PRODUCTION OF VIOLACEIN NANOPARTICLES VIA SONICATION TECHNIQUE WITH THE AID OF SURFACTANTS AS STABILIZER

MOHD AMIR ASYRAF BIN MOHD HAMZAH

A thesis submitted in fulfilment of the requirements for the award of the degree of Master of Philosophy

Faculty of Science Universiti Teknologi Malaysia

JUNE 2018

DEDICATION

This thesis is dedicated to my beloved father and mother, Mohd Hamzah bin Hassan and Norimah binti Salatin who have been very supportive through thick and thin for this meaningful two years. Thanks for all the prayers, advices and guidance.

Special thanks to all my wonderful friends who brighten my life with all the joys, sadness, smiles and laughters. This friendship has been very great experience in my life.

ACKNOWLEDGEMENT

IN THE NAME OF ALLAH THE MOST GRACIOUS THE MOST MERCIFUL

I would like to express my deepest gratitude to the almighty Allah for He has been my pillar of support through out this study. I am also grateful for He has granted me the opportunity to do my Master and allowed me to finish my research in the allocated time.

I would like to express my appreciation to both my supervisors, Dr. Siti Aminah Setu and Prof. Dr. Wan Azlina Ahmad for all your guidance, support, encouragement, patient and helps during the work course of this research. I gained many knowledge and experience during my two years and I believed these things will surely be helpful in my life onwards.

I am indebted to Ministry of Education and Universiti Teknologi Malaysia for the grants that they have provided. These grants helped me carrying out my research smoothly.

I am also thankful for all the biotechnology lab members that supported me through thick and thin. Your supports are one of the reasons I am able to finish my research.

ABSTRACT

Violacein, a violet pigment produced from *Chromobacterium violaceum* UTM5, has gained interest due to its biodegradability and pharmacological properties. However, its high production cost and limited solubility in water have become the major stumbling blocks for the pigment to be applied in different industries. In this study, liquid pineapple waste was used as an alternative inexpensive growth medium for bacteria cultivation instead of expensive synthetic nutrient broth, thus reducing the production cost of this pigment. The cultivation of C. violaceum in 50 L bioreactor gave a crude yield of 11846 ± 925 mg L⁻¹, which was comparable to the yield obtained using commercial growth medium. The crude pigment was successfully extracted using ethyl acetate. The presence of violacein, the major active compound of the crude pigment, was confirmed using high performance liquid chromatography (HPLC), infrared transform spectroscopy (FTIR) and ultraviolet-visible spectrophotometry (UV-Vis). Thermal gravimetric analysis was used to determine crystallinity and thermal degradation while Zetasizer analyzer was used to identify the isoelectric point, stability at various pHs, and particle size of violacein. Violacein nanoparticles were produced via sonication technique, with the aid of surfactants (Tween 80, Triton X-100, sodium dodecyl sulfate and dodecyltrimethylammonium bromide) as solubilizing and stabilizing agent, to address the violacein's poor solubility in water. The violacein nanoparticles were characterized using UV-Vis spectrophotometry, FTIR, thermal analysis and Zetasizer analysis. Water soluble violacein nanoparticles were produced at surfactant concentration greater than its critical micelle concentration, as indicated by FTIR. Zetasizer analysis showed the smallest violacein nanoparticle, which was 131.5 ± 2.001 nm, with polydispersity index (PDI) of 0.180 ± 0.018 , which indicated a monodispersed violacein nanoparticle distribution. The thermal analysis showed that violacein nanoparticles were in amorphous state and stable upon dispersion in water, with a zeta potential of -49.8 \pm 3.49 mV. The violacein nanoparticles have better solubility than the crude violacein pigment. The solubilized violacein nanoparticles remained well-dispersed upon storage in 28 days at different temperatures. In addition, the violet color of the violacein nanoparticles was maintained at pH range of 3 to 11, temperatures of up to 60°C, and under dark condition, despite its nanoscale size. Higher degradation rate was observed at high temperature and upon light illumination, with $k = 6.51 \times 10^{-3} \text{ h}^{-1}$ 1 , $t_{1/2} = 148$ h and $k = 6.75 \times 10^{-4}$ h⁻¹, $t_{1/2} = 1027$ h, respectively, following the firstorder kinetics. In conclusion, this study confirmed the feasibility of using liquid pineapple waste as cheap growth medium for cultivation of C. violaceum UTM5 in pilot scale (50-L bioreactor) while production of water-soluble violacein nanoparticles via sonication method with the aid of surfactants as stabilizers would increase its usefulness, especially in pharmaceutical industry.

ABSTRAK

Violasin, pigmen ungu yang terhasil daripada Chromobacterium violaceum UTM5, telah menarik banyak perhatian kerana sifat keterbiodegradan dan farmakologinya. Walau bagaimanapun, kos penghasilan yang tinggi keterlarutannya dalam air yang terhad menjadi penghalang utama untuk pigmen ini digunakan dalam pelbagai industri. Dalam kajian ini, sisa nenas cecair telah digunakan sebagai medium pertumbuhan alternatif yang murah bagi pemeliharaan bakteria untuk menggantikan kaldu nutrien sintetik yang mahal, seterusnya mengurangkan kos pengeluaran pigmen ini. Pemeliharaan C. violaceum di dalam bioreaktor bersaiz 50 L menghasilkan pigmen mentah sebanyak 11846 ± 925 mg L⁻¹, yang setanding dengan hasil yang diperoleh menggunakan medium pertumbuhan komersial. Pigmen mentah telah berjaya diekstrak menggunakan etil asetat. Kehadiran violasin, komponen aktif utama di dalam pigmen mentah, telah disahkan menggunakan kromatografi cecair berprestasi tinggi (HPLC), spektroskopi inframerah transformasi Fourier (FTIR) dan spektrofotometri ultra ungu-cahaya nampak (UV-Vis). Analisis gravimetri terma telah digunakan untuk menentukan kehabluran dan degradasi haba manakala penganalisis Zetasizer telah digunakan untuk mengenalpasti titik isoelektrik, kestabilan pada pelbagai pH, dan saiz zarah violasin. Nanopartikel violasin telah dihasilkan menggunakan teknik sonikasi, dengan bantuan beberapa surfaktan (Tween 80, Triton X-100, sodium dodekil sulfat dan dodekiltrimetilammonium bromida) sebagai agen pelarut dan penstabil untuk menangani keterlarutan violasin yang rendah dalam air. Nanopartikel violasin telah dicirikan menggunakan spektrofotometri UV-vis, FTIR, analisis terma dan analisis Zetasizer. Nanopartikel violasin yang mudah larut dalam air telah berjaya dihasilkan pada kepekatan surfaktan melebihi kepekatan kritikal misel, seperti yang ditunjukkan oleh FTIR. Analisis Zetasizer menunjukkan saiz nanopartikel violasin yang terkecil iaitu 131.5 ± 2.001 nm, dengan indeks kepoliserakan (PDI) 0.180 ± 0.018 , yang menunjukkan taburan nanopartikel violasin yang sekata. Analisis terma menunjukkan nanopartikel violasin berada dalam keadaan amorfus dan stabil apabila terserak di dalam air dengan potensi zeta -49.8 ± 3.49 mV. Nanopartikel violasin mempunyai keterlarutan yang lebih baik berbanding pigmen violasin mentah. Nanopartikel violasin yang larut ini kekal terserak apabila disimpan selama 28 hari pada suhu yang berbeza. Tambahan pula, warna ungu nanopartikel violasin masih kekal pada julat pH antara 3 dengan 11, suhu sehingga 60°C, dan dalam keadaan gelap walaupun saiznya berskala nano. Kadar degradasi violasin yang lebih tinggi telah dilihat pada suhu tinggi dan apabila terdedah pada cahaya, masing-masing dengan $k = 6.51 \times 10^{-3} \text{ h}^{-1}$, $t_{1/2} = 148 \text{ h}$ dan $k = 6.75 \times 10^{-4} \text{ h}^{-1}$, $t_{1/2} = 1027 \text{ h}$, mengikut kinetik tertib pertama. Kesimpulannya, kajian ini mengesahkan kebolehlaksanaan dalam menggunakan sisa nenas cecair sebagai medium pertumbuhan yang murah untuk pembiakan C. violaceum UTM5 pada skala perintis (50-L bioreaktor) manakala penghasilan nanopartikel violasin yang mudah larut dalam air menggunakan teknik sonikasi dengan bantuan surfaktan sebagai penstabil akan meningkatkan kegunaannya terutamanya dalam industri farmasi.

TABLE OF CONTENTS

CHAPTER	TITLE	PAGE
	DECLARATION	ii
	DEDICATION	iii
	ACKNOWLEDGEMENT	iv
	ABSTRACT	v
	ABSTRAK	vi
	TABLE OF CONTENTS	vii
	LIST OF TABLES	xiii
	LIST OF FIGURES	xv xix
	LIST OF SYMBOLS AND ABBREVIATIONS	
	LIST OF APPENDICES	XX
1	INTRODUCTION	1
	1.1 Background of the Study	1
	1.2 Problem Statement	3
	1.3 Objectives	4
	1.4 Scope of Study	4
	1.5 Significance of Study	5
2	LITERATURE REVIEW	7
	2.1 Pigment as Coloring Agent	7
	2.1.1 Synthetic Pigment	8
	2.1.2 Natural Pigments	11
	2.1.2.1 Plant Pigments	13
	2.1.2.2 Microbial Pigments	14
	2.2 Violacein Pigment	15

		2.2.1	Pineapp	ble Waste as Cheap Growth	16
			Medium	1	
		2.2.2	Structur	re and Derivatives of Violacein	17
		2.2.3	Physica	l Properties of Violacein Pigment	18
		2.2.4	Antibac	terial Activity of Violacein	19
			Pigmen	t	
		2.2.5	Violace	in Nanoparticles	23
	2.3	Techn	iques to	Increase Violacein Solubility in	23
		Water			
		2.3.1	Cosolve	ency	24
		2.3.2	Additio	n of Surfactants	25
			2.3.2.1	Types of Surfactants	26
			2.3.2.2	Critical Micelle Concentration	28
				(CMC)	
			2.3.2.3	Roles of Surfactant in	30
				Antibacterial Drug Formulation	
		2.3.3	Particle	Size Reduction	32
			2.3.3.1	Sonication Technique	32
3	MA	TERI <i>A</i>	LS ANI	O METHODS	34
	3.1			gents and Apparatus	34
	3.2	Instru	Ū	, 11	34
	3.3	Produ	ction of C	Crude Violacein Pigment	36
		3.3.1		n of <i>Chromobacterium violaceum</i>	36
			UTM5		
		3.3.2	Prepara	tion of Bacterial Growth Media	36
			3.3.2.1	Preparation of Nutrient Agar	36
			3.3.2.2	Preparation of Nutrient Broth	37
			3.3.2.3	Preparation of 0.1% Peptone	37
				Water Medium	
			3.3.2.4	Preparation of L-Tryptophan	37

		3.3.2.5	Preparation of Agricultural-	37
			Based Medium-Liquid Pineapple	
			Waste (LPW)	
	3.3.3	Mainten	nance of C. violaceum Stock	38
		Culture		
		3.3.3.1	Short-Term Bacterial Culture	38
		3.3.3.2	Long-Term Bacterial Culture	38
	3.3.4	Cultivat	ion of C. violaceum in 5 L and 50	38
		L Biore	actor	
		3.3.4.1	Bacterial Growth in Liquid	38
			Pineapple Waste (LPW)	
		3.3.4.2	Bacterial Growth in Nutrient	39
			Broth	
		3.3.4.3	Bacterial Growth without L-	39
			Tryptophan	
	3.3.5	Extracti	on of Crude Violacein	39
	3.3.6	Charact	erization of Crude Violacein	40
		Pigmen	t	
		3.3.6.1	Separation using Thin Layer	40
			Chromatography (TLC) and	
			High-Performance Liquid	
			Chromatography (HPLC)	
		3.3.6.2	Attenuated Total Reflectance -	41
			Fourier Transform Infrared	
			Spectroscopy (ATR-FTIR)	
			Analysis	
		3.3.6.3	UV-Visible Spectrophotometer	41
			Analysis	
		3.3.6.4	Determination of Particle Size,	42
			Polydispersity Index (PDI) and	
			Zeta potential of Crude Violacein	
		3.3.6.5	Thermal Analysis	42
3.4	Produ	ction of V	Violacein Nanoparticles	43

		3.4.1	Compar	ison between Stirring and	43
			Sonicati	on Methods	
		3.4.2	Product	ion of Violacein Nanoparticles	43
			using C	osolvency Approach	
		3.4.3	Effect o	f Type and Concentration	44
		3.4.4	Effect o	f Violacein Concentration	45
		3.4.5	Study or	n Sonication Parameters	46
			3.4.5.1	Study on the Effect of Pulse	46
				Time	
			3.4.5.2	Study on the Effect of Sonication	46
				Time	
		3.4.6	Charact	erization of Violacein	47
			Nanopa	rticles	
			3.4.6.1	Particle Size, PDI and Zeta	47
				Potential Analysis	
			3.4.6.2	ATR-FTIR Analysis	47
			3.4.6.3	Thermal Analysis	47
		3.4.7	Study or	n Dispersion Stability of Violacein	48
			Nanopa	rticles	
			3.4.7.1	Storage Time	48
			3.4.7.2	Temperature	48
		3.4.8	Study or	n Color Stability of Violacein	48
			Nanopa	rticles	
			3.4.8.1	Color Analysis of Pigment	48
			3.4.8.2	Storage Time	50
			3.4.8.3	рН	50
			3.4.8.4	Temperature	50
			3.4.8.5	Light Illumination	51
	3.5	Statist	ical Anal	ysis	52
4	R	ESULT	S AND I	DISCUSSION	53
	4.1	Produ	ction of C	Crude Violacein Pigment	53

	4.1.1	Product	ion and Extraction of Crude	53		
		Violace	in Pigment			
	4.1.2	Charact	erization of Crude Violacein	55		
		4.1.2.1	Separation using Thin Layer	56		
			Chromatography (TLC) and			
			High Performance Liquid			
			Chromatography (HPLC)			
		4.1.2.2	Attenuated Total Reflectance –	58		
			Fourier Transform Infrared			
			Spectroscopy (ATR-FTIR)			
			Analysis			
		4.1.2.3	UV-Visible Analysis	60		
		4.1.2.4	Size, Particle Distribution Index	64		
			(PDI) and Zeta potential of			
			Crude Violacein			
		4.1.2.5	Thermal Analysis	66		
4.2	Produ	luction of Violacein Nanoparticles				
	4.2.1	Compai	rison between Stirring and	67		
		Sonicat	ion Methods			
	4.2.2	Product	ion of Violacein Nanoparticles	69		
		Using C	Cosolvency Approach			
	4.2.3	Effect o	of Types and Concentration of	71		
		Surfacta	ants			
		4.2.3.1	Location of Violacein	75		
			Nanoparticles in Surfactant			
			Micelle			
	4.2.4	The Eff	ect of Violacein Pigment	77		
		Concen	tration			
	4.2.5	The Eff	ect of Sonication Parameters	80		
		4.2.5.1	Pulse Time	80		
		4.2.5.2	Sonication time	82		
	4.2.6	Charact	erization of Violacein	85		
		Nanopa	rticles			

			4.2.6.1	ATR-FTIR Analysis	85
			4.2.6.2	Thermal analysis	87
		4.2.7	Dispers	ion Stability of Violacein	88
			Nanopa	rticles	
		4.2.8	Color st	ability of Violacein Nanoparticles	89
			4.2.8.1	The Effect of Storage Time	89
			4.2.8.2	The Effect of pH	90
			4.2.8.3	The Effect of Temperature	91
			4.2.8.4	The Effect of Light Illumination	92
5	CO	NCLUS	SIONS A	ND FUTURE WORKS	94
	5.1	Concl	usion		94
	5.2	Future	e Works		95
REFEREN(CES				97
Appendices	A - D				112

LIST OF TABLES

TABLE	TITLE	PAGE
NO.		
2.1	Classification of synthetic pigments and their uses. (Lomax and Learner, 2006)	9
2.2	The list of natural pigments from plants, animals and microbes. (Malik <i>et al.</i> , 2012; Kumar <i>et al.</i> , 2015; Rajendran and Selvi, 2014; Dufossé, 2006; Kirti <i>et al.</i> , 2014; Tuli <i>et al.</i> , 2015).	11
2.3	Comparison among natural pigments from different sources in terms of production rate, yield, extraction step, cultivation cost and seasonal variation (Mata-Gómez <i>et al.</i> , 2014; Joana Gil-Chávez <i>et al.</i> , 2013; Seyedin <i>et al.</i> , 2015).	13
2.4	The list of pharmacological properties of violacein.	15
2.5	List of pathogenic bacterial strains inhibited by the violacein	19
2.6	The list of surfactants and their respective chemical structure used in this study.	27
3.1	List of instruments used in this study.	35
4.1	The yield of crude violacein pigment, expressed in mg L ⁻¹ , using different growth media (nutrient broth or LPW), with or without the addition of L-tryptophan.	54
4.2	Retention time, peak purity and quantification of violacein and deoxyviolacein compounds in crude pigment.	58
4.3	FTIR data for standard violacein, violacein fraction and crude violacein pigment.	60

4.4	Molar absorption coefficient of crude violacein ($\varepsilon_{violacein}$),	63
	in unit of L mol ⁻¹ cm ⁻¹ and L mg ⁻¹ cm ⁻¹ , determined using	
	the Beer Lambert's law.	
4.5	Production of violacein nanoparticles with (a) addition of	68
	1% (w/v) SDS and (b) addition of 40% (v/v) DMSO via (1)	
	stirring and (2) sonication techniques.	
4.6	Comparison between the CMC obtained from this study	73
	and other sources for each surfactant; Tween 80, Triton X-	
	100, SDS and DTAB.	
4.7	Solubilization power of each surfactant for violacein	74
4.8	The λ_{max} of violacein in different solvents and surfactants	76
	solutions.	
4.9	The effect of sonication time (pulse on in 5 s, pulse off in	84
	10 s) on the average size, PDI and zeta potential of	
	violacein nanoparticles in the presence of surfactant	
	(Tween 80 and SDS).	
4.10	The effect of pulse time on the average size, PDI and zeta	81
	potential of violacein nanoparticles.	
4.11	FTIR spectral data of crude violacein, SDS and violacein	86
	nanoparticle.	
4.12	Degradation constant (k) and half-life ($t_{1/2}$) of violacein	92
	nanoparticles at different temperatures; 10, 25, 40, 60 and	
	80°C.	
4.13	Degradation constant (k) and half-life ($t_{1/2}$) of violacein	93
	nanoparticles in the presence or absence of light.	

LIST OF FIGURES

FIGURE	TITLE	PAGE
NO.		
2.1	Relative energy of electronic transition for unconjugated	8
	(acetaldehyde and ethylene) and conjugated (1,3-butadiene	
	and acrolein) systems. Reproduced from Yadav (2005)	
2.2	Molecular structure of violacein and its derivatives	17
2.3	Schematic diagram of mode of action of violacein against	22
	S. aureus ATCC 29213 and MRSA ATCC 43300 cells,	
	with the permission from Aruldass (2016).	
2.4	The surfactant molecular structure and arrangement at the	26
	water-air interface.	
2.5	Micelle formation. (a) The micelle formation is explained	29
	by the change in surface tension behavior, as a function of	
	surfactant concentration (b) The steps in the micelle	
	formation (c) Micelle and reverse micelle structure in polar	
	and non-polar solvent system, respectively.	
3.1	The CIELAB color wheel. The formula to calculate the	49
	color angle depends on the a* and b* values (McLellan et	
	al., 1995)	
4.1	Biosynthesis pathway of violacein, as illustrated by	55
	(Hoshino, 2011). The pathway involves both enzymatic	
	reaction and nonenzymatic reaction. L-tryptophan acts as	
	the precursor.	
4.2	TLC analysis of (a) standard violacein, (b) violacein	56
	fraction and (c) crude violacein pigment. Two spots of	

	violacein and deoxyviolacein were observed under longwave UV (365 nm) exposure.	
4.2		-7
4.3	HPLC chromatograms of (a) standard violacein, (b)	57
	violacein fraction and (c) crude pigment. Violacein and	
	deoxyviolacein were eluted at 4.4 min and 6.9 min,	
	respectively.	
4.4	FTIR spectra of (a) crude violacein pigment, (b) violacein	59
	fraction and (c) standard violacein at wavenumber of 4000-	
	650 cm ⁻¹ .	
4.5	UV-Vis spectra of (a) crude violacein and (b) violacein	61
	fraction in different solvents; acetone (Yellow), decanol	
	(Red), DMSO (Blue), ethyl acetate (Green), ethanol	
	(Black) and methanol (Purple).	
4.6	Schematic diagram of conjugation system of the violacein	62
	molecule, which results to the violet appearance.	
4.7	Zeta potential of crude violacein pigment in 10% (v/v)	64
	DMSO/water solution at different pH. Results are	
	expressed in mean \pm standard deviation (n = 3).	
4.8	Schematic illustration of the surface charge of crude	65
	violacein at different pH. The isoelectric point (net charge	
	equals to zero) of crude violacein pigment at $pH = 3.3$. The	
	surface charge becomes positive when pH is below the	
	isoelectric point and becomes negative when pH is above	
	the isoelectric point.	
4.9	Thermal analysis of crude violacein at temperature from 30	66
	to 600°C.	
4.10	The production of violacein nanoparticles in various	70
	cosolvent systems using sonication (a) water: DMSO (b)	
	water: ethanol and (c) water: acetone.	
4.11	Production of violacein nanoparticles as a function of	72
	concentration of (a) Tween 80 (b) Triton X-100 (c) SDS	
	and (d) DTAB surfactants. The concentration of violacein	

	(n=3).	
4.12	The effect of SDS concentration on the size of violacein	75
	nanoparticles (nm). Values are presented as mean ±	
	standard deviation. a $p < 0.001$ as compared to the control	
	(1.0 % (w/v) SDS) using one-way ANOVA test using	
	Tukey for post-hoc analysis (n=3).	
4.13	The proposed location of violacein nanoparticles in the	77
	micelle of (a) nonionic and anionic surfactants and (b)	
	cationic surfactant.	
4.14	Concentration of violacein nanoparticles, in µg mL ⁻¹ (•)	78
	and entrapment efficiency of violacein nanoparticles (a) by	
	the surfactant SDS as a function of violacein pigment	
	concentration ($\mu g \ mL^{-1}$). Values are presented as mean \pm	
	standard deviation (n=3).	
4.15	Effect of varying concentration of violacein (µg mL ⁻¹) on	79
	the average size of violacein nanoparticles ($ullet$) and zeta	
	potential (v). Values are presented as mean ± standard	
	deviation (n=3).	
4.16	Effect of pulse time on the production (concentration) of	80
	violacein nanoparticles (•) and temperature of solution (°).	
4.17	Effect of sonication time (min) on the production and	82
	degradation of violacein nanoparticles (µg mL-1) in the	
	presence of different surfactants; Tween 80 (•), Triton X-	
	100 (\blacktriangle), SDS (\diamondsuit) and DTAB (\triangledown). Values for optical density	
	are presented as mean \pm standard deviation (n=3).	
4.18	FTIR spectra of (a) crude violacein, (b) SDS and (c)	85
	violacein nanoparticles at wavenumber of 4000-650 cm ⁻¹ .	
4.19	Thermal analysis of (a) SDS, (b) crude violacein and (c)	87
	violacein nanoparticles.	
4.20	The dispersion stability of violacein nanoparticles (a) upon	88
	storage within 28 days at room temperature and (b) at	

	different temperatures after 14 days. Absorbance values are	
	presented as mean \pm standard deviation (n=3).	
4.21	Color stability of violacein nanoparticles throughout 28	90
	days. The color was measured based on hue angle and	
	chroma values, which were detected using ColorFlex.	
4.22	The effect of pH on color stability at 0 h and 24 h. The hue	91
	and chroma values were measured using ColorFlex.	
4.23	The effect of temperature on the color stability of violacein	92
	nanoparticles, which was measured using color meter, after	
	28 days of storage time.	
4.24	The effect of light illumination on the color stability of	93
	violacein nanoparticles represented in 2D plot of hue and	
	chroma values, detected using color meter	

LIST OF SYMBOLS AND ABBREVIATIONS

ATR-FTIR : Attenuated Total Reflectance - Fourier Transform Infrared

Spectroscopy

CMC : Critical micelle concentration

HPLC : High-performance liquid chromatography

LPW : Liquid pineapple waste

MHA : Muller-Hinton agar

MHB : Muller-Hinton broth

NA : Nutrient agar

NB : Nutrient broth

PDI : Polydispersity index

TLC : Thin layer chromatography

UV-vis : UV-visible spectroscopy

mg : Milligram

mg L⁻¹ : Milligram per litre

°C : Degree celcius

% w/v : Percentage of weight in 100 mL of solvent/solution

% v/v : Percentage of volume in 100 mL of total solvent/solution

LIST OF APPENDICES

APPENDIX	TITLE	PAGE
A	Wavelength scan of violacein and deoxyviolacein compounds using HPLC.	112
В	Standard curves of violacein and deoxyviolacein,	113
	derived from HPLC data.	
С	Fitted absorbance line at 575 nm as a function of	114
	violacein concentration in (a) mol L ⁻¹ and (b) mg L ⁻¹ ,	
	for determination of violacein molar absorption	
	coefficient in different solvents.	
D	The slope of violacein in different surfactant solutions	115
	using regression analysis.	

CHAPTER 1

INTRODUCTION

1.1 Background of the Study

In 2010, tartrazine, quinoline yellow and carmoisine are the synthetic colorants that are banned in the United Kingdom and European Union as they trigger hyperactive behavior amongst children (Fusaro, 2010). Increase of awareness in regard to the danger of artificial (synthetic) colorants to human safety and environment leads to the increase in the use of natural colorants, known as biological colorants. These pigment, extracted from flora and fauna are found to be non-toxic, non-carcinogenic, and biodegradable (Venil *et al.*, 2013). On top of that, pharmacological properties exhibited by natural colorants have better advantages over synthetic pigments. For example, chlorophylls found in green plants exhibit anticancer properties as they can bind with cancer-causing chemicals to form complex structure, thus minimizing the absorption of potential carcinogens via gastrointestinal tract (İnanç, 2011).

Besides extracting the pigments from animals and plants, microbial pigments are also chosen due to their wide strain selection, shorter fermentation period, gene manipulability, lesser downstream processing (involves simple liquid-liquid extraction step) and cheaper growth medium availability (Venil *et al.*, 2013; Tuli *et al.*, 2015). Also, microorganisms can produce unique pigments such as violacein, prodigiosin and flexirubin, which are non-synthesizable by animals and plants. For example, bacterial strain of genus *Chromobacterium* is known for its ability to produce violacein pigment, with pharmacological properties as an antioxidant, and serves as

antimicrobial, antiprotozoal, and antipyretic compound (Durán *et al.*, 2016). The multiresistant *S. aureus* (MRSA) poses a challenge in dealing with antibiotics due to multidrug resistance behavior. The use of violacein to treat MRSA has been reported by Aruldass *et al.* (2015). As a colorant, the intense violet exhibited by violacein, even at low concentration, this is useful to formulate solvent-based ink in plastic application (Venil *et al.*, 2017; Durán *et al.*, 2007). In addition, violacein has been tested and the findings show its application as food colorant in yogurt and jelly (Venil *et al.*, 2015).

However, the challenge in manufacturing bacterial pigments is the need to produce the pigments in a large quantity at low cost (Malik *et al.*, 2012). Nutrient-rich agricultural waste medium obtained from brown sugar, rice bran, pineapple and sugar cane are increasingly popular due to its availability, low cost, and renewability (Ahmad *et al.*, 2012). In addition, the use of agricultural waste residues in bioprocess helps to reduce environmental pollution. Pineapple waste is the choice for growth medium in this study due to its high glucose content and other nutrients such as esters, ketones, alcohols, aldehydes, and acids, which are required for bacterial growth (Hemalatha and Anbuselvi, 2013). The use of pineapple waste as growth medium for the production of *Lactobacilli* sp. has been demonstrated by Pyar *et al.* (2014).

Furthermore, like other natural pigments, violacein has poor solubility in water, thus limits its usage in industrial application. The common organic solvents for violacein are dimethyl sulfoxide (DMSO), acetone, methanol and ethyl acetate, which are harmful to health and environment upon emission. Thus, particle size reduction which includes mechanical nanosization can improve solubility and dissolution rate of violacein in water due to the increase of surface area to volume ratio (Khadka *et al.*, 2014). Besides, violacein nanoparticles allow better membrane penetration and increase its pharmacological activities in drug delivery. To achieve nanosization, sonication is one of the effective top-down particle size reduction approaches. Sonication employs non-interaction vibration energy to disagglomerate and overcome bonding forces in dispersing the nanomaterials.

Nevertheless, agglomeration is a common issue in nanoparticle production. Agglomeration occurs when substances prefer to interact with the same molecules instead of interacting with solvent molecules to lower the kinetic energy and achieve a more stable structure (Mohd Hamzah *et al.*, 2017). Thus, the presence of surfactants as stabilizing agent is important to prevent agglomeration by providing steric or electrostatic repulsion. For example, baicalein nanocrystals which have potent antioxidant, antitumor, and anticancer properties were stable in water with the aid of a surfactant (Zhang *et al.*, 2011). Besides, surfactants can act as solubilizing agent by increasing the solubility in both organic and aqueous solvents. Tehrani-Bagha, Singh and Holmberg (2013) reported the increase of synthetic pigments' solubility when added with surfactant above its critical micelle concentration (CMC).

In this study, crude violacein extracted from *Chromobacterium violaceum* UTM5 was downsized to nanoparticles via sonication, with surfactants as stabilizing agent. The aim was to improve the violacein solubility in aqueous system with small particle size, narrower particle size distribution (low polydispersity index) and high zeta potential (high stability), besides retaining its violet color at different pH, temperature, time and light illumination.

1.2 Problem Statement

The major challenge to commercialize microbial pigments is to achieve high yield with cost-effective production (Malik *et al.*, 2012). The synthetic growth medium is expensive, thus hamper the production of the pigments at industrial scale, leading to its low usage in any applications where people prefer to use synthetic pigments. As for violacein, although exhibiting many pharmacological activities, it has not been utilized in commercial applications. Besides the high cost of production, another issue is its limited solubility in water, but can dissolve in methanol and DMSO (Durán *et al.*, 2007). The organic solvents are toxic even at low dosage (Galvao *et al.*, 2014). To reduce the production cost of violacein, liquid pineapple waste was used as a cheaper alternative growth medium. On the other hand, the solubility of violacein in

water can be increased by reducing its size into nanoscale, due to the increase of surface area to volume ratio. The high surface-to-volume ratio thus increases particle solubility in water system. This study focused on the use of sonication technique to produce the violacein nanoparticles. However, common issues with nanoparticles are poor solubility and dispersibility, this leads to aggregation and sedimentation process, which results in the loss of the bacterial pigment biological activities and reduces the pigment quality to be used as ink (Wu *et al.*, 2011). Thus, surface modification of the violacein nanoparticles with stabilizer molecules such as surfactants imparts the nanoparticles stability. The presence of surfactant as stabilizer acts as barrier, preventing agglomeration via two protection mechanisms, which are steric repulsion and electrostatic repulsion. In short, this research aims to produce a low cost and stable violacein nanoparticles with high dispersibility/solubility in aqueous system.

1.3 Objectives

- 1) To produce, extract and characterize crude violacein pigment from *C. violaceum* UTM5 using liquid pineapple waste as the growth medium.
- 2) To produce and characterize violacein nanoparticles via sonication technique with the aid of surfactants as stabilizers.
- 3) To test the solubility, stability and color performance of the violacein nanoparticles.

1.4 Scope of Study

The crude violacein used in this study was extracted from *C. violaceum* UTM5 strain and used without further purification. Liquid pineapple waste (LPW) and nutrient broth were used as growth medium for *C. violaceum* UTM5. The production of crude violacein was compared from using nutrient broth (NB), LPW (with and without L-tryptophan) as growth media. The bacteria was grown using continuous

shaking condition and extracted via liquid-liquid extraction using ethyl acetate and acetone as the solvents.

The production of violacein nanoparticles was done using water as the medium. The effectiveness of the sonication technique was first compared with mechanical stirring method. By focusing on sonication technique, several parameters including surfactant concentration and violacein concentration, pulse and sonication time were further investigated. Several common industrial surfactants were used in this study such as Tween 80 and Triton X-100 (nonionic surfactant), sodium dodecyl sulfate (anionic surfactant) and dodecyl trimethylammonium bromide (cationic surfactant). The performance of each surfactant was analyzed in terms of particle size, polydispersity index, zeta potential value, and solubilizing power. Sonication parameters such as sonication time and pulse time was optimized to produce stable violacein nanoparticles. The violacein nanoparticles were characterized using attenuated total reflectance Fourier transform infrared, thermal and Zetasizer analyzer.

The dispersion stability of the violacein nanoparticles were tested as a function of time (28 days) and at different temperatures. The dispersion stability was measured via violet color intensity using UV-Vis spectroscopy for upper and lower portion of the violacein solution. The color stability of the violacein nanoparticles was tested as a function of time, pH and temperature and under light illumination using UV-Vis spectrophotometer and Colorflex color meter.

1.5 Significance of Study

The use of liquid pineapple waste as an alternative growth medium reduces the production cost of violacein. Besides that, the use of fruit wastes for the production of microbial pigments leads to lower waste generation, better waste management and fulfills the waste-to-wealth initiative as described in RMK-11. The development of violacein nanoparticles will improve the solubility of violacein in water, thus making them useful for various applications especially in pharmaceutical industry. The use of

REFERENCES

- Aberoumand, A., 2011. A Review Article on Edible Pigments Properties and Sources as Natural Biocolorants in Foodstuff and Food Industry. *World Journal of Dairy and Food Sciences*, 6(1), pp.71–78.
- Adekunte, A.O., Tiwari, B.K., Cullen, P.J., Scannell, A.G.M. and O'Donnell, C.P., 2010. Effect of Sonication on Colour, Ascorbic Acid and Yeast Inactivation in Tomato Juice. *Food Chemistry*, 122(3), pp.500–507.
- Adeoye, M.D., Obi-Egbedi, N.O. and Iweibo, I., 2017. Solvent Effect and Photo-Physical Properties of 2,3-Diphenylcyclopropenone. *Arabian Journal of Chemistry*, 10, pp.134–140.
- Adesina, S.K., Ezeonyebuchi, U. and Akala, E.O., 2015. The Effect of Formulation Variables on Drug Loading of Antitubercular Drugs in Nanoparticle Formulations. *Materials Research Express*, 2(9), p.95403.
- Aggarwal, S. and Goel, A., 2012. Solubility and Its Enhancement Techniques of Poorly Soluble Drugs. *International Journal of Universal Pharmacy and Life Sciences*, 2(1), pp.65–82.
- Ahmad, W.A., Wan Ahmad, W.Y., Zakaria, Z.A. and Yusof, N.Z., 2012. Isolation of Pigment-Producing Bacteria and Characterization of the Extracted Pigments. In: *Application of Bacterial Pigments as Colorant The Malaysian Perspective*. Springer, Berlin, Heidelberg, pp. 25–44.
- Alshatwi, A.A., Subash-Babu, P. and Antonisamy, P., 2015. Violacein Induces Apoptosis in Human Breast Cancer Cells through Up Regulation of BAX, p53 and Down Regulation of MDM2. *Experimental and Toxicologic Pathology*, 68(1), pp.89–97.
- Andrighetti-Fröhner, C.R., Antonio, R. V., Creczynski-Pasa, T.B., Barardi, C.R.M. and Simões, C.M.O., 2003. Cytotoxicity and Potential Antiviral Evaluation of Violacein Produced by *Chromobacterium violaceum*. *Memorias do Instituto Oswaldo Cruz*, 98(6), pp.843–848.

- Antônio, R.V. and Creczynski-Pasa, T.B., 2004. Genetic Analysis of Violacein Biosynthesis by *Chromobacterium violaceum*. *Genetics and Molecular Research*: *GMR*, 3(1), pp.85–91.
- Antonisamy, P. and Ignacimuthu, S., 2010. Immunomodulatory, Analgesic and Antipyretic Effects of Violacein Isolated from *Chromobacterium violaceum*. *Phytomedicine*, 17(3–4), pp.300–304.
- Antonisamy, P., Kannan, P. and Ignacimuthu, S., 2009. Anti-Diarrhoeal and Ulcer-Protective Effects of Violacein Isolated from *Chromobacterium violaceum* in Wistar Rats. *Fundamental and Clinical Pharmacology*, 23(4), pp.483–490.
- Arif, S., Batool, A., Khalid, N., Ahmed, I. and Janjua, H.A., 2017. Comparative Analysis of Stability and Biological Activities of Violacein and Starch Capped Silver Nanoparticles. *RSC Adv.*, 7(8), pp.4468–4478.
- Aruldass, C.A., 2016. Violet Pigment from Chromobacterium violaceum UTM5 Grown in Liquid Pineapple Waste and Its Antibacterial and Cytotoxicity Activities. Universiti Teknologi Malaysia.
- Aruldass, C.A., Masalamany, S.R.L., Venil, C.K. and Ahmad, W.A., 2017. Antibacterial Mode of Action of Violacein from *Chromobacterium violaceum* UTM5 against *Staphylococcus aureus* and methicillin-resistant *Staphylococcus aureus* (MRSA). *Environmental Science and Pollution Research*, pp.1–17.
- Aruldass, C.A., Rubiyatno, R., Venil, C.K. and Ahmad, W.A., 2015. Violet Pigment Production from Liquid Pineapple Waste by *Chromobacterium violaceum* UTM5 and Evaluation of Its Bioactivity. *RSC Adv.*, 5(64), pp.51524–51536.
- Asencio, G., Lavin, P., Alegría, K., Domínguez, M., Bello, H., González-Rocha, G. and González-Aravena, M., 2014. Antibacterial Activity of the Antarctic Bacterium *Janthinobacterium* sp. SMN 33.6 against Multi-Resistant Gram-Negative Bacteria. *Electronic Journal of Biotechnology*, 17(1), pp.1–5.
- August, P.R., Grossman, T.H., Minor, C., Draper, M.P., MacNeil, I. a, Pemberton, J.M., Call, K.M., Holt, D. and Osburne, M.S., 2000. Sequence Analysis and Functional Characterization of The Violacein Biosynthetic Pathway from *Chromobacterium violaceum*. *Journal of Molecular Microbiology and Biotechnology*, 2(4), pp.513–519.
- Babitha, S., 2009. Microbial Pigments. In: *Biotechnology for Agro-Industrial Residues Utilisation*. Springer Netherlands, Dordrecht, pp. 147–162.
- Berde, C. V and Berde, V.B., 2015. Vegetable Waste as Alternative Microbiological

- Media for Laboratory and Industry. World Journal Of Pharmacy And Pharmaceutical Sciences, 4 (5), 1488, 4(5), pp.1488 1494.
- Berni, E., Marcato, P.D., Nakazato, G., Kobayashi, R.K.T., Vacchi, F.I., Umbuzeiro, G.A. and Durán, N., 2013. Violacein/Poly(ε-Caprolactone)/Chitosan Nanoparticles Against Bovine Mastistis: Antibacterial and Ecotoxicity Evaluation. *Journal of Physics: Conference Series*, 429(1).
- Bhadoriya, S.S., Madoriya, N., Madoriya, N., Shukla, K. and MS, P., 2013. Biosurfactants: A New Pharmaceutical Additive for Solubility Enhancement and Pharmaceutical Development. *Biochemistry & Pharmacology: Open Access*, 02(02).
- Biswas, S., Vaze, O.S., Movassaghian, S. and Torchilin, V.P., 2013. Polymeric Micelles for the Delivery of Poorly Soluble Drugs. In: *Drug Delivery Strategies for Poorly Water-Soluble Drugs*. John Wiley & Sons Ltd, Oxford, UK, pp. 411–476.
- Bohrey, S., Chourasiya, V. and Pandey, A., 2016. Polymeric Nanoparticles Containing Diazepam: Preparation, Optimization, Characterization, In-Vitro Drug Release and Release Kinetic Study. *Nano Convergence*, 3(1), p.3.
- Borowitzka, M.A., 2013. High-Value Products from Microalgae-Their Development and Commercialisation. *Journal of Applied Phycology*, 25(3), pp.743–756.
- Boverhof, D.R., Bramante, C.M., Butala, J.H., Clancy, S.F., Lafranconi, W.M., West, J. and Gordon, S.C., 2015. Comparative assessment of nanomaterial definitions and safety evaluation considerations. *Regulatory Toxicology and Pharmacology*, 73(1), pp.137–150.
- Bromberg, N., Dreyfuss, J.L., Regatieri, C. V., Palladino, M. V., Durán, N., Nader, H.B., Haun, M. and Justo, G.Z., 2010. Growth Inhibition and Pro-Apoptotic Activity of Violacein in Ehrlich Ascites Tumor. *Chemico-Biological Interactions*, 186(1), pp.43–52.
- Buckow, R., Kastell, A., Terefe, N.S. and Versteeg, C., 2010. Pressure and Temperature Effects on Degradation Kinetics and Storage Stability of Total Anthocyanins in Blueberry Juice. *Journal of Agricultural and Food Chemistry*, 58(18), pp.10076–10084.
- Cazoto, L.L., Martins, D., Ribeiro, M.G., Durán, N. and Nakazato, G., 2011. Antibacterial Activity of Violacein against *Staphylococcus aureus* Isolated from Bovine Mastitis. *The Journal of Antibiotics*, 64(5), pp.395–397.

- Chatterjee, S., Salaün, F. and Campagne, C., 2014. The Influence of 1-Butanol and Trisodium Citrate Ion on Morphology and Chemical Properties of Chitosan-Based Microcapsules during Rigidification by Alkali Treatment. *Marine Drugs*, 12(12), pp.5801–16.
- Choi, S.Y., Yoon, K.H., Lee, J. Il and Mitchell, R.J., 2015. Violacein: Properties and Production of a Versatile Bacterial Pigment. *BioMed Research International*, 2015, pp.1–8.
- Cooper, D.L. and Harirforoosh, S., 2014. Effect of Formulation Variables on Preparation of Celecoxib Loaded Polylactide-Co-Glycolide Nanoparticles. *PLoS ONE*, 9(12), pp.1–22.
- Darshan, N. and Manonmani, H.K., 2015. Prodigiosin and Its Potential Applications. *Journal of Food Science and Technology*, 52(9), pp.5393–5407.
- Dhakar, R.C., 2012. From Formulation Variables To Drug Entrapment Efficiency of Microspheres: A Technical Review. *Journal of Drug Delivery and Therapeutics*, 2(6).
- Dhakar, R.C., Maurya, S.D., Sagar, B.P.S., Bhagat, S., Kumar, P.S. and Jain, C.P., 2010. Variables Influencing the Drug Entrapment Efficiency of Microspheres: A Pharmaceutical Review. *Der Pharmacia Lettre*, 2(5), pp.102–116.
- Dole, M.N., Patel, P.A., Sawant, S.D. and Shedpure, P.S., 2011. Advance Applications of Fourier Transform Infrared Spectroscopy. *International Journal of Pharmaceutical Sciences Review and Research*, 7(2), pp.159–166.
- Dufossé, L., 2017. Current Carotenoid Production Using Microorganisms. In: Om V. Singh, (ed.) *Bio-pigmentation and Biotechnological Implementations*. John Wiley & Sons, Inc., Hoboken, NJ, USA, pp. 87–106.
- Dufossé, L., 2006. Microbial Production of Food Grade Pigments. *Food Technology* and *Biotechnology*, 44(3), pp.313–321.
- Durán, M., Ponezi, A.N., Faljoni-Alario, A., Teixeira, M.F.S., Justo, G.Z. and Durán, N., 2012. Potential Applications of Violacein: A Microbial Pigment. *Medicinal Chemistry Research*, 21(7), pp.1524–1532.
- Durán, N., Justo, G.Z., Durán, M., Brocchi, M., Cordi, L., Tasic, L., Castro, G.R. and Nakazato, G., 2016. Advances in *Chromobacterium violaceum* and Properties of Violacein-Its Main Secondary Metabolite: A Review. *Biotechnology Advances*, 34(5), pp.1030–1045.
- Durán, N., Justo, G.Z., Ferreira, C. V., Melo, P.S., Cordi, L. and Martins, D., 2007.

- Violacein: Properties and Biological Activities. *Biotechnology and Applied Biochemistry*, 48(3), p.127.
- Fakhr, F.A., Khanafari, A., Baserisalehi, M., Yaghoobi, R. and Shahghasempour, S., 2012. An Investigation of Antileukemia Activity of Violacein-Loaded Dendrimer in Jurkat Cell Lines. *African Journal of Microbiology Research*, 6(33), pp.6235– 6242.
- Fang, M.-Y., Zhang, C., Yang, S., Cui, J.-Y., Jiang, P.-X., Lou, K., Wachi, M. and Xing, X.-H., 2015. High Crude Violacein Production from Glucose by *Escherichia coli* Engineered with Interactive Control of Tryptophan Pathway and Violacein Biosynthetic Pathway. *Microbial Cell Factories*, 14(1), p.8.
- Fariya, M., Jain, A., Dhawan, V., Shah, S. and Nagarsenker, M.S., 2014. Bolaamphiphiles: A Pharmaceutical Review. Advanced Pharmaceutical Bulletin, 4(Suppl 2), pp.483–491.
- Füller, J.J., Röpke, R., Krausze, J., Rennhack, K.E., Daniel, N.P., Blankenfeldt, W., Schulz, S., Jahn, D. and Moser, J., 2016. Biosynthesis of Violacein, Structure and Function of L-Tryptophan Oxidase VioA from *Chromobacterium violaceum*. *Journal of Biological Chemistry*, 291(38), pp.20068–20084.
- Fusaro, D., 2010. When It Comes to Synthetic Food Colors: Beware the "Southampton Six." *Food Processing*.
- Galvao, J., Davis, B., Tilley, M., Normando, E., Duchen, M.R. and Cordeiro, M.F., 2014. Unexpected Low-Dose Toxicity of the Universal Solvent DMSO. *FASEB Journal*, 28(3), pp.1317–1330.
- Gillis, M. and De Ley, J., 2006. The Genera *Chromobacterium* and *Janthinobacterium*. In: Dworkin, M., Falkow, S., Rosenberg, E., Schleifer, K.-H. and Stackebrandt, E., (eds.) *The Prokaryotes*. Springer New York, New York, NY, pp. 737–746.
- Gürses, A., Açıkyıldız, M., Güneş, K. and Gürses, M.S., 2016. Dyes and Pigments. In: *Dyes and Pigments*. Springer International Publishing, pp. 13–29.
- Hait, S.K. and Moulik, S.P., 2001. Determination of Critical Micelle Concentration (CMC) of Nonionic Surfactants by Donor-Acceptor Interaction with lodine and Correlation of CMC with Hydrophile-Lipophile Balance and Other Parameters of The Surfactants. *Journal of Surfactants and Detergents*, 4(3), pp.303–309.
- Hajar, N., Zainal, S., Nadzirah, K.Z., Roha, A.M.S., Atikah, O. and Elida, T.Z.M.T.,2012. Physicochemical Properties Analysis of Three Indexes Pineapple (Ananas

- Comosus) Peel Extract Variety N36. APCBEE Procedia, 4, pp.115–121.
- Hemalatha, R. and Anbuselvi, S., 2013. Physicohemical Constituents of Pineapple Pulp and Waste. *Journal of Chemical and Pharmaceutical Research*, 5(2), pp.240–242.
- Hoshino, T., 2011. Violacein and Related Tryptophan Metabolites Produced by Chromobacterium violaceum: Biosynthetic Mechanism and Pathway for Construction of Violacein Core. Applied Microbiology and Biotechnology, 91(6), pp.1463–1475.
- Humayun, H.Y., Shaarani, M.N.N.M., Warrior, A., Abdullah, B. and Salam, M.A., 2016. The Effect of Co-solvent on the Solubility of a Sparingly Soluble Crystal of Benzoic Acid. *Procedia Engineering*, 148, pp.1320–1325.
- İnanç, A.L., 2011. Chlorophyll: Structural Properties, Health Benefits and Its Occurrence in Virgin Olive Oils. *Akademik Gıdatr (A.L. İnanç)*, 9(2), pp.90–344.
- Joana Gil-Chávez, G., Villa, J.A., Fernando Ayala-Zavala, J., Basilio Heredia, J., Sepulveda, D., Yahia, E.M. and González-Aguilar, G.A., 2013. Technologies for Extraction and Production of Bioactive Compounds to be Used as Nutraceuticals and Food Ingredients: An Overview. Comprehensive Reviews in Food Science and Food Safety, 12(1), pp.5–23.
- Jódar-Reyes, A.B., Martín-Rodríguez, A. and Ortega-Vinuesa, J.L., 2006. Effect of the Ionic Surfactant Concentration on the Stabilization/Destabilization of Polystyrene Colloidal Particles. *Journal of Colloid and Interface Science*, 298(1), pp.248–257.
- Kandisa, R.V., Saibaba KV, N., Shaik, K.B. and Gopinath, R., 2016. Dye Removal by Adsorption: A Review. *Journal of Bioremediation & Biodegradation*, 07(06).
- Khadka, P., Ro, J., Kim, H., Kim, I., Kim, J.T., Kim, H., Cho, J.M., Yun, G. and Lee, J., 2014. Pharmaceutical Particle Technologies: An Approach to Improve Drug Solubility, Dissolution and Bioavailability. *Asian Journal of Pharmaceutical Sciences*, 9(6), pp.304–316.
- Khan, M.I., 2016. Plant Betalains: Safety, Antioxidant Activity, Clinical Efficacy, and Bioavailability. *Comprehensive Reviews in Food Science and Food Safety*, 15(2), pp.316–330.
- Kirti, K., Amita, S., Priti, S., Kumar, A.M. and Jyoti, S., 2014. Colorful World of Microbes: Carotenoids and Their Applications. *Advances in Biology*, 2014(1), pp.1–13.

- Klaessig, F., Marrapese, M. and Abe, S., 2011. Current Perspectives in Nanotechnology Terminology and Nomenclature. In: Murashov, V. and Howard, J., (eds.) *Nanotechnology Standards*. Springer-Verlag New York, pp. 21–52.
- Konzen, M., De Marco, D., Cordova, C.A.S., Vieira, T.O., Antônio, R. V. and Creczynski-Pasa, T.B., 2006. Antioxidant Properties of Violacein: Possible Relation on Its Biological Function. *Bioorganic and Medicinal Chemistry*, 14(24), pp.8307–8313.
- Kumar, A., Vishwakarma, H.S., Singh, J. and Kumar, M., 2015. Microbial Pigments: Production and Their Applications in Various Industries. *International Journal of Pharmaceutical, Chemical and Biological Sciences*, 5(1), pp.203–212.
- Kumar, S. and Singh, P., 2016. Various Techniques for Solubility Enhancement: An Overview. *The Pharma Innovation Journal*, 5(1), pp.23–28.
- Li, X., Qin, Y., Liu, C., Jiang, S., Xiong, L. and Sun, Q., 2016. Size-Controlled Starch Nanoparticles Prepared by Self-Assembly with Different Green Surfactant: The Effect of Electrostatic Repulsion or Steric Hindrance. *Food Chemistry*, 199, pp.356–363.
- Liu, R., 2008. Water-Insoluble Drug Formulation 2nd ed., CRC Press.
- Lomax, S.Q. and Learner, T., 2006. A Review of the Classes, Structures, and Methods of Analysis of Synthetic Organic Pigments. *Journal of the American Institute for Conservation*, 45(2), pp.107–125.
- Mahmood, M.E. and Al-koofee, D. a F., 2013. Effect of Temperature Changes on Critical Micelle Concentration for Tween Series Surfactant. *Global Journal of Science Frontier Research Chemistry*, 13(4), pp.1–7.
- Malik, K., Tokkas, J. and Goyal, S., 2012. Microbial Pigments: A Review. *International Journal of Microbial Resource Technology Accepted*, 41(4), pp.361–365.
- Manzo, G., Carboni, M., Rinaldi, A.C., Casu, M. and Scorciapino, M.A., 2013. Characterization of Sodium Dodecylsulphate and Dodecylphosphocholine Mixed Micelles through NMR and Dynamic Light Scattering. *Magnetic Resonance in Chemistry*, 51(3), pp.176–183.
- Martins, D., Costa, F.T.M., Brocchi, M. and Durán, N., 2011. Evaluation of the Antibacterial Activity of Poly-(d,l-Lactide-co-Glycolide) Nanoparticles Containing Violacein. *Journal of Nanoparticle Research*, 13(1), pp.355–363.
- Martins, D., Frungillo, L., Anazzetti, M.C., Melo, P.S. and Durán, N., 2010.

- Antitumoral Activity of L-Ascorbic Acid-Poly- D,L-(Lactide-Co-Glycolide) Nanoparticles Containing Violacein. *International Journal of Nanomedicine*, 5, pp.77–85.
- Masilamani, K. and Ravichandiran, V., 2012. Effect of Formulation and Process Variables on Drug Content and Entrapment Efficiency of Aceclofenac Nanosuspension. *International Research Journal of Pharmacy*, 3(3), pp.315–318.
- Mata-Gómez, L.C., Montañez, J.C., Méndez-Zavala, A. and Aguilar, C.N., 2014. Biotechnological Production of Carotenoids by Yeasts: An Overview. *Microbial Cell Factories*, 13(1), pp.1–11.
- McLellan, M.R., Lind, L.R. and Kime, R.W., 1995. Hue Angle Determinations and Statistical Analysis for Multiquadrant Hunter L,a,b Data. *Journal of Food Quality*, 18, pp.235–240.
- Mishra, M., Muthuprasanna, P. and Prabha, K., 2009. Basics and Potential Applications of Surfactants—A Review. *Int J PharmTech Res*, 1(4), pp.1354–1365.
- Moawad, H., Abd El-Rahim, W.M. and Khalafallah, M., 2003. Evaluation of Biotoxicity of Textile Dyes Using Two Bioassays. *Journal of Basic Microbiology*, 43(3), pp.218–229.
- Mohajeri, E. and Noudeh, G.D., 2012. Effect of Temperature on the Critical Micelle Concentration and Micellization Thermodynamic of Nonionic Surfactants: Polyoxyethylene Sorbitan Fatty Acid Esters. *E-Journal of Chemistry*, 9(4), pp.2268–2274.
- Mohd Hamzah, M.A.A., Aruldass, C.A., Ahmad, W.A. and Setu, S.A., 2017. Effects of Surfactants on Antibacterial Drugs A Brief Review. *Malaysian Journal of Fundamental and Applied Sciences*, 13(2), pp.118–123.
- MPIB, 2016. Data Pengeluaran Nanas Malaysia Tahun 2016, Johor Bahru, Johor.
- Muhammad Khan, A. and Shah, S.S., 2008. Determination of Critical Micelle Concentration (Cmc) of Sodium Dodecyl Sulfate (SDS) and the Effect of Low Concentration of Pyrene on its Cmc Using ORIGIN Software. *Journal of Chemistry Society Pakistan*, 30(2), pp.186–191.
- Nakamura, Y., Asada, C. and Sawada, T., 2003. Production of Antibacterial Violet Pigment by Psychrotropic Bacterium RT102 Strain. *Biotechnology and Bioprocess Engineering*, 8(1), pp.37–40.

- Namazkar, S., Garg, R., Ahmad, W.Z. and Nordin, N., 2013. Production and Characterization of Crude and Encapsulated Prodigiosin Pigment. *International Journal of Chemical Sciences and Applications*, 4(3), pp.2278–6015.
- Natalia, C.-O., Mayra-Alexandra, C.-C., Vanessa, C.-A. and Luis-Daniel, P.-S., 2017. Influence of Environmental Factors on the Production of Violacein Synthesized By *Janthinobacterium lividum*. *The International Journal of Engineering and Science*, 06(01), pp.76–83.
- Nayak, A.K. and Panigrahi, P.P., 2012. Solubility Enhancement of Etoricoxib by Cosolvency Approach. *ISRN Physical Chemistry*, 2012, pp.1–5.
- Ngamwonglumlert, L., Devahastin, S. and Chiewchan, N., 2017. Natural Colorants: Pigment Stability and Extraction Yield Enhancement via Utilization of Appropriate Pretreatment and Extraction Methods. *Critical Reviews in Food Science and Nutrition*, 57(15), pp.3243–3259.
- Nobbmann, U. and Morfesis, A., 2008. Characterization of Nanoparticles by Light Scattering. *MRS Proceedings*, 1074(April 2016), pp.1074-I10-45.
- Nogueira, D.R., Mitjans, M., Infante, M.R. and Vinardell, M.P., 2011. The Role of Counterions in the Membrane-Disruptive Properties of pH-Sensitive Lysine-Based Surfactants. *Acta Biomaterialia*, 7(7), pp.2846–2856.
- Novotný, Č., Dias, N., Kapanen, A., Malachová, K., Vándrovcová, M., Itävaara, M. and Lima, N., 2006. Comparative Use of Bacterial, Algal and Protozoan Tests to Study Toxicity of Azo- and Anthraquinone Dyes. *Chemosphere*, 63(9), pp.1436–1442.
- O'Neill, C., Hawkes, F.R., Hawkes, D.L., Lourenço, N.D., Pinheiro, H.M. and Delée, W., 1999. Colour in Textile Effluents Sources, Measurement, Discharge Consents and Simulation: A Review. *Journal of Chemical Technology and Biotechnology*, 74(11), pp.1009–1018.
- Olorunsola, E.O. and Adedokun, M.O., 2014. Surface Activity as Basis for Pharmaceutical Applications of Hydrocolloids: A Review. *Journal of Applied Pharmaceutical Science*, 4(10), pp.110–116.
- Orna, M.V., 2013. Discovery of the Physics of Color. In: *The Chemical History of Color*. Springer-Verlag Berlin Heidelberg, pp. 11–28.
- Pereira, L. and Alves, M., 2012. Dyes-Environmental Impact and Remediation. In: Environmental Protection Strategies for Sustainable Development. Springer Netherlands, Dordrecht, pp. 111–162.

- Pongpeerapat, A., Itoh, K., Tozuka, Y., Moribe, K., Oguchi, T. and Yamamoto, K., 2004. Formation and Stability of Drug Nanoparticles Obtained from Drug/PVP/SDS Ternary Ground Mixture. *Journal of Drug Delivery Science and Technology*, 14(6), pp.441–447.
- Pyar, H., Liong, M.T. and Peh, K.K., 2014. Potentials of Pineapple Waste as Growth Medium for *Lactobacillus* species. *International Journal of Pharmacy and Pharmaceutical Sciences*, 6(1), pp.142–145.
- Rahman, F., 2015. The Treatment of Industrial Effluents for the Discharge of Textile Dyes Using by Techniques and Adsorbents. *Journal of Textile Science & Engineering*, 06(01), pp.1–9.
- Rahman, M., Bashar, A., Khalipha, R., Azad, A.K., Hossain, S. and Haque, S., 2014.
 Methods of Solubility and Dissolution Enhancement for Poorly Water Soluble
 Drugs: a Review. World Journal of Pharmacy and Pharmaceutical Sciences,
 3(5), pp.107–130.
- Rajendran, R. and Selvi, B.T., 2014. Natural Dyeing of Cotton Fabrics with Pigment Extracted from Roseomonas Fauriae. *Universal Journal of Environmental Research and Technology*, 4(1), pp.54–59.
- Rathi, P.B., Kale, M., Soleymani, J. and Jouyban, A., 2018. Solubility of Etoricoxib in Aqueous Solutions of Glycerin, Methanol, Polyethylene Glycols 200, 400, 600, and Propylene Glycol at 298.2 K. *Journal of Chemical & Engineering Data*.
- Ratna, P.B.S., 2012. Pollution due to Synthetic Dyes Toxicity & Carcinogenicity Studies and Remediation. *International Journal of Environmental Sciences*, 3(3), pp.940–955.
- Reshmi, S.K., Aravindhan, K.M. and Devi, P.S., 2012. The Effect of Light, Temperature, pH on Stabulity of Betacyanin Pigments in *Basella Alba* Fruit. *ASian Journal of Phatmaceutical and Clinical Research*, 5(4), pp.5–8.
- Rettori, D. and Durán, N., 1998. Production, Extraction and Purification of Violacein: an Antibiotic Pigment Produced by *Chromobacterium violaceum*. *World Journal of Microbiology and Biotechnology*, 14, pp.685–689.
- Reynolds, D.M., 2014. The Principles of Fluorescence. In: Coble, P., Lead, J., Baker, A., Reynolds, D.M. and Spencer, R.G.M., (eds.) *Aquatic Organic Matter Fluorescence*. Cambridge University Press, Cambridge, pp. 3–34.
- Rodrigo-Baños, M., Garbayo, I., Vílchez, C., Bonete, M.J. and Martínez-Espinosa, R.M., 2015. Carotenoids from Haloarchaea and Their Potential in Biotechnology.

- Marine Drugs, 13(9), pp.5508-5532.
- Rodrigues, A.L., Göcke, Y., Bolten, C., Brock, N.L., Dickschat, J.S. and Wittmann, C., 2012. Microbial Production of the Drugs Violacein and Deoxyviolacein: Analytical Development and Strain Comparison. *Biotechnology Letters*, 34(4), pp.717–720.
- Samani, B.H. and Lorigooini, Z., 2015. Effects of Ultrasonic on Microorganisms and Enzymes. *International Science and Investigation Journal*, 4(January), pp.106–113.
- Scheeren, L.E., Nogueira, D.R., Macedo, L.B., Vinardell, M.P., Mitjans, M., Infante,
 M.R. and Rolim, C.M.B., 2016. PEGylated and Poloxamer-Modified Chitosan
 Nanoparticles Incorporating a Lysine-Based Surfactant for pH-Triggered
 Doxorubicin Release. *Colloids and Surfaces B: Biointerfaces*, 138, pp.117–127.
- Seedher, N. and Agarwal, P., 2009. Various Solvent Systems for Solubility Enhancement of Enrofloxacin. *Indian Journal of Pharmaceutical Sciences*, 71(1), pp.82–7.
- Seedher, N. and Kanojia, M., 2009. Co-Solvent Solubilization of Some Poorly-SolubleAntidiabetic Drugs Solubilization Antidiabetic Drugs. *Pharmaceutical Development and Technology*, 14(2), pp.185–192.
- Sekhon, B.S., 2013. Surfactants: Pharmaceutical and Medicinal Aspects. *Journal of Pharmaceutical Technology, Research and Management*, 1(1), pp.43–68.
- Seyedin, A., Hatamian-zarmi, A., Rasekh, B. and Mir-derikvand, M., 2015. Natural Pigment Production by *Monascus purpureus*: Bioreactor Yield Improvement through Statistical Analysis. *Applied Food Biotechnology*, 2(2), pp.23–30.
- Shah, M., Agrawal, Y.K., Garala, K. and Ramkishan, A., 2012. Solid Lipid Nanoparticles of a Water Soluble Drug, Ciprofloxacin Hydrochloride. *Indian Journal of Pharmaceutical Sciences*, 74(5), pp.434–42.
- Sharma, D., Maheshwari, D., Philip, G., Rana, R., Bhatia, S., Singh, M., Gabrani, R., Sharma, S.K., Ali, J., Sharma, R.K. and Dang, S., 2014. Formulation and Optimization of Polymeric Nanoparticles for Intranasal Delivery of Lorazepam Using Box-Behnken Design: In Vitro and In Vivo Evaluation. *BioMed Research International*, 2014.
- Sharma, N., Madan, P. and Lin, S., 2016. Effect of Process and Formulation Variables on the Preparation of Parenteral Paclitaxel-Loaded Biodegradable Polymeric Nanoparticles: A Co-Surfactant Study. *Asian Journal of Pharmaceutical*

- Sciences, 11(3), pp.404–416.
- Shid, R.L., Dhole, S.N., Kulkarni, N. and Shid, S.L., 2014. Formulation and Evaluation of Nanosuspension Formulation for Drug Delivery of Simvastatin. *Int J Pharm Sci Nanotech Vol*, 7(4), pp.2650–2665.
- Singh, K. and Arora, S., 2011. Removal of Synthetic Textile Dyes from Wastewaters:

 A Critical Review on Present Treatment Technologies. *Critical Reviews in Environmental Science and Technology*, 41(9), pp.807–878.
- Singh, O., Kaur, R. and Mahajan, R.K., 2017. Flavonoid-Surfactant Interactions: A Detailed Physicochemical Study. *Spectrochimica Acta Part A: Molecular and Biomolecular Spectroscopy*, 170, pp.77–88.
- Singh, R., 2012. *Solubilization of Organic Dyes in Surfactant Micelles*. Chalmers University of Technology.
- Suslick, K.S., 1998. Sonochemistry. *Kirk-Othmer Encyclopedia of Chemical Tecnology*, pp.516–541.
- Suslick, K.S., 1997. Sonoluminescence and Sonochemistry. 1997 IEEE Ultrasonics Symposium Proceedings. An International Symposium (Cat. No.97CH36118), 1, pp.523–532.
- Tadros, T.F., 2005. Physical Chemistry of Surfactant Solutions. In: Applied Surfactants. Wiley-VCH Verlag GmbH & Co. KGaA, Weinheim, FRG, pp. 19–51.
- Tallury, P., Randall, M.K., Thaw, K.L., Preisser, J.S. and Kalachandra, S., 2007. Effects of Solubilizing Surfactants and Loading of Antiviral, Antimicrobial, and Antifungal Drugs on Their Release Rates from Ethylene Vinyl Acetate Copolymer. *Dental Materials: Official Publication of the Academy of Dental Materials*, 23(8), pp.977–82.
- Taurozzi, J.S., Hackley, V.A. and Wiesner, M.R., 2010. Protocol for Preparation of Nanoparticle Dispersions From Powdered Material Using Ultrasonic Disruption. CEINT, National Institute of Standars and Technology, pp.1–10.
- Tayade, P. and Modi, A., 2007. A Comparative Solubility Enhancement Profile of Valdecoxib with Different Solubilization Approaches. *Indian Journal of Pharmaceutical Sciences*, 69(2), p.274.
- Tehrani-Bagha, A.R. and Holmberg, K., 2013. Solubilization of Hydrophobic Dyes in Surfactant Solutions. *Materials*, 6(2), pp.580–608.
- Tehrani-Bagha, A.R., Singh, R.G. and Holmberg, K., 2013. Solubilization of Two

- Organic Dyes by Anionic, Cationic and Nonionic Surfactants. *Colloids and Surfaces A: Physicochemical and Engineering Aspects*, 417, pp.133–139.
- Tinoi, J. and Rakariyatham, N., 2015. Utilization of Pineapple Waste Hydrolysate for Lipid Production by Oleaginous Yeast *Rhodoturula glutinis*. *International Journal of Advanced Research*, 3(3).
- Toraman, O.Y., 2017. Experimental Investigations of Preparation of Calcite Particles by Ultrasonic Treatment. *Physicochemical Problems of Mineral Processing*, 53(2), pp.859–868.
- Torchilin, V.P., 2001. Structure and Design of Polymeric Surfactant-Based Drug Delivery Systems. *Journal of Controlled Release*, 73(2–3), pp.137–172.
- Trivedi, J.S. and Wells, M.L., 2008. Solubilization Using CoSolvent Approach. In: Liu, R., (ed.) *Water-Insoluble Drug Formulation*. CRC Press, pp. 141–168.
- Tuli, H.S., Chaudhary, P., Beniwal, V. and Sharma, A.K., 2015. Microbial Pigments as Natural Color Sources: Current Trends and Future Perspectives. *Journal of Food Science and Technology*, 52(8), pp.4669–4678.
- Tzintzun-Camacho, O., Sánchez-Segura, L., Minchaca-Acosta, A.Z., Rosales-Colunga, L.M., Hernández-Orihuela, A.L. and Martínez-Antonio, A., 2016. Development of a Bacterial Culture Medium from Avocado Seed Waste. *PeerJ Preprints*, 4, p.e2104v1.
- Upadhyay, A., Lama, J.P. and Tawata, S., 2013. Utilization of Pineapple Waste: A Review. *Journal of Food Science and Technology Nepal*, 6(0).
- Venil, C.K., Ainuddin, M., Wahidin, B., Arul, C., Wan, A.& and Ahmad, A., 2017.
 Production of Bacterial Pigments in Low Cost Medium and Formulation of Biodegradable Ink. *Indian Journal of Experimental Biology*, 55(July), pp.441–447.
- Venil, C.K., Aruldass, C.A., Abd Halim, M.H., Khasim, A.R., Zakaria, Z.A. and Ahmad, W.A., 2015. Spray Drying of Violet Pigment from *Chromobacterium* violaceum UTM 5 and Its Application in Food Model Systems. *International* Biodeterioration and Biodegradation, 102, pp.324–329.
- Venil, C.K. and Lakshmanaperumalsamy, P., 2009. An Insightful Overview on Microbial Pigment, Prodigiosin. *Electronic Journal of Biology*, 5(3), pp.49–61.
- Venil, C.K., Zakaria, Z.A. and Ahmad, W.A., 2013. Bacterial Pigments and Their Applications. *Process Biochemistry*, 48(7), pp.1065–1079.
- Ventura-camargo, B.D.C. and Marin-morales, M.A., 2013. Azo Dyes:

- Characterization and Toxicity A Review. *Textiles and Light Industrial Science and Technology*, 2(2), 2(2), pp.85–103.
- Vivian, J.T. and Callis, P.R., 2001. Mechanisms of Tryptophan Fluorescence Shifts in Proteins. *Biophysical Journal*, 80(5), pp.2093–2109.
- Wang, H., Jiang, P., Lu, Y., Ruan, Z., Jiang, R., Xing, X.H., Lou, K. and Wei, D., 2009. Optimization of Culture Conditions for Violacein Production by a New Strain of *Duganella* sp. B2. *Biochemical Engineering Journal*, 44(2–3), pp.119–124.
- Wu, L., Zhang, J. and Watanabe, W., 2011. Physical and Chemical Stability of Drug Nanoparticles. *Advanced Drug Delivery Reviews*, 63(6), pp.456–469.
- Wu, W., Ichihara, G., Suzuki, Y., Izuoka, K., Oikawa-Tada, S., Chang, J., Sakai, K., Miyazawa, K., Porter, D., Castranova, V., Kawaguchi, M. and Ichihara, S., 2014. Dispersion Method for Safety Research on Manufactured Nanomaterials. *Industrial Health*, 52(1), pp.54–65.
- Yadav, L.D.S., 2005. Ultraviolet (UV) and Visible Spectroscopy. *Organic Spectroscopy*, pp.7–51.
- Zhang, J., Lv, H., Jiang, K. and Gao, Y., 2011. Enhanced Bioavailability after Oral and Pulmonary Administration of Baicalein Nanocrystal. *International Journal of Pharmaceutics*, 420(1), pp.180–188.