

**SIGN AND COSTING OF SMALL SCALE CHROME
ELECTROPLATING PLANT.**

BY

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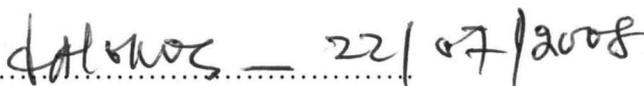
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CERTIFICATION PAGE

This thesis entitled "DESIGN AND COSTING OF A SMALL SCALE CHROME ELECTROPLATING PLANT", by BABALOLA RASHEED meets the regulation governing the award of degree of master of Engineering (M. Eng. Chem.) at Federal University of Technology, Minna and is approved for its contribution to knowledge and literary presentation.

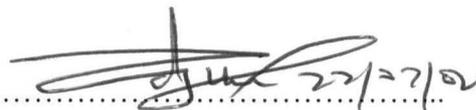
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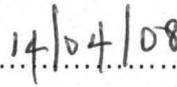
DECLARATION

by declare that this work was carried out by me and it is a record of my research work.

not been presented in any previous application for higher degree. All the sources of information are duly acknowledged by means of references.



Iola Rasheed



Date

DEDICATION

work is dedicated to the memory of my late mother Mrs. Sariyu Olalonpe Ayoka
ola and to my daughter Maryam Omomayowa Babalola and my yet unborn children.

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ABSTRACT

This project work was based on the designing of 2000kg/day of chrome electroplating plant for wrist watch chains. The design was carried out with the aid of an Engineering software and the various results of the equipment designed are shown in tables 4.1 to 4.4. A summary of the overall economic analysis has been presented in tables 4.6. The cost of equipment was \$21,220 (Twenty One Thousand Two Hundred and Twenty Dollars). The profit after tax (PAT) was \$61, 851 (Sixty One Thousand Eight Hundred and Fifty One Dollars); the pay back period (PBP) was 3 years and six months and the rate of return on investment was 27.28%.

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CHAPTER ONE

1.0

INTRODUCTION

This project work titled Design and Costing of small scale chrome electroplating plant is fully discussed accordingly.

Chrome electroplating is the covering of thin layer of chrome metal on a material usually a metal, plastic or other material in order to provide resistance to corrosion, wear, special appearance such as colour or luster or increase dimension, e.t.c. Thus, chrome electroplating is an electrolytic process whereby a chrome metal is cathodically deposited into another metal or a surface that has been made conductive. This process is done to import certain desirable properties while avoiding the prohibited cost of fabrications of the part entirely from chrome metal.

1.1 Background

Electroplating started as far back as 1800 when a chemist, Luigi Brugnatelli, carried out electroplating by making some metallic material surface to be coated with gold.

In 1805 a more refined electroplating process was conducted by a group of scientists in Britain and Russia when a printing press plate was electroplated with copper. By 1840 the process was refined and used for gold and silver plating.

In 1850 electroplating of bright nickel, brass, tin, and zinc on engineering material and other specific purposes was carried out by

scientists in great Britain. The years 1870 to 1940 were a quiet period for electroplating as no new scientific discoveries were made until the year 1945 when heavy gold plating for electronic components was discovered. In 1956 the "User friendly" plating bath that is based on formulae was developed for large-scale commercial use. (<http://electrochem2002>).

Nickel was the most widely used metal for corrosion resistance and shining property but the metal does not retain its luster for long.

Chrome metal thus quickly replaces nickel because it combined all other properties with the ability to retain luster for long.

Electroplating is the covering of a thin layer of metal on a material usually a metal plastic or other material in order to provide resistance to corrosion, special appearance such as colour or luster or increase dimension. Thus electroplating is an electrolytic process whereby a metal is cathodically deposited onto another metal or a surface that has been made conductive. This process is done to import certain desirable properties while avoiding the prohibitive cost of fabrications the part entirely from the metal used for coating other desirable characteristics of electroplating includes improved solder ability of the base metal (substrate), wear resistance, electrical conductivity, contact resistance and lubricity depending on their use. (Fowke et. al 1962).

Electroplated coating may be classified as either decorative or engineering but some fall into both.

Usually in electroplating a cheaper material is used as the basis material and plating technique is used to cover the surface of the material with an expensive material.

A typical example of the technique used to have expensive material is gold plating which is often found on religious articles.

Conventionally iron was and is still the most popular basis material for plating. We can therefore understand rust preventive technique easily when we concentrate on metal material, such as copper, brass, lead, zinc, as it's variations. Zinc plating is the most popular rust preventive plating method, followed by copper-nickel-chromium or nickel-chrome plating.

There are also alloys like brass, bronze plating as well as gold and silver plating.

Summarily, plating technique can be defined as applying thin coating and evenly deposited on the surface of base material. (Fowke et. al 1962).

1.2 Statement Of The Problems

Modern living conveniences such as wrist watches, spoons, openers, key holders, oven trays and transportation equipment such spare parts, car bumpers, bicycles and motorbikes parts are made

up of iron and steel, which are liable to corrosion and this can be prevented by painting, or electroplating.

In the recent years, the output of chrome plated product is increasing tremendously as paints peel off, Nickel does not retain its luster for long from the base material yet, only few industries are into chrome electroplating. Out of these few, most are not servicing the public, thus the need to have more chrome electroplating plant in Nigeria; Most people are in doubt what chrome plating plant entails and its financial feasibility. This project work is therefore aimed at:

- a. Carrying out a design of 2000kg/day of a small-scale chrome electroplating plant that is friendly to both human life and environment.
- b. To chrome plate wrist watch chains and other related materials.
- c. Prepare a financial feasibility profile for the plant operation.

1.3 Scope of Design work

- i. **Material and Energy balance**
- ii. **Equipment Design, Type and Specification.**
- iii. **Equipment Cost and Economy Analysis.**
- iv. **Cost Analysis**

1.4 Limitations

Findings of previous researchers were adopted

1.5 Approach

The approaches used in meeting these aims are as follows:

1. Source for local and international material on chrome electroplating.
2. Produce a flow diagram.
3. Carrying out material balance of the plant using MathCAD.
4. Carrying out energy balance on the plant using MathCAD.
5. Costing the plant using HYSYS/

CHAPTER TWO

2.0

LITERATURE REVIEW

2.1 Brief History

The early history of electroplating may be traced back to around 1800. A University Professor, Luigi Brugnatelli, is considered as the first person to apply electrodeposition process to electroplate gold. Brugnatelli was a friend of Alessandro Volta (after whom the electric unit "volt" has been named) who discovered the chemical principles that would make possible the development of electrical cells.

Volta's first actual demonstration of that was called. As a consequence of his development, Brugnatelli's early work using voltaic electricity enable him to experiment with various plating solutions.

By 1805 he had refined his process enough to plate a fine layer of gold over large silver metal objects. He wrote in a letter to the Belgian Journal of Physics and Chemistry (later reprinted in Britain), which reads:

"I have lately gilt in a complete manner two large silver metals, by bringing them into communication by means of a steel wire, with a negative pole of a voltaic pile, and keeping them one after the other immersed in amount of gold newly made and well saturated".(<http://electrochem,2002>).

Unfortunately for Brugnatelli, a disagreement or falling out with the French Academy of Science, the leading scientific body of Europe at the time, prevented Brugnatelli's work from being published in the scientific journals of his day. His work remained largely unknown outside of his native Italy except for a small group of associates. By 1839, however, group of scientist in Britain and Russia had independently devised metal deposition processes similar to those of Brugnatelli's for copper electroplating of printing press plates. By 1840, this discovery was adapted and refined by Henry and George Elkington of Birmingham, England for gold and silver plating.

Collaborating with their partner John Wright and using formulae developed by the later potassium cyanide plating baths, the Elkingtons were able to have the first viable patents for gold and silver electroplating issued on their name. From Great Britain the electroplating process for gold and silver quickly spread throughout the rest of Europe and later to United States. By the 1850's electroplating method of bright nickel, brass, tin, and zinc were commercialized and were applied for engineering and specific commercial purposes. In time, the industrial age and financial capital had expanded from Great Britain to the rest of the world. As a result, electrodepositing process expanding in scope and found more and more usage in the production of a variety of goods and services.

While this expansion was taking place, no significant scientific discoveries were made until the emergency of the electronic industry in the mid forties of the last century. The years from 1870 to 1940 were a quiet period as far as electroplating was concerned, significant only in gradual improvement in larger scale manufacturing processes, and reaction principles and plating bath formulae. During the later years of the forties, rediscovery of heavy gold plating for electronic components took place. In comparison to that, during the mid to later fifties the usage of new and more “user friendly” plating baths based on formulae were developed and introduced for large scale commercial use.([http:// electrochem 2002](#)).

2.2 An Over View Of Electroplating

Electroplating is electrodeposition of metals on metals alloys and non-metals. The objects of electroplating are as follows: -

- i. To change the surface properties of metals and non-metals.
- ii. To get improved appearance on basic metals.
- iii. To obtain improved resistance to corrosion, tarnish, chemical attack and wear.

Electroplating is performed in a liquid solution called the “plating bath”. The bath is a specially designed chemical solution that contains the desired metals (such as gold, copper, or nickel) dissolved in a form of submicroscopic metallic particles introduce in the bath to obtain smooth and bright deposits. The object that is to

be plated is submerged into the electrolyte (plating bath). Place usually at the center of the bath, the object that is to be plated acts as the Cathode, and the Anode placed at the opposite edges of the plating tank. It is then connected to the power supply (rectifier) thus causing film deposit on both sides of the cathode. In the bath, the electric current is carried largely by the positively charged ions from the anode(s) toward the negatively charged cathode. This movement makes the metals ions in the bath, to migrate toward extra electrons, that are located at or near the cathodes surface outer layer. (Adegoke R. 2001).

2.3 Theory Of Electroplating

If we have a solution of metallic salt and a potential difference is applied to this salt solution by means of two electrodes, it is ionized thus the metals ions migrate to cathode and are deposited there. If the anode is of the same metal of which the salt in solution is the anode metals passing into the solution in ionic form reform the salt. In this way there is a continuous deposition of metal on the cathode. (Fowke D. G. 1962).

For $ZnCl_2$ solution in water it is ionized as given below



Zinc will go to cathode and will get deposited there. Cathode ion will go to anode metals and will react with it to form zinc chloride.

2.3.1 Principles of chrome electroplating

The article to be electroplated is immersed in a solution containing dissolved chrome metal and made the cathode by connecting it to the negative lead of a low voltage D.C. supply(rectifier). The circuit is completed by immersing the Anodes e.g. wrist watch, into the solution and these are connected to the positive lead. Dissolved chrome ions are driven by a D.C. electrical current of low voltage and high amperage. The ions gain electrons at the cathode, causing the wrist watches to be plated. The potential difference applied between anode and cathode which is usually between 2 and 16 volts D.C. is the driving force for the transfer of charge ions across the metal solution interfaces.(Joe I. et al 1979).

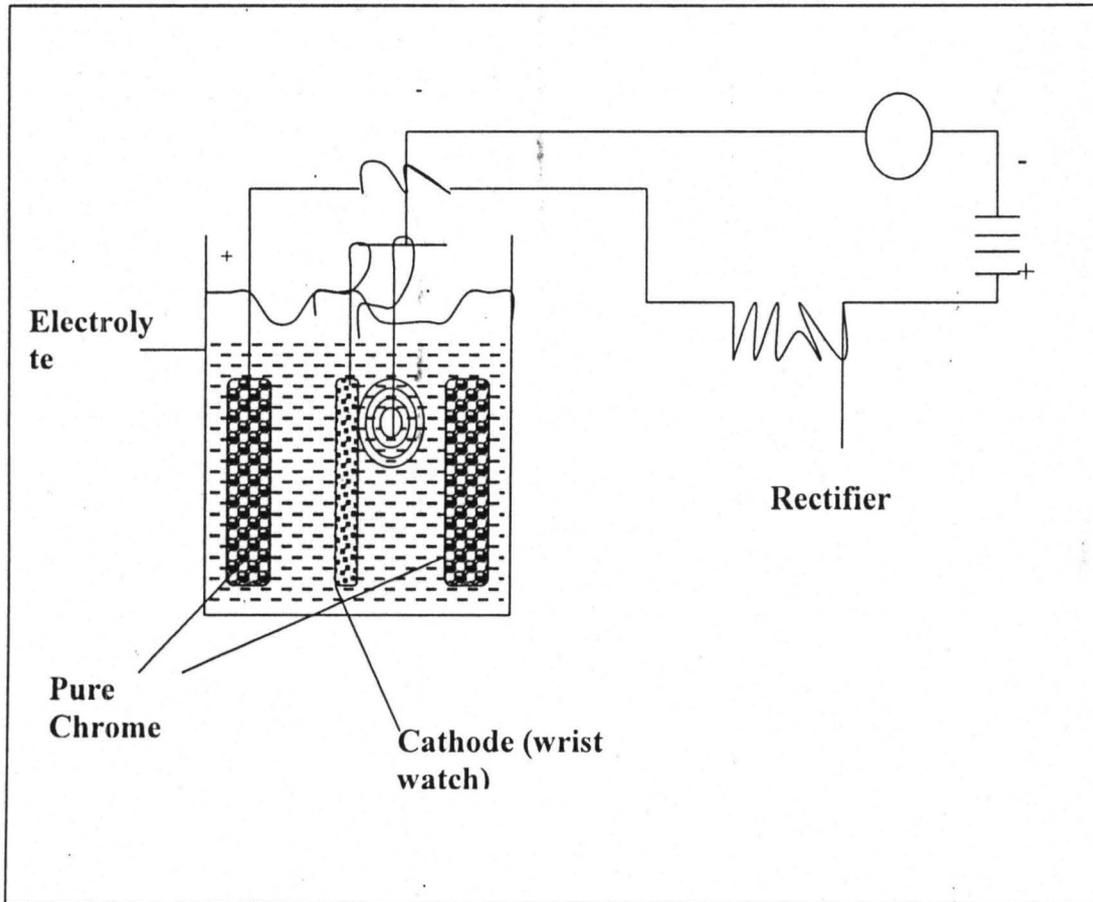


Fig 2.1 Schematic diagram of chrome electroplating of wrist watch chains

2.3.2 Table 2.1 Electrochemical series

Manganese.....	-1.18	ELECTRONEGATIVE METALS
Zinc.....	0.76	More cathodic potentials
Chromium.....	0.56	less noble potential
Iron (ferrous).....	0.44	more noble potentials
Cadium.....	0.40	
Indium.....	0.44	ELETROPOSITIVE
METALS		
Cobalt.....	0.28	
Nickel.....	0.25	
Tin (stannous).....	0.14	
Lead hydrogen.....	0.12	
Copper (cupric)	+0.34	
Copper (cuprous).....	+0.52	
Silver.....	+1.50	
Gold.....	+1.6	

The electrochemical series lists the element in the order of their standard electrode potentials and to a degree, indicates the ease with which they ionize. As displacement series; usually a metals with a higher negative value will displace one lower in the series.

(M. M. Upper 1995). For example, if a piece of zinc is immersed in a solution of copper sulphate, copper is immediately deposited upon the zinc by chemical action without the assistance of applied current. Electropositive metals are deposited in preference to more electronegative metals.

2.3.3 Quantity of metal deposited

Faraday's laws of electrolysis guide quantity of metals deposited. According to these laws, the quantity deposited depends on the quantities of electricity passed per second and the chemical equivalent of these substances. (Joe L. et al 1979).

2.3.4 Rate of deposition

The minimum quantity of electricity require to deposit the equivalent weight in gramme of metals is 96500 coulombs. For instance, 96500 coulombs will discharge 1 mole of hydrogen; this quantity of electricity is called Faraday. One faraday is equal to one mole of electron since, the discharge of one mole of single charge ions involved the transfer of one mole of electron which contains Avogadro's number i.e. 6.02×10^{23} of electrons

Thus: $W=ItA/ZF$2.2

Where A is the atomic weight of metals deposited, ZF is a faraday constant. The relation between atomic weight and chemical equivalent is atomic weight=equivalent weight x valency.

The valency may be defined as being the number of atoms of hydrogen with which one atom of the element combines, or which it replaces. The figure is always a whole number, for example silver is monovalent and cadmium divalent, or in other words, have valencies of one and two respectively.

2.3.4.1 electrolytes

The electrolyte is the medium that carries the current by means of ions. It is the ability of a solvent especially water to ionize substances, i.e. split them into two components which carries negative and positive charge that make electroplating possible.

2.3.5 Presence of ion and electrical conductance

When an electric field is applied to metals, according to the direction of applied field, electrons start moving from negatively charged end of metal to its positive end, this way current flows in metals. Thus, a substance conducts electricity when an electron or some other electrically charged particles like ion has to move through it. In all electrically conducting solution, the solute molecules give rise to two types of electrically charged particles or

ions. The negatively charged types of ions are called anions and the positively charged ones are called cations.

2.3.6 Electrolysis

When a direct electric current is passed through an electrolyte (such as a molten salt or an aqueous solution of a salt, acid or base), chemical reactions take place at the contacts between the circuit and the solution. This process is called electrolysis. Electrolysis takes place in an electrolytic cell.

The components, which make contact with the electrolyte, are called electrodes. The electrode which is attached to the negative pole of the battery, and which supplies electrons to the electrolyte, is called cathode. Reduction takes place at the cathode.

The electrode which is attached to the positive pole of the battery, and which accepts electrons from the electrolyte is called the anode.

Various reactions take place at the electrodes during electrolysis. In general, reduction takes place at the cathode, and oxidation takes place at the anode. (Ollord et .al 1974).

2.3.7 Faraday's laws

In 1883, Michael Faraday explained the nature of electrolysis. Faraday's law is basic to the understanding of all electrolytic processes and may be summarized by the two following statements:

1. The amount of chemical change (metals deposited) that is produced by an electric current passing through solution in a cell is directly related to the quantity of electricity that flows.
2. The amount of substance liberated (i.e. metal deposited) by a given quantity of electricity is directly related to the gram equivalent weight of the substance. More specifically, 96,500-ampere seconds (or coulombs) reduce one gram equivalent of metals ions to metals atoms.

Faraday's law is used by the plater to determine the plating time and current required to deposit a specified weight (and considering the specific gravity and surface area, also a specified thickness) of a metal. Table 2.1 gives the electrochemical equivalents for some typically electroplated metals. The amount of time required to deposit a certain thickness at a specific current density can be calculated from this data. $Q = IT$ 2.3

where Q is the qty of electricity passed, I is current and T is time.(upper, 1995).

2.3.7.1 Cathode efficiency

Faraday's law states that the total amount of chemical change at an electrode is directly related to the quantity of electricity (that is,

the current flowing). In plating, however, the main concern is the quantity of metal deposited. Any other reactions, such as the liberation of hydrogen at the cathode, decrease the cathode efficiency. The average cathode efficiency percentage is determined by dividing the actual weight of metals deposited by the theoretical weight and then multiplying by 100. Change in solution composition and other external factors can improve cathode efficiencies to some degree; however the solution used is the major factor for determining efficiency.

2.3.7.2 current distribution

Current density is the applied current divided by the total surface area of the work piece.

**Table 2.2: Electrochemical Equivalent And Deposit Metal Weight
Calculated from Faraday's Law (Uppal, 1995)**

METAL	VALENCE	ATOMIC SPECIFIC WEIGHT		THICK COATING	WEIGHT IN 1 MIL (Oz/ft ²)	WEIGHT GRAVITY
Cadmium (Cd)	2	112.40	8.65	0.074 (2.097)	0.71 (8.659)	9.73 (4.125)
Chromium (Cr)	6	52.01	7.1	0.011 (0.323)	0.59 (7.146)	51.8 (22.110)
Cobalt	3		7.1	0.023 (0.646)	0.59 (7.146)	25.9 (11.060)
Copper (Cu)	2	58.93	8.9	0.039 (1.099)	0.74 (8.719)	19.0 (7.926)
Gold (Au)	1	63.54	8.96	0.084 (2.370)	0.74 (8.935)	8.84 (3.770)
Indium (In)	2	63.54	8.96	0.042 (1.186)	0.74 (8.935)	17.8 (7.540)
Iron (Fe)	1	197.0	19.3	0.236 (7.348)**	1.47 (19.32)**	6.2 (2.631)
Lead (Pb)	3	197.0	19.3	0.079 (2.449)**	1.47 (19.32)**	18.6 (7.887)
Nicket (Ni)	3	114.82	7.31	0.045 (1.428)**	0.56 (7.278)**	12.0 (5.092)
Palladium (Pd)	2	55.85	7.86	0.037 (1.042)	0.65 (7.868)	17.9 (7.54)
Platinum (Pi)	2	207.19	11.34	0.136 (3.865)	0.94 (11.350)	6.9 (2.936)
Rhodium (Rh)	2	58.71	8.90	0.039 (1.095)	0.74 (8.880)	19.0 (8.044)
Silver (Ag)	2	106.4	12.0	0.064 (1.985)**	0.86 (24.34)**	13.5 (6.045)
Tin (Sn)	4	195.09	21.41	0.058 (1.819)**	1.60**	27.9 (11.77)
Zinc (Zn)	3	102.9	12.4	0.041 (1.280)**	0.95 (8.29)**	22.9 (9.73)
	1	107.87	10.5	0.129 (4.042)**	0.79 (10.5)**	62 (2.605)
	2	118.69	7.30	0.078 (2.214)	0.61 (7.30)	7.8 (3.30)
	4	118.69	7.30	0.039 (1.106)	0.61 (7.30)	15.6 (6.6040)
	2	65.38	7.14	0.043 (1.219)	0.59 (7.15)	14.3 (5.8630)

***Assume 100% cathode efficiency for metal deposition**

Current Density is usually measured in A/ft^2 or A/dm^2 . Depending on the system of units.

In Electroplating, current is concentrated at edges and points as well as in area closer to the opposite electrode (anode) and these latter areas consequently receive a greater deposit thickness (see fig 2.1). Normally, a part is plated to meet a minimum thickness specification. Since the excess metal thickness in high current density areas is usually not desired, the throwing power is defined as the electrolyte's ability to minimize the difference in deposit thickness between high and low current density areas.

Plating solution in which the cathode efficiency decreases with an increase in current density, generally displays better throwing power because less metal will be deposited in the less efficient high current density areas. Cyanide zinc is an example of such a bath. Chromium plating, on the other hand, becomes more efficient with an increase in current density and consequently has poor throwing power.

2.3.8 Introduction to basic electroplating process

In the simplest terms, plating transfers metals from positive Anode to the negative Cathode following Faraday's law.

Objects are immersed into chemical baths or vats to change their surface condition. Every plating is UNIQUE and the number of

tanks and their chemical make up differ based on desired result.

Metal slabs or balls (Anode) are placed in an electrolyte solution.

Dissolved metal ions are driven by a DC electrical current of low voltage and high amperage (Rectifier). During electroplating pure metal dissolving from the anode is deposited on the cathode by an electron gain – transfer method which can be represented thus: -



M^+ represents metallic ion required Cu^+ and Ni^+ represent copper and nickel ions, and Cu and Ni represent deposit of copper and nickel.

Objects to be plated are placed on racks or perforated barrels and immersed in a series of chemical solutions to prepare them for plating. The objects to be plated must be "SURGICALLY" clean in order to be successfully plated.

2.4 Plating Process Line

Plating process can be divided into 3 basic units namely:-

- i. Pre-treatment
- ii. Plating
- iii. Post-plating

2.4.1 Pre-treatment

Each material has its own treatment procedure. It is therefore necessary to know the nature of material or basis metal. Aluminum for example, cannot be cleaned in solutions formulated for cleaning steel. The nature of contamination and an incorrect identification may result in destruction or damage of the parts. Cleaning affects adhesion, appearance, composition and corrosion resistance of final deposit. The plater must therefore, be informed about pre-existing contamination. Such as inclusions in the base materials. Information supplied often help a customer – client relation resulting in successful metal finishing. The material is therefore, derusted or cleaned using necessary acid and depending on the base material. Acids used are sulfuric, hydrochloric, nitric or a combination of these. This removes rust, scale or smuts.

Depending on the material and configuration, it is polished or buffed/mopped. After proper rinsing, it is transferred to the degreaser tank where oils and greases are removed from the surface, then rinsed and ready for plating. Sandpapering is another manual method of polishing the surface of materials. These pre-treatment processes of pickling and degreasing can also be achieved by electrolysis called Electrolytic degreasing or pickling. The composition of materials or chemical used is based on the material being cleaned and it is unique for each base material. It is

commonly accepted and often quoted by electroplaters that can make a poor coating performed with excellent pretreatment, but one cannot make an excellent coating perform with poor pre-treatment. Surface pretreatment by chemical and or mechanical means is important not only in the case of preparations for electroplating but is also required in preparation for painting. In either of these, methods are designed to ensure good adhesion of the coating the processes include solvent degreasing, alkali cleaning and acid dipping which are described below (Adegoke R. 2001)

2.4.1.1 polishing

This is mechanical cleaning of the substrate to remove surface defects (scratches, etc) by polishing and buffing.

2.4.1.2 swirling

This is a process of rinsing the pretreated articles in cold distilled water in order to avoid contamination of the electrolyte.

2.4.1.3 solution heating

For many of the processes associated with electroplating it is necessary to heat the solution. The available methods of heating include steam, hot water, gas and electricity.

2.4.1.4 solution cooling

A number of processes in which heavy current densities are employed or are operate at fairly low temperature may required

cooling to prevent rise in solution. The cooling media employed are well water, recirculated water chilled water and brine.

2.4.2 Engineering application

Plating for appearance (decorative plating). For Decorative Plating Metals Used are gold, Nickel, brass/radius and copper. For special decorative effects tin, zinc, cadmium, lead, palladium and platinum, etc are also used.

2.4.2.1 plating for protection

Steel must be protected from rusting chemical attack and so it is electroplated. Protective metals applied are zinc, cadmium and tin, etc. where both protection and good appearance are desired steel is given successive coating layers of copper, nickel and chromium. Copper and Nickel provide protection against rusting and chemical attack. The chromium plate above all plate, gives clean metal appearance. Nickel dulls by tarnish when exposed to atmosphere and chromium does not.

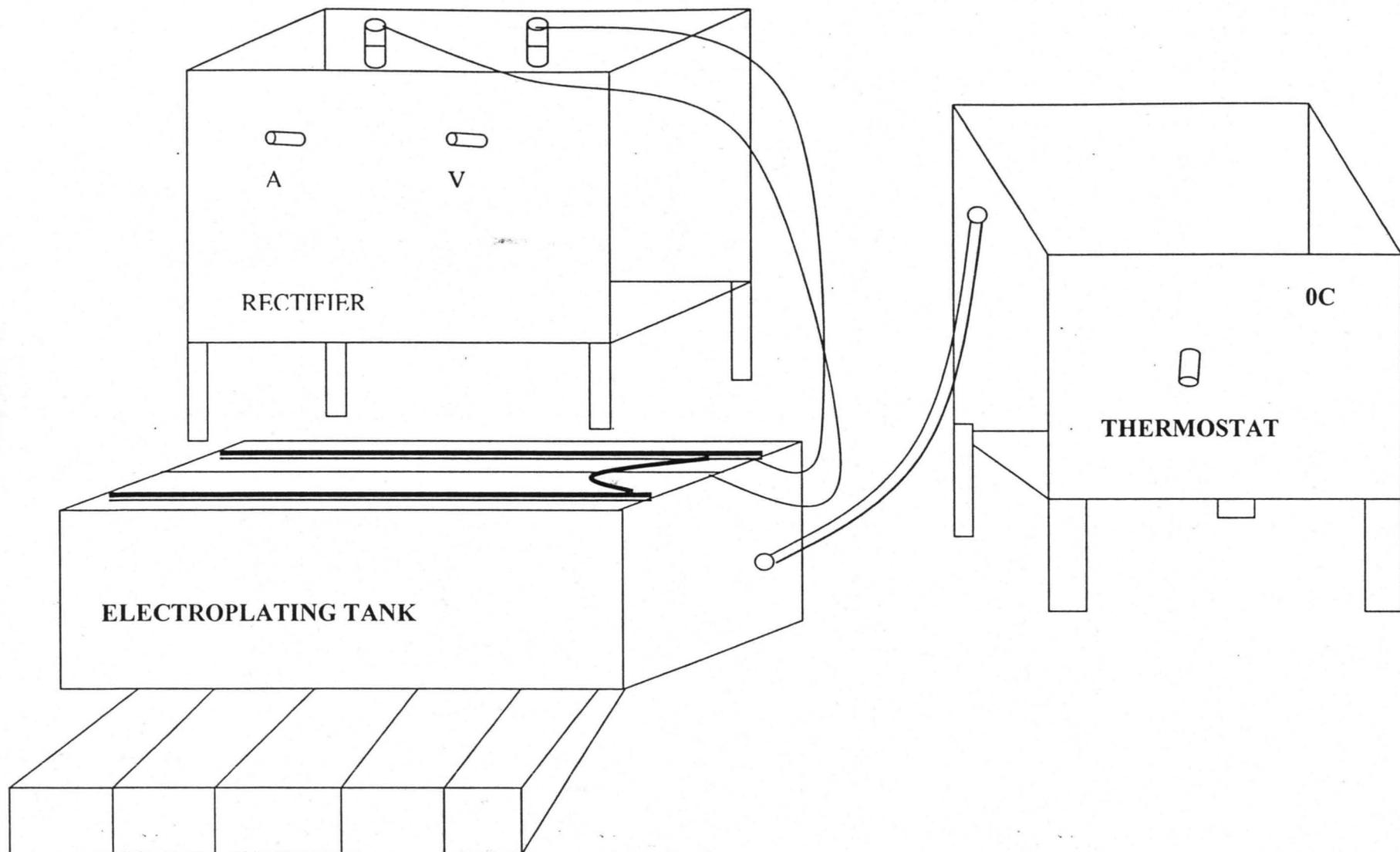


Fig. 2.2: Schematic Diagram of Electroplating Equipments.

2.4.2.2 plating bath constituent

Although it is often possible to electrodeposit metal from a simple solution of a metallic salt alone, it is rarely a practicable commercial process. Modern plating baths are somewhat complex and may contain as many as six or eight constituents, all which play an important part. Chlorides, for example, are added to nickel sulphate solutions to promote anode dissolution. Boric acid acts as 'buffer' and maintain p^H equilibrium. Colloids to refine the texture of the deposit, other metals and organic Brightener to give luster, and anti-pitting agent to prevent 'pin-holing' due to hydrogen bubbles and Leveler allow even deposit on irregular object.

Operating Condition

For optimum plating some parameters such as p^H , temperature, current etc. P^H should be in range of 3-4, temperature from room temperature to $60^{\circ}c$ depending on anode metal and a battery or rectifier to supply low voltage of 2-8v.

2.4.2.3 solution agitation

With many processes it is necessary to keep the solution well mixed in order to prevent local variation in concentration. Unless the solution is agitated there will be a reduction in the metal content of the solution surrounding the cathode, which will result in the production of a rough or "burnt" deposit.

2.4.2.4 electroforming

By electro-forming is meant formation of articles by electro-deposition of metal. In electroforming the layer of electrodeposited

metal is quite thick. Automobile, aircraft, radio, radar and ammunition industries use parts made entirely by electroplating. Mandrels are used for depositing metals over it. The electro-form is removed without damage from its mandrel after obtaining the desired thickness

2.4.2.5 electroless plating

Electroless plating is a process of plating by chemical reduction. For example, nickel can be plated by the addition of a reducing agent, such as sodium hypophosphate, a solution, to nickel salts. The nickel ion is reduced to the metals, which under proper conditions is deposited as a coherent and adhesive plate rather than simple as a powder throughout the solution. It is used, where electroplating is impossible, example plating the insides of pipes.

2.4.2.6 displacement plating

Displacement plating occurs when the metal deposited is nobler than the substrate, and the substrate dissolves. The reaction ceases when the substrate is completely covered by a pore-free deposit.

2.4.3 Electroplating equipment

Electroplating may be divided into barrel, rack and strip continuous (reel-to-reel) plating. Barrel plating is used for plating smaller parts in some electrolytes. Rack plating is used for larger parts and for chromium plating. (Adegoke R. 2001).

2.4.3.1 barrel plating

Barrel plating is usually performed in either horizontal or oblique barrels constructed of polypropylene or other suitable plastics. The walls of the barrels are perforated, and the barrel is plated during

plating. Electrical contact is obtained via a flexible conductor known as a dangler. Some barrels are not perforated, but contain the plating solution and an anode.

Mixed loads are not recommended in barrel plating unless the plater is not too concerned about plating thickness and distribution, or unless the parts require just a flash plate. Longer or larger parts in the mixture will usually receive the greater deposit and buildup on high current density areas will be increased. In addition, sorting the parts after plating is time consuming.

Larger pieces weighing more than 1 lb (0.5kg) and containing sharp edges should not be barrel plated because they will be damaged in the tumbling action of the barrel and the barrel itself may be damaged. Flat and lightweight parts should not be barrel plated because they tend to stick together and do not tumble properly. Wire forms are more easily rack plated than barrel plated.

2.4.3.2 rack plating

Rack plating is usually employed in the processing of parts that are too heavy, too large or too complex in shape to be barrel plated. The parts can vary from a small knob that is to be nickel-chromium plated, to a large roller, weighing a ton or more for hard chromium plating. Rack plating is used with manual, semiautomatic and fully automatic machines.

One of the most important considerations in designing a plating rack is that the rack must have adequate current carrying capacity. Racks are usually constructed of copper because of its high current

carrying capacity, ease of fabrication, and relatively high strength. After fabrication, the racks are covered with an inert insulating material to protect them and prevent metal plating on them, and to keep the plating solutions from becoming contaminated. Parts are hung or dipped to the insulated rack. The contact to the part should be made on a non critical area of the part such as the back, inside, or through a hole.

Another important factor in designing a plating rack is the position of the work on the rack. It must be positioned to obtain the most uniform current distribution possible and to prevent entrapment of air or gas in holes or pockets, which will restrict deposition of metal in these areas, resulting in non uniform thicknesses. The position of the plating rack in the processing track is also important for obtaining as uniform a plating deposit as possible. If the work is located too close to the anodes, excessive buildup of plating thicknesses, known as burning occurs in the high current density areas of the part. The parts are usually tracked so that they tend to shield the edges of adjoining parts, and are frequently racked back to back to increase the capacity of the rack and at the same time reduce the deposition of the metal in non critical areas. If a particular part has a deep recess, an auxiliary anode can be used to ensure the deposition of the required thickness in the recessed area.

2.4.3.3 strip plating

Strip plating is a plating process whereby the work piece is a continuous strip being pulled through each process station (tank) by a take-up null. Wire and lead frames are commonly strip plated with tin,

tin lead, nickel, and precious metals. Steel sheet may be continuously zinc, tin, chromium, copper, brass, nickel iron or nickel zinc plated. The strip may be plated at specific points as it goes through the cycle, such selective plating is very common with precious metals.

2.4.4 Process tank

Tanks may be fabricated of hot rolled low-carbon steel and protected with a rubber or other suitable liner rated to withstand the bath's corrosive action and to keep the steel tank from contaminating the solution. Polypropylene and other high-strength plastics may be used for smaller tanks. Alkaline-cleaner tanks and some plating solutions do not require liners. Rinse tanks may be equipped with spray nozzles and the water counter-flowed back into the previous rinse tank to reduce water usage.

2.4.5 Temperature control

Temperature control is important in most plating solutions. Some require cooling heating or both. For cooling, cold water piped through reeling coils of suitable construction may be sufficient or heaters dangers may be required. Heat may be provided by steam through heating coils or by electric immersion heaters. Instruments are commonly used to automatically control plating bath temperatures.

2.5 Power Supplies

Alternating current is converted to direct current by means of a rectifier or motor generator set, with rectifiers being preferred. Regulated and unregulated power supplies are available that provide filtered direct current with good reliability. A regulated power supply

may be of the silicon-controlled or saturable reactor type. These power supplies maintain constant current or voltage under varied loads and are generally used in large operations. The more common unregulated types include the tap switch and manual powerstat. Both air and liquid cooled rectifiers are available. The major advantage of the more expensive liquid-cooled type is that the unit is completely sealed so that rectifier components are not exposed to the atmosphere, which may be very corrosive in a plating shop. As a general rule, a rectifier should be operated at a minimum of 50% of its maximum rated output current to provide low-ripple direct current.

2.5.1 Anodes

The anode carries a positive charge and completes the electrolytic circuit. In most cases, the anode is fabricated of the metal being plated and serves to replenish the solution with metal ions. However, insoluble (metal) anodes are used in some applications, such as chromium plating and with precious metals. Anodes may be fabricated metal slabs or balls, or chips contained in an inert basket. The metallurgical structure and composition are important considerations for proper anode performance. Titanium is often used for the baskets because of its inert oxide film.

Most modern nickel plating installations use nickel chips in titanium baskets, sheets, or cathode-deposited shapes. Zinc is usually plated from ball anodes, and copper from ball or bar anodes.

2.5.2 Automated control

Electroplating is subject to a wide variety of variables that frequently change. Automatic controllers add stability and consistency to the operation and should be used wherever possible. Automatic ampere hour feeders monitor the plating time and may be used to approximate thickness as well as automatically feed addition agents. Solutions pH may be controlled automatically with a relatively inexpensive piece of equipment, such control is especially useful in electrolytes where the pH tends to rise due to generation of the hydroxyl ion, such as in nickel and chloride zinc solutions. Rinse waters may be controlled by automatic conductivity meters connected to solenoid valves, these ensure good rinsing with reduced water consumption.

2.6 Substrate Preparation

In preparing the substrate for plating, it is important to properly select the correct pretreatment method. Pretreatment influences the adhesion, appearance, composition and corrosion resistance of the deposit.

Some of the factors that should be considered when selecting the pretreatment cycle are type of substrate, nature of the contamination, how the part is used, and part geometry. Each basis metal may require a different pretreatment. Aluminum, for example, cannot be properly processed in solutions formulated for steel. Even variations in alloy may cause the finisher to change pretreatment. (Adegoke R. 2001)

Typical Processing Cycles for Substrate Preparation for

Electroplating.

I(a) Low-carbon steel-oily, not rusted

1. Soak clean, 180°F (82°C), 2-3 minutes.
2. Cold water rinse, 30 seconds.
3. Anodic electro clean, 180°F (82°C).
4. Rinse
5. Acid dip, 30 seconds
6. Cold water rinse, 30 seconds
7. Plate

I(b) Low-carbon steel - rusted

1. Soak clean, (180°C), 2-3 minutes.
2. Cold water rinse
3. Periodic reverse electroclean, room temperature, 30-60 A/H² (3.2 & 4A/din²).
4. Plate.

II. Stainless steel and high-nickel alloys.

1. Soaked clean, 1-5 minutes
2. Cold water rinse, 30 seconds
3. Periodic reverse electroclean (10 seconds anodic, 7 seconds cathodic), room temperature, 30-60 A/H² (3.2-6.4 A/dm²). 3-5 minutes
4. Cold water rinse, 30 seconds.
5. Hydrochloric acid dip. 30% by volume, room temperature, one minute.

6. Wood's nickel strike, 20-60 A/H² (2.2-6.4 A/dm²). 1-3 minutes
7. Cold water rinse, 30 seconds
8. Plate.

III Cast Iron

1. Soak clean, 103 minutes
2. Cold water rinse.
3. Periodic reverse electro clean (7 seconds anodic, 5 seconds cathodic), 5 minutes
4. Warm water rinse.
5. Cold water rinse
6. Hydrochloric acid, 30% by volume, room temperature in seconds.
7. Cold water rinse
8. Plate

IV. Cast iron alternate method

1. Cathodic electro clean, 1-3 minutes
2. Warm water rinse
3. Cold water rinse
4. Hydrochloric acid dip. 10% by volume, 5-10 seconds
5. Cold water rinse
6. Plate.

V. Wrought aluminum alloys most alloys

1. Soak clean, nonetch cleaner, 1-1 minute
2. Cold water rinse
3. Alkaline etch clean

4. Cold water rinse
5. Desmut in a solution containing 50% by volume nitric acid, 25% by volume sulfuric acid, 25% by volume water, and 1 lb/gal (120 g/l) fluoride salt at room temperature for 20-45 seconds.
6. Cold water rinse.
7. Zincate, 5-30 seconds.
8. Rinse.
9. Plate, preferably first in a cyanide copper strike.

VI. Wrought aluminium alloys 5000 and 6000 series

1. Soak clean, nonetch cleaner, 1-3 minutes.
2. Cold water rinse.
3. Acid etch, 160°F (70°C) 1-3 minutes.
4. Cold water rinse.
5. Nitric acid, 50% by volume, room temperature, 30 seconds.
6. Cold water rinse.
7. Zincate, 5-30 seconds.
8. Cold water rinse.
9. Plate, preferably first in a cyanide copper strike.

VII. Sand or die-cast aluminium alloys

1. Nonetch soak clean, 1-2 minutes
2. Cold water rinse
3. Alkaline etch clean.
4. Cold water rinse
5. Desmut in a solution containing 50% by volume nitric acid,

5. Desmut in a solution containing 50% by volume nitric acid, 25% by volume sulfuric acid, 25% by volume water, and 1 lb/gal (120 g/l) fluoride salt at room temperature for 20-45 seconds.
6. Cold water rinse
7. Zincate, 5-30 seconds
8. Rinse
10. Plate, preferably first in a cyanide copper strike.

VIII. Copper and Copper alloys

1. Soak clean, 3-5 minutes
2. Cold water rinse
3. Electro clean (cathodic or anodic depending on formulation), 30-90 seconds.
4. Cold water rinse.
5. Acid dip, 10% sulfuric acid or dry acid salt, 30 seconds.
6. Cold water rinse.
7. Plate.

IX Zinc-based die castings

1. Soak clean, 3-5 minutes.
2. Cold water rinse.
3. Spray alkaline clean, 30-60 seconds.
4. Cold water rinse.
5. Anodic electro clean, 10-25 A/H² (1.1-2.7/dm²), 25-50 seconds.
6. Cold water rinse.
7. Acid dip. 30 seconds

8. Rinse
9. Plate, preferably first in a cyanide copper strike.

2.6.1 Product finishing

- The acid dip may be 10% sulfuric acid at 122-180°F (50 ox/gal (120 g/l) dry acid salt at 77-140°F (25-60°C).
- The two most widely used processes for the pretreatment of aluminium alloys before electroplating are the zincate process and the stannate process.
- On difficult to plate alloys, use double zincate with an intermediate nitric acid dip to improve deposit adhesion.
- Acid dip may be 1% hydrofluoric acid, 1% fluoboric acid or a dry acid salt.

Table 2.3: American Society for Testing and Materials Recommended Policies for Preparation of Substrates to be Electroplated.

(Fowke, 1962)

Metal Substrate	Standard Number
Low-carbon steel	B.183
High-carbon steel	B.242
Zinc alloy disc castings	B.252
Aluminium alloys	B.253
Stainless steel	B.254
Copper and copper-based alloys	B.281
Lead and lead alloys	B.319
Iron castings	B.320
Nickel	B.343
Magnesium and magnesium alloys	B.480
Titanium and titanium alloys	B.481
Tungsten and Tungsten alloys	B.482
Nickel alloys	B.558

Several stages are generally required to provide adequate cleaning of the substrate and activation pre-cleaning, intermediate alkaline cleaning and electro cleaning. Pre-cleaning is designed to remove a large excess of soil, especially deposits of butting compound or grease. It is also useful in reducing the viscosity of waxes and heavy oils to enable later cleaning stages to be more effective, or to surround fingerprints and dry dust with an oily matrix to facilitate removal by alkaline cleaners.

Intermediate alkaline cleaning removes solvent residues and residual soil that has been softened or conditioned by precleaning. Spray or soak alkaline cleaning may also be used as a precleaning stage, followed by additional alkaline cleaning, if the soil and metal lend themselves to this treatment. Electro-cleaning is soak cleaning with agitation provided by the upward movement of bubbles of hydrogen or oxygen formed by the electrolytic decomposition of water in the solution.

Some parts cannot be etched because surface finish must be maintained, just as part used in structural applications should not be subjected to pretreatments that may cause hydrogen embrittlement. The design of the work may require special handling and surface treatment. For example, a large part may require external manual finishing or parts with deep recesses or blind holes may require special handling and drainage techniques to avoid excessive drag-out and cross-contamination.

2.6.2 Operating parameters

The four main concerns in electroplating are temperature, pH, and chemistry of the plating bath as well as current density. Most plating solutions have an optimum temperature range for producing best results, and close control of temperature is important for proper current control. As the temperature of the solution increases, conductivity increases, and therefore the current increase for a fixed applied voltage, the converse is also true. Over plating or under plating occurs if the temperature is not maintained properly.

The pH control of plating solutions is necessary to maintain the acidity or alkalinity that has been determined to produce the best results. Appearance, stress, leveling, electrode efficiency and coating hardness are influenced by the pH of the solution. Current density is a very important variable in all electroplating operations. The character of the deposit, its distribution, the current efficiency, and perhaps whether a deposit forms at all may depend on the current density employed.

2.7 Comparison of product with its competitors

The competitors of chrome are: Nickel, zinc, gold, tin, and copper. Amount all the competitors chrome stands out as the most durable of all and that which give the best satisfaction on Metal because of its shining and wear resistance properties any chrome Plated materials do not loose their appearance and never break, these make It has higher quality than its competitors e.g. Chrome plated jewelries, Automobile, motorcycle and bicycle part last longer and regains its shining ability.

2.8 Quality Control Aspect of Electroplating Technology

Since the main purpose of electroplating is to get a rust-free, lustrous finish on the object, apart from other engineering application of increasing surface thickness, a deviation from this target, is a deviation from excreted result.

Quality control therefore in planting starts from the beginning to the end to get customer fully satisfying and satisfied.

Material to be plated must be specified since some finishes cannot be applied to certain materials. Metals fabricators and spare parts manufactures must clearly understand the coating needed rather

than what they just thought they wanted.

While designers are expected to know what the conditions the plated product will survive, engineers must also know and understand coating's limitation.

Since each of the coating; zinc, nickel, copper, chrome has unique properties with respect to use and environment, selection of a single coating or a combination for a given application is never a matter of chance, aesthetic nature or serious economic of financial consideration. Expected coating and selection require specialized knowledge as to use, application and shelf life.

To become a better customer and quality control conscious, the quotation that "the requirement of a supplier must be established long before he produces any material", by W Edwards Deming, out of the Crisis of quality control theorem must be followed.

Therefore, nickel plating is the best for food equipment/ material like oven trays, and the like, while chrome is acceptable for car bumpers.(Ollord, E.A. 1974)

In effect, quality specification and control is to make both plater and client a synergy of successful ventures. From the start therefore fabricators, engineers and benefactors of plating process will not choose high carbon steel, instead of low carbon steel, defective, pin-holed and scratched, pitted, scrapped metal, under the guise or disguise of "no-fund" will be a penny wise, pound foolish situation.

From quality assurance stand point, platers will use his science of

materials knowledge to help client in the right choice of materials, dimensions, tolerance, and even in designing for improved palatability. Product starts receiving good control and specification from inception or conceptualization stage.

Plating quality is also controlled if the following salient point and actions are taken from the shop floor. Plating solutions are given preventive maintenance approach which makes plating to be cost effective, efficient, interesting and satisfying. Platers therefore tend towards zero control commandments to make plating profitable.

Tanks or Vats must be designed to specification with regard to lining. Most of the plating tanks are lined with polypropylene, rubber or fibre-glass materials with no outlet taps or valves.

There is need to periodically filter the solutions. Nickel solutions are regularly filtered or ultra filtered.

Air agitation and cathode rod movement are needed plating solutions like nickel to reduce hydrogen-oxygen embrittlement (covering) on the plated objects.

Some good plating will not take place unless at temperature above room temperature. Nickel plating and some cleaning (degreasing) processes are few examples.

During plating, operation conditions are religiously adhered to. The current voltage, current density, pH, temperature are closely monitored for good plating to take place.

Some shop floor problems reduced to the bearest minimum as a result of these quality control measures.

Constant addition (little) as the plating is done daily. Such daily addition of individual or formulated "propriety" salt are expected to keep platers out of trouble. Large addition of salt or additives like levelers, brighter, conducting salts are regarded in modern plating as poisonous additions.

Bus bars, anodes must be periodically wire brushed, cleaned and kept in a conducting state.

Plating time (duration) must be strictly adhered to, to avoid burning and uneconomical coating. (Lesile,1986)

Equipment must be given periodic servicing and on-line daily, weekly, and monthly maintenance.

To maintain uniform coating, chemical composition of the electrolyte's as well as electrochemistry of the bath must be maintained.

This is not least exhausted but proper adherence to these as well as good house keeping will lead to trouble-free plating.

2.9 Health and Safety Aspect

The very nature of Electroplating processes, products, and by products makes health and safety a primary consideration in setting up an electroplating plant of this nature. Most important fundamental principal is that the safe handling of chemicals, operation of plant and processes is a management responsibility.(Lesile,W.Flott 1986)

Proper facilities, and wares provided must be used judiciously, appropriately and correctly. General precautions must be adhered to properly.

In addition to the wok shop and laboratory design which must

comply with health and safety regulation and requirements, a well organized medical service plays an important role in having efficient and effective workforce. A periodic medical examination is necessary in order to maintain the general good health of employees,(Adegoke R.,2002)

Enforcement of safety rules and regulations, mixing, and preparation technique of plating solutions, proper operating technique of machines and laboratory equipments and safe handling of chemical and waste are all necessary for a productive and profitable plating shop.

In addition, employees should be encouraged to be interested, alert and observing. Demonstration of respect for safety measures against fire, explosion and other hazards does much to endanger the same attitude amongst those who later carry out electroplating processes as operatives.

Chemical used in electroplating are toxic, carcinogenic and harmful. Cyanides must be stored away from acids and careful storage technique must be emphasized.

It is necessary that one of the supervisors should be trained in first aid or as a safety officer and there is a need to train all workers on matters of health and safety.

2.10 Important Uses and Application of Electroplating Technology

Modern living conveniences and transportation equipment are made up of coated iron since iron is corrosive to air, moisture and fumes. With the remarkable increase of iron products, in recent years, the output of plated products is also increasing tremendously. As

electroplating technique has made great advances, the importance, uses and application of electroplating in industries has widely come to be recognized.

Although plating is applied commercially to copper, brass, nickel-brass, zinc, zinc-base die-casting the greatest proportion is applied to steel.

A small amount of plating is applied to non metals such as plastics, but these must first be coated with some electrically conductive materials. The plating materials are tin, zinc, silver. Nickel and chromium.

However, except in connection with production of tin plate for can industry most plating is done with nickel and chromium. Nickel plating is widely used for corrosion resistance, but since it does not retain its luster for long, it has been replaced to a great extent by chromium. However, chrome is seldom used alone and most chrome plating is a three layer process consisting first of a layer of copper, then one of nickel and finally, a thin layer of chrome 0.0001 inch or 0.00025cm.

The purpose of electroplating is to protect, beautify and also used to build up worn parts to larger dimension. It is also to coat the face of cutting tools to reduce friction and wear and to resist wear on the surface or gauges.

Generally, plating is used to save resources as new functions are added to base materials. This can enhance higher conductivity, improve resistance to corrosion economic and artistic losses. In printed and integrated circuit which is a semi conductor, the chip is surface

treated and gold plated on its lead frame.

The terminal of lead frame is always tin plated or solder plated.

Plating is basic industry that supports various other industries. It is not gainsaying or an exaggeration to say that modern industries cannot exist or thrive without plating techniques. The main uses and applications of plating with examples are shown in tables (A) and (B).

Table 2.4A Main Uses Of Plating

(Fowke D .G 1986)

PURPOSE	AIM	EXAMPLE OF PLATING	EXAMPLES OF PLATED PRODUCT
DECORATION	Beautify the surface and prevent rust at the same time	Gold, silver, copper, nickel, chromium, zinc (chromate treated)	Bicycles, automobiles, motorbikes, sewing machines, clocks and watches, electric irons, toasters, personal accessories
Corrosion	Prevent rust and discoloration. To serve as undercoat plating for paints	Zinc, cadmium, lead (chromate treated) thodium	Electrical appliances, bolts, nuts, washers
Surface Hardening	Harden the surface to prevent abrasion and enhance duration	Chromium nickel	Chafing parts of engines, various types of rollers
Easing friction	Improve fitting between surface and prevent scratches and disfigurement	Tin	Pistons, piston rings, engine cylinders
Bonding	Ensure close adhesion to prevent exfoliation	Brass	Parts of engines
Preventing cementation and nitriding	Prevent hardening	Copper, tin	Various machine tools
Building up	Match measurements	Chromium nickel, copper	Same as above
Reproducing the surface	Reproduce the same surface (known as electroforming)	Copper, nickel	Matrices for gramophone records
Coloring	After plating, coloration in preferred colors is carried out (sometimes referred to as bronzing)	Copper, brass, zinc, silver, gold, (iron), (stainless steel), (aluminum)	Personal accessories, furniture.

Table 2.4B: Classification Of Main Electronic And Application Examples:

CLASSIFICATION	EXAMPLES	EXAMPLES OF APPLICATION
Super-Conductive Materials	Lead, niobium tantalum	General, motor, magnetic levitation train
Photoelectron material	Quartz, glass, silicon, germanium, indium-gallium arsenic	Fiber optic communications, light measurement video disc computer terminal
Conducive material	Copper, aluminum silver gold sodium	Wire print circuit terminal
Semiconductor material	Silicon, germanium selenium, gallium arsenic, cadmium sulfide	Transistor, diode, electronic copying machine, IC
Insulating material	Nitrogen, freon, mica, ceramics, glass, bakelite, silicon oil	Insulator, condenser, transformer, breaker
Resistance material	Nichrome, tantalum, manganin, constantan, graphite	Resistor, potentiometer, electric furnace, electric heater, transducer
Magnetic core material	Pure iron, perm alloy, silicon steel	Transformer, motor, generator, relay.
Magnet material	KS steel, MK steel, fernate, alnico	Magnetic head, speaker, cartridge
Temperature measuring material	Platinum, nickel, platinum rhodium	Thermocouple, temperature regulator, temperature measurement
Joining material	Solder, silver solder, aluminum solder	Wining of parts, assembling of parts, joining of wires
Fuse material	Tin lead, silver, brass	Power fuse, temperature fuse
Contact material	Gold silver, platinum, tungsten	Switch, jack, relay, switchboard
Electron tube material	Tungsten, molybdenum, nickel	Vacuum tube, discharge tube, X-ray tube.

2.11 General Description Of Electrodepositing Process

The main points of electrophoresis painting are:

- It is a process by dipping: It needs a container or bath of paint the volume of which being large enough to completely immerse the part to be coated: from few liters in a laboratory for painting small

panels, the volume may reach several tens of m³ for cap-parts (roof, bonnet) till several hundred m³ for cars.(PAN,1997)

Dipping into the paint tank is proceeded automatically either vertical and discontinuous dip, or continuously by the means of a conveyor which carries the car-bodies into the tank inlet, each one after the other and get it out after needed time at the outlet of the tank. That way the paint is progressively consumed during the dipping of parts; the consumption has to be compensated by addition of equivalent quantity and quality of fresh product to maintain constant paint quality all time along its exploitation.

- It is a process involving electricity: A direct current generator is necessary, the negative terminal of which is connected to the part to be painted (cathode) and positive one to anodes+.

2.11.1 Electrochemical Reactions

Electro deposition process may occur if:

- The paint to be coated is electrically conductive.
- Its viscosity is low enough to let the part to be printed completely submerged.
- The part itself is metallic.
- Another metallic contrary electrode is also submerged in the paint bath.
- The part to be painted and the contrary-electrode are connected to a direct current generator

The hereunder plan represents an electrodeposition installation:

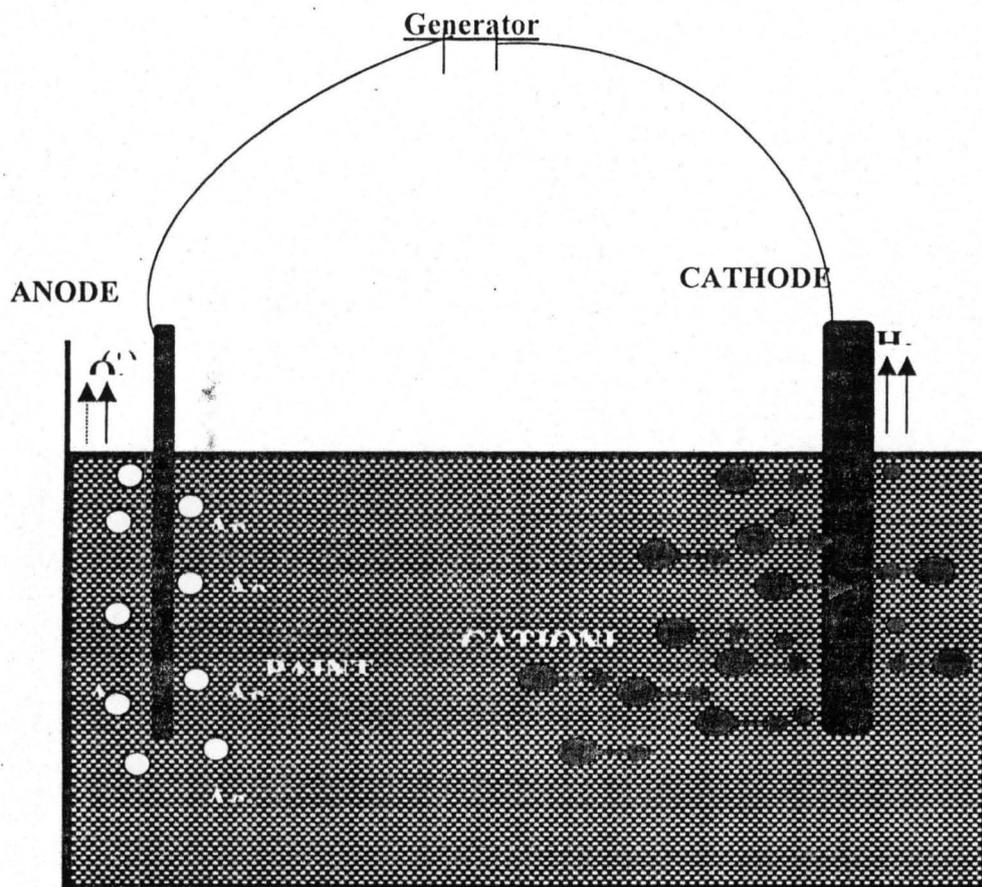


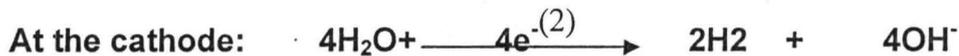
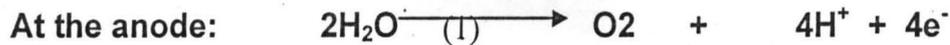
Fig. 2.3: Electro deposition of Paints

Paint is in fact polyelectrolyte in water and behaves, as it is when electrical current passes through. When a voltage input is applied between anode and cathode previously immersed in electrolytes aqueous solution (Paint), electrical field is created from electrons motion.(PAN, 1997)

The electrolyte milieu is here the ionized polymer previously described: Ac^- ; water electrolysis takes place together with ions migration (it is the actual electrophoresis) towards the electrodes: anions Ac^- to the anode, cations H^+ to the cathode.

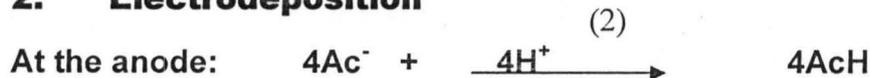
Reactions at the electrodes occur as soon as a potential difference is applied on terminals of current generator; there are 3 types.

1. Water Electrolysis

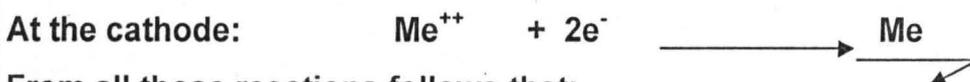
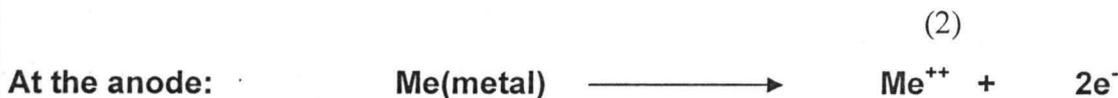


Obviously Oxygen gassing is produced at the anode, Hydrogen gassing, twice the volume of previous one, at the cathode. Reaction (1) is an oxidation with acidification at the interface anode-paint: reaction (2) is a reduction with alkalisation at the interface paint-cathode. Both are essential to realize the second type of reactions:

2. Electrodeposition



The main side reaction is promoted by the acid created in the anode neigourgh; if this last is made of corrodable material; it is oxidized and constituent metal dissolves. Dissolution promotes metallic cations going to the cathode:



From all those reactions follows that:

- At the same time the paint lays over the steel is produced hydrogen gas form. If this gassing becomes excessive, deposition is obstructed; then too powerful electrical field privileges water electrolysis to the detriment of paint electrodeposition, rupture is occurring.
- If the resin is highly ionized, rupture becomes worse because NH^+ groups on a given molecule need identical number of OH^- groups.

To create OH-, same number of voltage electrons is consumed and proportional quantity of hydrogen escapes.

Moreover it is not necessary to neutralize all amine groups 'n' in NH+' to make the cationic polymer water soluble; minimum neutralization reaction is related to the polymer molecular weight. Excess of electricity consumption (electron volt 'e' = 96.500 Coulombs) + risk of rupture even with low powerful electrical field + enough solubility in water are the 3 reasons why cationic binder is always under-neutralized: neutralization reaction (number of equivalents of Ac-/number of equivalent of N) is generally managed between 40 and 60%. Control of this ratio is mentioned.(PAN,1997)

There is a strong risk of anode corrosion, the reason why it is necessary to use non-corrodable material for anode, like graphite stainless steel.

Hence depositing of paint is the result of 4 connected events:

- Water Electrolysis
- Cations migration; is electrophoresis.
- Polymer coagulation on the cathode-liquid interface; is electrodeposition.
- Water shrinkage from that interface by osmotic pressure to let an homogeneous coat of coagulated polymer be deposited; is electrolysis.

2.12 Description Of The Product

Chrome is a blue white metal that takes on a high polish. It is hard, malleable, ductile, somewhat ferromagnetic and a fair conductor of electricity and heat.

The atomic mass of natural chrome is 24 and mass no of 51.996g, a member of the block and a group II of the periodic table of elements. It has melting point Of 187⁰C and boiling point of 2672⁰C. Chrome is of moderate strength and Hardness, when viewed as very

small particles chrome appears black as do Metals in general.

Chrome has the following physical properties tensile strength of 600N/mm^2 , Proof stress of 132 N/mm^2 , modulus of electricity of 220kn/mm^2 , hardness Between 80 – 150 and specific gravity of 8.80 chrome plating is applied over underplated layers of copper and nickel for decorative purposes and directly on the base metal for engineering purpose.

Chrome may be either shiny or dull and often tends to highlight imperfections In the base metal only very thin amount of chrome is coated on the metal. The usual thickness for bright chromium deposits is 0.25 to 0.8 microns (0.00001 to 0.00003 In^2). hardchrome is applied for wear resistance, or to restore an old warn out part to its original dimensions. It is general applied directly onto the base metal.

Chrome plated products are wrist watches chains, screw drivers, spanners, steel (car bombers), key Holders, keys, jewelry, padlocks, wheels, rim, bicycle part etc.(JOE L. et al 1981).

Table 2.5 Trouble Shooting Guide for Chrome Electroplating

PROBLEM	POSSIBLE CAUSES	SUGGESTED SOLUTION
Poor chromium coverage	i. Decorative Chromium Rectifier failure.	Check rectifier meters for stable operation. AC ripple should not exceed 5%. Rectifier should operate at 38% traced amperage.
	ii. Poor electrical connection Inactive anodes Incorrect bath temperature. Incorrect bath concentration.	Check electrical connections for lightness and contact. Check coating on anodes. A yellowish lead chromate coating indicates that the anode is inactive. A black-brown lead peroxide coating is on active anodes. Make sure proper current is used. When plating inside diameters. Use largest anode possible. Check bath temperature. Check bath agitation. Mix bath thoroughly and take bath sample. Analyze chromium and sulfate content. Adjust accordingly.
	iii. Improper racking Poor cleaning or rinsing. Copper or nickel roughness Passive nickel Chloride contamination Improper current density.	Check rack-to-fixtue connection. Incorporate shields or rubbers in racking to direct current to specific areas. Check temperature of raising water and maintain between 70 and 750F (21 and 240C). Check copper or nickel plate for roughness. Dip nickel-plated part in 50% by volume solution of hydrochloric acid. If part brightness increases, correct nickel-plating bath. Dammy with high anode current density and agitation or treat with silver oxide. Add sulfate acid to lower ratio and

	<p>Incorrect chronic acid to sulfate catalyst ratio.</p> <p>Impurities.</p>	<p>barium carbonate to increase ratio.</p> <p>Limit metallic impurities to less than 0.5 oz/gal (3.75 g/l). increase chronic acid concentration.</p> <p>Add sulfuric acid to lower ratio and barium carbonate to increase ratio.</p> <p>Limit metallic impurities to less than 0.5 oz/gal (3.75g/l). increase chronic acid concentration.</p>
Burned chromium deposits	<p>iv. Passive nickel deposit</p> <p>Incorrect bath temperature</p> <p>Improper bath concentration.</p> <p>Inactive anodes.</p> <p>Incorrect anode length.</p> <p>Current interruption.</p> <p>Improper Racking.</p> <p>High current density.</p> <p>Sulfate de-concentration or other catalyst too high.</p> <p>High or low concentration</p>	<p>Current nickel brightener concentration increase density in nickel bath.</p> <p>Correct bath temperature.</p> <p>Analyze chromium and sulfate content. Correct if necessary.</p> <p>Check anode color. Inactive anodes are coated with a yellowish lead chromate.</p> <p>Anode should be 2-4⁰ (50-100mm) shorter than cathode. Mask anodes that are too long with suitable insulator or cut to proper length.</p> <p>Check racking of parts and electrical contacts.</p> <p>Check rack to fixture connection. Incorporate shields or rubbers in racking to direct current to specific areas.</p> <p>Place work 3⁰ (75mm) below surface of plating solution.</p> <p>Add barium carbonate to bath to lower sulfate concentration. Ratio should be 2.1.</p> <p>Check ratio and adjust accordingly.</p>

	ration.	
Dull chromium deposit.	v. Dull or passive nickel.	Correct nickel brightener concentration. Increase current density in nickel bath.
	Excessive ripple in current.	Check rectifier operation. Ripple should not exceed 50%.
	Current interruption.	Check racking of parts and electrical contracts.
	Incorrect bath temperature.	Correct bath temperature.
	Low current density.	Adjust current density to bath temperature and concentration. Check rinsing procedures.
	Impurities.	Limit metallic impurities to less than 0.5 oz/gal (3.75 g/l). increase chronic acid concentration.
	Excess fluoride catalyst or fluoride contamination.	Add boric acid to reduce catalyst or use high anode current density dummy.
	High or low concentration ratio.	Check ratio and adjust accordingly.
White Motching	vi. Film of immersion nickel on work piece.	Rinse work piece thoroughly.
	Passive Nickel.	Check nickel plating bath or proper concentration.
	Excessive ripple in current.	Check rectifier operation. Ripple should not exceed 5%.
	Bipolarity caused by rapid work movement in automatic machines.	Reduce speed of work movement through machine.
	Bipolarity caused by dead entry into chromium.	Use live lead at reduce current.
	Bipolarity caused by dead entry into chromium.	Use live exit-lead from nickel.

	Bipolarity caused by dead exit from nickel tanks. Impurities especially chloride.	Check rinsing procedures.
Deposit roughness.	vii. Poor basis metal finish. Copper or nickel roughness.	Reject parts with rough finish. Check copper or nickel-plating for roughness.
Poor Chromium adhesion.	viii. Poor nickel adhesion.	Refers to solutions for bipolarity conditions under "white blotching".
No Chromium Plate.	ix. Rectifier failure. Poor electrical connections. Missing anodes. Passive nickel. Low current density. Incorrect catalyst concentration.	Check operation of rectifier. Rectifier should operate at no less than 75% rated amperage. Check connections for good contact. Check for replacement of anodes after they have been removed. Check nickel bath. Adjust current density to bath temperature and concentration. Analyze chromium-plating solution and adjust accordingly.
Poor Chromium brightness.	x. Hard Chromium.	
	xi.	
	xii. Poor basis metal finish. Excessive reverse etching. Improper bath temperature. Improper bath concentration. Improper current density. High concentration ratio.	Correct bath temperature. Check bath agitation. Take Bauman reading of bath and correct. Adjust current density to bath temperature and concentration. Check ratio and adjust accordingly.
Chromium roughness.	xiii. High trivalent chromium.	Check color of chromium solution; a black color indicates trivalent chromium contamination. Electrolyze chromium-plating solution.

	<p>Magnetic particles.</p> <p>Excessive reverse etching.</p> <p>High current density.</p> <p>Improper concentration ratio.</p>	<p>Attach a magnet to a piece of wood and then drag through solution to remove particles.</p> <p>Adjust current density to bath temperature and concentration.</p> <p>Check ratio and adjust accordingly.</p>
Bauer Chromium deposit.	<p>xiv. Improper concentration ratio.</p> <p>xv.</p>	<p>Check ratio and adjust accordingly.</p> <p>Check rack design and contacts.</p>
	<p>xvi. Poor fixturing.</p> <p>xvii.</p> <p>xviii.</p> <p>xix.</p> <p>xx.</p>	<p>Use shield to direct current to specific areas.</p> <p>Anode should be shorter than cathode. Mask anodes that are too long with suitable insulator.</p>
	<p>xxi. Incorrect Anode length.</p> <p>xxii.</p> <p>xxiii.</p> <p>xxiv.</p> <p>xxv.</p> <p>xxvi. Improper anode-cathode relationship.</p> <p>xxvii.</p>	<p>For outside diameter plating. Anode cathode distance should be approximately 4" (100mm). For inside diameter plating. The distance should be from ½ or 1 (12-25mm).</p>
Poor Chromium adhesion.	<p>xxviii. Improper cleaning or rinsing.</p> <p>Improper reverse etching.</p> <p>Current interruption.</p>	<p>Follow recommended cleaning procedure.</p> <p>Do not reverse etch in plating bath. Current used in reverse etching should be proportionated to work areas. Work should be at bath temperature before applying current.</p> <p>Check racking of parts and electrical contacts.</p>
Burned Chromium deposits.	<p>xxiv. Etching in alkaline cleaner</p> <p>Improper fixturing.</p>	<p>Check cleaner concentration.</p> <p>Racks should be made from copper for best results. Check rack design and contact.</p>

	<p>Improper anode cathode relationship.</p> <p>Improper solution level.</p> <p>Improper anode length.</p> <p>Inadequate stop-off anodes.</p> <p>Improper bath temperature.</p> <p>Low bath concentration.</p> <p>High current density.</p> <p>Impurities.</p> <p>Excess catalyst or chloride contamination.</p>	<p>See "uneven chromium deposit".</p> <p>Work should be 3" (75mm) below surface of plating solution.</p> <p>Anode should be shorter than cathode.</p> <p>Correct bath temperature.</p> <p>Mix bath thoroughly and take sample to obtain baume reading. Adjust accordingly.</p> <p>Adjust current density to bath temperature and concentration.</p> <p>Adjust current density to bath temperature and concentration.</p>
Lack of chromium hardness.	<p>xxx. Incorrect bath temperature.</p> <p>Improper bath concentration.</p> <p>High ratio.</p> <p>Insufficient deposit thickness.</p>	<p>Check bath temperature and agitation.</p> <p>Take Baume reading of bath and correct.</p> <p>Check ratio and adjust.</p> <p>Check deposit thickness.</p>
Fitting	<p>xxxi. Poor basis metal and rework.</p> <p>Improper cleaning.</p> <p>Incorrect catalyst suppressant.</p> <p>Incorrect fume suppressant.</p>	<p>Check metal prior to plating.</p> <p>Check cleaning procedures.</p> <p>Analyze chromium plating solution and adjust accordingly.</p> <p>Eliminate use of fume suppressant that lower surface tension.</p>
No Chromium plate.	xxxii. Rectifier failure.	Check operation of rectifier.

	Poor electrical connection. Missing anodes. Inactive anodes. Incorrect catalyst concentration.	Check connections for good contact. Check if anodes are properly placed. Check anode colour reactive anodes and coated with a yellowish lead chromate. Analyze chromium plating solution and adjust accordingly.
PROBLEM	POSSIBLE CAUSES	SUGGESTED SOLUTION
Poor chromium coverage	i. Decorative Chromium Rectifier failure.	Check rectifier meters for stable operation. AC ripple should not exceed 5%. Rectifier should operate at 38% traced amperage.
	ii. Poor electrical connection Inactive anodes Incorrect bath temperature. Incorrect bath concentration.	Check electrical connections for tightness and contact. Check coating on anodes. A yellowish lead chromate coating indicates that the anode is inactive. A black-brown lead peroxide coating is on active anodes. Make sure proper current is used. When plating inside diameters. Use largest anode possible. Check bath temperature. Check bath agitation. Mix bath thoroughly and take bath sample. Analyze chromium and sulfate content. Adjust accordingly.
	iii. Improper racking Poor cleaning or rinsing. Copper or nickel roughness Passive nickel	Check rack-to-fixture connection. Incorporate shields or rubbers in racking to direct current to specific areas. Check temperature of rinsing water and maintain between 70 and 75°F (21 and 24°C). Check copper or nickel plate for roughness. Dip nickel-plated part in 50% by volume solution of hydrochloric acid.

	<p>Chloride contamination</p> <p>Improper current density.</p> <p>Incorrect chronic acid to sulfate catalyst ratio.</p> <p>Impurities.</p>	<p>If part brightness increases, correct nickel-plating bath. Dampy with high anode current density and agitation or treat with silver oxide.</p> <p>Add sulfate acid to lower ratio and barium carbonate to increase ratio.</p> <p>Limit metallic impurities to less than 0.5 oz/gal (3.75 g/l). increase chronic acid concentration.</p> <p>Add sulfuric acid to lower ratio and barium carbonate to increase ratio.</p> <p>Limit metallic impurities to less than 0.5 oz/gal (3.75g/l). increase chronic acid concentration.</p>
Burned chromium deposits	<p>iv. Passive nickel deposit</p> <p>Incorrect bath temperature</p> <p>Improper bath concentration.</p> <p>Inactive anodes.</p> <p>Incorrect anode length.</p> <p>Current interruption.</p> <p>Improper Racking.</p> <p>High current density.</p>	<p>Current nickel brightener concentration increase density in nickel bath.</p> <p>Correct bath temperature.</p> <p>Analyze chromium and sulfate content. Correct if necessary.</p> <p>Check anode color. Inactive anodes are coated with a yellowish lead chromate.</p> <p>Anode should be 2-4⁰ (50-100mm) shorter than cathode. Mask anodes that are too long with suitable insulator or cut to proper length.</p> <p>Check racking of parts and electrical contacts.</p> <p>Check rack to fixture connection. Incorporate shields or rubbers in racking to direct current to specific areas.</p> <p>Place work 3⁰ (75mm) below surface</p>

	<p>Sulfate de-concentration or other catalyst too high.</p> <p>High or low concentration ration.</p>	<p>of plating solution.</p> <p>Add barium carbonate to bath to lower sulfate concentration. Ratio should be 2.1.</p> <p>Check ratio and adjust accordingly.</p>
Dull chromium deposit.	<p>v. Dull or passive nickel.</p> <p>Excessive ripple in current.</p> <p>Current interruption.</p> <p>Incorrect bath temperature.</p> <p>Low current density.</p> <p>Impurities.</p> <p>Excess fluoride catalyst or fluoride contamination.</p> <p>High or low concentration ratio.</p>	<p>Correct nickel brightener concentration. Increase current density in nickel bath.</p> <p>Check rectifier operation. Ripple should not exceed 50%.</p> <p>Check racking of parts and electrical contracts.</p> <p>Correct bath temperature.</p> <p>Adjust current density to bath temperature and concentration. Check rinsing procedures.</p> <p>Limit metallic impurities to less than 0.5 oz/gal (3.75 g/l). increase chronic acid concentration.</p> <p>Add boric acid to reduce catalyst or use high anode current density dummy.</p> <p>Check ratio and adjust accordingly.</p>
White Motching	<p>vi. Film of immersion nickel on work piece.</p> <p>Passive Nickel.</p> <p>Excessive ripple in current.</p> <p>Bipolarity caused by rapid work movement in automatic machines.</p>	<p>Rinse work piece thoroughly.</p> <p>Check nickel plating bath or proper concentration.</p> <p>Check rectifier operation. Ripple should not exceed 5%.</p> <p>Reduce speed of work movement through machine.</p>

	<p>Bipolarity caused by dead entry into chromium.</p> <p>Bipolarity caused by dead entry into chromium.</p> <p>Bipolarity caused by dead exit from nickel tanks.</p> <p>Impurities especially chloride.</p>	<p>Use live lead at reduce current.</p> <p>Use live exit-lead from nickel.</p> <p>Check rinsing procedures.</p>
Deposit roughness.	<p>vii. Poor basis metal finish.</p> <p>Copper or nickel roughness.</p>	<p>Reject parts with rough finish.</p> <p>Check copper or nickel-plating for roughness.</p>
Poor Chromium adhesion.	viii. Poor nickel adhesion.	Refers to solutions for bipolarity conditions under "white blotching".
No Chromium Plate.	<p>ix. Rectifier failure.</p> <p>Poor electrical connections.</p> <p>Missing anodes.</p> <p>Passive nickel.</p> <p>Low current density.</p> <p>Incorrect catalyst concentration.</p>	<p>Check operation of rectifier. Rectifier should operate at no less than 75% rated amperage.</p> <p>Check connections for good contact.</p> <p>Check for replacement of anodes after they have been removed.</p> <p>Check nickel bath.</p> <p>Adjust current density to bath temperature and concentration.</p> <p>Analyze chromium-plating solution and adjust accordingly.</p>
Poor Chromium brightness.	x. Hard Chromium.	
	xi.	
	<p>xii. Poor basis metal finish.</p> <p>Excessive reverse etching.</p> <p>Improper bath temperature.</p> <p>Improper bath concentration.</p> <p>Improper current density.</p> <p>High concentration ratio.</p>	<p>Correct bath temperature. Check bath agitation.</p> <p>Take Bauman reading of bath and correct.</p> <p>Adjust current density to bath temperature and concentration.</p> <p>Check ratio and adjust accordingly.</p>

Chromium roughness.	<p>xiii. High trivalent chromium.</p> <p>Magnetic particles.</p> <p>Excessive reverse etching.</p> <p>High current density.</p> <p>Improper concentration ratio.</p>	<p>Check color of chromium solution; a black color indicates trivalent chromium contamination. Electrolyze chromium-plating solution.</p> <p>Attach a magnet to a piece of wood and then drag through solution to remove particles.</p> <p>Adjust current density to bath temperature and concentration.</p> <p>Check ratio and adjust accordingly.</p>
Bauer Chromium deposit.	<p>xiv. Improper concentration ratio.</p> <p>xv.</p>	<p>Check ratio and adjust accordingly.</p> <p>Check rack design and contacts.</p>
	<p>xvi. Poor fixturing.</p> <p>xvii.</p> <p>xviii.</p> <p>xix.</p> <p>xx.</p>	<p>Use shield to direct current to specific areas.</p> <p>Anode should be shorter than cathode. Mask anodes that are too long with suitable insulator.</p>
	<p>xxi. Incorrect Anode length.</p> <p>xxii.</p> <p>xxiii.</p> <p>xxiv.</p> <p>xxv.</p> <p>xxvi. Improper anode-cathode relationship.</p> <p>xxvii.</p>	<p>For outside diameter plating. Anode cathode distance should be approximately 4" (100mm). For inside diameter plating. The distance should be from ½ or 1 (12-25mm).</p>
Poor Chromium adhesion.	<p>xxviii. Improper cleaning or rinsing.</p> <p>Improper reverse etching.</p> <p>Current interruption.</p>	<p>Follow recommended cleaning procedure.</p> <p>Do not reverse etch in plating bath. Current used in reverse etching should be proportionated to work areas. Work should be at bath temperature before applying current.</p> <p>Check racking of parts and electrical contacts.</p>

Burned Chromium deposits.	<p>xxiv. Etching in alkaline cleaner</p> <p>Improper fixturing.</p> <p>Improper anode cathode relationship.</p> <p>Improper solution level.</p> <p>Improper anode length.</p> <p>Inadequate stop-off anodes.</p> <p>Improper bath temperature.</p> <p>Low bath concentration.</p> <p>High current density.</p> <p>Impurities.</p> <p>Excess catalyst or chloride contamination.</p>	<p>Check cleaner concentration.</p> <p>Racks should be made from copper for best results. Check rack design and contact.</p> <p>See "uneven chromium deposit".</p> <p>Work should be 3" (75mm) below surface of plating solution.</p> <p>Anode should be shorter than cathode.</p> <p>Correct bath temperature.</p> <p>Mix bath thoroughly and take sample to obtain baume reading. Adjust accordingly.</p> <p>Adjust current density to bath temperature and concentration.</p> <p>Adjust current density to bath temperature and concentration.</p>
Lack of chromium hardness.	<p>xxx. Incorrect bath temperature.</p> <p>Improper bath concentration.</p> <p>High ratio.</p> <p>Insufficient deposit thickness.</p>	<p>Check bath temperature and agitation.</p> <p>Take Baume reading of bath and correct.</p> <p>Check ratio and adjust.</p> <p>Check deposit thickness.</p>
Fitting	<p>xxxii. Poor basis metal and rework.</p> <p>Improper cleaning.</p> <p>Incorrect catalyst suppressant.</p>	<p>Check metal prior to plating.</p> <p>Check cleaning procedures.</p> <p>Analyze chromium plating solution and adjust accordingly.</p>

	Incorrect fume suppressant.	Eliminate use of fume suppressant that lower surface tension.
No Chromium plate.	xxxii. Rectifier failure. Poor electrical connection. Missing anodes. Inactive anodes. Incorrect catalyst concentration.	Check operation of rectifier. Check connections for good contact. Check if anodes are properly placed. Check anode colour reactive anodes and coated with a yellowish lead chromate. Analyze chromium plating solution and adjust accordingly.

2.13 Small Scale Industries

2.13.1 Definitions

It will be vague to offer a precise definition of the terms “small scale” business because the expression covers a wide range of activities that are related to social and economic development of a given nation, even among researchers and practitioners, There has been no consensus as regards what constitutes a small scale business. The multiplicity of definition relates to the culture and peculiar circumstances of the person(s) attempting the definition. (Adama et al, 2003).

Hence, there is no single yardstick for classifying a business unit as small scale. The classification essentially varied according to time and setting. Since attempts in the past to produce a universally accepted definition of what business unit can rightly be referred to a small scale, industry had caused some confusion and protracted debate among discussant. So the solution to this multiplicity of definition or criteria employed would seem to be in the revision of whatever definition is used from time to time based on specific objective as to what these micro economic unit are meant to achieve (Blunt et at, 1990).

The central Bank of Nigeria in its 1998 operational guidelines defined small-scale business with reference to two Financial Institutions; the Merchant Bank, a small scale business is the “one with a capital investment of more than N200,000 excluding the cost of land and annual turnover of N500,000”. But for Commercial Banks, a small scale business is “one with maximum annual turnover not exceeding

N500,000(Koh,1988).

The center for management science development at the University of Netherlands, defines a small-scale business as a business in which manager personally perform at the function of management. These centers classifies small scale business as an establishment employing not more than 99 persons, while the employment of 1-9 persons is regarded as cottage industry (Blunt et al, 190).

The United Nations industrial development organization (UNIDO) classified small scale enterprises as those with less than 150 employees on its pay roll. Such enterprises can be classified into two:

- i. The traditional craftsmen and artisan and
- ii. The small manufacturing enterprises which produce a variety of consumer and simple producer goods.

The Nigeria Bank for commerce and industry (NBCI) in 1987, defined small enterprises as those with total cost of not more than 750,000.

This definition which is expected to be valid until 1990 before it is reviewed has been accepted by the government (Central Bank of Nigeria, 1995).

However, the definitions of small scale enterprises as those whose total cost, excluding cost of land but including working capital is above N1,000,000 but does not exceed N10,000,000 has irraddvertently left out credit-worthy cottage industries. This has created problems for the category of enterprises and banks that are other-wise disposed to granting credit to them. For this

purpose therefore, small scale enterprises shall include cottage industries and defined as enterprises whose total cost, excluding cost of land but including working capital does not exceed N10,000,000 (Oladeepo, 1998).

Finally, according to National Council of Industrial/Federal Ministry of Industry, 2002 gives the following definition as regards various sectors of industries.

- i. **Cottage Industry:** This is any firm with total cost including working capital but excluding cost of land below N1.5 million and labour size below 11 workers.
- ii. **Small Scale Industry:** This is any firm with total cost including working capital but excluding cost of land above N1.5million but not exceeding N50 million and labour size between 11 and 35 workers.
- iii. **Medium Scale Industry:** This is any firm with total cost including working capital but excluding cost of land above N50,000,000 but not exceeding N200,000,000 and labour size of between 36 and 100 workers.
- iv. **Large Scale Industry:** This refers to any firm with total cost including working capital but excluding cost of land above N200 million and labour size above 100 workers (Onyeneke, 2002).

Furthermore, the revision of these definitions are done from time to time to enable many more unit of small scale industry to modernize their operation and following the trend of change within the sector.

2.13.2 Cost indices

A cost index is merely an index value for a given point in time, showing the cost at that time relative to a certain base time. If the cost at some time in the past is known, the equivalent cost at the present time can be determined by multiply the original cost by the ratio of the present index value to the index value applicable when the original cost was obtained (Peters and Timmerhaus, 1991).

$$\text{Present Cost} = (\text{Original Cost}) \times (\text{Index value at preset time}) / (\text{Index value at time Original cost was obtained}).$$

Statements of cost are based on the value of the dollar at a given year, and a cost index is needed to compare cost, which are presented on different bases. The uses of cost index to express the changing purchasing power of the dollar dates back to 1913 with the ENR cost index which is a weighted average of the cost of steel, lumber, cement and common labour (Rudd and Watson, 1968).

Cost Indices can be used to give a general estimate, although, no index can take into account all factors, such as special technological advancement, or local conditions.

The common index permits fairly accurate estimates if the time period involved is less than 10 years.

Many different types of cost Indices are published regularly. The most common of these Indices which can be used for estimating equipment cost are the Marshall and Smith Equipment Cost Index, and the Chemical Engineering Plant Cost Index. For the purpose of this work, the latter shall be used.

2.13.3 Chemical engineering plant cost index

Construction cost for chemical engineering plants form the basis of the Chemical Engineering Plant Cost Index. The four major components of this index are weighted by percentage in the following manner: equipment, machinery and supports, 61; erection and installation labour, 22; building materials and labour, 7' and engineering and supervision, 10. The major component, equipment, is further subdivided and weighted as follows:

Fabricated equipment, 37; process machinery, 14; pipes , valves and fillings, 20; process instruments and control, 7; pumps and compressors, 7; electrical equipment and materials, 5; structural supports, paints and instrumentation, 10. All index components are based on 1957-1959 = (Peter and Timmerhaus).

2.14 Process Simulation

This is an act of representing some aspects of the real world by symbols that may be easily manipulated to facilitate their study. The important steps of process simulation are therefore, description of the part of the "real world" that needs to be simulated, representation of this part of the 'real world' in terms of a model (mathematical or symbolic), and finally, solution of the mathematical model to obtain numbers or symbols. With respect to chemical engineering, the real world is a chemical process described by a process flow sheet. Typically, process simulation is needed to solve problems related to process design, process analysis, process control and many more. Depending on the type of the problem, different types of process

simulation problems are formulated. For example, verification of design or for inventory control (mass and energy balance) requires the solution of steady state simulation problems while verification of the control system or analysis of the effect of disturbances requires the solution of dynamic simulation problems.

Obviously, each simulation problem is associated to a corresponding mathematical model. In steady state simulation, the mathematical model is usually represented by a mixed set of algebraic equations and ordinary differential equation. For complex or rigorous simulation problems, the mathematical model may even be represented by a mixed set of algebraic equations. Each mathematical model, therefore, is associated to a corresponding method of solution.

Since there can be many way to solve a set of equations representing a mathematical model, it is necessary to have a simulation strategy which ensures that the simulation problem is solves efficiently and that the simulation results are reliable. Reliability also depends on the model accuracy. Use of inappropriate models or modes parameters may result in erroneous simulation results. (floudas, 1995).

As described above, process simulation requires a number of important steps that are linked to each other. Each step, however, can be large enough to justify the use of a computer to solve the problem. A computer problem that combines the important steps into a single program package is called a process simulator. The process simulator is designed to help the user in performing the important steps without the need to develop new computer programs for solving new

problems.

In the solution of problems related to process synthesis or design, control or operation, analysis etc., the user needs to provide sufficient information to properly specify the simulation problem. Typically, a simple steady state simulation of a process flow sheet requires a description of the process flow sheet (including an identification of the compounds present in the system and unit operations present in the flow sheet), choice of method of solution, specification of equipment parameters and input streams and many more. Solution of a process design problem may require many solution of the process simulation problem (for each trial, a different simulation problem may need to be solved). User does not have all the information necessary to specify the simulation problems, in order to assist the user; the concept of an integrated simulation system has been introduced.

The objective of integrated system is to help the user to generate the necessary information so that a simulation problem can be properly defined and solved. Usually, the time spent on collecting the information necessary to defined a simulation problem is much greater than the time spent by the process simulator in finding the solution. By reducing the time spent by the user in defining the simulation problem, the integrated system helps to improve the work efficiency or user. In the integrated system helps to improve the efficiency of the user. In the integrated system, the user has available, computational tools that can be used to generate the missing information.

2.14.1

Structure of a process simulator

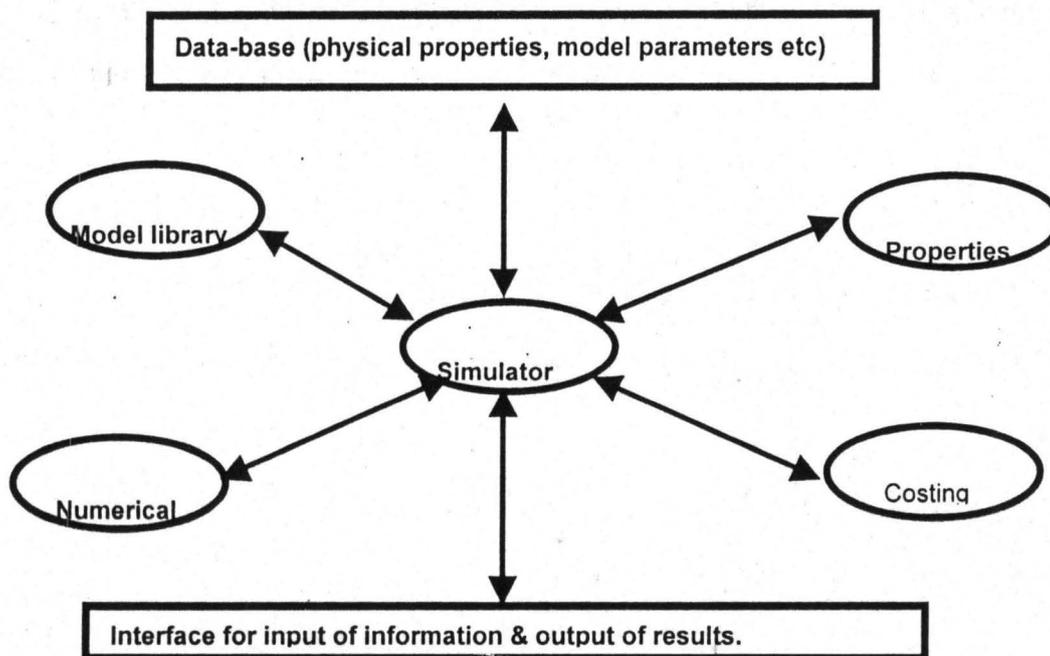


Figure 2.4 Main features of a process simulation

2.14.2

Commercial process simulators

The three common commercial process simulator are;

ASPEN Plus - Aspen Technology, USA.

Pro-II/Provision – Simulation Sciences, USA (Seibe, UK)

HYSIS-Hypro tech, Canada (AEA Technologies, UK).

ASPEN and Pro-II are steady state process simulators with process optimization feature. Both of them employ a sequential modular approach. HYSIS provides both steady and unsteady simulation features. It employs a mixed simulation strategy. ASPEN has the largest number of users followed by pro-II and then HYSIS. All the three simulators have windows-based PC-versions. All have extensive data for compound and physical properties, it has a large collection of unit operation models and thermodynamic models.

2.14.3 Computer application to chemical engineering

Computer can be applicable to the following chemical engineering subject: Heat transfer, process dynamic and control, introduction to Chemical Engineering, , chemical reaction engineering, fluid mechanics, mathematical methods, thermodynamics, separation process, process design and electrochemical processes.

2.14.4 Hysys

This simulator provides you with an intuitive and interactive approach toward process modeling, simulation and optimization.

This software lets you to create detailed high-fidelity plant simulations for analyzing and optimizing your plant's operation. Through the completely interactive HYSYS interface, you have access to a fine level of equipment geometry, performance detail as well as the ability to completely customize your simulation using its extensibility capability.

This program is used by industry to design and simulate process plants such as oil and gas refineries, chemical and pharmaceuticals production and electrochemical facilities. (HYSIS Simulation Manual, 2004).

2.14.5 MathCAD

MathCAD provides a rich problem-solving environment that gives you a wide choice of tools and supports a variety of analysis and visualization techniques. MathCAD is used by hundreds of thousands of engineers, scientists, and students across a broad range of technical disciplines to perform calculations, to interface with other engineering,

drawing and business applications and to publish their "live" documents on the web.

In the most general sense, you can think of MathCAD as a combination of;

(i) A powerful technical computing environment centered on real math notation.

And

(ii) A flexible, full-featured technical word processor.

This means that with MathCAD the tasks of performing computations and documenting them are integrated into one seamless process, resulting in substantial increases in productivity.

2.14.5.1 features of the mathCAD window

Before you move on to learning more about MathCAD's features and especially about building math and text in your own MathCAD worksheets, take a moment to look at the MathCAD application window in front of you.

(a) The Main Menu

File Edit View Insert Format
Math Symbolic Window Help

This gateway to math, graphics, and functions provides the commands that handle the details of editing and managing your worksheets. Click on the menu to see the array of computational and formatting functionally available to you.

2.14.5.2 the math toolbar

Choose Toolbars Math from the View menu to see a tear-away bar whose buttons bring up toolbars of math operators. You can click the bar under the main menu. If you let your mouse hover over each of the buttons on the bar in turn, you will see a tool tip telling you what menu each button brings up:

Calculator toolbar

Graph toolbar

Boolean toolbar

Programming toolbar

Matrix toolbar

Greek toolbar

Calculus toolbar

Click on one of these buttons in the bar to bring up the associated operator toolbar. You can then use the operator toolbar to insert math symbols right into your MathCAD worksheet.

2.15 Costing

It is an economic fact that the purchasing power of the monetary unit, no matter what the currency is, erodes and depreciates with time, and there is little if any reason, to suggest that this trend will change in the future. This depreciation in the value of money is termed inflation. Inflation is the rise in price level or fall in the purchasing power, or the rate of increase in some national price level. The official rate of inflation is based on a specified mixture of goods required in cash outlay required to purchase this basket of goods is the official rate of inflation.

Due to inflation and other factors, the economic environment in which a chemical plant (Chrome Electroplating Plant) operates is a dynamic and not static one, and it undergoes continuous change, as will

the factors that determine its profitability, e.g. labour, raw materials and utilities. The overall cost of establishing such a plant will equally change upwards, as the years slowly go by. One of the most important factors contributing to the changes above is inflation, which will also seriously affect the cost of equipment.

2.15.1 Equipment cost estimation

There are methods of estimation of equipment costs of the Chrome Electroplating Plant amongst are the method of Cost Index, the Six-Tenth-Factor Rule Method, and method of Cost Index and scaling.

2.15.2 The method of cost index

If the cost of each piece of equipment at sometime in the past is known, then the equivalent cost at the present time can be determined by multiplying the original or historical cost by the ratio of the present index value to the historical index value applicable when the historical cost was obtained (See Table 3.1 for cost data).

Let historical cost (Original) cost of each equipment be HC, say; the historical cost index be HI; the current cost index be CI, and the current cost of the current cost of the equipment be CC.

Therefore, the current cost of any equipment was given by the following equation:

$$CC = HC (CI/HI) \dots \dots \dots 2.7$$

2.15.3 Six-tenths factor rule method

It is often necessary to estimate the cost of a piece of equipment when no cost data are available for the particular size of

operational capacity in question. Good results can be obtained by using the logarithmic relation known as the six-tenth-factor the cost data are available. According to this rule, if the cost of a given equipment at one capacity is known, the cost of a desired similar unit with X times the capacity of the first equipment is approximately $(X)^{0.6}$ times the cost of the original equipment.

Let HCB – historical cost of equipment B (desired).

HCA – historical cost of equipment A.

QA – known capacity of equipment A.

QB – capacity of equipment B.

Therefore, the historical cost of equipment B that is not available in literature is given by:

$$HCB = HCA (QB/QA)^{0.6} \dots\dots\dots 2.8$$

The current cost of equipment A is given by the following cost index equation:

$$CCA = HCA (CIA/HIA) \dots\dots\dots 2.9$$

From equation 3.2, it follows that the current cost of the desired equipment B is given by:

$$CCB = HCA (CIA/HIA)(QB/QA)^{0.6} \dots\dots\dots 2.10$$

Combining equation 3.1 and equation 3.4 obtain.

$$CCB = HCA (CIA/HIA)(QB/QA)^{0.6} \dots\dots\dots 2.11$$

2.15.4 Method of cost index and scaling.

This method is similar to the Six-Tenth-Factor Rule method, but different in the fact that whereas the index to which the equipment capacities are raised was constant (0.6) for the former method, it varied

for this method depending on the equipment. Secondly this method is used when cost data for a particular size of operational capacity of the same equipment is available, but lack for the desired size.

The current cost of a piece of equipment of desired size B was given by the following equation:

$$CCB = HCA (CIA/HIA) (QB/QA)^{EI} \dots\dots\dots 2.12$$

Cost data for this equation is given in Table 3.1.

2.15.5 Estimation of the delivered purchased equipment cost

The cost of delivered purchased equipment is the basis for estimating Capital Investment. This was computed here as a summation of the individual equipment costs determined by any of the three equipment cost estimation methods discussed in 3.2.1.

Therefore,

$$PECT = \sum_{i=1}^n CC_1 \dots\dots\dots 2.13$$

Cost data for this equation is given in Table 3.1.

2.15.6 Estimation of the fixed capital investment

This consists of manufacturing fixed-capital investment, which represents the capital necessary for the installed process equipment with all auxiliaries that are needed for complete process operation are typical examples of costs included in the manufacturing fixed-capital investment.

Fixed capital required for construction of overheads and for all plant components that are not directly related to the process operation is designated as the non-manufacturing fixed-capital investment. These

plant components included the land, processing buildings, administrative and other offices, warehouses, laboratories, transportation, shipping and receiving facilities, utility and waste disposal facilities, shops and other permanent parts of the plant. The construction overhead costs consist of field-office and supervision expenses, home-office expenses, engineering expenses, miscellaneous construction cost, contractor's fees, and contingencies (Peters and Timmerhaus 1991).

The fixed-capital investment shall be determined by means of the factored estimate method, which is a method by which the investment cost in a completed system can be extrapolated from the delivered cost of the major items of processing equipment (Rudd and Watson, 1968).

It has been observed that the cost of the other essential items needed to complete the process system can be correlated with the investment cost in major items of equipments, and the capital investment can be estimated by the application of experience factors to the base investment of delivered purchased equipment cost (Rudd and Watson 1968).

Thus, the factored estimate equation below results. The experience factors f are obtained from a study of many similar processing systems. Values for the direct and indirect factors are given in Table 3.2.

$$FCAPN = [PECT + (\sum f_i PEECT)] f_1 \dots \dots \dots 2.14$$

Where $f_1 = (1 + \sum f_i)$.

2.15.7 Computation of the total and working capital investments.

Total capital investment consists of fixed capital and working capital investment.

The fixed-capital investment was taken as eighty five percent of the total capital investment, while the remaining fifteen percent became the working capital investment.

Thus, the total capital investment given by;

$$\text{TCAPN} = \text{FCAPN}/0.85 \dots\dots\dots 2.15$$

And, the working capital investment was given by:

$$\text{WCAPN} = 0.15 * \text{TCAPN} \dots\dots\dots 2.16$$

Cost estimation of equipment and economic acceptable plant design must present a process that is capable of operating under conditions which will yield profit when the cost for any type of commercial process is to be determined, sufficient accuracy has to be provided for reliable decision.

There are many factors affecting investment and production cost.

These are:

- i. Company Policies
Source of equipment
- ii. Price fluctuation
- iii. Operating and rate of production
- iv. Government Policies

(Peters et al, 1991)

Before an industrial plant can be put into operation, a large sum of money must be supplied to purchase and install the necessary machinery and equipment. (Gaeul, 1977).

Hysis cost estimation and Marshal and Smith cost estimation index were used and the summary of result for the cost can be seen in chapter four.

Cost of equipment from hysis online cost.

Simulator and cost are in Dollar.

Exchange rate ER is 150 Naira = 1 Dollar

This cost estimation is based on cost data of 2006 and is available in dollar.

Exchange rate (ER) 130 Naira = 1 Dollar

Data for cost of Acid pickling tank' Degreasing Tank, Electroplating tank and washing tanks are estimated using Marshall and Smith cost index. However this cost estimation method cannot be used for some equipment e.g. cooler, rectifier, tank mixers, heaters, and air dryer. direct online costs were used.

$P_c = M.S (101.9) 1.066$

Where D = Diameter of the tank/column in ft.

H = Height of the tank/column in ft.

M.S = Marshall and Smith index = 110

$F_c = 1.00 + F_m + F_p$

(Gealu, etal, 2006)

The material of construction selected for most equipment is carbon steel because of its corrosive resistance.

Hence, $F_m = 1.00$

$F_p = 1/00$

$F_c = 1 + F_m + F_p$

$F_c = 3$

Diameter = D

Volume = V

Height = H

CHAPTER THREE

3.0 METHODOLOGY

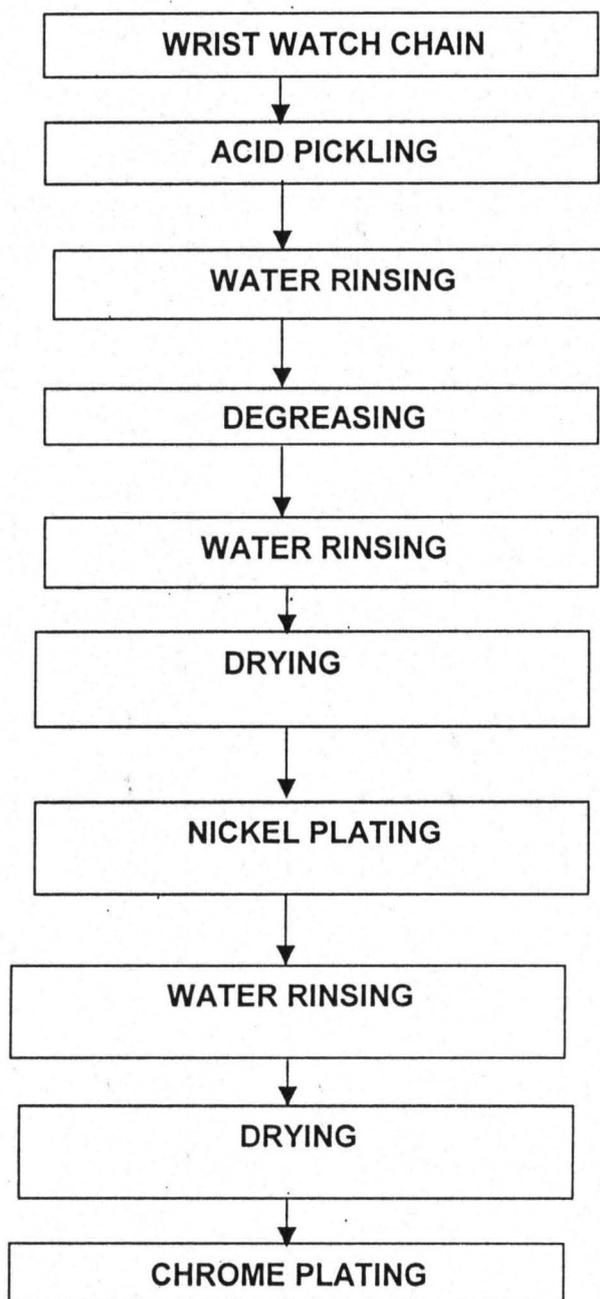
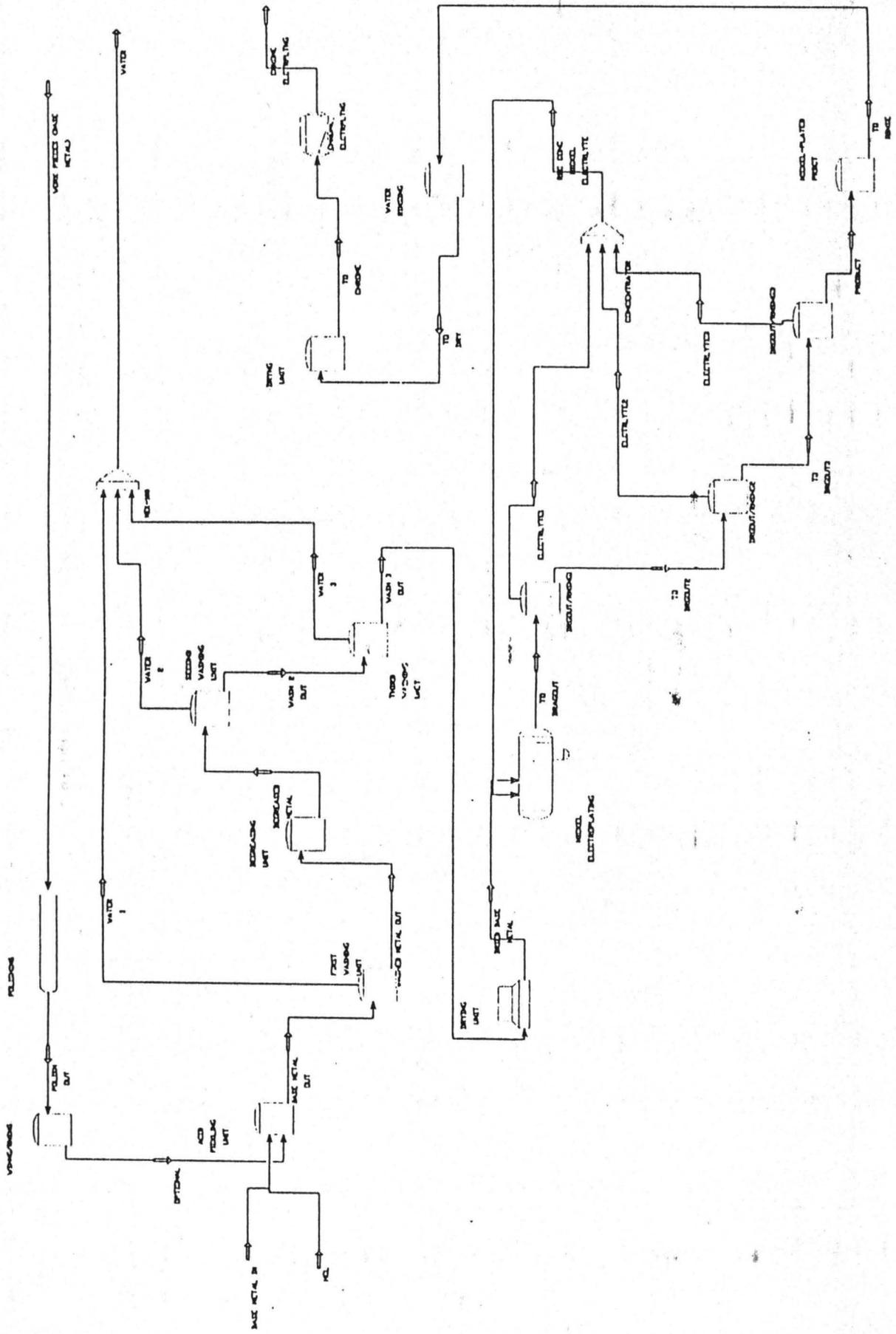


Fig. 3.0 Block Diagram Of Chrome Electroplating.

FLOW DIAGRAM OF CATHODIC



3.1 Process description

3.1.1 Acid pickling

In this process, oxides are removed from surface of the basis metal for steel, warm, dilute sulfuric acid is used in large-scale operations because it is inexpensive; but room temperature, dilute hydrochloric acid is also used for pickling of steel, attack of the metal is retarded while the oxide is being dissolved. In addition to rough pickling acid treatment to activate the surface just prior to plating are often used. When an inhibitor is added to acids, it covers the exposed iron material and thus inhibits excessive corrosion by acids.

The following reaction took place



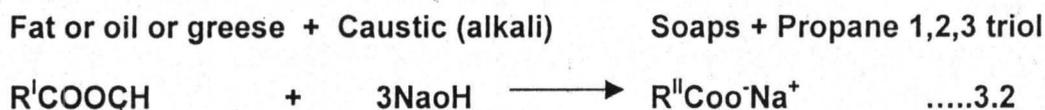
3.1.2 Water rinsing

Cold distilled water is used for rinsing of the product from Acid pickling to avoid contamination of the electrolyte, by removing the iron III chloride formed and at the same time reduces the traces of acid on the work pieces before degreasing.

3.1.3 Degreasing (alkaline soak cleaning)

This process generally involves the use of chlorinated hydrocarbons in either the heated or vapour form. The solvents commonly used are Trichloroethylene, perchloroethylene and 1,1,1-trichloroethylene. This process is basic, and therefore a very important process in the plating process. The degreasing salt is dissolved in water and the solution is heated to a temperature of between 50⁰c to 80⁰c.

After ward the object is suspended in the solution and left for about 15 minutes by which time all the dirt and grease would have been completely removed. The degreasing speed tend to become faster when the product to be plated is stained by an highly viscous oil, since the oil will soften faster under a higher temperature. Saponification process is the basic reaction here



3.1.4 Water rinsing

Similarly cold distilled water is used to rinse off the soaps, emulsified products and left over sodium hydroxide on the product.

3.1.5 Drying

The water on the base metal is dried to avoid further dilution of content of the electroplating bath .

3.1.6 Nickel electroplating

Nickel plating solutions, nickel anodes are used to provide the necessary anode surface to secure as possible distribution of the electro deposit on the work and to maintain the nickel content for the plating solution. A typical electrolyte for Nickel plating can be made by dissolving the following salts

Nickel Sulphate	225 – 357g/L,	Mean 291g/L
Nickel Chloride	30 – 75g/L,	Mean 52.5g/L
Boric acid	30 – 55g/L,	Mean 42.5g/L

2-3 droplets of H₂SO₄ at a P^H Of 3.6.

The nickel anodes are enclosed in cotton polypropylene or

terylene bags.

3.1.7 Operating condition for Nickel plating solutions.

Cathode Current density	4.0 to 6.5amp/dm ² (40 to 60 amp/ft ²)
P ^H Value	4.0 to 4.8
Voltage	3 to 4 volts
Temperature	55 to 65 ⁰ c
Anode	nickel

3.1.8 Chrome electroplating

In Chrome Electroplating, chrome anodes are used to provide the necessary anode surface to secure distribution of the electro deposit on the work and to maintain the nickel content for the plating solution. A typical electrolyte for chromium decorative plating can be made by dissolving the following salts

Chrome Sulphate	200 – 300g/L,	Mean 250g/L
Ratio of chromic acid to sulfate		100:1 to 125:1
chrome Chloride	30 – 75g/L,	Mean 52.5g/L

3.1.9 Operating condition for chromium plating solutions.

Cathode Current density	8.0 to 12amp/dm ² (75 to 110 amp/ft ²)
Voltage	4 to 5 volts
Temperature	35 to 40 ⁰ c
Anode:	lead/antimony.

3.1.10 Dragout/Water rinsing tanks

The dragout/water rinsing tank does two things;

To capture the dragout volume and also rinse the works pieces(wristwatches) from concentrated Nickel solution and subsequently from concentrated chrome solution. The tanks are connected to each other and Nickel and chromium plating tank by drop board.

3.1.11 Concentrator

Concentrator helps to recover Nickel and chrome metal from plating rinse waters (Dragout tanks). However, chemical precipitation, solvent extraction, ion exchange or reverse Osmosis will recover metal as concentrated solutions or sludges, which can be recycle back for reused. Also, where the metal from plating rinse water is required as pure metal then an electrolytic. Extraction may be used for this design ion exchange with reverse Osmosis concentrator is used.

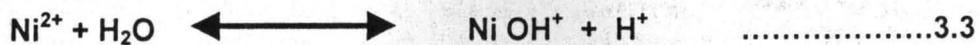
3.1.12 Air drying

The work piece (base metal) after electroplating was spread to dry under atmospheric air before packaging.

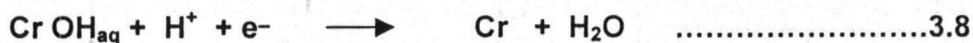
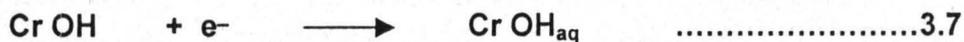
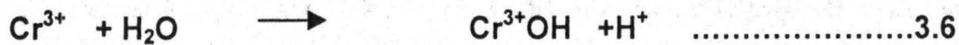
3.2 Equation of The Reactions

The equation of the reaction is obtained based on the anodic and cathodic half reactions. Electrochemical deposition of nickel and chrome from typical electrolyte is a relatively slow process compared to the speed at which their ions moved in the bath. This fact is possible to use very high current densities at direct current conditions, and even higher current densities using pulse plating.

The deposition of nickel takes place through a number of intermediate steps, as indicated below,



Similarly the deposition of chrome takes place as follows:



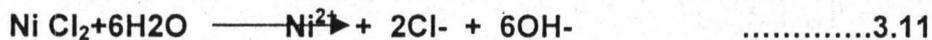
From the reaction above, it can be seen that that pH value in the bath is of great importance. Since free hydrogen ions play an important role in the deposition mechanism. The bath temperature is also an important parameter, since the speed of the reaction is all dependence on temperature (but some more than others).

Also the concentration of nickel and chrome ions at the cathode, and the current density (available electrons) has influence on the deposition process .

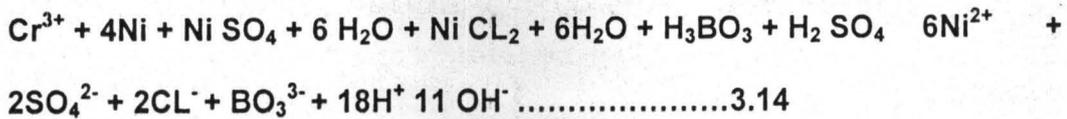
Also, noted is the equation of reaction at the positive and negative poles respectively.



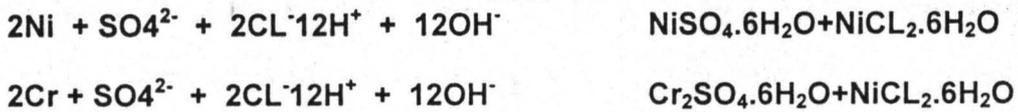
In the electrolyte, the dissociation is



The net reaction is given as (overall equation), this involve the combination of equation 3.6 – 3.10



3.2.1 Anodic reaction



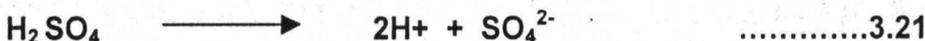
3.2.2 Cathodic reaction



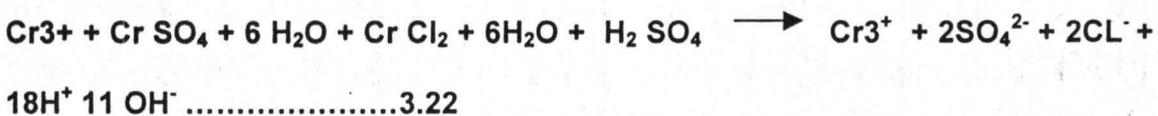
Similarly, for chrome:



In the electrolyte, the dissociation is



The net reaction is given as (overall equation), this involve the combination of equation 3.18 – 3.21



3.3 Material Balance

A computer aided design was used for this plant, MathCAD was chosen to carry out the material balance . The basis chosen was 2000kg/day of wrist watches, Rust of 0.05% and grease of 0.02% were used. The mass flow rates of the materials were calculated: rust (Fe₂O₃) was 1.0kg/day and grease was 0.4kg/day and reacting mass of rust was 6.25 x 10⁻³kg.

The volume of the metal base was 254.0L and for the acid pickling vessel was 304.8L and acid concentration was 0.148mol/m³. These were followed with the evaluation of material balances for the various pre plating and plating units starting from acid pickling vessel to dragout tanks where the input stream of previous unit is made to serve as an input stream to the next unit finally, 1998.6kg/day of wrist watches goes for electroplating and 351.62kg of nickel metal was required and 187.888kg of chrome metal was required for this task.

Overall Material balance

$$\text{Basis} = 2000 \frac{\text{kg}}{\text{day}}$$

$$1 \text{ day} = 8 \text{ hours}$$

$$\text{Rust} := 0.05\% \cdot \text{Basis}$$

$$\text{Mass_flowrate_of_Rust} = 1 \text{ kg /day}$$

$$\text{Grease} := 0.02\% \cdot \text{Basis}$$

$$\text{Iron} := \text{Basis} - (\text{Rust} + \text{Grease})$$

$$\text{Mass_flowrate_of_Rust} := \text{Rust}$$

$$\text{Mass_flowrate_of_Grease} := \text{Grease}$$

$$\text{Mass_flowrate_of_Grease} = 0.4 \text{ kg}$$

3.4.2 Material balance around the acid pickling unit

3.4.2.1 input streams

SPECIES₁ :=

Fe ₂ O ₃
HCl
Fe
FeCl ₃
Grease
Water

Mwt₁ =

160
36.5
56
162.2
287.27
18

kg·mol⁻¹

$$\text{Mole_of_Rust} := \frac{\text{Rust}}{160 \frac{\text{kg}}{\text{mol}}}$$

$$\text{Mole_of_Rust} = 6.25 \times 10^{-3} \text{ mol}$$

$$\text{Mole_of_HCl} := 6 \cdot \text{Mole_of_Rust}$$

$$\text{Mole_of_HCl} = 0.038 \text{ mol}$$

$$\text{HCl} := \text{Mole_of_HCl} \cdot 36.5 \frac{\text{kg}}{\text{mol}}$$

SPECIES₁ :=

Fe ₂ O ₃
HCl
Fe
FeCl ₃
Grease
Water

Inlet_Mass_1₁ :=

Rust
HCl
Iron
0
Grease
0

$$\text{Unit_one_input} := \sum_{I=1}^6 \text{Inlet_Mass_1}_I$$

$$\text{Unit_one_input} = 2 \times 10^3 \text{ kg}$$

3.4.2.2 evaluation of HCl concentration

$$\text{Mass_of_metal_base} := \text{Basis}$$

$$\text{Density_of_metal_base} := 7874 \frac{\text{kg}}{\text{m}^3}$$

$$\text{Volume_of_metal_base} := \frac{\text{Mass_of_metal_base}}{\text{Density_of_metal_base}}$$

$$\text{Volume_of_metal_base} = 254.001 \text{ L}$$

$$\text{Tank_Volume} := 1.2 \cdot \text{Volume_of_metal_base}$$

$$\text{Tank_Volume} = 304.801 \text{ L}$$

$$\text{Conc}_{\text{HCl}} := \frac{\text{Mole_of_HCl}}{\text{Volume_of_metal_base}}$$

$$\text{Conc}_{\text{HCl}} = 0.148 \frac{\text{mol}}{\text{m}^3}$$

3.4.2.3 output stream

$$\text{Mole_of_Water} := 3 \cdot \text{Mole_of_Rus}$$

$$\text{Mole_of_FeCl}_3 = 0.013 \text{ mol}$$

$$\text{Mole_of_Iron} := \frac{\text{Iron}}{56 \frac{\text{kg}}{\text{mol}}}$$

$$\text{Mole_of_Water} = 0.019 \text{ mol}$$

$$\text{Outlet_Mass_1}_j := \text{Outlet_Mole}_j \cdot \text{Mwt}_j$$

$$\text{Mole_of_Grease} := \frac{\text{Grease}}{287.27 \frac{\text{kg}}{\text{mol}}}$$

$$\text{Mole_of_Grease} = 1.392 \times 10^{-3} \text{ mol}$$

$$\text{Mole_of_Iron} = 35.689 \text{ mol}$$

$$\text{SPECIES}_1 :=$$

Fe ₂ O ₃
HCl
Fe
FeCl ₃
Grease
Water

$$\text{Outlet_Mole}_1$$

0.0
0.0
Mole_of_Iron
Mole_of_FeCl ₃
Mole_of_Grease
Mole_of_Water

$$\text{SPECIES}_1 :=$$

Fe ₂ O ₃
HCl
Fe
FeCl ₃
Grease
Water

$$\text{Outlet_Mass}_1 =$$

0	kg
0	
1.9986710 ³	
2.0275	
0.4	
0.3375	

$$\text{Unit_one_output} := \sum_{I=1}^6 \text{Outlet_Mass}_1$$

$$\text{Unit_one_output} = 2 \times 10^3 \text{ kg}$$

3.4.3 Unit two

Estimation of the amount of washing Water

From solubility curve; solubility of FeCl₃ at 25°C

$$\text{Solubility} := 4.3 \cdot 10^{-3} \frac{\text{mol}}{\text{m}^3}$$

$$\text{Volume_of_Water_Required} := \frac{\text{Mole_of_FeCl}_3}{\text{Solubility}}$$

$$\text{Volume_of_Water_Required} = 2.907 \times 10^3 \text{ L}$$

$$\rho_{\text{water}} = \text{Density_of_water}$$

$$\rho_{\text{water}} := 997.99 \frac{\text{kg}}{\text{m}^3}$$

$$\text{Mass_of_water} := \rho_{\text{water}} \cdot \text{Volume_of_Water_Required}$$

$$\text{Mass_of_water} = 2.901 \times 10^3 \text{ kg}$$

3.4.3.1 unit two inlet

$$\text{Unit_Two_inlet} := \text{Outlet_Mass_1}$$

SPECIES₁ :=

Fe ₂ O ₃
HCl
Fe
FeCl ₃
Grease
Water

Unit_Two_inlet =

0	kg
0	
1.9986710 ³	
2.0275	
0.4	
0.3375	

$$\text{Total_mass_inlet} := \left(\sum_{I=1}^6 \text{Unit_Two_inlet}_I \right) + \text{Mass_of_water}$$

$$\text{Total_mass_inlet} = 4.9 \times 10^3 \text{ kg}$$

3.4.3.2 unit two output

SPECIES₁ :=

Fe ₂ O ₃
HCl
Fe
FeCl ₃
Grease
Water

Outlet_Mass_2a₁ :=

0.0
0.0
1998.6
0.0
0.4
0.0

Outlet_Mass_2b₁ :=

0.0
0.0
0.0
2.027
0.0
2.901·10 ³

$$\text{Total_mass_outlet} := \sum_{i=1}^6 [(\text{Outlet_Mass_2a}_i + \text{Outlet_Mass_2b}_i) \cdot \text{kg}]$$

$$\text{Total_mass_outlet} = 4.9 \times 10^3 \text{ kg}$$

3.4.4 Unit three

Degreasing unit



$$\text{Mass_of_Grease} := \text{Grease}$$

$$\text{Mwt}_{\text{Grease}} := 287.27 \frac{\text{kg}}{\text{mol}}$$

$$\text{Grease_Reacting_Mole} := \frac{\text{Mass_of_Grease}}{\text{Mwt}_{\text{Grease}}}$$

$$\text{Grease_Reacting_Mole} = 1.392 \times 10^{-3} \text{ mol}$$

$$\text{Mass_of_NaOH} := \text{Grease_Reacting_Mole} \cdot 40 \frac{\text{kg}}{\text{mol}}$$

$$\text{Mass_of_NaOH} = 0.056 \text{ kg}$$

3.4.4.1 inlet stream

Species_κ :=

Fe
NaOH
Grease
Water

Unit_3_Inlet_κ =

1.9986710 ³	kg
0.056	
0.4	
0	

$$\text{Unit_3_inlet} := \sum_{\kappa=1}^4 \text{Unit_3_Inlet}_{\kappa}$$

$$\text{Unit_3_inlet} = 1.9991 \times 10^3 \text{ kg}$$

3.4.4.2 outlet stream

$$\text{Mass_of_Soap_formed} := \text{Grease_Reacting_Mole} \cdot \text{Mwt}_{\text{soap}}$$

$$\text{Mass_of_Soap_formed} = 0.427 \text{ kg}$$

$$\text{Mwt}_{\text{soap}} = 100 \frac{\text{kg}}{\text{mol}}$$

$$\text{Mass_of_Water_formed} := \text{Grease_Reacting_Mole} \cdot \text{Mwt}_{\text{water}}$$

$$Mwt_{\text{water}} \equiv 18 \frac{\text{kg}}{\text{mol}}$$

$$\text{Mass_of_Water_formed} = 0.0251 \text{ kg}$$

3.4.4.3 unit three outlet

Species_ω :=

Fe
NaOH
Grease
Water
Soap

Unit_3_Outlet_ω =

1.9986710 ³	kg
0	
0	
0.02506	
0.42672	

$$\text{Unit_3_outlet} := \sum_{\omega=1}^5 \text{Unit_3_Outlet}_{\omega}$$

$$\text{Unit_3_outlet} = 1.9991 \times 10^3 \text{ kg}$$

3.4.5 Unit four

Water rinsing (Soap Removal)

Calculation of Amount of Water Required for Rinsing

Solubility of Sodium Stearate at 25° C

$$\text{Solubility}_{\text{soap}} = \frac{0.1 \text{ g}_{\text{soap}}}{100 \text{ g}_{\text{water}}} \quad (\text{Shreves et.al})$$

$$\text{Mass_of_water_required} := \frac{\text{Mass_of_Soap_formed}}{\text{Solubility}_{\text{soap}}}$$

Using 35% Excess Water for Rinsing

$$\text{Mass_of_water_required} = 426.721 \text{ kg}$$

$$\text{Actual_water} := \text{Mass_of_water_required} + 35\% \cdot \text{Mass_of_water_required}$$

$$\text{Actual_water} = 576.073 \text{ kg}$$

3.4.5.1 input summary

Species _ω :=	Mass_Input_4 _ω =
Fe	1.9986710 ³ kg
NaOH	0
Grease	576.0977
Water	0.4267
Soap	

$$\text{UNIT}_4_Input := \sum_{\omega=1}^5 \text{Mass_Input}_4_{\omega}$$

$$\text{UNIT}_4_Input = 2.5751 \times 10^3 \text{ kg}$$

3.4.5.2 out summary

Species _ω :=	Mass_Output_4a _ω	Mass_Output_4b _ω
Fe	1.999710 ³ kg	
NaOH	0	0 kg
Grease	0	0
Water	3.5	0
Soap	0	572.598
		0.427

$$\text{UNIT}_4_OUTPUT := \sum_{\omega} (\text{Mass_Output}_4b_{\omega} + \text{Mass_Output}_4a_{\omega})$$

$$\text{UNIT}_4_OUTPUT = 2.575 \times 10^3 \text{ kg}$$

3.4.6 Unit five

drying of base metal

3.4.6.1 inlet stream

(water removal)

$$\text{Mass_Inlet}_5_{\omega} := \text{Mass_Output}_4a_{\omega}$$

Species _ω :=	Mass_Inlet_5 _ω =
Fe	1.9986710 ³ kg
NaOH	0
Grease	0
Water	3.5
Soap	0

3.4.6.2 output stream

Species_ω :=

Fe
NaOH
Grease
Water
Soap

Mass_Output_5a_ω :=

1.9986710 ³	kg
0	
0	
0	
0	

Mass_Output_5b_ω

0	kg
0	
0	
3.5	
0	

3.4.7 Calculation of the mass of nickel required per metal base

density of nickel

$$\rho_{\text{nickel}} := 8908 \frac{\text{kg}}{\text{m}^3}$$

mass of nickel per metal base

$$M_{\text{nickel}} := \rho_{\text{nickel}} \cdot \text{VOLUME_OF_NICKE}$$

$$M_{\text{nickel}} = 0.021 \text{ kg}$$

$$\text{Chain_Unit_Mass} := 0.1200 \text{ kg}$$

$$\text{Chain_Total_Number} := \frac{\text{METAL_BASE}}{\text{Chain_Unit_Mass}}$$

$$\text{Chain_Total_Number} = 1.665 \times 10^4$$

$$\text{TOTAL_NICKEL_REQUIRED} := \text{Chain_Total_Number} \cdot M_{\text{nickel}}$$

$$\text{TOTAL_NICKEL_REQUIRED} = 351.62 \text{ kg}$$

3.4.7.1 input summary

Species_q :=

Base Meta
Nickel

Unit_6_Input_q =

1.999710 ³	kg
351.62	

$$\text{UNIT_6_INPUT} := \sum \text{Unit_6_Input}_q$$

$$\text{UNIT_6_INPUT} = 2.35 \times 10^3 \text{ kg}$$

$$\text{Amount_of_solution} := 1.2 \cdot \text{METAL_BASE}$$

$$\text{Amount_of_solution} = 2.398 \times 10^3 \text{ kg}$$

3.4.7.2 output stream

$$\text{Associated_Solution} := 0.95\% \cdot \text{Amount_of_solution}$$

$$\text{Associated_Solution} = 22.784 \text{ kg}$$

$$\text{Species}_{\phi} :=$$

Base_Metal
Nickel
Solution

$$\text{Unit_6_Output}_{\phi} =$$

1.999710 ³	kg
351.62	
22.784	

$$\text{UNIT_6_OUTPUT} := \sum_{\phi=1}^3 \text{Unit_6_Output}_{\phi}$$

$$\text{UNIT_6_OUTPUT} = 2.373 \times 10^3 \text{ kg}$$

3.4.8 Unit seven

DRAG OUT (WATER RINSING USING FOUR CASCADE TANK)

$$\text{Unit_7_Input}_{\phi} := \text{Unit_6_Output}_{\phi}$$

$$\text{Species}_{\phi} :=$$

Base_Metal
Nickel
Solution

$$\text{Unit_7_Input}_{\phi} =$$

1.9986710 ³	kg
351.6197	
22.784	

$$\text{Solution_Solubility} = \frac{1.25 \text{ g}}{100 \text{ g}_{\text{water}}}$$

$$\text{Amount_of_Water_Required} := \frac{\text{Associated_Solution}}{\text{Solution_Solubility}}$$

$$\text{Amount_of_Water_Required} = 1.823 \times 10^3 \text{ kg}$$

$$\text{UNIT_7_INPUT} = \text{Amount_of_Water_Required} \sum_{\phi} \text{Unit_7_Input}_{\phi}$$

$$\text{UNIT_7_INPUT} = 4.196 \times 10^3 \text{ kg}$$

3.4.8 Output streams

$$\text{Associated_Water} := 0.50\% \cdot \text{Amount_of_Water_Required}$$

$$\text{Associated_Water} = 9.114 \text{ kg}$$

Species $\phi :=$

Base_Metal
Nickel
Water
Solution

EXIT_7A $_{\tau} =$

1.999710 ³	kg
351.62	
9.114	
0	

EXIT_7B $_{\tau} =$

0	kg
0	
1.814710 ³	
22.784	

$$\text{UNIT_7_OUTPUT} := \sum_{\tau} (\text{EXIT_7A}_{\tau} + \text{EXIT_7B}_{\tau})$$

$$\text{UNIT_7_OUTPUT} = 4.196 \times 10^3 \text{ kg}$$

Air dryer

3.4.9 Unit 8

3.4.9.1 input

Species $\phi :=$

Base_Metal
Nickel
Water
Solution

Unit_8_Input $_{\tau} =$

1.9986710 ³	kg
351.6197	
9.1136	
0	

3.4.9.2 output

3.4.10 Chrome plating unit

3.4.10.1 determination of the amount of chrome needed for plating using the classical oxide method.

Species $\phi :=$

Base_Metal
Nickel
Water
Solution

Unit_8a_Output $_{\tau} =$

1.9986710 ³	kg
351.6197	
0	
0	

Unit_8b_Output $_{\tau} =$

0	kg
0	
9.114	
0	

$$\text{Total_Surface_Area} = 7.9 \times 10^{-3} \text{ m}^2$$

$$\text{CHROME_THICKNESS} := 0.02 \text{ m}$$

$$\text{VOLUME_OF_CHROME} := \text{Total_Surface_Area} \cdot \text{CHROME_THICKNESS}$$

(Steven S. Zumdahl, 1998.)

3.4.10.2 calculations of the volume of chrome required per metal base

VOLUME OF CHROME = 1.58×10^{-3} L

3.4.10.3 calculation of the mass of chrome required per metal base

density of chrome

$$\rho_{\text{chrome}} := 7140 \frac{\text{kg}}{\text{m}^3}$$

mass of chrome per metal base

$$M_{\text{chrome}} := \rho_{\text{chrome}} \cdot \text{VOLUME_OF_CHROME}$$

$$M_{\text{chrome}} = 0.011 \text{ kg}$$

$$\text{Chain_Unit_Mass} = 0.12 \text{ kg}$$

$$\text{Chain_Total_Number} = 1.665 \times 10^4$$

$$\text{TOTAL_CHROME_REQUIRED} = 187.888 \text{ kg}$$

$$\text{TOTAL_CHROME_REQUIRED} = \text{Chain_Total_Number} \cdot M_{\text{chrome}}$$

3.4.10.4 input summary

Species ϕ :=

Base_Metal
Nickel
Water
Chrome

Unit_9a_Input $_{\tau}$ =

1.9986210 ³	kg
351.6197	
0	
0	

Unit_9b_Input $_{\tau}$ =

0	kg
0	
0	
187.888	

$$\text{Unit_9_Output}_{\tau} := \sum_{\tau} (\text{Unit_9a_Input}_{\tau} + \text{Unit_9b_Input}_{\tau})$$

$$\text{Unit_9_Output} = 2.5381 \times 10^3 \text{ kg}$$

3.4.10.5 output summary

$$\text{Unit_9_Output}_{\tau} := (\text{Unit_9a_Input}_{\tau} + \text{Unit_9b_Input}_{\tau})$$

Species ϕ :=

Base_Metal
Nickel
Water
Chrome

Unit_9_Output $_{\tau}$ =

1.9986210 ³	kg
351.6197	
0	
187.8884	

$$\text{Unit_9_Input} := \sum_{\tau} (\text{Unit_9a_Input}_{\tau} + \text{Unit_9b_Input}_{\tau})$$

3.5 Energy Balance

MathCAD was Similarly used to carry out the energy balance, knowing the temperature of each of the unit; from program tool bar and thermodynamic package the different change in enthalpy of formation of chemical species involved were selected and consequently, for each species the product of mass and enthalpy(ΔH) multiplied by 1000J/kg gives energy balance for each component and the MathCAD automatically calculate. This is done for each unit starting from acid pickling tank to drying unit. Finally, the time for nickel plating was 3.56hrs energy spent was 20.89×10^5 j and power was 22.5 watts while time for chrome plating was 1.42hrs energy spent was 2.557×10^7 j and the power was 20.5 watts.

3.5.1 Energy balance

3.5.1.1 energy balance around unit one at 25°C

$$Q_{\text{one.IN}} = \text{mass_in_1} \Delta H_1$$

$$Q_{\text{IN}} = Q_{\text{OUT}}$$

SPECIES _{ω} :=

Fe ₂ O ₃
HCl
FeCl ₃
Grease
Water

Mass_Out_1 _{ω} :=

0
0
2.0275
0.4
2.025

ΔH_1 _{ω} :=

7950.63
2534.2
2999.5
3150.1
15808

(Steven S. Zumdahl, 1998.)

$$Q_{\text{one.IN}} := \sum_{\omega} \left[\left(\text{mass_in_1}_{\omega} \cdot \Delta H_{1\omega} \right) \frac{1000\text{J}}{\text{kg}} \right]$$

$$Q_{\text{one.IN}} = 1.268 \times 10^7 \text{ J}$$

$$Q_{\text{one.OUT}} = \text{Mass_out_1} \cdot \Delta H_2$$

SPECIES_ω :=

Fe ₂ O ₃
HCl
FeCl ₃
Grease
Water

mass_in_1_ω :=

Rust
HCl
0
Grease
0

ΔH_{2ω} :=

7950.63
2534.2
2999.5
3150.1
15808

$$Q_{\text{one.OUT}} := \sum_{\omega} \left[\left(\text{Mass_Out_1}_{\omega} \cdot \Delta H_{2\omega} \right) \cdot 1000 \right]$$

$$Q_{\text{one.OUT}} = 3.935 \times 10^7 \text{ J}$$

3.5.1.2 calculation of energy required

Q_{RQD} = Quantity_of_Energy_Required

$$Q_{\text{RQD}} := Q_{\text{one.OUT}} - Q_{\text{one.IN}}$$

$$Q_{\text{RQD}} = 2.667 \times 10^7 \text{ J}$$

3.5.2 Energy balance around unit two at 25^oc

3.5.2.1 Input energy

$$Q_{\text{two.IN}} = \text{Mass_Out_1}_{\omega} \cdot \Delta H_{2\omega}$$

$$Q_{\text{two.IN}} = Q_{\text{two.OUT}}$$

SPECIES_ω :=

Fe ₂ O ₃
HCl
FeCl ₃
Grease
Water

Mass_Out_1_ω :=

0
0
2.0275
0.4
2.025

ΔH_{2ω} :=

7950.63
2534.2
2999.5
3150.1
15808

$$Q_{\text{two.IN}} := \sum_{\omega} \left[\left(\text{Mass_Out}_{-1\omega} \cdot \Delta H_{2\omega} \right) \cdot 1000 \right]$$

$$Q_{\text{two.IN}} = 3.935 \times 10^7 \text{ J}$$

$$Q_{\text{two.OUT}} := Q_{\text{two.IN}}$$

$$Q_{\text{two.OUT}} = 3.935 \times 10^7 \text{ J}$$

3.5.3 Energy balance around unit three

Input Energy At 25°C

Species_κ :=

Fe
NaOH
Grease
Water

Unit_3_Inlet_κ =

1.9986710 ³	kg
0.056	
0.4	
0	

ΔH_{3κ} :=

0
-4555.1
-3150.1
-15808

$$Q_{\text{three.IN}} := \sum_{\kappa} \left[\left(\text{Unit_3_Inlet}_{\kappa} \cdot \Delta H_{3\kappa} \right) \cdot 1000 \frac{\text{J}}{\text{kg}} \right]$$

Output Energy At 50°C

Species_ω :=

Fe
NaOH
Grease
Water
Soap

Unit_3_Outlet_ω =

1.9986710 ³	kg
0	
0	
0.0251	
0.4267	

ΔH_{3ω} :=

0
-4327.8
-3098.4
-15707
-2513.9

$$Q_{\text{three.OUT}} := \sum_{\omega} \left[\left(\text{Unit_3_Outlet}_{\omega} \cdot \Delta H_{3\omega} \right) \cdot 1000 \frac{\text{J}}{\text{kg}} \right]$$

3.5.3.1 Energy balance equation

$$Q_{\text{three.IN}} + Q_{\chi} = Q_{\text{three.OUT}}$$

$$Q_{\chi} = \text{ENERGY_REQUIREI}$$

$$Q_{\chi} := Q_{\text{three.OUT}} - Q_{\text{three.IN}}$$

$$Q_{\chi} = 4.872 \times 10^4 \text{ J}$$

3.5.4 Energy balance around unit four

Energy Input At 50°C

Species ω :=	Unit_3_Outlet ω =	$\Delta H_{4\omega}$:=
Fe	1.9986710 ³ kg	0
NaOH	0	-4327.8
Grease	0	-3098.4
Water	0.0251	-15707
Soap	0.4267	-2513.9

$$Q_{\text{four.IN}} := \sum_{\omega} \left[\left(\text{Mass_Output_4a}_{\omega} \cdot \Delta H_{4\omega} \right) \cdot 1000 \frac{\text{J}}{\text{kg}} \right]$$

Energy Output At 25°C

Species ω :=	Mass_Output_4a ω =	$\Delta H_{4A\omega}$:=
Fe	1.9986710 ³ kg	0
NaOH	0	-4555.1
Grease	0	-3150.1
Water	3.5	-15808
Soap	0	-2534.2

$$Q_{4A_{\text{out}}} := \sum_{\omega} \left(\text{Mass_Output_4a}_{\omega} \cdot \Delta H_{4A\omega} \cdot 1000 \frac{\text{J}}{\text{kg}} \right)$$

Species ω :=	Mass_Output_4b ω	$\Delta H_{4B\omega}$:=
Fe	0 kg	0
NaOH	0	-4555.1
Grease	0	-3150.1
Water	572.598	-15808
Soap	0.427	-1024.2

$$Q_{4B_{\text{out}}} := \sum_{\omega} \left[\left(\text{Mass_Output_4b}_{\omega} \cdot \Delta H_{4B\omega} \right) \cdot 1000 \frac{\text{J}}{\text{kg}} \right]$$

$$Q_{\text{four.OUT}} := Q_{4A_{\text{out}}} + Q_{4B_{\text{out}}}$$

$$Q_{\Psi} := Q_{\text{four.IN}} - Q_{\text{four.OUT}}$$

$$Q_{\Psi} = 9.052 \times 10^9 \text{ J}$$

Q_{Ψ} = Amount_of_Energy_Removed

3.5.5 Energy balance around unit five

$$\text{ENERGY_IN} = \text{ENERGY_OUT}$$

Input Energy At 25°C

Species _ω :=	Mass_Inlet _{5_ω} =	ΔH _{5_ω} :=
Fe	1.999710 ³ kg	0
NaOH	0	-4555.1
Grease	0	-3150.1
Water	3.5	-15808
Soap	0	-2534.2

$$Q_{\text{five.IN}} := \sum_{\omega} \left[\left(\text{Mass_Inlet}_{5_{\omega}} \cdot \Delta H_{5_{\omega}} \right) \cdot 1000 \frac{\text{J}}{\text{kg}} \right]$$

$$Q_R = \text{ENERGY_REQUIRED_TO_DRIVE_AWAY_WATER_MOLECUL}$$

$$Q_R = M \cdot C_p \cdot \Delta T$$

$$M := 3.5 \text{ kg}$$

$$C_p := 4185 \frac{\text{J}}{\text{kg} \cdot \text{K}}$$

$$\Delta T := (303 - 298) \text{ K}$$

$$Q_R = 7.324 \times 10^4 \text{ J}$$

$$\text{UNIT_5_EXIT_ENERGY} := Q_R - Q_{\text{five.IN}}$$

$$\text{UNIT_5_EXIT_ENERGY} = 5.54 \times 10^7 \text{ J}$$

3.5.6 Energy balance around unit six

Species _q :=	Unit_6_Input _q =
Base_Metal	1.9986710 ³ kg
Nickel	351.6197



Applying Faraday's Law

$$W_{\text{Ni}} = \text{MOLECULAR_WEIGHT_OF_NICKE}$$

$$W_{\text{Ni}} := 58.69 \frac{\text{kg}}{\text{mol}}$$

$$M_{\text{Ni}} = \text{MASS_OF_NICKEL_DEPOSITEI}$$

$$M_{\text{Ni}} := \text{TOTAL_NICKEL_REQUIRE}$$

$$n_{\text{Ni}} = \text{MOLE_OF_NICKEL_DEPOSITEI}$$

$$n_{\text{Ni}} := \frac{M_{\text{Ni}}}{W_{\text{Ni}}}$$

$$n_{\text{Ni}} = 5.991 \text{ mol}$$

$$2 \text{ mol} \cdot e^{-} = 96486 \text{ coulomb} \quad (\text{Faraday's Law})$$

$$M_{\text{Ni}} = E \cdot I \cdot T \quad \dots \text{eqn}_1$$

E is the Electrochemical Equivalence

I is the Current Density

T is the time required for Plating

Calculation of Amount of Columb Required

$$A_{\text{columb}} = \text{Amount_of_Columb}$$

$$A_{\text{columb}} := n_{\text{Ni}} \cdot \frac{96486 \text{ C}}{2 \text{ mol}}$$

$$A_{\text{columb}} = 2.89 \times 10^5 \text{ C}$$

$$\text{From Literature,} \quad I := 22.5 \frac{\text{C}}{\text{s}}$$

$$\text{From eqn}_1, \quad \frac{M_{\text{Ni}}}{E} = I \cdot T$$

$$\frac{M_{Ni}}{E} = A_{columb}$$

Then,

$$A_{columb} = I \cdot T$$

$$\text{Time_Required} := \frac{A_{columb}}{I}$$

$$\text{Time_Required} = 1.285 \times 10^4 \text{ s}$$

$$\text{Time_Required} = 3.56 \text{ hr}$$

$$Q_{\text{Plating}} = \text{Energy_Required_for_Plating}$$

$$Q_{\text{Plating}} = I \cdot V \cdot \text{Time_Required}$$

$$V = \text{Voltage_Required}$$

$$V := 220 \text{ volt}$$

$$Q_{\text{Plating}} = 2.89 \times 10^5 \text{ J}$$

$$\text{Power} := \frac{Q_{\text{Plating}}}{\text{Time_Required}}$$

$$\text{Power} = 22.5 \text{ W}$$

3.5.7 Chrome plating unit



3.5.7.1 Applying Faraday's Law

$$W_{Cr} = \text{MOLECULAR_WEIGHT_OF_CHROMIUM}$$

$$M_{Cr} = \text{MASS_OF_CHROME_DEPOSITE}$$

$$W_{Cr} := 51.9961 \frac{\text{kg}}{\text{mol}}$$

$$M_{Cr} := \text{TOTAL_CHROME_REQUIRE}$$

$$n_{Ni} = \text{MOLE_OF_CHROME_DEPOSITED}$$

$$n_{Cr} := \frac{M_{Cr}}{W_{Cr}}$$

$$n_{Cr} = 3.614 \text{ mol}$$

$$3 \text{ mol} \cdot e^- = 96486 \text{ coulomb}$$

$$M_{Ni} = E \cdot I \cdot T$$

$$A_{\text{coulomb}} = \text{Amount_of_Coulomb}$$

Calculation of Amount of Coulomb Required

$$A_{\text{coulomb}} := n_{Cr} \cdot \frac{96486 \text{ C}}{3 \text{ mol}}$$

$$A_{\text{coulomb}} = 1.162 \times 10^5 \text{ C}$$

$$\text{Time_Required} := \frac{A_{\text{coulomb}}}{I}$$

$$\text{Time_Required} = 5.165 \times 10^3 \text{ s}$$

$$\text{Time_Required} = 1.243 \text{ hr}$$

$$Q_{\text{plating}} = \text{Energy_Required_for_Plating}$$

$$Q_{\text{plating}} = 2.557 \times 10^7 \text{ J}$$

$$\text{Power} = \frac{Q_{\text{Plating}}}{\text{Time Required}}$$

$$\text{Power} = 20.5$$

CHAPTER FOUR

4.0 EQUIPMENT DESIGN

The result of the equipment that were designed are given in Table 4.1 to 4.3 and the result of the cost can be seen in Table 4.4. Detail calculation shown in Appendix i.

Table 4.1: Design Summary Vessel Parameter and Sizing for Electroplating Equipment

Design Parameters	Acid Pickling	Washing Tank 1	Degreasing Unit	Washing Tank 2	Washing Tank 3	Nickel Plating Unit	Drag Out Tank 1
M/TYPE	Carbon Steel	Carbon steel	Carbon steel	Carbon steel	Carbon steel	Carbon steel	Carbon steel
Diameter (m)	1.200	1.198	0.3048	1.203	1.200	-	0.9405
Height (m)	1.799	1.797	1.804	1.799	1.799	-	1.411
Length (m)	4.801	3.535	1.676	1.804	2.515	3.353	0.9800
Volume (m ³)	2.024	2.024	2.583	2.050	2.034	2.583	1.676
Shell thickness	6.350	6.350	6.350	6.350	6.350	6.350	6.350
Corrosion thickness (mm)	3.175	3.175	3.175	3.175	3.175	3.175	3.175
Vessel Pressure (Kpa)	100	100	100	100	100	90.00	90.00
Joint efficiency	1.000	1.000	1.000	1.000	1.000	1.000	1.000
L/D Ratio	3.000	5.000	5.000	5.000	5.000		
Allowance stress (Kpa)	9.446 e + 004	9.446 e + 004	9.446 e + 004	9.446 e + 004	9.446 e + 004	9.446 e + 004	9.446 e + 004

Design Parameter	Dragout Tank 2	Dragout Tank 3	Concentrator	Chrome Plating Tank
M/TYPE	Carbon Steel	Carbon steel	Carbon steel	Carbon steel
Diameter (m)	0.9405	0.9405	0.9405	-
Height (m)	1.411	1.411	1.411	3.535
Length (m)	1.676	1.676	1.676	2.583
Volume (m3)	0.9800	0.9800	0.1223	-
Shell thickness	6.350	6.350	-	6.350
Corrosion thickness (mm)	3.175	3.175	3.175	3.175
Vessel Pressure (Kpa)	90	90	90	90.00
Joint efficiency	1.000	1.000	-	1.000
L/D ratio	-	5.000	-	
Allowance stress (Kpa)	9.446 e + 004	9.446 e + 004	9.446 e + 004	9.446 e + 004

Table 4.2 Design Summary of Pumps Parameters and Sizing for The Entire Electroplating Process.

I. Design	Pump 1	Pump 2	Pump 3	Pump 4	Recycling Pump
Power	1.978 kJ/h	1.045 kJ/h	2.090 kJ/h	16.09 kJ/h	53.71 kJ/h
Pressure	0.6897m	1.012m	1.012m	1.012m	1.016m
Heat Duty	5.498e ⁻⁰⁸⁸ kw	2.903e ⁻⁰⁰⁴ kw	5.305e ⁻⁰⁰⁴ kw	4.470e ⁻⁰⁰³ kw	1.480e ⁻⁰⁰² kw
Capacity	1.560m ³	0.1002 m ³ /h	0.2004 m ³ /h	1.543 m ³ /h	5.131 m ³ /h
Heat Velocity	017m	001m	90m	009m	007m
Head DP	-1.48e ⁻⁰¹⁷ 10kpa	-5.557e ⁻ 10kpa	-2.223e ⁻ 10kpa	-1.318e ⁻ 10.00kpa	-1.457e ⁻⁰⁰⁷ 10.00kpa
Radius of Gyration	0.1000	0.1000	0.1000	0.1000	

Table 4.3 Design Summary of Parameter for Heater, Pre Heater Cooler.

Design parameter	Pre heater	Heater	Cooler
Duty	5.466e+005kJ/h	6.319kJ/h	-5.251e+005kJ/h
Pressure drops	0.0000k/a	0.0000k/a	2.000k/a
Area (cm)	4.320m ²	4.650m ²	
Overall heat Loose (kJ/h)			

Table 4.4 Design Summary of Parameter for Air Dryer

Design parameter	Sizing
Volumetric Air flow	7.200e+005 (ACT m ³ /h)
UA	-1625Kj/c-h
Working fluid duty	1.225kJ/h
Mass Air Flow	8.491e+005kg/h
LMTD	1.865 C

4.1 Result of Equipment Costing

Exchange rate ER as at 2006 is 130 Naira for 1 Dollar as at 2006.

Table 4.5 Costing of Equipment Using Marshall and Smith Cost Estimation Index

FM		FP	
Carbon steel CS	1.00	< 50 P SIG	1.00
Stainless Steel SS	3.67	200	1.15
Moniel Steel	6.34	400	1.35
Titanium	7.89	600	1.60

Table 4.6 Cost of Raw Material Using Direct Cost Estimation

Raw Material	Quantity	COST	IN	Equivalent Cost in Naira
Cr (So ⁴) ₃		DOLLAR		
Ni SiO ₄ . 6 H ₂ O	3295 kg	234		30,500
NiCl ₂ . 6H ₂ O	10.71kg	230		21,800
H ₂ BO ₄	7.03 kg	167		1,200
H ₂ SO ₄	1.0 x 10-2L	19.2		2,500
Dissolvable Nickel Anode	40.679Kg	98.5		12,800
HCL	8.4635 X 10-2 Kg	13.8		1800
NaOH	3.402 x 10-2 Kg	6.9		900
Water required for entire process	6.5874 x 10 3L	65.4		8,500
Wrist watches	1999.0kg	153.8		19,900

4.2 Summary OF Result

Table 4.7 Result of Cost Estimation and Economic Analysis Using Marshal and Smith Cost Index Data

Items	Economic Analysis Marshal and Smith Cost Index)	
	DOLLARS	(Naira
Purchase cost of equipment	21,220.6	2,758,677
Total fixed capital cost (TFC)	212,205.9	27,586,770
Working capital (WC)	106,103	13,793,385
Total land cost (TLC)	21,220.6	2,768,677
Total fixed cost invested (Tinv)	339,529.5	44,138.832
Labour cost	12,732.4	1,655,206
Plant maintenance and repairs (MC)	16,976.5	2,206,941
Insurance InSc	2,122.1	275,867
Local Taxes (Ltxc)	4,244.1	551.735.4
Royalties (Royc)	2,122.1	275.867
Laboratory cost (Labc)	848.8	110,347.08
Supervision (Sc)	169.8	22,069.4
Plant overhead cost (POHc)	2,122.1	275,867
Administrative cost (ADMc)	1,061	137,933.85
Reaserch and development cost (RADc)	3,183.1	413,801.55
Total fixed operating cost (TFOc)	45,412	5,903,567
Total annual fixed operating cost TFO	2,270.6	295,178
Cost of raw material (CRMc)	1,134.9	147,540
Annual cost of raw material (ACRMc)	340,476.9	44,262,000
Miscellaneous (Msc)	212.2	27,586.77
Utilities Cost (utc)	84.8	11,034.7
Packaging Pac	4.2	551.7
Total variable operating cost (Tvoc)	340,778.3	44,301,173
Annual variable cost (Tvo) annual	17,038.9	2,215.058
Total annual operating cost (AOCc)	19,309.5	2,510,236
Annual production rate (APR)	-	181440000kg/yr.
Production cost per kg (PC)	0.015	2.0
Annual product sales (APS)	167,483	21,772,800
Selling price (SP)	0.09	12
Profit before tax (PBT)	148,174	19,262,563
Depreciation annual (Dep)	15,385	2,000,040
Tax payable (TP)	86,312.6	11,220,639
Profit after tax (PAT)	61,851	8,041,923
Net Income (NIN)	77,245.8	10,041,964
Pay Back Period (PBP)	-	3.5 years
Rate of return on investment ROR	-	27.28%

4.3 Discussion Of Results

The small scale chrome electroplating plant was designed with the aid of computer program and process simulator. MathCAD was used to carry out the material and energy balance, Hysys was used to obtain the equipment design and process flow sheet .

The results of the equipment designed shows that the Acid Pickling unit, washing tanks, degreasing unit, Nickel plating unit, dragout unit concentrator and chrome plating unit are all rectangular shape tanks as shown in Table 4.1 to 4.13 and the corresponding result of the costing were shown in Tale 4.14 to 4.26. Marshall and Smith Cost Estimation Index approach was used for the cost estimation and economic analysis for the plant and are has shown in table 4.28. Details are shown in Appendix III.

The purchase cost of equipment was \$21,220 (Twenty One Thousand Two Hundred and Twenty Dollars) and the total fixed cost to be invested on the plant was \$339,529 (Three hundred and Thirty Nine Thousand Five Hundred and Twenty Nine Dollars).

Also the annual cost of raw materials was \$340,476 (Three Hundred and Fourty Thousand Four Hundred and Seventy Six Dollars), the annual production rate was estimated as 181440000kg/yr and was as shown in table 4.28. The cost of chrome electroplating of 0.12kg of iron wrist watch chain was N2 and the selling price was N12 each respectively.

The plant has an annual operating cost of \$2,270 (Two Thousand Two Hundred and Seventy Dollars) and the profit after tax (PAT) was

\$61,851 (Six One Thousand Eight Hundred and Fifty One Dollars) while the profit before tax (PBT) was \$148,174 (One Hundred and Forty Eight Thousand One Hundred and Seventy Four Dollars). The pay back period of the capital invested on the plant was 3 ½ years and the rate of return on investment (ROR) was 27.28% . However, in this design work attention was given to the process aspect while datas are provided in Appendix I for Mechanical Technicians and Fabricators. The general observation of table 4.28 shows that electroplating is a very lucrative business the any intending business man can venture into and make handsome profit.

CHAPTER FIVE

5.0 CONCLUSION

A powerful technical computer environment (MATHCAD) was used to carry out the material and energy balance of this design work while process simulator (HYSYS) was used for the equipment design and flow sheet construction. The Marshal & Smith cost index estimation approach was used. This design of 2000kg/day of chrome electroplating of a wrist watch chain plant, was made to develop a cost effective, user and environmental friendly plant, that takes into consideration the technological development of a viable economy and market structure for an electroplating plant .

The economic analysis as shown in table 4.28 shows that the cost of equipment was \$21,220 (Twenty One Thousand Two and Twenty Dollar). the profit after tax of \$61,851 (Sixty One Thousand Eight Hundred and Fifty One Dollars) with pay back period of 3 ½ years and rate of return (R)OR of 27.28%. In view of all these it can be concluded that this project is economically viable.

5.1

Recommendation

The under listed recommendation are of paramount importance for any potential businesspersons or government bodies. Interested in electroplating business.

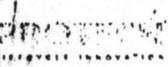
1. The design of the plant should be supervised by a technical personnel.
2. An adequate safety measure should be put into consideration.
3. A good electroplating bath maintenance culture ensured.
4. A Proper statistic of bath technological parameters should be kept to ensure quality product.
5. Further work remains to be done on optimization of electroplating plant.

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APPENDIX I EQUIPMENT DESIGN DETAILS

 <p>TEAM LND Calgary, Alberta CANADA</p>	Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc
	Unit Set: SI
	Date/Time: Thu Dec 21 16:45:40 2006

Tank: ACID PICKLING UNIT

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
METAL	

Outlet Stream

Stream Name	To Unit Operation
R METAL	Tank: WASHING UNIT 1

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Volume:	2.034 m3	Level SP:	50.00 %	Liquid Volume	1.017 m3
Pressure:	100.0 kPa	Pressure Drop:	0.0000 kPa	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heating

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot:	No
2.034 m3	Diameter	1.200 m	Height
			1.799

Nozzles

Elevation Relative to Ground Level	0.0000 m	Diameter	1.200 m	Height	1.799
		BASE METAL	HCl		WATER
er	(m)	8.997e-002	8.997e-002		8.997e-002
n (Base)	(m)	0.8997	0.8997		1.799
n (Ground)	(m)	0.8997	0.8997		1.799
n (% of Height)	(%)	50.00	50.00		100.00
		BASE METAL			
er	(m)	8.997e-002			
n (Base)	(m)	0.0000			
n (Ground)	(m)	0.0000			
n (% of Height)	(%)	0.00			

Detailed Heat Loss Parameters

Heat Loss (kJ/h)	0.0000	Area (m2)	6.782
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Temperature Profile

0.0000	Inner Wall (C)	25.00	Outer Vessel (C)	25.00	Outer Insulation (C)	25.00	Ambient (C)	25.00
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Conduction

Thickness	Metal	Insulation	
(m)	1.000e-002	3.000e-002	
(kJ/kg-C)	0.4730	0.8200	
(kg/m3)	7801	520.0	
ivity	(W/m-K)	45.00	0.1500

Convection

Cap Phase U (kJ/h-m2-C)	7200	Inside Lq Phase U (kJ/h-m2-C)	-	Outside U (kJ/h-m2-C)	54.00
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Level Taps: Level Tap Specification

Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Tap	Liquid Level	Aqueous Level
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Options



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Thu Dec 21 16:45:40 2006

Tank: ACID PICKLING UNIT (continued)

PV Work Term Contribution (%) 100.00 *

DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	2.034 *	Level Calculator	Vertical cylinder
Vessel Diameter (m)	1.200	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	1.799	Feed Delta P (kPa)	0.0000 *
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	100.0

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	-	-	0.0000
Liquid	-	-	0.0000
Aqueous	-	-	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000 *	0.0000
Liquid	0.0000	0.0000 *	0.0000
Aqueous	0.0000	0.0000 *	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc
Unit Set: SI
Date/Time: Fri Feb 09 09:17:03 2007

Vessel Sizing: Vessel Sizing-8

Vertical Separator Properties: CHROMEPLATING TANK

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	300.0 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7396 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7068 *

Costing Results

Base Cost (\$US)	7476	Associated Cost (\$US)	1658	Total Cost (FOB \$US)	9134
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Sizing Results

Diameter (m)	0.6096	Liquid Residence Time (seconds)	300.0
Total Length (m)	3.353	Liquid Surge Height (m)	1.521
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	0.6644	Liquid Res. Time at LLSD (seconds)	82.94
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	1.070	Ellipsoidal Head (m)	0.1524
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.832



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:29:17 2007

Tank: TEMPORARY STORAGE TANK

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
CHROME SOLUTION	Separator CHROME SOLUTION CONCENTRATOR

Outlet Stream

Stream Name	To Unit Operation

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume: --	Level SP: 50.00 %	Liquid Volume: --	
Vessel Pressure: 90.00 kPa	Pressure Drop: 0.0000 kPa *	Duty: 0.0000 kJ/h	Heat Transfer Mode: Heating

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot: No	
Volume --	Diameter --	Height --	

Nozzles

Vessel Elevation Relative to Ground Level	0.0000 m *	Diameter	Height

Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options

Work Term Contribution	(%)	100.00 *	
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	--	Level Calculator	Vertical cylinder
Vessel Diameter (m)	--	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	--	Feed Delta P (kPa)	0.0000 *
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	90.00

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:29:17 2007

Tank: TEMPORARY STORAGE TANK (continued)

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:21:17 2007

Conversion Reactor: CHROME PLATING TANK

CONNECTIONS

Inlet Stream Connections

Stream Name	From Unit Operation
SOLUTION	
PLATED METAL	Tank DRAGOUT TANK.3

Outlet Stream Connections

Stream Name	To Unit Operation
CHROMED METAL	Tank: DRAGOUT TANK.4
L.SOLN	Mixer: CHROME SOLUTION MIXER

Energy Stream Connections

Stream Name	From Unit Operation
Q.TH	

PARAMETERS

Physical Parameters		Optional Heat Transfer: Heating	
Delta P	Vessel Volume	Duty	Energy Stream
0.0000 kPa	--	6.641e+006 kJ/h	Q.TH

User Variables

RATING

Sizing

Cylinder	Vertical	Reactor has a Bool:	No
Volume --	Diameter --	Height	--

Nozzles

Base Elevation Relative to Ground Level	0.0000 m *	Diameter	--	Height	--
	SOLUTION	PLATED METAL	CHROMED METAL		
Diameter (m)	5.000e-002	5.000e-002	5.000e-002		
Elevation (Base) (m)	0.0000	0.0000	0.0000		
Elevation (Ground) (m)	0.0000	0.0000	0.0000		
Elevation (% of Height) (%)	--	--	--		
	L.SOLN				
Diameter (m)	5.000e-002				
Elevation (Base) (m)	0.0000				
Elevation (Ground) (m)	0.0000				
Elevation (% of Height) (%)	--				

DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	--	Level Calculator	Vertical cylinder
Vessel Diameter (m)	--	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	--	Feed Delta P (kPa)	0.0000 *
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	90.00

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:21:17 2007

Conversion Reactor: CHROME PLATING TANK (continued)

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000

Duty Valve Source : Direct_Q

SP (kJ/h)	Min. Available (kJ/h)	Max. Available (kJ/h)
6.641e+006	--	--

Liquid Heater Height as % of Vessel Volume

Top of Heater : 5.00 %	Bottom of Heater : 0.00 %
------------------------	---------------------------

Heat Flow into the PFR: Heating



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:28:27 2007

Separator: CHROME SOLUTION CONCENTRATOR

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
L	Heater PRE_HEATER

Outlet Stream

Stream Name	To Unit Operation
WATER VAPOUR	
CHROME SOLUTION	Tank: TEMPORARY STORAGE TANK

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	0.9966 m3	Level SP:	50.00 %	Liquid Volume:	0.4983 m3
Vessel Pressure:	90.00 kPa	Pressure Drop:	0.0000 kPa *	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heating

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot:	No
Volume 0.9966 m3	Diameter 0.8700 m *	Height	1.676 *

Nozzles

Base Elevation Relative to Ground Level	0.0000 m *	Diameter	0.8700 m *	Height	1.676 *
	L	WATER VAPOUR	CHROME SOLUTION		
Diameter (m)	8.382e-002	8.382e-002	8.382e-002		
Elevation (Base) (m)	0.8382	1.676	0.0000		
Elevation (Ground) (m)	0.8382	1.676	0.0000		
Elevation (% of Height) (%)	50.00	100.00	0.00		

Detailed Heat Loss Parameters

Overall Heat Loss (kJ/h)	0.0000	Area (m2)	4.582
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Temperature Profile

Liquid (C)	0.0000 *	Inner Wall (C)	25.00	Outer Vessel (C)	25.00	Outer Insulation (C)	25.00	Ambient (C)	25.00
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Conduction

	Metal	Insulation
Thickness (m)	1.000e-002	3.000e-002
ρ (kJ/kg-C)	0.4730	0.8200
Density (kg/m3)	7801	520.0
Conductivity (W/m-K)	45.00	0.1500

Convection

Inside Vap Phase U (kJ/h-m2-C)	7200	Inside Liq Phase U (kJ/h-m2-C)	--	Outside U (kJ/h-m2-C)	54.00
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Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low

Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level

Options

Work Term Contribution (%)	100.00 *
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DYNAMICS

Vessel Parameters: Initialize from Product



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:28:27 2007

Separator: CHROME SOLUTION CONCENTRATOR (continued)

Vessel Volume	(m3)	0.9966	Level Calculator	Vertical cylinder
Vessel Diameter	(m)	0.8700	Fraction Calculator	Use levels and nozzles
Vessel Height	(m)	1.676	Feed Delta P	(kPa) 0.0000
Liquid Level Percent	(%)	50.00	Vessel Pressure	(kPa) 90.00

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc
Unit Set: SI
Date/Time: Wed Feb 07 12:27:16 2007

Mixer: CHROME SOLUTION MIXER

CONNECTIONS

Inlet Stream

STREAM NAME	FROM UNIT OPERATION
SOLN	Conversion Reactor
SOLN.2	Tank
SOLN.3	Tank
SOLN.4	Tank

Outlet Stream

STREAM NAME	TO UNIT OPERATION
CHROME SOLUTION	Heater

PARAMETERS

User Variables

NOZZLE PARAMETERS

Case Elevation Relative to Ground Level	0.0000 m *		
	L.SOLN	L.SOLN.2	L.SOLN.3
meter (m)	5.000e-002	5.000e-002	5.000e-002
levation (Base) (m)	0.0000	0.0000	0.0000
levation (Ground) (m)	0.0000	0.0000	0.0000
	L.SOLN.4	CHROME SOLUTION	
meter (m)	5.000e-002	5.000e-002	
levation (Base) (m)	0.0000	0.0000	
levation (Ground) (m)	0.0000	0.0000	

DYNAMICS

Pressure Specification: Set Outlet to Lowest Inlet

Holdup Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:25:42 2007

Cooler: COOLER 1

CONNECTIONS

Inlet Stream

STREAM NAME	FROM UNIT OPERATION
CONCENTRATED NICKEL SOLUTION	Separator CONCENTRATOR

Outlet Stream

STREAM NAME	TO UNIT OPERATION
	Pump P-100

Energy Stream

STREAM NAME	TO UNIT OPERATION

PARAMETERS

Pressure Drop:	2.000 kPa *	Duty:	1.235e+006 kJ/h	Volume:	0.1000 m ³
Zone:	Not Selected	Zones:	1		

User Variables

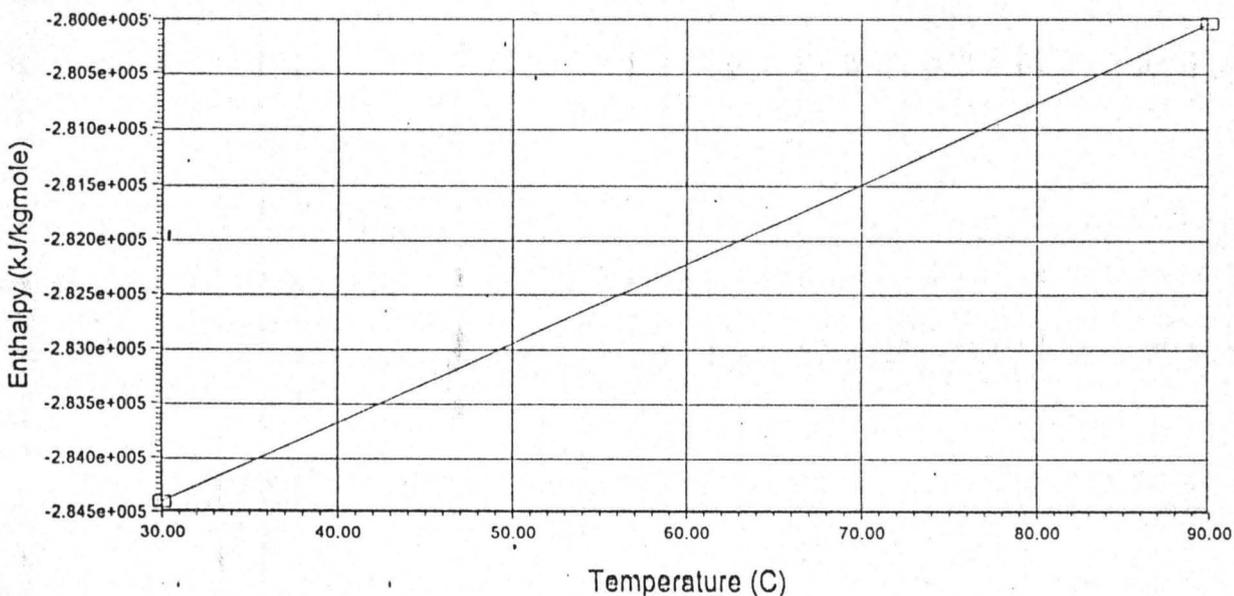
NOZZLE PARAMETERS

Elevation Relative to Ground Level				0.0000 m *
	CONCENTRATED NICKEL SOLUTION	CNS		
Height (m)	5.000e-002	5.000e-002		
Height (Base) (m)	0.0000	0.0000		
Height (Ground) (m)	0.0000	0.0000		

PERFORMANCE PROFILES

Zone	Pressure (kPa)	Temperature (C)	Vapour Fraction	Enthalpy (kJ/kgmole)
Inlet	90.00	90.00	0.0000	-280029.47
0	88.00	30.00	0.0000	-284409.78

PERFORMANCE PLOT





TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:25:42 2007

Cooler: COOLER 1 (continued)

PERFORMANCE TABLE

Temperature (C)	Pressure (kPa)	Heat Flow (kJ/h)	Enthalpy (kJ/kgmole)	Vapour Fraction
90.00	90.00	0.00	-280029.47	0.0000
30.00	88.00	-1234767.49	-284409.78	0.0000

DYNAMICS

Model Details: Supplied Duty

Zone	1 *	Delta P (kPa)	2.000 *
Volume (m3)	0.1000 *	Overall K (kg/hr/sqrt(kPa-kg/m3))	117.4 *
Duty (kJ/h)	1.235e+006		

Holdup Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000

Individual Zone Holdups: Zone 0

Delta P Specs and Duties

Zone	dP Value (kPa)	dP Option	Duty (kJ/h)
0	2.000	not specified	-6084

Zone Conductance Specifications

Zone	k (kg/hr/sqrt(kPa-kg/m3))	Specification
0	117.4	Disabled



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Thu Dec 21 17:08:03 2006

Air cooler: AIR DRIER

CONNECTIONS

Inlet Stream

STREAM NAME	FROM UNIT OPERATION
W.B.M	Tank WASHING UNIT.3

Outlet Stream

STREAM NAME	TO UNIT OPERATION
DRIED BASE METAL	Conversion Reactor ELECTROPLATING TANK

DESIGN PARAMETERS

Pressure Drop:	10.00 kPa *	UA:	-97.27 kJ/C-h
Inlet Air Temp:	25.00 C	Outlet Air Temp:	25.00 C
Configuration:	one tube row, one pass		

PERFORMANCE

Working Fluid Duty:	147.2 kJ/h	Correction Factor:	0.8111
UA:	-97.27 kJ/C-h	LMTD:	1.865 C *
Feed Temp:	25.43 C	Prod Temp:	30.00 C *
Air Feed Temp:	25.00 C	Air Prod Temp:	25.00 C
Volumetric Air Flow:	7.200e+005 ACT m ³ /h *	Mass Air Flow:	8.491e+005 kg/h *

User Variables

SIZING

Fan	Fan 0	Fan 1
Number of Fans	2 *	
Speed (rpm)	60.00	60.00
Speed (rpm)	60.00	60.00
Max Acceleration (rpm)	--	--
Design Speed (rpm)	60.00	60.00
Design airflow (ACT m ³ /h)	3.600e+005	3.600e+005
Current airflow (ACT m ³ /h)	3.600e+005	3.600e+005

NOZZLE PARAMETERS

Base Elevation Relative to Ground Level	0.4500 m *		
	W B M	DRIED BASE METAL	
Diameter (m)	5.000e-002	5.000e-002	
Elevation (Base) (m)	0.0000	0.0000	
Elevation (Ground) (m)	0.4500	0.4500	

NOTES



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\WR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Thu Dec 21 16:52:36 2006

Pump: PUMP.1

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
NaOH	

Outlet Stream

Stream Name	To Unit Operation
NaOH	Heater HEATER

Energy Stream

Stream Name	From Unit Operation
QP.1	

PARAMETERS

Adiabatic Efficiency (%):	75.00 *	Delta P:	10.00 kPa *	Duty:	6.964e-008 kW
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CURVES

Delta P:	10.00 kPa *	Duty:	6.964e-008		
Coefficient A:	0.0000 *	Coefficient B:	0.0000 *	Coefficient C:	0.0000
Parameter Preferences	Units for Delta P: m	Flow Basis	ActVolFlow	Units for Flow:	m3/h

User Variables

RATING

Characteristic Curves

Speed:		
Flow	Head	Efficiency (%)

NPSH

NPSH Required	--	NPSH Available	--	Enable NPSH Curves:	No
NPSH Curves					

Nozzle Parameters

Base Elevation Relative to Ground Level				0.0000 m *
		NaOH	NaOH	
Diameter	(m)	5.000e-002	5.000e-002	
Elevation (Base)	(m)	0.0000	0.0000	
Elevation (Ground)	(m)	0.0000	0.0000	

Inertia

Rotational inertia (kg-m2)	0.5000	Radius of gyration (m)	0.1000 *	Mass (kg)	50.00 *	Friction loss factor (rpm)	6.000 *
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Start Up

Design Flow Typical Operating Capacity	10.00 m3/h
--	------------

PERFORMANCE

Results

Total Head	--	Velocity Head	-1.548e-017 m
Pressure Head	0.6897 m	Delta P excluding Static Head Results	--

DYNAMICS

Dynamic Specifications

Duty	(m)	--	Not Active	Power	(kJ/h)	2.507e-004	Not Active
Adiabatic Efficiency	(rpm)	--	Not Active	Capacity	(m3/h)	1.560e-005	
Polytropic Efficiency	(%)	75.00 *	Active	Use Characteristic Curves			Not Active
Pressure Increase	(kPa)	10.00 *	Active	Pump is Acting as a Turbine			Not Active



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Thu Dec 21 16:52:36 2006

Pump: PUMP.1 (continued)

Holdup Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc
Unit Set: SI
Date/Time: Thu Dec 21 16:56:32 2006

Heater: HEATER

CONNECTIONS

Inlet Stream

STREAM NAME	FROM UNIT OPERATION
NaOH.	Pump PUMP 1

Outlet Stream

STREAM NAME	TO UNIT OPERATION
SODIUM HYDROXIDE	Tank DEGREASING UNIT

Energy Stream

STREAM NAME	FROM UNIT OPERATION
Q.HEATER1	

PARAMETERS

Pressure Drop:	0.0000 kPa	Duty:	6.319 kJ/h	Volume:	0.8000 m ³
Function:	Not Selected	Zones:	1		

User Variables

RATING

NOZZLE PARAMETERS

Base Elevation Relative to Ground Level			0.7620 m
	NaOH.	SODIUM HYDROXIDE	
Diameter (m)	5.000e-002	5.000e-002	
Elevation (Base) (m)	0.0000	0.0000	
Elevation (Ground) (m)	0.7620	0.7620	

DETAILED HEAT LOSS PARAMETERS

Overall Heat Loss (kJ/h)	--	Area (m ²)
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Temperature Profile

Fluid (C)	--	Inner Wall (C)	25.00	Middle Wall (C)	25.00	Outer Wall (C)	25.00	External (C)	25.00
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Conduction

	Metal	Insulation
Thickness (m)	1.000e-002	3.000e-002
Cp (kJ/kg-C)	0.4730	0.8200
Density (kg/m ³)	7801	520.0
Conductivity (W/m-K)	45.00	0.1500

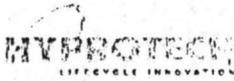
Convection

Outside U (kJ/h-m ² -C)	54.00	Inside U (kJ/h-m ² -C)	7200
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PERFORMANCE PROFILES

Zone	Pressure (kPa)	Temperature (C)	Vapour Fraction	Enthalpy (kJ/kgmole)
Inlet	120.00	25.00	0.0000	-182180.76
0	120.00	50.00	0.0000	-173089.36

PERFORMANCE PLOT



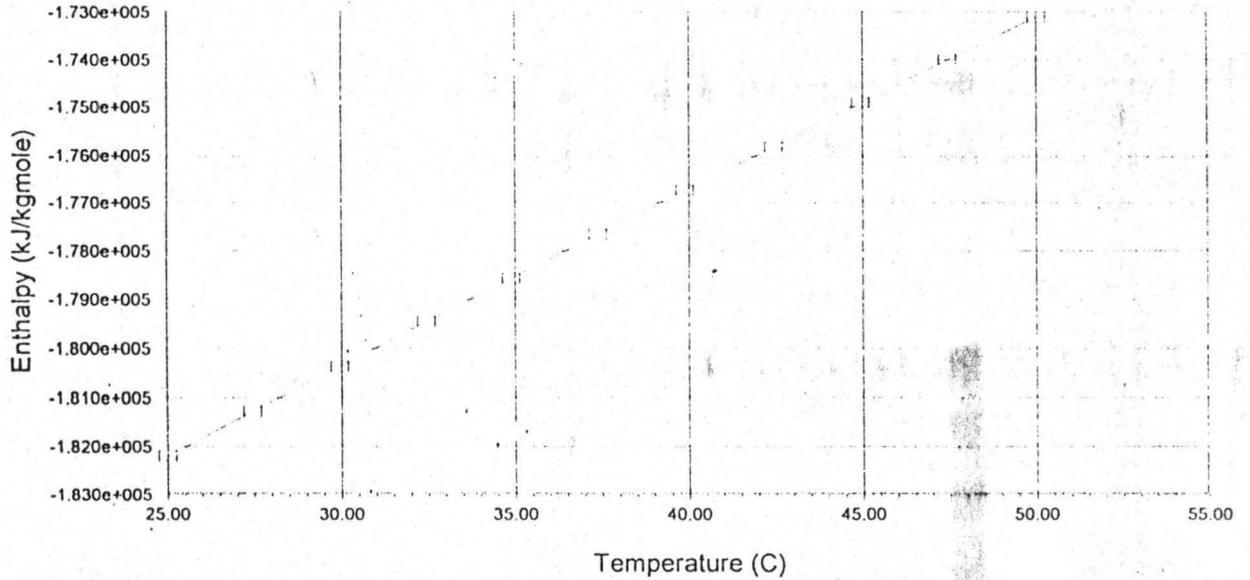
TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Thu Dec 21 16:56:32 2006

Heater: HEATER (continued)



PERFORMANCE TABLE

Temperature (C)	Pressure (kPa)	Heat Flow (kJ/h)	Enthalpy (kJ/kgmole)	Vapour Fraction
25.00	120.00	0.00	-182180.76	0.0000
27.47	120.00	0.63	-181271.62	0.0000
29.96	120.00	1.26	-180362.48	0.0000
32.44	120.00	1.90	-179453.34	0.0000
34.93	120.00	2.53	-178544.20	0.0000
37.43	120.00	3.16	-177635.06	0.0000
39.93	120.00	3.79	-176725.92	0.0000
42.44	120.00	4.42	-175816.78	0.0000
44.96	120.00	5.06	-174907.64	0.0000
47.48	120.00	5.69	-173998.50	0.0000
50.00	120.00	6.32	-173089.36	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:25:04 2007

Pump: P-100

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
CNS	Cooler COOLER 1

Outlet Stream

Stream Name	To Unit Operation
P.CNS	Recycle RCY-1

Energy Stream

Stream Name	From Unit Operation
Q.P	

PARAMETERS

Adiabatic Efficiency (%): 95.00 * Delta P: 10.00 kPa * Duty: 1.480e-002

CURVES

Delta P: 10.00 kPa * Duty: 1.480e-002

Coefficient A: 0.0000 * Coefficient B: 0.0000 * Coefficient C: 0.00

Parameter Preferences Units for Delta P: m Flow Basis ActVolFlow Units for Flow: m

User Variables

RATING

Characteristic Curves

Flow	Head	Efficiency (%)
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NPSH

NPSH Required -- NPSH Available -- Enable NPSH Curves: No

NPSH Curves

Nozzle Parameters

Base Elevation Relative to Ground Level 0.0000 m

	CNS	P.CNS
Diameter (m)	5.000e-002	5.000e-002
Elevation (Base) (m)	0.0000	0.0000
Elevation (Ground) (m)	0.0000	0.0000

Inertia

Rotational inertia (kg-m2) 0.5000 Radius of gyration (m) 0.1000 * Mass (kg) 50.00 * Friction loss factor (rpm) 6.000 *

Start Up

Design Flow Typical Operating Capacity 10.00 m3/h

PERFORMANCE

Results

Total Head 1.016 m * Velocity Head -1.457e-007 m
Pressure Head 1.016 m Delta P excluding Static Head Results --

DYNAMICS

Dynamic Specifications

Duty (m)	1.016	Not Active	Power (kJ/h)	53.27	Not Active
Adiabatic Efficiency (rpm)	--	Not Active	Capacity (m3/h)	5.089	
Polytropic Efficiency (%)	95.00	Active	Use Characteristic Curves		Not Active



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR, BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:25:04 2007

Pump: P-100 (continued)

Pressure Increase (kPa)	10.00	Active	Pump is Acting as a Turbine	Not Active
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Holdup Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc
Unit Set: SI
Date/Time: Fri Feb 09 09:10:03 2007

Vessel Sizing: DEGREASING UNIT

Vertical Separator Properties: CONCENTRATOR

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	300.0 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7396 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7068 *

Costing Results

Base Cost (\$US)	7476	Associated Cost (\$US)	1658	Total Cost (FOB \$US)	9134
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Sizing Results

Diameter (m)	0.6096	Liquid Residence Time (seconds)	300.0
Total Length (m)	3.353	Liquid Surge Height (m)	1.567
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	1.000	Liquid Res. Time at LLSD (seconds)	80.44
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	1.024	Ellipsoidal Head (m)	0.1524
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.786



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:26:38 2007

Separator: CONCENTRATOR

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
P	Heater PRE-HEATER

Outlet Stream

Stream Name	To Unit Operation
WATER VAPOUR	
CONCENTRATED NICKEL SOLUTION	Cooler: COOLER 1

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	0.1223 m ³	Level SP:	50.00 %	Liquid Volume:	6.116e-002 m ³
Vessel Pressure:	90.00 kPa	Pressure Drop:	0.0000 kPa *	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heating

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot:	No
Volume 0.1223 m ³	Diameter 0.3048 m *	Height	1.676 *

Nozzles

Base Elevation Relative to Ground Level	0.0000 m *	Diameter	0.3048 m *	Height	1.676 *
		P	WATER VAPOUR	CONCENTRATED NICKEL SOLUTION	
Diameter (m)	8.382e-002		8.382e-002	8.382e-002	
Elevation (Base) (m)	0.8382		1.676	0.0000	
Elevation (Ground) (m)	0.8382		1.676	0.0000	
Elevation (% of Height) (%)	50.00		100.00	0.00	

Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low

Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level

Options

PV Work Term Contribution (%)	100.00 *
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m ³)	0.1223	Level Calculator	Vertical cylinder
Vessel Diameter (m)	0.3048 *	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	1.676 *	Feed Delta P (kPa)	0.0000 *
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	90.00

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m ³)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:26:38 2007

Separator: CONCENTRATOR (continued)

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:22:05 2007

Vessel Sizing: Vessel Sizing-11

Vertical Separator Properties: DRAGOUT TANK.6

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	3537 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7396 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7068 *

Costing Results

Base Cost (\$US)	5205	Associated Cost (\$US)	608.5	Total Cost (FOB \$US)	5814
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Sizing Results

Diameter (m)	0.3048	Liquid Residence Time (seconds)	3537
Total Length (m)	1.676	Liquid Surge Height (m)	0.4572
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	1.000	Liquid Res. Time at LLSD (seconds)	3537
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.5334	Ellipsoidal Head (m)	7.620e-002
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.219



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\WR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:24:03 2007

Tank: DRAGOUT TANK 6

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
WCM.2	Tank DRAGOUT TANK.5
WATER.10	Tee TEE-100

Outlet Stream

Stream Name	To Unit Operation
WCM.3	
L.SOLN.4	Mixer: CHROME SOLUTION MIXER

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	--	Level SP:	50.00 %	Liquid Volume:	
Vessel Pressure:	90.00 kPa	Pressure Drop:	0.0000 kPa	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	

User Variables

RATING

Sizing

Cylinder	--	Vertical	Separator has a Boot:	No
Volume	--	Diameter	Height	

Nozzles

Base Elevation Relative to Ground Level	0.0000 m	Diameter	--	Height
		WCM.2	WATER.10	WCM.3
Diameter (m)		5.000e-002	5.000e-002	5.000e-002
Elevation (Base) (m)		0.0000	0.0000	0.0000
Elevation (Ground) (m)		0.0000	0.0000	0.0000
Elevation (% of Height) (%)		--	--	--
		L.SOLN.4		
Diameter (m)		5.000e-002		
Elevation (Base) (m)		0.0000		
Elevation (Ground) (m)		0.0000		
Elevation (% of Height) (%)		--		

Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options

PV Work Term Contribution (%)	100.00
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	--	Level Calculator	Vertical cylinder
Vessel Diameter (m)	--	Fraction Calculator	Use levels and nozzle
Vessel Height (m)	--	Feed Delta P (kPa)	0.0000
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	90.00

Holdup: Vessel Levels



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:24:03 2007

Tank: DRAGOUT TANK 6 (continued)

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:21:23 2007

Vessel Sizing: Vessel Sizing-10

Vertical Separator Properties: DRAGOUT TANK.5

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	1.047e+004 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1: 0.4000 *	A4: 0.8116 *	A5: 8.600 *	A8: 1017 *	A6: -0.2165 *	A9: 0.7396 *	A7: 4.580e-002 *	A10: 0.7068 *
A2: 2.000 *							
A3: 0.2000 *							

Costing Results

Base Cost (\$US)	5205	Associated Cost (\$US)	608.5	Total Cost (FOB \$US)	5813.5
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Sizing Results

Diameter (m)	0.3048	Liquid Residence Time (seconds)	1.047e+004
Total Length (m)	1.676	Liquid Surge Height (m)	0.4572
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	1.000	Liquid Res. Time at LLSD (seconds)	1.047e+004
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.5334	Ellipsoidal Head (m)	7.620e-002
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.211



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:20:30 2007

Vessel Sizing: Vessel Sizing-9

Vertical Separator Properties: DRAGOUT TANK.4

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq Res Time (seconds)	1.042e+004 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1: 0.4000 *	A4: 0.8116 *	A5: 8.600 *	A8: 1017 *				
A2: 2.000 *		A6: -0.2165 *	A9: 0.7396 *				
A3: 0.2000 *		A7: 4.580e-002 *	A10: 0.7068 *				

Costing Results

Base Cost (\$US)	5205	Associated Cost (\$US)	608.5	Total Cost (FOB \$US)	5814
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Sizing Results

Diameter (m)	0.3048	Liquid Residence Time (seconds)	1.042e+004
Total Length (m)	1.676	Liquid Surge Height (m)	0.4572
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	1.000	Liquid Res. Time at LLSD (seconds)	1.042e+004
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.5334	Ellipsoidal Head (m)	7.620e-002
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.219



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc
 Unit Set: SI
 Date/Time: Wed Feb 07 12:21:49 2007

Tank: DRAGOUT TANK.4

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
CHROMED METAL	Conversion Reactor CHROME PLATING TANK
WATER.8	Tee TEE-100

Outlet Stream

Stream Name	To Unit Operation
WCM.1	Tank: DRAGOUT TANK.5
L.SOLN.2	Mixer: CHROME SOLUTION MIXER

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume: 1.216 m3	Level SP: 50.00 %	Liquid Volume: 0.6081 m3
Vessel Pressure: 90.00 kPa	Pressure Drop: 0.0000 kPa *	Duty: 0.0000 kJ/h
Heat Transfer Mode: Heating		

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot: No
Volume: 1.216 m3	Diameter: 0.7620 m *	Height: 2.667 *

Nozzles

Base Elevation Relative to Ground Level: 0.0000 m *	Diameter: 0.7620 m *	Height: 2.667 *	
	CHROMED METAL	WATER.8	WCM.1
Diameter (m)	0.1334	0.1334	0.1334
Elevation (Base) (m)	1.333	1.333	2.667
Elevation (Ground) (m)	1.333	1.333	2.667
Elevation (% of Height) (%)	50.00	50.00	100.00
	L.SOLN.2		
Diameter (m)	0.1334		
Elevation (Base) (m)	0.0000		
Elevation (Ground) (m)	0.0000		
Elevation (% of Height) (%)	0.00		

Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options

PV Work Term Contribution (%)	100.00 *
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	1.216	Level Calculator	Vertical cylinder
Vessel Diameter (m)	0.7620 *	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	2.667 *	Feed Delta P (kPa)	0.0000 *
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	90.00

Holdup: Vessel Levels



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:21:49 2007

Tank: DRAGOUT TANK.4 (continued)

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:13:17 2007

Vessel Sizing: Vessel Sizing-3

Vertical Separator Properties: DRAGOUT TANK.3

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	3537 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7395 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7067 *

Costing Results

Base Cost (\$US)	5205	Associated Cost (\$US)	608.5	Total Cost (FOB \$US)	5813.5
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Sizing Results

Diameter (m)	0.3048	Liquid Residence Time (seconds)	3537
Total Length (m)	1.676	Liquid Surge Height (m)	0.4572
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	1.000	Liquid Res. Time at LLSD (seconds)	3537
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.5334	Ellipsoidal Head (m)	7.620e-002
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.2100



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:19:29 2007

Tank: DRAGOUT TANK.3

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
P.IRON. WATER.7	Tank: DRAGOUT TANK.2 Tee: WATER SPLITTER

Outlet Stream

Stream Name	To Unit Operation
PLATED METAL WASH WATER.3	Conversion Reactor: CHROME PLATING TANK Mixer: MIXER.2

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	0.9800 m3 *	Level SP:	50.00 %	Liquid Volume:	0.4900 m3
Vessel Pressure:	90.00 kPa	Pressure Drop:	0.0000 kPa *	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heating

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot:	No
Volume: 0.9800 m3 *	Diameter: 0.9405 m	Height:	1.411

Nozzles

Base Elevation Relative to Ground Level	*0.0000 m *	Diameter	0.9405 m	Height	1.411
		P.IRON.	WATER.7	PLATED METAL	
Diameter (m)		7.054e-002	7.054e-002	7.054e-002	
Elevation (Base) (m)		0.7054	0.7054	1.411	
Elevation (Ground) (m)		0.7054	0.7054	1.411	
Elevation (% of Height) (%)		50.00	50.00	100.00	
		WASH WATER.3			
Diameter (m)		7.054e-002			
Elevation (Base) (m)		0.0000			
Elevation (Ground) (m)		0.0000			
Elevation (% of Height) (%)		0.00			

Detailed Heat Loss Parameters

Overall Heat Loss (kJ/h)	0.0000	Area (m2)	4.168
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Temperature Profile

Fluid (C)	0.0000 *	Inner Wall (C)	25.00	Outer Vessel (C)	25.00	Outer Insulation (C)	25.00	Ambient (C)	25.00
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Conduction

	Metal	Insulation
Thickness (m)	1.000e-002	3.000e-002
Cp (kJ/kg-C)	0.4730	0.8200
Density (kg/m3)	7801	520.0
Conductivity (W/m-K)	45.00	0.1500

Convection

Inside Vap Phase U (kJ/h-m2-C)	7200	Inside Liq Phase U (kJ/h-m2-C)	--	Outside U (kJ/h-m2-C)	54.00
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Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:19:29 2007

Tank: DRAGOUT TANK.3 (continued)

Options

PV Work Term Contribution	(%)	100.00 *	
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume	(m3)	0.9800 *	Level Calculator	Vertical cylinder
Vessel Diameter	(m)	0.9405	Fraction Calculator	Use levels and nozzles
Vessel Height	(m)	1.411	Feed Delta P	(kPa) 0.0000 *
Liquid Level Percent	(%)	50.00	Vessel Pressure	(kPa) 90.00

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:12:35 2007

Vessel Sizing: Vessel Sizing-2

Vertical Separator Properties: DRAGOUT TANK.2

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	1.047e+004 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m ³)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7396 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7068 *

Costing Results

Base Cost (\$US)	5205	Associated Cost (\$US)	608.5	Total Cost (FOB \$US)	5814
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Sizing Results

Diameter (m)	0.3048	Liquid Residence Time (seconds)	1.047e+004
Total Length (m)	1.676	Liquid Surge Height (m)	0.4572
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	1.000	Liquid Res. Time at LLSD (seconds)	1.047e+004
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.5334	Ellipsoidal Head (m)	7.620e-002
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.219



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:18:54 2007

Tank: DRAGOUT TANK.2

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
P.IRON	Tank DRAGOUT TANK.1
WATER.6	Tee WATER SPLITTER

Outlet Stream

Stream Name	To Unit Operation
P.IRON.	Tank DRAGOUT TANK.3
WASH WATER.2	Mixer: MIXER.2

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	0.9800 m3 *	Level SP:	50.00 %	Liquid Volume:	0.4900 m3
Vessel Pressure:	90.00 kPa	Pressure Drop:	0.0000 kPa *	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heat

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Bool:	No
Volume	0.9800 m3 *	Diameter	0.9405 m
		Height	1.411

Nozzles

Base Elevation Relative to Ground Level	0.0000 m *	Diameter	0.9405 m	Height	1.411
		P.IRON	WATER.6	P.IRON.	
Diameter (m)		7.054e-002	7.054e-002	7.054e-002	
Elevation (Base) (m)		0.7054	0.7054	1.411	
Elevation (Ground) (m)		0.7054	0.7054	1.411	
Elevation (% of Height) (%)		50.00	50.00	100.00	
		WASH WATER.2			
Diameter (m)		7.054e-002			
Elevation (Base) (m)		0.0000			
Elevation (Ground) (m)		0.0000			
Elevation (% of Height) (%)		0.00			

Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options

PV Work Term Contribution (%)	100.00 *
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	0.9800 *	Level Calculator	Vertical cylinder
Vessel Diameter (m)	0.9405	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	1.411	Feed Delta P (kPa)	0.0000 *
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	90.00

Holdup: Vessel Levels



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:18:54 2007

Tank: DRAGOUT TANK.2 (continued)

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:11:46 2007

Vessel Sizing: Vessel Sizing-1

Vertical Separator Properties: DRAGOUT TANK.1

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	1.042e+004 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7396 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7068 *

Costing Results

Base Cost (\$US)	5205	Associated Cost (\$US)	608.5	Total Cost (FOB \$US)	5814
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Sizing Results

Diameter (m)	0.3048	Liquid Residence Time (seconds)	1.042e+004
Total Length (m)	1.676	Liquid Surge Height (m)	0.4572
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	1.000	Liquid Res. Time at LLSD (seconds)	1.042e+004
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.5334	Ellipsoidal Head (m)	7.620e-002
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.219



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc
Unit Set: SI
Date/Time: Wed Feb 07 12:18:07 2007

Tank: DRAGOUT TANK.1

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
PLATED IRON	Conversion Reactor
WATER.5	Tee

Outlet Stream

Stream Name	To Unit Operation
P.IRON	Tank: DRAGOUT TANK.2
WASH WATER.1	Mixer: MIXER.2

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	0.9800 m3	Level SP:	50.00 %	Liquid Volume:	0.4900 m3
Vessel Pressure:	90.00 kPa	Pressure Drop:	0.0000 kPa	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heating

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot:	No
Volume	0.9800 m3	Diameter	0.9405 m
		Height	1.411

Nozzles

Base Elevation Relative to Ground Level	0.0000 m	Diameter	0.9405 m	Height	1.411
		PLATED IRON	WATER.5	P.IRON	
Diameter	(m)	7.054e-002	7.054e-002	7.054e-002	
Elevation (Base)	(m)	0.7054	0.7054	1.411	
Elevation (Ground)	(m)	0.7054	0.7054	1.411	
Elevation (% of Height)	(%)	50.00	50.00	100.00	
		WASH WATER.1			
Diameter	(m)	7.054e-002			
Elevation (Base)	(m)	0.0000			
Elevation (Ground)	(m)	0.0000			
Elevation (% of Height)	(%)	0.00			

Detailed Heat Loss Parameters

Overall Heat Loss (kJ/h)	0.0000	Area (m2)	4.168
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Temperature Profile

Liquid (C)	0.0000	Inner Wall (C)	25.00	Outer Vessel (C)	25.00	Outer Insulation (C)	25.00	Ambient (C)	25.00
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Conduction

	Metal	Insulation
Thickness (m)	1.000e-002	3.000e-002
Thermal Property (kJ/kg-C)	0.4730	0.8200
Density (kg/m3)	7801	5200
Conductivity (W/m-K)	45.00	0.1500

Convection

Inside Vap Phase U (kJ/h-m2-C)	7200	Inside Liq Phase U (kJ/h-m2-C)	--	Outside U (kJ/h-m2-C)	54.00
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Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\WR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:18:07 2007

Tank: DRAGOUT TANK.1 (continued)

Options

PV Work Term Contribution	(%)	100.00 *	
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume	(m3)	0.9800 *	Level Calculator	Vertical cylinder
Vessel Diameter	(m)	0.9405	Fraction Calculator	Use levels and nozzles
Vessel Height	(m)	1.411	Feed Delta P	(kPa) 0.0000 *
Liquid Level Percent	(%)	50.00	Vessel Pressure	(kPa) 90.00

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc
Unit Set: SI
Date/Time: Fri Feb 09 09:14:00 2007

Vessel Sizing: Vessel Sizing-4

Vertical Separator Properties: ELECTROPLATING TANK

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	300.0 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1: 0.4000 *	A4: 0.8116 *	A5: 8.600 *	A8: 1017 *	A6: -0.2165 *	A9: 0.7396 *	A7: 4.580e-002 *	A10: 0.7068 *
A2: 2.000 *							
A3: 0.2000 *							

Costing Results

Base Cost (\$US)	7476	Associated Cost (\$US)	1658	Total Cost (FOB \$US)	9134
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Sizing Results

Diameter (m)	0.6096	Liquid Residence Time (seconds)	300.0
Total Length (m)	3.353	Liquid Surge Height (m)	1.521
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	0.6644	Liquid Res. Time at LLSD (seconds)	82.94
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	1.070	Ellipsoidal Head (m)	0.1524
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.832



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:15:33 2007

Conversion Reactor: ELECTROPLATING TANK

CONNECTIONS

Inlet Stream Connections

Stream Name	From Unit Operation
RCY	Recycle RCY-1
DRIED BASE METAL	Air cooler AIR DRIER

Outlet Stream Connections

Stream Name	To Unit Operation
PLATED IRON	Tank: DRAGOUT TANK.1
NICKEL SOLUTION	Mixer: MIXER.2

Energy Stream Connections

Stream Name	From Unit Operation
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PARAMETERS

Physical Parameters		Optional Heat Transfer:	
Delta P	Vessel Volume	Duty	Energy Stream
0.0000 kPa	2.583 m ³	0.0000 kJ/h	

User Variables

RATING

Sizing

Cylinder	Vertical	Reactor has a Boot:
Volume: 2.583 m ³	Diameter: 1.299 m	No
	Height: 1.949 m	

Nozzles

Base Elevation Relative to Ground Level	Diameter	Height
0.0000 m	1.299 m	1.949 m
RCY		
Diameter (m)	9.743e-002	
Elevation (Base) (m)	0.9743	
Elevation (Ground) (m)	0.9743	
Elevation (% of Height) (%)	50.00	
DRIED BASE METAL		
Diameter (m)	9.743e-002	
Elevation (Base) (m)	0.0000	
Elevation (Ground) (m)	0.0000	
Elevation (% of Height) (%)	0.00	
PLATED IRON		
Diameter (m)	9.743e-002	
Elevation (Base) (m)	0.9743	
Elevation (Ground) (m)	0.9743	
Elevation (% of Height) (%)	50.00	
NICKEL SOLUTION		
Diameter (m)	9.743e-002	
Elevation (Base) (m)	0.0000	
Elevation (Ground) (m)	0.0000	
Elevation (% of Height) (%)	0.00	

Detailed Heat Loss Parameters

Overall Heat Loss (kJ/h)	0.0000	Area (m ²)	7.953
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Temperature Profile

Fluid (C)	0.0000	Inner Wall (C)	25.00	Outer Vessel (C)	25.00	Outer Insulation (C)	25.00	Ambient (C)	25.00
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Conduction

	Metal	Insulation
Thickness (m)	1.000e-002	3.000e-002
Cp (kJ/kg-C)	0.4730	0.8200
Density (kg/m ³)	7801	520.0
Thermal Conductivity (W/m-K)	45.000	0.1500

Convection

Outside U (kJ/h-m ² -C)	54.00	Inside U (kJ/h-m ² -C)	7200
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m ³)	2.583	Level Calculator	Vertical cylinder
Vessel Diameter (m)	1.299	Fraction Calculator	Use levels and nozzles



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc

Unit Set: SI

Date/Time: Wed Feb 07 12:15:33 2007

Conversion Reactor: ELECTROPLATING TANK (continued)

Vessel Height	(m)	1.949	Feed Delta P	(kPa)	0.0000 *
Liquid Level Percent	(%)	50.00	Vessel Pressure	(kPa)	90.00

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:10:43 2007

Vessel Sizing: DEGREASING UNIT

Vertical Separator Properties: DEGREASING UNIT

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	1.360e+006 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7396 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7058 *

Costing Results

Base Cost (\$US)	5205	Associated Cost (\$US)	608.5	Total Cost (FOB \$US)	5813.5
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Sizing Results

Diameter (m)	0.3048	Liquid Residence Time (seconds)	1.360e+006
Total Length (m)	1.676	Liquid Surge Height (m)	0.4572
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	0.8129	Liquid Res. Time at LLSD (seconds)	1.360e+006
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.5334	Ellipsoidal Head (m)	7.620e-002
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.219



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc
Unit Set: SI
Date/Time: Thu Dec 21 16:58:45 2006

Tank: DEGREASING UNIT

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
WASHED B.M.	Tank WASHING UNIT 1
SODIUM HYDROXIDE	Heater HEATER

Outlet Stream

Stream Name	To Unit Operation
DEGREASED B.M	Tank: WASHING UNIT2
SOAP + WATER	Mixer: MIXER

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	--	Level SP:	50.00 %	Liquid Volume:	--
Vessel Pressure:	100.0 kPa	Pressure Drop:	0.0000 kPa	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heater

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot:	No
Volume --	Diameter --	Height	--

Nozzles

Base Elevation Relative to Ground Level	0.0000 m	Diameter	--	Height	--
		WASHED B.M.	SODIUM HYDROXIDE	DEGREASED B.M	
Diameter (m)		5.000e-002	5.000e-002	5.000e-002	
Elevation (Base) (m)		0.0000	0.0000	0.0000	
Elevation (Ground) (m)		0.0000	0.0000	0.0000	
Elevation (% of Height) (%)		--	--	--	
		SOAP + WATER			
Diameter (m)		5.000e-002			
Elevation (Base) (m)		0.0000			
Elevation (Ground) (m)		0.0000			
Elevation (% of Height) (%)		--			

Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options

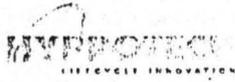
PV Work Term Contribution (%)	100.00
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	--	Level Calculator	Vertical cylinder
Vessel Diameter (m)	--	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	--	Feed Delta P (kPa)	0.0000
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	100.0

Holdup: Vessel Levels



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\WR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Thu Dec 21 16:58:45 2006

Tank: DEGREASING UNIT (continued)

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc
Unit Set: SI
Date/Time: Fri Feb 09 09:15:38 2007

Vessel Sizing: Vessel Sizing-7

Vertical Separator Properties: WASHING UNIT.3

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	300.0 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1: 0.4000 *	A4: 0.8116 *	A5: 8.600 *	A8: 1017 *	A2: 2.000 *	A6: -0.2165 *	A9: 0.7396 *	
A3: 0.2000 *		A7: 4.580e-002 *	A10: 0.7068 *				

Costing Results

Base Cost (\$US)	6297	Associated Cost (\$US)	1094	Total Cost (FOB \$US)	7391
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Sizing Results

Diameter (m)	0.4572	Liquid Residence Time (seconds)	300.0
Total Length (m)	2.515	Liquid Surge Height (m)	0.8147
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	0.5607	Liquid Res. Time at LLSD (seconds)	161.9
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.9760	Ellipsoidal Head (m)	0.1143
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.700



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.3.hsc
Unit Set: SI
Date/Time: Thu Dec 21 17:04:59 2006

Tank: WASHING UNIT.3

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
WASHED.BM	Tank WASHING UNIT2
P.WATER.4	Pump PUMP.4

Outlet Stream

Stream Name	To Unit Operation
W.B.M	Air cooler: AIR DRIER
WATER+LIGTH SOAP	Mixer: MIXER

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	2.034 m ³ *	Level SP:	50.00 %	Liquid Volume:	1.017 m ³
Vessel Pressure:	100.0 kPa	Pressure Drop:	0.0000 kPa *	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heat

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot:	No
Volume	2.034 m ³ *	Diameter	1.200 m
		Height	1.799

Nozzles

Base Elevation Relative to Ground Level	0.0000 m *	Diameter	1.200 m	Height	1.799
		WASHED BM	P WATER.4		W.B.M
Diameter (m)		8.997e-002	8.997e-002		8.997e-002
Elevation (Base) (m)		0.8997	0.8997		1.799
Elevation (Ground) (m)		0.8997	0.8997		1.799
Elevation (% of Height) (%)		50.00	50.00		100.00
		WATER+LIGTH SOAP			
Diameter (m)		8.997e-002			
Elevation (Base) (m)		0.0000			
Elevation (Ground) (m)		0.0000			
Elevation (% of Height) (%)		0.00			

Detailed Heat Loss Parameters

Overall Heat Loss (kJ/h)	0.0000	Area (m ²)	6.782
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Temperature Profile

Fluid (C)	0.0000 *	Inner Wall (C)	25.00	Outer Vessel (C)	25.00	Outer Insulation (C)	25.00	Ambient (C)	25.00
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Conduction

	Metal	Insulation
Thickness (m)	1.000e-002	3.000e-002
Cp (kJ/kg-C)	0.4730	0.8200
Density (kg/m ³)	7801	520.0
Conductivity (W/m-K)	45.00	0.1500

Convection

Inside Vap Phase U (kJ/h-m ² -C)	7200	Inside Liq Phase U (kJ/h-m ² -C)	-	Outside U (kJ/h-m ² -C)	54.00
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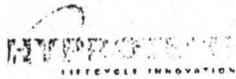
Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\WR. BABALOLA.3.hsc
 Unit Set: SI
 Date/Time: Thu Dec 21 17:04:59 2006

Tank: WASHING UNIT.3 (continued)

PV Work Term Contribution (%)	100.00
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DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	2.034	Level Calculator	Vertical cylinder
Vessel Diameter (m)	1.200	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	1.799	Feed Delta P (kPa)	0.0000
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	100.0

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:14:39 2007

Vessel Sizing: Vessel Sizing-5

Vertical Separator Properties: WASHING UNIT2

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	564.7 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7396 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7068 *

Costing Results

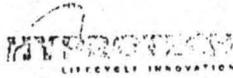
Base Cost (\$US)	5205	Associated Cost (\$US)	608.5	Total Cost (FOB \$US)	5814
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Sizing Results

Diameter (m)	0.3048	Liquid Residence Time (seconds)	564.7
Total Length (m)	1.676	Liquid Surge Height (m)	0.4572
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	1.000	Liquid Res. Time at LLSD (seconds)	564.7
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.5334	Ellipsoidal Head (m)	7.620e-002
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.219



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\WR. BABALOLA.2.hsc
Unit Set: SI
Date/Time: Thu Dec 21 17:01:19 2006

Tank: WASHING UNIT2

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
DEGREASED.B.M	Tank: DEGREASING UNIT
P.WATER.3	Pump: PUMP.3

Outlet Stream

Stream Name	To Unit Operation
WASHED.BM	Tank: WASHING UNIT.3
WATER+SOAP	Mixer: MIXER

Energy Stream

Stream Name	From Unit Operation

PARAMETERS

Vessel Volume:	2.050 m3 *	Level SP:	85.00 % *	Liquid Volume:	1.742 m3
Vessel Pressure:	100.0 kPa	Pressure Drop:	0.0000 kPa *	Duty:	0.0000 kJ/h
				Heat Transfer Mode:	Heating

User Variables

RATING

Sizing

Cylinder	Horizontal	Separator has a Boot:	No
Volume: 2.050 m3 *	Diameter: 1.203 m	Length:	1.804

Nozzles

Base Elevation Relative to Ground Level	0.0000 m *	Diameter	1.203 m	Length	1.804
		DEGREASED.B.M	P.WATER.3	WASHED.BM	
Diameter (m)		6.014e-002	6.014e-002	6.014e-002	
Elevation (Base) (m)		0.6014	0.6014	1.203	
Elevation (Ground) (m)		0.6014	0.6014	1.203	
Elevation (% of Height) (%)		50.00	50.00	100.00	
		WATER+SOAP			
Diameter (m)		6.014e-002			
Elevation (Base) (m)		0.0000			
Elevation (Ground) (m)		0.0000			
Elevation (% of Height) (%)		0.00			

Detailed Heat Loss Parameters

Overall Heat Loss (kJ/h)	0.0000	Area (m2) *	6.817
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Temperature Profile

Fluid (C)	0.0000 *	Inner Wall (C)	25.00	Outer Vessel (C)	25.00	Outer Insulation (C)	25.00	Ambient (C)	25.00
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Conduction

	Metal	Insulation
Thickness (m)	1.000e-002	3.000e-002
Cp (kJ/kg-C)	0.4730	0.8200
Density (kg/m3)	7801	520.0
Conductivity (W/m-K)	45.00	0.1500

Convection

Inside Vap Phase U (kJ/h-m2-C)	7200	Inside Liq Phase U (kJ/h-m2-C)	--	Outside U (kJ/h-m2-C)	54.00
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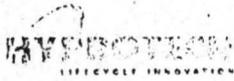
Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Thu Dec 21 17:01:19 2006

Tank: WASHING UNIT2 (continued)

PV Work Term Contribution (%) 100.00 *

DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	2.050 *	Level Calculator	Horizontal cylinder
Vessel Diameter (m)	1.203	Fraction Calculator	Use levels and nozzles
Vessel Length (m)	1.804	Feed Delta P (kPa)	0.0000 *
Liquid Level Percent (%)	85.00 *	Vessel Pressure (kPa)	100.0

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	-	-	0.0000
Liquid	-	-	0.0000
Aqueous	-	-	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:15:08 2007

Vessel Sizing: Vessel Sizing-6

Vertical Separator Properties: WASHING UNIT.1

L/D Ratio	5.000 *	Demister to Top (m)	0.3048 *
Liq. Res. Time (seconds)	300.0 *		

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1: 0.4000 *	A4: 0.8116 *	A5: 8.600 *	A8: 1017 *				
A2: 2.000 *		A6: -0.2165 *	A9: 0.7396 *				
A3: 0.2000 *		A7: 4.580e-002 *	A10: 0.7068 *				

Costing Results

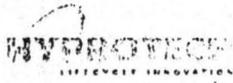
Base Cost (\$US)	7476	Associated Cost (\$US)	1658	Total Cost (FOB \$US)	9134
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Sizing Results

Diameter (m)	0.6096	Liquid Residence Time (seconds)	300.0
Total Length (m)	3.353	Liquid Surge Height (m)	1.719
L/D Ratio	5.000	LLSD (m)	0.4572
Max. Allowable Vapour Velocity (m/s)	0.8989	Liquid Res. Time at LLSD (seconds)	73.10
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	0.8721	Ellipsoidal Head (m)	0.1524
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	1.634



TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Thu Dec 21 16:50:06 2006

Tank: WASHING UNIT.1

CONNECTIONS

Inlet Stream

Stream Name	From Unit Operation
BASE.METAL	Tank ACID PICKLING UNIT
P.WATER.2	Pump PUMP.2

Outlet Stream

Stream Name	To Unit Operation
WASHED.B.M.	Tank DEGREASING UNIT
FeCl3 +WATER	

Energy Stream

Stream Name	From Unit Operation
QEEE	

PARAMETERS

Vessel Volume:	2.024 m ³	Level SP:	50.00 %	Liquid Volume:	1.012
Vessel Pressure:	100.0 kPa	Pressure Drop:	0.0000 kPa	Duty:	4.962e+005 kJ/h
				Heat Transfer Mode:	Heating

User Variables

RATING

Sizing

Cylinder	Vertical	Separator has a Boot:	No
Volume	2.024 m ³	Diameter	1.198 m
		Height	1.797

Nozzles

Base Elevation Relative to Ground Level	0.0000 m	Diameter	1.198 m	Height	1.797
		BASE METAL	P. WATER.2	WASHED B.M	
Diameter (m)		8.983e-002	8.983e-002	8.983e-002	
Elevation (Base) (m)		0.8983	0.8983	1.797	
Elevation (Ground) (m)		0.8983	0.8983	1.797	
Elevation (% of Height) (%)		50.00	50.00	100.00	
		FeCl3 +WATER			
Diameter (m)		8.983e-002			
Elevation (Base) (m)		0.0000			
Elevation (Ground) (m)		0.0000			
Elevation (% of Height) (%)		0.00			

Detailed Heat Loss Parameters

Overall Heat Loss (kJ/h)	0.0000	Area (m ²)	6.759
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Temperature Profile

Fluid (C)	0.0000	Inner Wall (C)	25.00	Outer Vessel (C)	25.00	Outer Insulation (C)	25.00	Ambient (C)	25.00
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Conduction

	Metal	Insulation
Thickness (m)	1.000e-002	3.000e-002
Cp (kJ/kg-C)	0.4730	0.8200
Density (kg/m ³)	7801	520.0
Conductivity (W/m-K)	45.00	0.1500

Convection

Inside Vap Phase U (kJ/h-m ² -C)	7200	Inside Liq Phase U (kJ/h-m ² -C)	--	Outside U (kJ/h-m ² -C)	54.00
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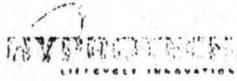
Level Taps: Level Tap Specification

Level Tap	PV High	PV Low	OP High	OP Low
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Level Taps: Calculated Level Tap Values

Level Tap	Liquid Level	Aqueous Level
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Options



TEAM LND
 Calgary, Alberta
 CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.2.hsc

Unit Set: SI

Date/Time: Thu Dec 21 16:50:06 2006

Tank: WASHING UNIT.1 (continued)

PV Work Term Contribution (%) 100.00 *

DYNAMICS

Vessel Parameters: Initialize from Product

Vessel Volume (m3)	2.024 *	Level Calculator	Vertical cylinder
Vessel Diameter (m)	1.198	Fraction Calculator	Use levels and nozzles
Vessel Height (m)	1.797	Feed Delta P (kPa)	0.0000 *
Liquid Level Percent (%)	50.00	Vessel Pressure (kPa)	100.0

Holdup: Vessel Levels

Phase	Level (m)	Percent (%)	Volume (m3)
Vapour	--	--	0.0000
Liquid	--	--	0.0000
Aqueous	--	--	0.0000

Holdup: Details

Phase	Accumulation (kgmole/h)	Moles (kgmole)	Volume (m3)
Vapour	0.0000	0.0000	0.0000
Liquid	0.0000	0.0000	0.0000
Aqueous	0.0000	0.0000	0.0000
Total	0.0000	0.0000	0.0000

Duty Valve Source : Direct_Q

SP (kJ/h)	Min. Available (kJ/h)	Max. Available (kJ/h)
4.962e+005	--	--

Liquid Heater Height as % of Vessel Volume

Top of Heater : 5.00 % Bottom of Heater : 0.00 %

Heat Flow into the PFR: Heating

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TEAM LND
Calgary, Alberta
CANADA

Case Name: C:\Program Files\Hyprotech\HYSYS 3.1\Cases\MR. BABALOLA.hsc

Unit Set: SI

Date/Time: Fri Feb 09 09:09:22 2007

Vessel Sizing: DEGREASING UNIT

Vertical Separator Properties: ACID PICKLING UNIT

Max. Vap. Velocity (m/s)	0.5343 *	Liq. Res. Time (seconds)	300.0 *
L/D Ratio	3.000 *	Demister to Top (m)	0.3048 *

Construction Information

Chem Eng Fab Index	252.5	Allowable Stress (kPa)	9.446e+004
Material Type	Carbon Steel	Shell Thickness (mm)	6.350
Mass Density (kg/m3)	7861	Corrosion Thickness (mm)	3.175
FMC	1.000	Joint Efficiency	1.000

Costing Coefficients

Shell Thick		Shell Mass		Base Cost		Accessories	
A1:	0.4000 *	A4:	0.8116 *	A5:	8.600 *	A8:	1017 *
A2:	2.000 *			A6:	-0.2165 *	A9:	0.7396 *
A3:	0.2000 *			A7:	4.580e-002 *	A10:	0.7068 *

Costing Results

Base Cost (\$US)	1.200e+004	Associated Cost (\$US)	3894	Total Cost (FOB \$US)	1.589e+004
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Sizing Results

Diameter (m)	1.372	Liquid Residence Time (seconds)	300.0
Total Length (m)	4.801	Liquid Surge Height (m)	0.4300
L/D Ratio	3.000	LLSD (m)	0.7620
Max. Allowable Vapour Velocity (m/s)	0.5343	Liquid Res. Time at LLSD (seconds)	615.5
Demister Thickness (mm)	0.0000		

Vapour Space

Sump To Inlet Nozzle (m)	0.3048	Demister To Head (m)	0.3048
Inlet Nozzle to Demister (m)	3.418	Ellipsoidal Head (m)	0.3048
Demister Thickness (mm)	0.0000	Total Vapour Height (m)	4.0316

APPENDIX II

Cathodic balance

Nickel deposited is 40.679kg.

Chrome deposited is 351.62kg.

Anodic Balance

Since it is assumed that what is removed from Anode is deposited on the cathode. It implies that 351.62kg and 187.89kg of nickel and chrome were what is removed from Anode and consequently deposited at the cathode.

Electrolyte balance

Volume of Electroplating tank is 2.583m^3

$$\begin{aligned} \text{Volume of Bath} &= 0.6 \times \text{volume of the Electroplating Tank.} \\ &= 0.6 \times 2.583 \\ &= 1.550\text{m}^3 \end{aligned}$$

It implies total quantity of water required to make the Electrolyte is 1.550m^3 and quantity of salt to dissolve can be calculated thus.

$\text{NiSO}_4 \cdot 6\text{H}_2\text{O}$ 291g	1000L	1kg
L	1m^3	1000g
$= 291\text{kg}/\text{m}^3$		
$\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$ 52.5g	1000L	1kg
L	1m^3	1000g
$= 52.5\text{kg}/\text{m}^3$		

H_3BO_3 (Boric Acid) 42.5g	1000L	1kg
L	$1m^3$	1000g

$Cr_2(SO_4)_3 \cdot 6H_2O$ 300g	1000L	1Kg
L	$1m^3$	1000g

$$= 300kg/m^3$$

Sulphuric Acid tetraoxosulphate (vi) Acid

10ml is required it implies

0.01L is required

$$1000L \longrightarrow 1M^3$$

$$1L = \frac{1}{1000m^3}$$

$$0.01L = \frac{1 \times 0.01m^3}{1000}$$

$$0.01L = 1.0 \times 10^{-5} m^3$$

Density of H_2SO_4 is $1184kg/m^3$

$$\text{Density} = \frac{\text{mass}}{\text{volume}}$$

$$\begin{aligned} \text{Mass} &= \text{Density} \times \text{Volume} \\ &= \frac{1184kg \times 1.0 \times 10^{-5}m^3}{m^3} \\ &= 0.01184kg \end{aligned}$$

Volume Equivalent of constituent of the electrolyte

$$\frac{\text{NiSO}_4 \cdot 6\text{H}_2\text{O}}{386.01} \times 291 \times 1.550 = 1.168\text{m}^3$$

$$\frac{\text{NiCl}_2 \cdot 6\text{H}_2\text{O}}{386.01} \times 52.5 \times 1.550 = 0.211\text{m}^3$$

$$\frac{\text{Cr}_2(\text{SO}_4)_3 \cdot 6\text{H}_2\text{O}}{386.01} \times 300 \times 1.550 = 1.205$$

$$\frac{\text{Boric Acid H}_3\text{B}_2}{386.01} \times 42.5 \times 1.550 = 0.171\text{m}^3$$

$$\frac{\text{Sulphuric Acid H}_3\text{SO}_4}{386.01} = 0.01184 \times 1.550 = 4.754 \times 10^{-5}\text{m}^3$$

Quantity of each constituent that will be require to make 1.502 titles are

$$\text{NiSO}_4 \cdot 6\text{H}_2\text{O} \quad \frac{291\text{kg}}{\text{m}^3} \times 1.168\text{m}^3 = 339.89\text{kg}$$

$$\text{NiCl}_2 \cdot 6\text{H}_2\text{O} = \frac{52.5\text{KG}}{\text{M}^3} \times 0.211\text{m}^3 = 11.08\text{kg}$$

$$\text{H}_3\text{B}_2 = \frac{42.5\text{kg}}{\text{M}^3} \times 0.171\text{m}^3 = 7.27\text{kg}$$

$$\text{H}_2\text{SO}_4 = 0.01184\text{kg}$$

Direct evaluation of cost of material

$$\text{NiSO}_4 \cdot 6\text{H}_2\text{O} \quad 1\text{kg} \longrightarrow \text{N}400.00$$

$$339.89\text{kg} \longrightarrow \text{N}400 \times 339.89 = \text{N} 135,956$$

$$\text{NiCl}_2 \cdot 6\text{H}_2\text{O} \quad 1\text{kg} \longrightarrow \text{N}340.00$$

$$11.08\text{Kg} \longrightarrow \text{N}340 \times 11.08$$

$$= \text{N}3,767$$

Boric Acid 1 kg → N630.00

7.27kg → N620 x 7.27

= N4.580

10ml H₂SO₄ l is needed

10ml = 0.01L

1L = N500

Cr₂(SO₄)₃·6H₂O 1Kg → N380

189.89Kg → N380 x 189.89 = N72,158

0.01L = $\frac{0.01 \times N500}{1} = N5$

Anode 1kg → N720

351.62kg → N720 x 351.62 = N253.159

Water required in dissolution of salts.

25 litres → N5.00

1.550m³ → 1550 litres

Cost of water required in dissolution 5 x 1550 = N310.00

25

Total water required for washing

Post acid pickling washing 2901kg

Post decreasing unit washing 576.073kg

Post electroplating washing (Dragout) 1823kg

Total water required is (2901 + 576.07 + 1823) kg = 5,300.07kg

Density = $\frac{\text{Mass}}{\text{Volume}}$

Volume = $\frac{\text{Mass}}{\text{Density}}$

$$\text{Volume } \frac{5300.07}{1000} \frac{\text{Kg}}{\text{Kg/m}^3} = 5.3\text{m}^3$$

$$5.3\text{m}^3 \longrightarrow 5,300\text{Litres}$$

Total water required for the entire process =

Water required for dissolution + water required for post (Acid pickling, Decreasing and Electroplating) + water required in Acid pickling tank + water required in dragout tank =

$$(1550 + 5300 + 1220 + 1215) \text{ litres.}$$

$$= 9,285 \text{ litres}$$

$$\text{cost of this Water} = \frac{\text{N}5 \times 9,285 \times 10^3}{20}$$

$$= \text{N}46,425$$

$$\text{Actual mass of HCL required} = 0.6935\text{kg/m}^3 \times 1.2204\text{m}^3 = 0.846\text{kg}$$

$$\text{Actual mass of NaOH require} = 0.028\text{kg/m}^3 \times 1.215\text{m}^3 = 0.034\text{kg}$$

$$\text{Volume of Acid} = 0.6 \times 1.550\text{m}^3$$

$$\text{Pickling unit} = 0.6 \times 1.550 \times 1000\text{L} = 930\text{L}$$

$$\text{Volume of Degreasing} = 0.6 \times 2.583$$

$$\text{Unit} = 0.6 \times 2.583 \times 100\text{L}$$

$$= 154.98\text{L}$$

Costing of NaOH and HCl required

NaOH

$$1\text{kg} \longrightarrow \text{N}5,00$$

$$0.034 \longrightarrow 0.039 = \text{N}17.00$$

HCl

$$2.795\text{kg} \longrightarrow \text{N}300$$

$$1\text{kg} \longrightarrow \frac{\text{N}300}{2.795} = \text{N}107.33$$

$$0.8464\text{kg} = \frac{300 \times 0.8464}{2.795}$$

$$= \text{N}90.85$$

Cost of wrist watch 1kg of raw wrist watch chain \longrightarrow N10

Each wrist watch weight of 0.25kg

$$0.25\text{kg} \longrightarrow 0.25 \times 10 = \text{N}2.50$$

$$1999.0\text{kg} \longrightarrow 1999.0 \times 10 = 19,990$$

$$= \text{N}1.2 \times 10^4$$

APPENDIX III

COST ESTIMATION OF EQUIPMENT AND ECONOMIC ANALYSIS USING COST INDEX ESTIMATION (Marshall and Smith Cost Index)

Cost equipment

The cost are based on cost data of 2006 which are available in dollars.

Exchange rate ER as at 2006: 130 naira = 1 dollar

Therefore ER: 130 naira

Purchased Cost Data

The purchased cost data for process equipment is given below as:

Purchase Cost Data for Reactors, Degreasing Tank, Acid Pickling, tank electroplating tank and washing tanks.

$$PC = \frac{M \cdot S}{280} (101 \cdot D^{1.066} H^{0.802}) ER$$

Where D = Diameter of the column (ft)

H = Height of the column (ft)

M.S = Marshall and Smith index

Fc = 1.00 + Fm + Fp

Fm		Fp	
CS	1.00	<50Psia	1.00
SS	3.67	200	1.15
Monel Steel	6.34	400	1.35
Titanium	7.89	600	1.60

Note: CS = Carbon Steel

SS = Stainless Steel

Monel Steel

Purchase cost of the acid pickling unit

Purchase cost PC is given by the relation below:

Marshall and Smith index is Ms: = 110

The material of construction selected is

Carbon Steel:

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + f_m + f_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3 \quad \text{Note: } 1m^3 \text{ } 3.281$$

Diameter of the tank D: 1.200m or 3.937 ft

Height H: 1.799m or 5.903ft

$$PC_{\text{Acidpickling tank}} = \frac{M/s}{280} \left[101.9(D)_{\text{Acid pickling tank}}^{1.066} (H_{\text{Acidpickling tank}}) 0.802 F_c ER \right]$$

$$= \frac{11.0}{280} \left[101 \times 9(3.907)^{1.066} \times (5.903) 0.802 \times 3 \right] 130$$

$$= \text{N}277,191$$

Purchased cost of the Washing Tank 1

Purchase cost Pc is given by the relation below:

Marshall and Smith index is Ms: = 110

The material of construction is carbon steel

For carbon steel

$$F_m = 1.00 \quad f_p = 1.00$$

$$F_c = 1 + f_m + f_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

Diameter of the tank	1.198m	→	2.9306ft
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Height of the tank	1.979m	→	5.896ft
--------------------	--------	---	---------

$$P_{C_{\text{washing tank}}} = \frac{M_s}{280} \left[101.9(D_{\text{washing tank}})^{1.066} (H_{\text{washing tank}})^{0.802} F_c \right] ER$$

$$= \frac{110}{280} 101.9(3.9306)^{1.066} \times (5.896)^{0.802} \times 3 \times 130$$

$$= \text{N}278,711$$

Degreasing Unit

Purchasing cost of the degreasing tank

Purchase cost P_c is given by the relation below:

Marshall and Smith index is $M_s: 110$

The material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + f_m + f_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

Diameter of the tank	D =	1.198m or 3.931 ft
----------------------	-----	--------------------

	H =	1.797m or 5.896 ft
--	-----	--------------------

$$\begin{aligned}
 PC_{\text{Degreasing tank}} &= \frac{Ms \left[101.9 (D_{\text{Degreasing tank}})^{1.066} (H_{\text{Degreasing tank}})^{0.802} Fc \right]}{280} ER \\
 &= \frac{110 \left[101.9 (3.931)^{1.066} \times (5.896)^{0.802} \times 3 \right]}{280} 130 \\
 &= \text{N}278,741
 \end{aligned}$$

Purchasing Cost of Washing Tank 2

Purchasing cost P_c is given by the relation below

Marshall and Smith index is $Ms = 110$

The material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

$$\text{Diameter of the tank} \quad D = 1.203\text{m or } 3.947 \text{ ft}$$

$$H = 1.804\text{m or } 5.919 \text{ ft}$$

$$\begin{aligned}
 PC_{\text{washing tank 2}} &= \frac{Ms \left[101.9 (D_{\text{washing tank 2}})^{1.066} (H_{\text{washing tank 2}})^{0.802} Fc \right]}{280} ER \\
 &= \text{N}297,500 \\
 &= \frac{110 \left[101.9 (3.947)^{1.066} \times (3.919)^{0.802} \times 3 \right]}{280} 130 \\
 &= \text{N}201,753
 \end{aligned}$$

Purchasing cost of the washing tank 3

Purchase cost P_c is given by the relation below

Marshall and Smith index is $Ms = 110$

The material of construction selected is carbon steel.

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

$$\text{Diameter of the tank} \quad D = 1.299 \text{ or } 4.262$$

$$H = 1.949 \text{ or } 6.395$$

$$P_{C_{\text{washing tank 3}}} = \frac{M_s}{280} \left[101.9 (D_{\text{washing tank 3}})^{1.066} (H_{\text{washing tank 3}})^{0.802} F_c \right] ER$$

$$= \frac{110}{280} \left[101.9 (4.262)^{1.066} \times (6.395)^{0.802} \times 3 \right] 130$$

$$= \text{N}324,284$$

Purchasing Cost of the Electroplating Tank

Purchase Cost P_c is given by the relation below:

Marshall and Smith Index is $M_s = 110$

The material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

Diameter of the tank

$$D = 1.299\text{m or } 4.262$$

$$H = 1.949\text{m or } 6.395$$

$$P_{C_{\text{Electroplating tank}}} = \frac{Ms}{280} \left[101.9 (D_{\text{electroplating tank}})^{1.066} (H_{\text{electroplating tank}})^{0.802} F_c \right] ER$$

$$= \frac{110}{280} \left[101.9 (4.262)^{1.066} \times (6.395)^{0.802} \times 3 \right] 130$$

$$= \text{N}324,284$$

Purchasing cost of the Dragout tank 1

Purchase cost Pc is given by the relation below

Marshall and Smith index is : Ms = 110

the material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

$$\text{Diameter of the tank} \quad D = 0.9405\text{m or } 3.08616\text{ft}$$

$$H = 1.411 \text{ or } 4.629\text{ft}$$

$$P_{C_{\text{Dragout tank 1}}} = \frac{Ms}{280} \left[101.9 (D_{\text{dragout tank 1}})^{1.066} (H_{\text{dragout tank 1}})^{0.802} F_c \right] ER$$

$$= \frac{110}{280} \left[101.9 (3.086)^{1.066} \times (4.629)^{0.802} \times 3 \right] 130$$

$$= \text{N}177,374$$

Purchasing cost of the Dragout Tank 2

Purchase Cost P_c is given by the relation below:

Marshall and Smith index is : $M_s = 110$

The material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

$$\text{Diameter of the tank} \quad D = 0.9405\text{m or } 3.086\text{ft}$$

$$H = 1.411\text{m or } 4.629\text{ft}$$

$$\begin{aligned} P_{C_{\text{Dragout tank 2}}} &= \frac{M_s}{280} [101.9 (D_{\text{dragout tank 2}})^{1.066} (H_{\text{Dragout tank 2}})^{0.802} F_c] ER \\ &= \frac{110 [101.9 (3.086)^{1.066} \times (4.629)^{0.802} \times 3] 130}{280} \\ &= \text{N}177,374 \end{aligned}$$

Purchasing cost of the Dragout Tank 3

Purchase Cost is given by the relation below:

Marshall and Smith index is $M_s = 110$

The material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

$$\text{Diameter of the tank} \quad D = 0.9405\text{m or } 3.086\text{ft}$$

$$H = 1.411\text{m or } 4.629\text{ft}$$

$$\begin{aligned}
 & \frac{PC_{\text{Dragout tank 3}}}{280} = \frac{Ms [101.9 (D_{\text{Dragout tank 3}})^{1.066} (H_{\text{Dragout tank 3}})^{0.802} Fc] ER}{280} \\
 & = \frac{110 [101.9 (3.086)^{1.066} \times (4.629)^{0.802} \times 3] \times 130}{280} \\
 & = \text{N177,374}
 \end{aligned}$$

Purchasing cost of the Dragout Tank 4

Purchase Cost is given by the relation below:

Marshall and Smith index is $Ms = 110$

The material of construction selected is carbon steel

$$Fm = 1.00 \quad Fp = 1.00$$

$$Fc = 1 + Fm + Fp$$

$$Fc = 1 + 1 + 1$$

$$Fc = 3$$

$$\text{Diameter of the tank} \quad D = 0.762\text{m or } 2.500\text{ft}$$

$$H = 2.667\text{m or } 8.750\text{ft}$$

$$\begin{aligned}
 & \frac{PC_{\text{Dragout tank 3}}}{280} = \frac{Ms [101.9 (D_{\text{Dragout tank 3}})^{1.066} (H_{\text{Dragout tank 3}})^{0.802} Fc] ER}{280} \\
 & = \frac{110 [101.9 (2.500)^{1.066} \times (8.750)^{0.802} \times 3] \times 130}{280} \\
 & = \text{N236,139}
 \end{aligned}$$

Purchasing cost of the Dragout Tank 5

Purchase Cost is given by the relation below:

Marshall and Smith index is $Ms = 110$

The material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

$$\text{Diameter of the tank} \quad D = 0.762\text{m or } 2.500\text{ft}$$

$$H = 2.667\text{m or } 8.750\text{ft}$$

$$P_{C_{\text{Dragout tank 3}}} = M_s [101.9 (D_{\text{Dragout tank 3}})^{1.066} (H_{\text{Dragout tank 3}})^{0.802} F_c] \text{ ER}$$

$$= \frac{110 [101.9 (2.500)^{1.066} \times (8.750)^{0.802} \times 3]}{280} \times 130$$

$$= \text{N}236,139$$

Purchasing cost of the Dragout Tank 6

Purchase Cost is given by the relation below:

$$\text{Marshall and Smith index is} \quad M_s = 110$$

The material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

$$\text{Diameter of the tank} \quad D = 0.762\text{m or } 2.500\text{ft}$$

$$H = 2.667\text{m or } 8.750\text{ft}$$

$$P_{C_{\text{Dragout tank 3}}} = M_s [101.9 (D_{\text{Dragout tank 3}})^{1.066} (H_{\text{Dragout tank 3}})^{0.802} F_c] \text{ ER}$$

$$= \frac{110 [101.9 (2.500)^{1.066} \times (8.750)^{0.802} \times 3]}{280} \times 130$$

$$= \text{N}236,139$$

Purchasing cost of the Concentrator

Purchase Cost P_c is given by the relation below:

Marshall and Smith index is : $M_s = 110$

The material of construction selected is carbon steel

$$F_m = 1.00 \quad F_p = 1.00$$

$$F_c = 1 + F_m + F_p$$

$$F_c = 1 + 1 + 1$$

$$F_c = 3$$

$$\text{Diameter of the tank} \quad D = 0.3048\text{m or } 1.000\text{ft}$$

$$H = 1.676\text{m or } 5.499\text{ft}$$

$$P_{C_{\text{concentrator}}} = \frac{M_s}{280} [101.9 (D_{\text{concentrator}})^{1.066} (H_{\text{Dragout tank}})^{0.802} F_c] \text{ ER}$$

$$= \frac{110 [101.9 (1.000)^{1.066} \times (5.499)^{0.802} \times 3]}{280} \times 130$$

$$= \text{N}61,260$$

Purchasing cost of Pump

$$\text{Cost Pump} = C_2 \left(\left(\frac{Q_2}{Q_1} \right)^n \right) \left(\frac{M-s \text{ 2006}}{M-s \text{ 1979}} \right) \text{ ER Naira}$$

Where Q = Volumetric Flow rate gal/min

C = Cost of pump and

$M-s$ = Marshall and Smith index

n = Exponent

Note that all flow rates are converted into gallon/minutes

From Timmerhaus, Cost of pump of 10gpm is 1×10^3

$$Q_1 = 10 \text{ gal/min} \quad C_1 = 1000$$

Flow rate of the pump 1 designed is:

Q_2	=	1.560m ³	1000L	4 gallon	1h
		h	1m ³	4L	60 min

$$Q_2 = 6.5 \text{ gal/min}$$

Marshall and Smith index

M. s 1979 645

$$n = 0.34$$

Accordingly Chebrony, Marshall and Smith index obtained by extrapolating equation for year 2006 is:

M. s_2006 is :

$$M. s - 2006 = 1098$$

Cost of pump 1 in year 2006 is:

$$C_2 = \left[\left(\frac{C_1 Q_2}{Q_1} \right)^n \right] \frac{M - s \text{ 2006}}{M - s \text{ 1979}} \text{ ER Naira } C^2 \left[1000 \left(\frac{6.5}{10} \right)^{0.34} \right] \frac{1098}{645} 130$$

$$C_2 = \left[\left(\frac{C_1 Q_2}{Q_1} \right)^n \right] \frac{1098}{645} 130$$

$$= \text{N}2,002$$

Cost of pump 2 in year 2006 is

$$Q_1 = 10 \text{ gal/min.} \quad C_1 = 1000$$

$$M - s - 1979 = 645$$

$$M - s \text{ 2006} = 1098 \quad n = 0.34$$

Q^2	=	0.1002m ³	1000L	1 gallon	1h
		h	1m ³	4 litres	60 min

$$Q_2 = 0.4175 \text{ gallon/min}$$

$$C_2 = \left[\left(C_1 \frac{Q_2}{Q_1} \right)^n \right] \frac{M - s \text{ 2006}}{M - s \text{ 1979}} \text{ ER naira}$$

$$C_2 = \left[\left(1000 \frac{0.4175}{10} \right)^{0.34} \right] \frac{1098}{645} \times 130$$

$$= \text{N787}$$

Cost of pump 3 in year 2006 is

$$Q_1 = 10 \text{ gal/min.} \quad C_1 = 1000$$

$$M - s - 1979 \quad 645$$

$$M - s - 2006 \quad 1098 \quad n = 0.34$$

$Q_2 =$	0.2 044	1000L	1 gallon	1h
	H	1m ³	4 litres	60 min

$$Q_2: = 0.852 \text{ gallon/min}$$

$$C_2 = \left[\left(C_1 \frac{Q_2}{Q_1} \right)^n \right] \frac{M - s \text{ 2006}}{M - s \text{ 1979}} \text{ ER naira}$$

$$C_2 = \left[1000 \left(\frac{0.852}{10} \right)^{0.34} \right] \frac{1098}{645} \times 130$$

$$= \text{N1,003}$$

Cost of pump in year 2006 is

$$Q_1 = 10 \text{ gal/min.} \quad C_1 = 1000$$

$$M - s - 1979 \quad 645$$

$$M - s - 2006 \quad 1098 \quad n = 0.34$$

$Q_2 =$	$\frac{1.543\text{m}^3}{H}$	$\frac{1000L}{1\text{m}^3}$	$\frac{1 \text{ gallon}}{4 \text{ litres}}$	$\frac{1\text{h}}{60 \text{ min}}$
---------	-----------------------------	-----------------------------	---	------------------------------------

$$Q_2: = 6.43 \text{ gallon/min}$$

Cost of pump 4 in year 2006 is

$$Q_2: = 0.835 \text{ gallon/min}$$

$$C_2 = \left(\frac{M - s \ 2006}{C_1 \left(\frac{Q}{Q_1} \right)^n} \right) \left(\frac{\text{ER naira}}{M - s \ 1979} \right)$$

$$C_2 = \left(1000 \left(\frac{6.43}{10} \right)^{0.34} \right) \left(\frac{1098}{645} \right) 130$$

$$= \text{N}4,261$$

Cost of recycling pump in year 2006 is

$$Q_1 = 10 \text{ gal/min.} \quad C_1 = 1000$$

$$M - s - 1979 \quad 645$$

$$M - s - 2006 \quad 1098 \quad n = 0.34$$

$Q_2 =$	$\frac{5.131\text{m}^3}{H}$	$\frac{1000L}{1\text{m}^3}$	$\frac{1 \text{ gallon}}{4 \text{ litres}}$	$\frac{1\text{h}}{60 \text{ min}}$
---------	-----------------------------	-----------------------------	---	------------------------------------

$$Q_2: = 21.38 \text{ gallon/min}$$

$$C_2 = \left(\left(C_1 \frac{Q_2}{Q_1} \right)^n \right) \left(\frac{M - s \text{ 2006}}{M - s \text{ 1979}} \right) \text{ER naira}$$

$$C_2 = \left[1000 \left(\frac{21.38}{10} \right)^{0.34} \left(\frac{1098}{645} \right) \right] 130$$

= N3000

Total cost of pump = cost of pump 1 + cost of pump 2 + cost of pump 3 +

Cost of pump 4 + cost of recycling pump

N(2,002 + 787 + 1003 + 4,261 + 3000)

Tc pumps N11,053

Operating Cost

Total purchased cost of equipment

$$P_{C_{\text{Equip}}} = P_{C_{\text{Acid pickling Tank}}} + P_{C_{\text{washing tank 1}}} + P_{C_{\text{Degreasing tank 2}}} + P_{C_{\text{washing tank 2}}} + P_{C_{\text{washing tank 3}}} + P_{C_{\text{Electroplating tank}}} + P_{C_{\text{Dragout tank 1}}} + P_{C_{\text{dragout tank 2}}} + P_{C_{\text{dragout tank 3}}} + P_{C_{\text{dragout tank 4}}} + P_{C_{\text{dragout tank 5}}} + P_{C_{\text{dragout tank 5}}} + P_{C_{\text{Concentrator}}} + T_{C_{\text{pumps}}}$$

$$+ P_{C_{\text{heater}}} + P_{C_{\text{air dryer}}} + P_{C_{\text{preheater}}} + P_{C_{\text{cooler}}} + P_{C_{\text{Mixer}}}$$

$$P_{C_{\text{Equip}}} = \text{N}(277,191 + 278,711 + 278,741 + 201,753 + 324,284 + 324,284 + 177,374 + 177,374 + 177,374 + 236,139 + 236,139 + 236,139 + 61,260 + 2002 + 787 + 1003 + 4261)$$

Pc = N2,758,677

ECONOMIC ESTIMATION OF TOTAL CAPITAL INVESTMENT

The total capital investment can be given by

$$T_{\text{in}} = T_{\text{FC}} + W_{\text{c}} + T_{\text{LC}}$$

Where TFC = Total fixed cost

Wc = Working capital

TLC = Total Land cost.

Total Fixed Cost

The factorial method can be used with the relationship below:
(Direct Cost)

$$\begin{aligned} \text{TFC:} &= 10\text{Pc equip} \\ &= 758,677 \\ &= \text{N}27,586,770 \end{aligned}$$

Working Capital

$$\begin{aligned} \text{WC} &= 0.5 \text{ TFC} \\ &= 0.5 \text{ N}27,586,770 \\ &= \text{N}13,793,385 \end{aligned}$$

Total land cost

$$\begin{aligned} \text{TLC} &= 0.1 \text{ TFC} \\ &= 0.1 \times \text{N}27,586,770 \\ &= \text{N}2,758,677 \end{aligned}$$

$$\begin{aligned} \text{Tinv} &= \text{TLC} + \text{WC} + \text{TFC} \\ &= \text{N}(2,758,677 + 13,793,385 + 27,586,770) \\ &= \text{N}44,138,832 \end{aligned}$$

Operating Cost

This is divided into fixed and variable operating cost.

Let the plant life be 20 years $n = 20$ years

Fixed Operating Cost

Direct Labour Cost

$$\begin{aligned} \text{Lbc} &= 0.06 \text{ TFC} \\ &= 0.06 \times \text{N}27,586,770 \\ &= \text{N}1,655,206.2 \end{aligned}$$

Plant Maintenance and Repair

$$Mc = 0.08 \text{ TFC}$$

$$Mc = 0.08 \times \text{N}27,586,770$$

$$Mc = \text{N}2,206,941.6$$

Insurance

$$\text{Insc} = 0.01 \text{ TFC}$$

$$= 0.01 \times \text{N}27,586,770$$

$$= \text{N}275,867.7$$

Local Taxes

$$\text{Lbc} = 0.02 \text{ TFC}$$

$$= \text{N}0.02 \times 27,586,770$$

$$= \text{N}551,735.4$$

Royalties and licensed fee

$$\text{Royc} = 0.01 \text{ TFC}$$

$$= 0.01 \times \text{N}27,586,770$$

$$= \text{N}275,867.7$$

Laboratory Cost

$$\text{Labc} = 0.21\text{bc}$$

$$= 0.2 \times \text{N}551,735.4$$

$$= \text{N}110,347.08$$

Plant overhead cost

$$\text{POHc} = 0.5 \text{ Lbc}$$

GENERAL EXPENSES

i. Administrative

$$\begin{aligned} \text{ADMc} &= 0.25 \text{ Lbc} \\ &= 0.25 \times \text{N}551,735.4 \\ &= \text{N}137,933.85 \end{aligned}$$

ii. Research and Development

$$\begin{aligned} \text{RADc} &= 0.015 \text{ TFC} \\ &= 0.015 \times \text{N}27,586,770 \\ &= \text{N}413,801.55 \end{aligned}$$

Total Fixed Operating Cost

$$\text{TFOc} = \text{RADc} + \text{ADMc} + \text{POHc} + \text{Labc} + \text{Royc} + \text{Labc} + \text{Insc} + \text{Mc} + \text{Lbc} + \text{Sc}$$

$$\text{TFOc} = \text{N}[413,801.55 + 137,933.85 + 275,867.7 + 110,347.08 + 275,867.7 + 551,735.4 + 275,867.7 + 2,206,941.6 + 655,206]$$

$$\text{TFOc} = \text{N}5,903,567.98$$

Annual fixed operating Cost

$$\text{TFO annum} = \frac{\text{TFOc}}{\text{N}}$$

$$\begin{aligned} \text{TFO annum} &= \frac{\text{N}5,903,567.98}{20} \\ &= \text{N}295,178.4/\text{Annum} \end{aligned}$$

Variable Operating Cost

Cost of raw materials CRMC

$$\begin{aligned} \text{Cost of Raw Material} &= \text{Chrome sulphate} + \text{Nickel Sulphate Cost} + \\ \text{Nickel Chloride Cost} &+ \text{Boric Acid Cost} + \text{Sulphoric Acid} + \text{Aniode} \\ &\text{Cost} + \text{Total cost of water for the electroplating} \\ &\text{system} + \text{Cost of Hydrochloric acid} + \text{Cost of} \\ &\text{Sodium Hydroxide} + \text{Cost of Base Metal} \end{aligned}$$

$$\text{CRMc} = \left(\begin{array}{l} 30,500 + 52,300 + 1,200 + 2,500 + 50 + 18,000 + 14,000 + \\ 9,000 + 19,990 \\ \text{N}147,540 \end{array} \right)$$

Annual cost of raw material

536,148	300 days
Day	1 year

$$\text{ACRM} = \text{N}44,262,000/\text{Annum}$$

Miscellaneous

$$\begin{aligned} \text{Msc} &= 0.05 \text{ Lbc} \\ &= 0.05 \times \text{N}551,735.4 \\ &= \text{N}27,586.77 \end{aligned}$$

Utilities cost

$$\begin{aligned} \text{Utc} &= 0.02 \text{ Lbc} \\ &= 0.02 \times \text{N}551,735.4 \\ &= \text{N}11,034.7 \end{aligned}$$

Packaging

$$\begin{aligned} \text{PAC} &= 0.001 \text{ Lbc} \\ &= 0.001 \times \text{N}551,735.4 \\ &= \text{N}551.7 \end{aligned}$$

Total Variable Operating Cost

$$\begin{aligned} \text{TVOc} &= \text{Pac} + \text{Msc} + \text{ACRM} + \text{Utc} \\ &= \text{N}(551.7 + 27,586.77 + 11,034.7 + 147,540) \\ &= \text{N}44,301,173 \end{aligned}$$

Annual Variable Operating Cost

$$\begin{aligned} \text{TVO annum} &= \frac{\text{TVOc}}{n} \\ \text{TVO annum} &= \frac{44,301,173}{20} \\ \text{TVO annum} &= \text{N}2,215,058 \end{aligned}$$

Total Annual Operating Cost

$$\begin{aligned} \text{AOCc} &= \text{TFO annual} + \text{TVO annual} \\ &= \text{N}295,178 + 2,215,058 \\ &= \text{N}2,510,236.7 \end{aligned}$$

Profit Analysis

Annual Production Rate (APR)

$$= 7 \times 3600 \times 20 \times 300 = 1814400\text{kg}$$

$$\begin{aligned} \text{Production Cost (PC) per kg} &= \text{AOCc/APR} \\ &= \text{N}2,501,236.7/181,440\text{kg} \\ &= \text{N}2.0 \end{aligned}$$

Selling Price

This is a function of total production cost demanded and market forces.

Distribution price of N12.00 per kg is assumed

$$= 12 \times 181440000$$

APS (Annual Product Sales)

$$= \text{N}21,772,800$$

Profit before tax (PBT)

$$\begin{aligned} \text{PBT} &= \text{APS} - \text{AOCc} \\ &= \text{N}[21,772,800 - 2,510,236.7] \end{aligned}$$

$$\text{PBT} = \text{N}19,262,563.3$$

Annual Depreciation (Depr)

Let S = Salvage value after n years of the plant

Assume plant life of 20 years

Then S:=

Let Vs after 20 years be taken as 15% of TFC in

$$\frac{15}{100} \times \text{N}27,586,770$$
$$S = \text{N}4,138,015.5$$
$$\text{Dep} = \frac{T_{in} - s}{n} = \frac{\text{N}[44,138.832 - 4,138,015.5]}{20}$$
$$= \text{N}2,000,040.8$$

Tax Payable (TP)

Assume tax ratio of 65% and depreciation is tax allowable, hence tax ratio = 0.65

$$\text{TP} = (\text{PBT} - \text{Depr}) \text{ Tax ratio}$$
$$= \text{N}[19,262,563.3 - 2,000,040.8]$$
$$\text{TP} = 17,262,522.5 \times 0.65$$
$$\text{TP} = \text{N}11,220,639.6$$

Profit after tax (PAT)

$$\text{PAT} = \text{PBT} - \text{TP}$$
$$= \text{N}[19,262,563.3 - 11,220,639.6]$$
$$= \text{N}8,041,923.7$$

Net Income (NIN)

$$\text{NIN} = \text{N}8,041,923.7 + 2,000,040.8/\text{year}$$
$$= \text{N}10,041,964.5$$

Pay back period (PBP)

$$\text{PBP} = \frac{T_{inv}}{\text{NIN}}$$
$$= \frac{\text{N}44,138.832}{\text{N}10,041,964.5}$$

$$\begin{array}{rcl}
 \text{PBP} & = & 4.39 \text{ year} \\
 & & \begin{array}{|l}
 \hline
 4.39 \text{ year} \quad | \quad 300 \text{ days} \\
 \hline
 \quad \quad \quad | \quad 1 \text{ year} \\
 \hline
 \end{array}
 \end{array}$$

$$\text{PBP} = 1317 \text{ days}$$

Rate of return on investment

$$\text{RORI} = \frac{\text{Nin} + \text{Dep}}{\text{Tinv}} = 100$$

$$\frac{\text{N}10,041,964.5 + 2,000,040.8}{\text{N}44,138,832} = 0.273 = 108.5\%$$

$$\text{N}44,138,832$$

$$= 0.273 \times 100 = 27.28\%$$