

ELECTRODEPOSITION OF NICKEL ALUMINIDE BASED ALLOY ON FE SUBSTRATE IN DEEP EUTECTIC SOLVENT

Submitted in accordance with the requirement of the Universiti Teknikal Malaysia Melaka (UTeM) for the Bachelor Degree of Manufacturing Engineering (Hons).

by

LIM YAN YING

B051510083

951214085462

FACULTY OF MANUFACTURING ENGINEERING

2019



ΙΙΝΙ\/FRSITI ΤΕΚΝΙΚΔΙ ΜΔΙ ΔΥSIΔ

BORANG PENGESAHAN STATUS LAPORAN PROJEK SARJANA MUDA

Tajuk:ELECTRODEPOSITION OF NICKEL ALUMINIDE BASED ALLOY
ON FE SUBSTRATE IN DEEP EUTECTIC SOLVENT

Sesi Pengajian: 2018/2019 Semester 2

Saya LIM YAN YING (951214-08-5462)

mengaku membenarkan Laporan Projek Sarjana Muda (PSM) ini disimpan di Perpustakaan Universiti Teknikal Malaysia Melaka (UTeM) dengan syarat-syarat kegunaan seperti berikut:

- 1. Laporan PSM adalah hak milik Universiti Teknikal Malaysia Melaka dan penulis.
- 2. Perpustakaan Universiti Teknikal Malaysia Melaka dibenarkan membuat salinan untuk tujuan pengajian sahaja dengan izin penulis.
- 3. Perpustakaan dibenarkan membuat salinan laporan PSM ini sebagai bahan pertukaran antara institusi pengajian tinggi.
- 4. *Sila tandakan ($\sqrt{}$)

SULIT(Mengandungi maklumat yang berdarjah keselamatan atau kepentingan
Malaysiasebagaimana yang termaktub dalam AKTA RAHSIA RASMI 1972)

TERHAD

(Mengandungi maklumat TERHAD yang telah ditentukan oleh organisasi/ badan di mana penyelidikan dijalankan)

√ TIDAK TERHAD

Disahkan oleh:

Alamat Tetap:

Cop Rasmi:

Tarikh: _____

Tarikh: _____

*Jika Laporan PSM ini SULIT atau TERHAD, sila lampirkan surat daripada pihak berkuasa/organisasi berkenaan dengan menyatakan sekali sebab dan tempoh laporan PSM ini perlu dikelaskan sebagai SULIT atau TERHAD.

DECLARATION

I hereby, declared this report entitled "Electrodeposition of Nickel Aluminide Based Alloy on Fe Substrate in Deep Eutectic Solvent" is the results of my own research except as cited in reference.

Signature:Author's Name: LIM YAN YINGDate: 3rd June 2019

APPROVAL

This report is submitted to the Faculty of Manufacturing Engineering of Universiti Teknikal Malaysia Melaka as a partial fulfillment of the requirements for the degree of Bachelor of Manufacturing Engineering (Hons.). The members of the supervisory committee are as follow:

(Dr. Muhammad Zaimi bin Zainal Abidin)

ABSTRAK

Penyiasatan penggunaan pelarut eutektik sebagai elektrolit dalam elektrodeposi Nickel Aluminide dijalankan dalam kajian ini kerana fabrikasi aloi binari dalam pelarut eutektik belum dieksplorasi dengan sepenuhnya. Tujuan kajian ini adalah untuk menghasilkan lapisan aloi berasaskan Nickel Aluminide pada substrat Fe dengan elektrodeposisi serta menggunakan pelarut eutektik sebagai elektrolit. Mula-mulanya, substrat Fe digilap sehingga permukaan mirip cermin. Seterusnya, substrat masing-masing degreased, dibilas menggunakan etanol dan air suling. Kemudian, substrat akan dipasang dengan resin epoksi. Akhirnya, sampel elektrodeposited dengan ketumpatan arus 0.001, 0.002, 0.004, 0.005, dan 0.01 A/dm², SERTA masa penyaduran dari 15, 30 dan 45 minit. Mandi plating terdiri daripada campuran Nikel (II) Klorida dan Aluminium (III) Klorida bercampur homogen dengan larutan ChCl: EG akan dihasilkan. Pengimbasan mikroskop elektron (SEM) dan X-ray difraksi (XRD) digunakan untuk menentukan morfologi permukaan dan komposisi elemen sampel bersalut. Kemudian, pengukuran elektrokimia dilakukan dalam 3.5% berat natrium klorida untuk mengkaji sifat kakisan aloi bersalut. Akhirnya, ujian kekerasan dan kekasaran permukaan sampel bersalut masing-masing dapat ditentukan dengan penguji mikro Vickers hardness dan profilometer.

i

ABSTRACT

The investigation of the use of deep eutectic solvent as the electrolyte in the electrodeposition of Nickel Aluminide is conducted in this study due to the fabrication of binary alloy in deep eutectic solvent has not been fully explored. The aim of this study is to produce a coating of Nickel Aluminide based alloy on Fe substrate using deep eutectic solvent as electrolyte by electrodeposition. Fe substrate is first polished until a mirror-like surface. Next, the substrate degreased, rinsed using the ethanol and distilled water respectively. Then, the substrate were mounted by epoxy resin. Finally, the sample was electrodeposited with current density of 0.001, 0.002, 0.004, 0.005, 0.01 A/dm², and plating time from 15, 30 and 45 minutes. The plating bath consists of Nickel (II) Chloride and Aluminium (III) Chloride were mixed homogenously with the ChCl:EG solution. Scanning electron microscope (SEM) and X-ray diffraction (XRD) were used to determine the surface morphology and the elemental composition of the coated samples. Then, electrochemical measurements were done in 3.5 wt.% of sodium chloride to study the corrosion behaviour of the coated alloy. Lastly, hardness tester and profilometer respectively.

DEDICATION

I would like to dedicate this work to my Beloved parents Appreciated friends Honourable supervisor and lecturers For giving me moral support, cooperation and encouragements. Thank you.

ACKNOWLEDGEMENT

I had receive the help and guideline from many respected parties throughout the project, who deserve my greatest gratitude. I would like to express my deep gratitude to Dr. Muhammad Zaimi bin Zainal Abidin, my respected supervisor, for his great mentoring, patient guidance, enthusiastic, encouragement and useful critiques throughout this project. His willingness to spare his time generously has been much appreciated.

Also, I would like to give a special thanks to my friends and family who gave me much motivation, encouragement and cooperation mentally while completing this report. They had given their precious comments and advice throughout my research. Thanks for being supportive all these while.

Finally, I would like to thank everyone who had directly or indirectly involve in this research, as well as expressing my apology that I could not mention personally to each and every one of you.

TABLE OF CONTENT

Abst	trak	i		
Abst	tract	ii		
Ded	ication	iii		
Ack	nowledgement	iv		
Tabl	e of content	v		
List	of tables	ix		
List	of figures	x		
List	of abbreviations	xii		
List	of symbols	xiv		
CHA	APTER 1: INTRODUCTION	1		
1.1	Background Of Study	1		
1.2	Problem Statement	2		
1.3	Objectives	3		
1.4	1.4 Scope			
CH	APTER 2: LITERATURE REVIEW	5		
2.1	Coating	5		
2.2	Types Of Coating Techniques	6		
	2.2.1 Vapour depositions	6		

	2.2.1.1 Physical vapour deposition	6
	2.2.1.2 Chemical vapour deposition	6
	2.2.2 Galvanizing	7
	2.2.4 Diffusion coating	7
	2.2.5 Conversion coating	8
	2.2.6 Thermal spraying	8
	2.2.7 Electrochemical techniques	8
	2.2.7.1 Electroless deposition	9
2.4	Nickel Aluminide Based Alloy	11
	2.4.1 Applications of <i>Ni</i> ₃ <i>Al</i> based alloys	13
2.5	Electrodeposition Of Alloys	15
	2.5.1 Microstructural evolutions of Nickel-Aluminide nanocomposites during possible synthesis and thermal spray process.	wder 16
	2.5.2 Microstructure of aluminide coating on the Nickel based superalloy MAR	M247
	deposited by CVD aluminizing process	18
	2.5.3 Electrodeposition of Aluminum-Nickel alloys in organic chloroaluminate	ionic
	liquids	20
2.6	Factors Affecting Electrodeposition	23
	2.6.1 Effect of current density and distribution	23
	2.6.2 Effect of ph	24
	2.6.3 Effect of bath temperature	26
	2.6.4 Effect of concentration bath	27
	2.6.5 Effect of electrodepositing time	28
2.7	Electrodeposition In Deep Eutectic Solvent	28
CHA	APTER 3: METHODOLOGY	30
3.1	Project Flow Chart	31

3.2	Chemicals 3				
3.3	Preparation Of Substrate	32			
3.4	Preparation Of Electrolyte				
3.5	Parameters Used In Electrodeposition	33			
3.6	Electrodeposition Of Nickel Aluminide Based Alloy On Fe Substrate In Deep Eutectic Solvent	33			
3.7	Analysis Techniques Of Coated Samples	34			
	3.7.1 Surface morphology of electrodeposited samples	35			
	3.7.2 Microstructure and phase identification of electrodeposited samples	35			
	3.7.3 Mechanical analysis of electrodeposited samples	37			
	3.7.3.1 Hardness testing	37			
	3.7.3.2 Surface roughness	37			
	3.7.4 Analysis of the electrochemical measurements of electrodeposited samples	38			
CHA	APTER 4: RESULT AND DISCUSSION	40			
CHA 4.1	APTER 4: RESULT AND DISCUSSION Characterization Of Electroplated Samples	40 40			
	Characterization Of Electroplated Samples	40			
4.1	Characterization Of Electroplated Samples 4.1.1 Scanning electron microscope	40 40			
4.14.2	Characterization Of Electroplated Samples 4.1.1 Scanning electron microscope X-ray Diffraction Analysis (XRD)	40 40 42			
4.14.2	Characterization Of Electroplated Samples 4.1.1 Scanning electron microscope X-ray Diffraction Analysis (XRD) Mechanical Analysis Of Nickel Aluminide Coating	40 40 42 45			
4.14.2	Characterization Of Electroplated Samples 4.1.1 Scanning electron microscope X-ray Diffraction Analysis (XRD) Mechanical Analysis Of Nickel Aluminide Coating 4.3.1 Hardness test	40 40 42 45 45			
4.14.24.34.4	 Characterization Of Electroplated Samples 4.1.1 Scanning electron microscope X-ray Diffraction Analysis (XRD) Mechanical Analysis Of Nickel Aluminide Coating 4.3.1 Hardness test 4.3.2 Surface roughness 	40 40 42 45 45 47			
4.14.24.34.4	 Characterization Of Electroplated Samples 4.1.1 Scanning electron microscope X-ray Diffraction Analysis (XRD) Mechanical Analysis Of Nickel Aluminide Coating 4.3.1 Hardness test 4.3.2 Surface roughness Corrosion Behaviour of Ni-Al Coating 	40 40 42 45 45 45 47 50			

	5.2.1 Effect of substrate surface preparation	
	5.2.2 Effect of plating bath	55
5.3	Sustainable Design And Development	56
5.4	Limitations	56
REF	ERENCES	58
APP	ENDICES	61

LIST OF TABLES

3.1	Fixed parameters for electrodeposition process.	32
3.2	Varying parameters for electrodeposition process.	32
3.3	Analysis techniques to analyse the coatings formed on Fe substrate.	33
3.4	Technical specifications of XRD machine.	35
4.1	SEM images of electrodeposition of Nickel Aluminide onto	
	Fe substrate with different plating parameters.	39
4.2	Average reading of hardness of coated samples.	43
4.3	Average reading of surface roughness of coated samples	46
4.4	Linear polarization curve measurement of chosen samples	50

LIST OF FIGURES

2.1	Schematic diagram for electroless plating (Sahoo et al., 2016).	9
2.2	Set-up for electrodeposition process.	10
2.3	Ni-Al system (Talaş, 2018).	12
2.4	Physical properties of nickel aluminides (Talaş, 2018).	12
2.5	Chemical composition of nickel aluminide based alloys selected for	
	commercial applications (Weber &Corp, 2016).	13
2.6	Water turbine rotor made of a nickel aluminide based alloy (Talaş, 2018).	14
2.7	XRD patterns of Ni-Al powder at different milling times	
	(Movahedi, 2014).	16
2.8	Cross-sectional of microstructure for HVOF coating (Movahedi, 2014).	17
2.9	XRD pattern of the HVOF coating (Movahedi, 2014).	17
2.10	Scheme of equipment for deposition of aluminide coatings	
	by CVD method (Zielińska et al., 2013).	18
2.11	Surface morphology of an aluminide coating fabricated by CVD	
	method at 1050°C/8h (Zielińska et al., 2013).	19
2.12	Cross sections of an aluminide coating fabricated by CVD	
	method at 1050°C/8h (Zielińska et al., 2013).	19
2.13	XRD pattern of aluminide coating deposited by CVD method	
	(Zielińska et al., 2013).	19
2.14	SEM images for pure Al deposit at -0.3 V in 1.5:1 M AlCl ₃ -EMIM	
	containing 0.026 M $NiCl_2$. and pure Ni deposit at 0.4 V in	
	1.5:1 M AlCl ₃ -EMIM containing 0.1 M NiCl ₂ (Waqar, 2014).	21
2.15	XRD pattern of the coated samples (Waqar, 2014).	21

2.16	Stress and strain curve at pH value of 2.8, 4.1 and 4.67	
	(Lu, Ou, Huang, Yan, &Yan, 2013).	24
2.17	SEM image for FeCo fils deposited at pH (a) pH=1; (b) pH=2;	
	(c) pH=3 ; (d) pH=4 and (e) pH=5 (Lu et al., 2013)	25
2.18	Electrodeposition of Zn-Co-Fe alloys on steel substrates at bath	
	temperature of (a) 20° C; (b) 40° C and (c) 60° C (Attia, 2017)	26
3.1	Process flow chart of project	30
3.2	Set-up of the electrodeposition of nickel aluminide based alloy	
	on Fe substrate in deep eutectic solvent.	33
3.3	SEM machine SEM EVO 50 (Carl Zeiss SMT, UK)	34
3.4	PW3040/60 X'PERT PRO X-ray diffraction system	
	(PANalytical, Netherland) operating via $CuK\alpha$ radiation	35
3.5	Schematic diagram of electrochemical analysis	37
4.1	XRD diagram of the coated samples for each parameters	43
4.2	Graph of average hardness for the coated samples	46
4.3	Graph of average surface roughness for the coated samples	48

xi

LIST OF ABBREVIATIONS

Cr	-	Chromium
Мо	-	Molybdenum
Zr	-	Zirconium
В	-	Boron
С	-	Carbon
Ti	-	Titanium
W	-	Tungsten
Si	-	Silicon
Ni	-	Nickel
Zn	-	Zinc
Co	-	Cobalt
Ag	-	Silver
Al	-	Aluminium
Fe	-	Iron
AgCl	-	Silver Chloride
KCl	-	Potassium Chloride
HCl	-	Hydrochloric acid
ChCl	-	Choline Chloride
NaCl	-	Sodium Chloride
H ₂	-	Hydrogen gas
Ni ₃ Al	-	Nickel Aluminide
Al_2O_3	-	Aluminium Oxide
TiO ₂	-	Titanium Oxide
AlCl ₃	-	Aluminium (III) Chloride
NiCl ₂	-	Nickel (II) Chloride
SEM	-	Scanning Electron Microscopy
XRD	-	X-ray Diffraction
PVD	-	Physical Vapour Deposition
CVD	-	Chemical Vapour Deposition

HVOF	-	High-velocity Oxy Fuel
DES	-	Deep eutectic solvent
EG	-	Ethylene Glycol
EMIM	-	1-ethyl-3-methylimiazolium chloride
FCC	-	Face centered cubic
R	-	Reducing agent
Me	-	Metal to be deposited
WE	-	Working electrode
CE	-	Counter electrode
RE	-	Reference electrode
E _{ocp}	-	Open-circuit potential
I _{corr}	-	Corrosion current density
E _{corr}	-	Corrosion potential

LIST OF SYMBOLS

Mm-MicrometreA-AreaA-Currentcm-Centermetred-DensityGPa-Giga PascalHV-Hardness VickerK-KelvinM-MolMm-MillimetreMpy-milli-inches per yearN-NewtonNm-Degree CelciusS-SecondV-Voltswt.%-ThetaQ-Charged passedI-Current passedt-the time the current is passedzM-mass for oxidized/reduced species	%	-	Percentage
A-Currentcm-Centermetred-DensityGPa-Giga PascalHV-Hardness VickerK-KelvinM-MolMm-MillimetreMpy-nilli-inches per yearNm-Degree CelciusS-SecondV-SecondV-VoltsMn%-ThetaQ-ThetaQ-Charged passedI-Current passedt-the time the current is passedm-change in oxidation statem-mass for oxidized/reduced species	Mm	-	Micrometre
cm- Centermetred- DensityGPa- Giga PascalHV- MHardness VickerK- MM- KelvinM- MolMm- MMpy- MillimetreNm- NewtonNm- NewtonNm- Degree CelciusS- MolV- SecondV- NetaQ- ThetaQ- Collage passedI- Current passedI- Current passedI- Current is passedz- Cual pass for oxidized/reduced species	А	-	Area
d-DensityGPa-Giga PascalHV-Hardness VickerK-KelvinM-MolMm-MillimetreMpy-milli-inches per yearN-NewtonNm-Degree CelciusS-SecondV-Voltswt.%-ThetaQ-ThetaQ-Current passedt-Current passedz-Change in oxidation statem-Secondition state	А	-	Current
GPa-Giga PascalHV-Hardness VickerK-KelvinM-MolMm-MillimetreMpy-milli-inches per yearN-NewtonNm-Degree CelciusS-SecondV-SecondV-Voltswt.%-ThetaQ-Carrent passedI-Carrent passedt-Carrent passedz-Change in oxidation statem-Sec oxidized/reduced species	cm	-	Centermetre
HV-Hardness VickerK-KelvinM-MolMm-MillimetreMpy-milli-inches per yearN-NewtonNm-Degree Celcius°C-Degree CelciusS-SecondV-Voltswt.%-VoltsQ-ThetaQ-Charged passedI-Lerrent passedz-Charge in oxidation statem-Secondized/reduced species	d	-	Density
K-KelvinM-MolMm-MillimetreMpy-milli-inches per yearN-NewtonNm-Nanometre°C-Degree CelciusS-SecondV-Voltswt.%-ThetaQ-Charged passedI-Charged passedI-Current passedt-Charge in oxidation statem-Mes for oxidized/reduced species	GPa	-	Giga Pascal
M-MolMm-MillimetreMpy-milli-inches per yearN-NewtonNm-Nanometre°C-Degree CelciusS-SecondV-Voltswt.%-ThetaQ-Charged passedI-current passedt-the time the current is passedz-Change in oxidation statem-mass for oxidized/reduced species	HV	-	Hardness Vicker
Mm- MillimetreMpy- Milli-inches per yearNm- NewtonNm- Nanometre°C- Degree CelciusS- SecondV- Voltswt.%- Veight PercentageQ- Sharged passedI- Charged passedI- SecondJ- SecondMu- SecondWt.%- SecondQ- SecondI- SecondQ- SecondI- Secon	Κ	-	Kelvin
Mpy-milli-inches per yearN-NewtonNm-Nanometre°C-Degree CelciusS-SecondV-Voltswt.%-Weight PercentageQ-ThetaQ-charged passedI-the time the current is passedz-Change in oxidation statem-mass for oxidized/reduced species	Μ	-	Mol
N-NewtonNm-Nanometre°C-Degree CelciusS-SecondV-Voltswt.%-Weight PercentageΘ-ThetaQ-charged passedI-current passedt-the time the current is passedz-change in oxidation statem-mass for oxidized/reduced species	Mm	-	Millimetre
Nm-Nanometre°C-Degree CelciusS-SecondV-Voltsvt.%-Weight PercentageΘ-ThetaQ-charged passedI-current passedt-the time the current is passedz-charge in oxidation statem-ses for oxidized/reduced species	Мру	-	milli-inches per year
°C-Degree CelciusS-SecondV-Voltswt.%-Weight PercentageΘ-ThetaQ-charged passedI-current passedt-the time the current is passedz-charge in oxidation statem-mass for oxidized/reduced species	Ν	-	Newton
S-SecondV-Voltswt.%-Weight PercentageΘ-ThetaQ-charged passedI-current passedt-the time the current is passedz-charge in oxidation statem-mass for oxidized/reduced species	Nm	-	Nanometre
V-Voltswt.%-Weight PercentageΘ-ThetaQ-charged passedI-current passedt-the time the current is passedz-charge in oxidation statem-mass for oxidized/reduced species	°C	-	Degree Celcius
wt.%-Weight PercentageΘ-ThetaQ-charged passedI-current passedt-the time the current is passedz-charge in oxidation statem-mass for oxidized/reduced species	S	-	Second
Θ-ThetaQ-charged passedI-current passedt-the time the current is passedz-change in oxidation statem-mass for oxidized/reduced species	V	-	Volts
Q-charged passedI-current passedt-the time the current is passedz-change in oxidation statem-mass for oxidized/reduced species	wt.%	-	Weight Percentage
I-current passedt-the time the current is passedz-change in oxidation statem-mass for oxidized/reduced species	Θ	-	Theta
t-the time the current is passedz-change in oxidation statem-mass for oxidized/reduced species	Q	-	charged passed
z-change in oxidation statem-mass for oxidized/reduced species	Ι	-	current passed
m - mass for oxidized/reduced species	t	-	the time the current is passed
-	Z	-	change in oxidation state
M - molar mass for oxidized/reduced species	m	-	mass for oxidized/reduced species
in inotal mass for ontailed reduced species	М	-	molar mass for oxidized/reduced species
F - faraday constant	F	-	faraday constant
n - amount of substance oxidized or reduced	n	-	amount of substance oxidized or reduced

CHAPTER 1 INTRODUCTION

A coating refers to a masking which can be applied onto the surface of certain object. Practically, the purpose of the coating is to provide protection, improve wear resistance, improve corrosion resistant property and improve appearance of an object. Coating processes can be classified into 7 main category namely, electrochemical technique, vapour deposition, galvanizing, powder coating, diffusion coating, conversion coating and thermal spraying. The study focused on one of the technique under the 'Electrochemical Techniques' called, electrodeposition. The study of electroplating metallic alloys and metals has been practiced for about a century and the prior endeavours are all well recorded. Electrodeposition was used to prepare metallic mirrors and create corrosion resistant surface on certain objects in the past. The exploration of electrodeposition tracked back to Michael Faraday and his laws of electrolysis. The principle law expressed that the total quantity of chemical change generated by an electrical current is always proportional total amount of charge passing through the electrolyte.

1.1 Background Of Study

Electrodeposition referred to a process whereby a layer of metal is deposited or coated on another electrode by electrolysis. This process can be achieved by an electrolytic cell, which involves passing electric current through a solution called electrolyte. In electrodeposition, the cathode which is the piece to be plated is connected to the negative terminal whereas the anode is connected to the positive terminal. Both electrodes are immersed in an electrolyte, which is a solution consisting a salt of the same metal as the anode. An example of copper plating onto brass will be discussed to clarify the basic mechanism of electrodeposition. The brass electrode (substrate to be plated) connected to the negative terminal is referred as cathode whereas the copper electrode connected to the positive terminal is called anode. The electrolyte chosen has to be a solution of a salt of the same metal as the anode, thus the electrolyte used in this example is copper sulphate solution. Both anode and cathode will be dipped into the copper sulphate solution after connected to the positive and negative terminal of the battery. When electric field is applied, the positively charged copper ions will be drawn to the negatively charged brass electrode and accumulate on its surface, slowly forming a thin layer of copper plate on the bass substrate.

In this study, Nickel Aluminide based alloy is used to coat on Fe substrate. Nickel Aluminide (NiAl) is known as an intermetallic alloy made up of aluminium and nickel which exhibits similar properties to both ceramic and metal. Nevertheless, intermetallic based alloys exhibit several specific features such as, high melting point, high temperature strength, but low ductility compare to those conventional metals and alloys. All these properties resulted intermetallic compounds possesses similar properties like ceramics but different from ceramics by having good thermal and electric conductivity.

1.2 Problem Statement

Nickel aluminide based alloys are known to have high tensile and compression strength, high corrosion resistance in atmospheres that are rich with oxygen and carbon, high corrosion resistance towards sodium chloride, acid and base solution as well as excellent high temperature wear resistance. Conventional electrodeposition method uses water as a solvent for cation and anion movement. Nickel deposition can be done easily in aqueous bath, but deposition of Aluminium is hard to conduct because Aluminium ions form oxides on the surface of the substrate when react with water. Unlike the nature of electrolytes used for plating processes which poses certain limitations such as, hydrogen evolution in specific conditions, reactive towards specific metals and narrow potential window, deep eutectic solvent possesses extensive potential windows and better chemical inertness. Also, current Nickel Aluminide based alloy coating is fabricated using other coating method like thermal spraying and CVD which consume a lot of energy, time and cost. Therefore, the deposition of Nickel Aluminide in deep eutectic solvent is less known and requires further investigation regarding plating bath composition, plating parameter and its coating various properties. Nevertheless, the study on electrodeposition of Nickel aluminide based alloys on Fe substrate is carried out due to several reasons mainly due to the limitations of iron substrate namely highly reactive, corrodes rapidly in moist air and elevated temperature. Besides, the surface roughness, surface morphological properties, composition and hardness after the coating of nickel aluminide based alloys on iron substrate is analysed as well.

1.3 Objectives

The objectives of this study are as per below:

- 1. To coat Nickel Aluminide based alloy on Fe substrate.
- To study the surface morphology, composition of Ni-Al based alloy coating on Fe substrate.
- To investigate the mechanical properties of Ni-Al based alloy coating on Fe substrate.
- To examine the effect of corrosion behaviour of Ni-Al based alloy coating on Fe substrate.

1.4 Scope

The focus of this study is the electrodeposition of Nickel Aluminide based alloy coating on Fe substrate in deep eutectic solvent. Also, the electrolyte used in this study is deep eutectic solvent instead of aqueous based solvent. Deep eutectic solvent provides the medium without the presence of hydroxide that will oxidize Al surface. The electrodeposition of Ni_3Al based alloy coating on Fe substrate in deep eutectic solvent using variant current density, bath temperature and plating time are studied. The current density used in this study are 0.001, 0.002, 0.004, 0.005 and 0.01 A/ dm^2 , whereas the plating time will ranged from 15, 30 and 45 minutes. After the electrodeposition process is completed, surface morphology and composition of Ni_3Al based alloy coating on Fe substrate are investigated using Scanning Electron Microscopy (SEM) and X-ray diffraction (XRD) analysis for phase identification. The hardness of Ni_3Al based alloy coating on Fe substrate

is determined by using Micro Vickers Hardness Tester and the surface roughness of Ni-Al based alloy coating on Fe substrate is analysed by profilometer. Lastly, the corrosion behaviour in is investigated using electrochemical measurements.

CHAPTER 2 LITERATURE REVIEW

2.1 Coating

A coating is a layer of covering spread over the surface of a substance, normally referred as substrate. The coating can be defined as a layer of material, deposited naturally or synthetically on the surface of a substrate made of another material, in order to obtain desired decorative or technical properties (Burakowski and Wierzchon, 1999). A material sometimes might fail to possess certain specifications or requirements, hence this is where coating plays a significant role. For instance, a rougher surface might needed to be flattened or polished for lower frictional resistance, whereas a smoother surface might need harder surface to strengthen the wear resistance of the substance. Also, an unappealing object might have to aesthetically enhanced to be improved in decorative intention (Audigié et al., 2018). In short, coating are mostly conducted for decorative or functional purposes. Depending on the demand of applications, surface coatings able to either improve the aesthetic appearance, surface roughness, corrosion resistance, frictional resistance and also wear resistance of a material. There are several coating processes exist which can be categorized into 7 categories namely vapour depositions, galvanizing, powder coatings, diffusion coatings, conversion coatings, thermal spraying and electrochemical techniques. Each technique has its own limitations and advantages. A suitable coating method can be selected based on the required specifications or applications.

2.2 Types Of Coating Techniques

2.2.1 Vapour depositions

Vapour deposition is a coating process whereby the materials in its vapour state are condensed either through conversion, condensation or chemical reaction to develop a solid material. Vapour depositions normally conducted in a vacuum chamber. The purpose of vapour depositions are applied to develop coatings in order to modify the electrical, optical, thermal, mechanical properties as well as enhancing wear and corrosion resistance of the substrate (Audigié *et al.*,2018). Vapour deposition process can be classified into Physical Vapour Deposition (PVD) and Chemical Vapour Deposition (CVD).

2.2.1.1 physical vapour deposition

In Physical Vapour Deposition, materials needed to be deposited are converted into vapour form, and then condensed onto the substrates' surface. Thickness of the coating formed will ranged from 0-20mm. The desired material is atomized and ionized in 2 different ways. It is either through physical evaporation or plasma sputtering. In physical evaporation, material is heated in high vacuum to develop a high vapour pressure which is adequate for the condensation to take place on the substrate, whereas in plasma sputtering involves the bombardment of the material surface (Kumar *et al.*, 2015). In this technique, cathode will be the coating material and an inert gas.

2.2.1.2 chemical vapour deposition

Chemical Vapour Deposition is one of the coating process whereby the deposits are formed due to the reaction between the hot substrate and its atmosphere containing gases or vapour of the coating material. The substrate is heated to temperature above 850° C (Audigié *et al.*, 2018). CVD are normally chosen to obtain hard coatings.