



UNIVERSITI PUTRA MALAYSIA

**TOTAL MERCURY AND METHYLMERCURY CONCENTRATION IN
FISH AND THEIR REDUCTION THROUGH PROCESSING**

PARVANEH HAJEB

FSTM 2009 15



**TOTAL MERCURY AND METHYLMERCURY CONCENTRATION IN
FISH AND THEIR REDUCTION THROUGH PROCESSING**

By

PARVANEH HAJEB

**Thesis Submitted to the School of Graduate Studies, Universiti Putra Malaysia,
in Fulfilment of the Requirements for the Degree of Doctor of Philosophy**

June 2009



DEDICTED TO MY BELOVED FAMILY



Abstract of thesis presented to the Senate of Universiti Putra Malaysia in fulfilment of the requirements for the degree of Doctor of Philosophy

**TOTAL MERCURY AND METHYLMERCURY CONCENTRATION IN
FISH AND THEIR REDUCTION THROUGH PROCESSING**

By

PARVANEH HAJEB

June 2008

Chairman : Jinap Selamat, PhD

Faculty : Food Science and Technology

This research has been conducted to study the levels of total mercury and methylmercury, and their correlation in different marine fish species available for consumption in Peninsular Malaysia. Artificial methods have been used to remove mercury from fish. Method for methylmercury determination in fish samples was optimized using response surface methodology (RSM). Total mercury and methylmercury levels were determined using Cold vapor atomic absorption spectrophotometry (CV-AAS) and Gas chromatography-micro electron capture detector (GC- μ ECD), respectively. Samples of twelve species of common marine fish consumed by Malaysians were collected from local wholesale market in Malaysia. On the basis of total mercury and methylmercury levels measured in commonly



consumed fish, two species, long tail tuna and short-bodied mackerel identified with high mercury contents were sampled from east and west coast of Peninsular Malaysia. Methods for elimination of mercury in raw fish fillet has been developed using acidic solutions containing mercury chelating agents. The optimum conditions for methylmercury extraction were found by using acid concentration of 12.118 M, cysteine concentration of 2.375%, solvent volume of 1.5 ml, and extraction time of 35 min. Total mercury and methylmercury levels in fish samples studied were in the range of not detected to 1.010 and not detected to 0.914 $\mu\text{g/g}$ wet wt, respectively. The methylmercury to total mercury ratio ranged from 49.1% to 87.5%, with the highest ratio was in predatory fishes. All of the fish species showed strong positive correlation between methylmercury and total mercury levels ($R^2 > 0.86$). High levels of total mercury and methylmercury were detected in short-bodied mackerel and long tail tuna. Samples of these two species from east coast of Peninsular Malaysia showed higher levels of mercury compared to those from west coast. In all of the locations, significant positive correlations were found between fish body weight and mercury content. The industrial optimized method produced a solution which can remove mercury from raw fish fillet up to 91%. The optimum conditions for mercury reduction was achieved using cysteine concentration of 1.25 %, EDTA of 275 (mg/L), NaCl of 0.5 (%), pH of 3.75 and exposure time of 18 min. The home-used optimized protocol produced a solution which can remove mercury from raw fish fillet up to 81%. The overall optimal condition resulting to the maximum mercury removal in fish fillet was obtained at combined level of pH of 2.79, 0.5% NaCl, and 13.5 (min) of exposure time.



Abstrak tesis yang dikemukakan kepada Senat Universiti Putra Malaysia sebagai memenuhi keperluan untuk ijazah Doktor Falsafah

PENCEMARAN JUMLAH MERKURI DAN METILMERKURI DI DALAM IKAN DAN PENGURANGAN NYA SEWAKTU PEMROSESAN

Oleh

PARVANEH HAJEB

Jun 2008

Pengerusi: Jinap Selamat, PhD

Fakulti: Sains Makanan dan Teknologi

Penyelidikan ini telah dijalankan bagi mengkaji tahap jumlah merkuri dan metilmerkuri serta kaitannya dengan pengambilan pelbagai spesis ikan laut yang didapati di Semenanjung Malaysia. Satu kaedah yang diubahsuai telah digunakan bagi membuang merkuri dari ikan. Pemoptimuman keadah bagi pengesanan metilmerkuri di dalam ikan telah dijalankan dengan menggunakan keadah 'Response surface methodology' (RSM). Manakala tahap jumlah merkuri dan metilmerkuri masing-masing dikesan dengan menggunakan spektrofotometri serapan atom wap sejuk dan kuromatografi gas-penangkapan elektron. Dalam kajian ini sebanyak 12 spesis ikan laut yang biasa dimakan oleh penduduk Malaysia telah diambil dari pasar borong tempatan. Tahap jumlah merkuri dan metilmerkuri, nisbah metilmerkuri



dengan total merkuri (%MeHg) dan kaitannya dengan isi ikan serta jantungnya telah dikesan. Berdasarkan kepada keputusan yang diperolehi, didapati 2 spesis ikan yang biasa dimakan iaitu tongkol dan kembong telah dikenalpasti mengandungi kandungan merkuri yang tertinggi, dimana sampel tersebut telah diambil dari pantai timur dan barat Semenanjung Malaysia. Kaedah bagi menyingkirkan merkuri dari filet ikan mentah telah dibangunkan dengan menggunakan larutan berasid yang mengandungi agen pengkelat merkuri. Keadaan optimum bagi pengekstrakan metilmerkuri adalah kepekatan asid 12.118M, kepekatan cysteine 2.375, isipadu pelarut 1.5 ml dan masa pengekstrakan 35 min. Tahap jumlah merkuri dan metilmerkuri dalam ikan yang masing-masing dikaji adalah di antara julat tidak dikesan-1.010 $\mu\text{g/g}$ dan tidak dikesan-0.914 $\mu\text{g/g}$ berat basah. Manakala julat nisbah metilmerkuri dengan jumlah merkuri (%MeHg) adalah di antara 49.1% dan 87.5%, yang mana nisbah yang tertinggi adalah dari ikan-ikan pemangsa. Semua spesis ikan menunjukkan korelasi yang kuat secara positif di antara metilmerkuri dan tahap jumlah merkuri ($R^2 > 0.86$). Tahap jumlah merkuri dan metilmerkuri dikesan paling tinggi di dalam ikan tongkol dan ikan kembong. Berdasarkan kepada kedua-dua spesis ikan tersebut didapati spesis ikan dari pantai timur Semenanjung Malaysia mempunyai merkuri lebih tinggi berbanding dari pantai barat. Data dari semua lokasi pensampelan menunjukkan terdapat korelasi positif di antara berat badan ikan dan kandungan merkuri. Untuk penggunaan peringkat industri, larutan yang digarakan pada keadaan optimum bagi kepadat menyingkirkan merkuri dari filet ikan mentah sehingga 91%. Keadaan optimum bagi penyingkiran merkuri adalah kepekatan cysteine (1.25%), kepekatan EDTA (275 (mg/L)), kepekatan NaCl (0.5

%), pH (3.75) dan tempuh pendedahan (18 min). Manakala untuk penggunaan di rumah, lamtan yang digunakan pada keadaan optimum dapat mengeluarkan merkuri dari filet ikan mentah sebanyak 81%. Keadaan optimum bagi penyingkiran merkuri secara maksima dari fillet ikan diperolehi dengan menggabungkan tahap pH (2.79), kepekatan NaCl (0.5%) dan masa pendedahan (13.5min).

AKNOWLEDGEMENTS

My full praise to our God for enabling me to complete my study.

My sincere appreciation to my supervisor and chair person of the supervisory committee, Professor Dr. Jinap Selamat, who was a great source of motivation, encouragement and scientific guidance throughout the period of my study. I am also deeply indebted to her for arranging of the necessary funding.

I would like to express my deep thanks to my supervisory committee members, Professor Dr. Jamilah Bakar, Associate Professor Dr. Fatimah Abu Bakar and Associate Professor Dr. Ahmad Ismail, for their valuable contribution and suggestions. Special gratitude to Associate Professor Dr. Ahmad Ismail who provided me with an opportunity to use his well-equipped laboratory.

Thanks to The Ministry of Science, Technology and Innovation of Malaysia for sponsoring this research under EScience Fund project No. UPM0002449.

My thankfulness to Mr. Zulkifli from LKIM for his kind helps to collect fish samples from fish landings, Mr. Halim, Mr. Hamizan, Ms. Liza (Faculty of Food Science and Technology) for their helps during laboratory experiments.



I am also very much indebted to my dear friends Gisia, Elham, Maimunah, Afidah, Khairulnisak, Fatimah, Safzan, Farzad, Shahram, Dr. Yazdan and Dr. Hamed, for their support and being my friend.

My deepest appreciation and gratitude to my dear family members for their spiritual, financial and moral support. All of you are respected and loved for being there for me.



I certify that an Examination Committee met on 19 / 06 / 2009 to conduct the final examination of Parvaneh Hajeb on his PhD degree of Food Science thesis entitled “Concentration of mercury in fish muscle and its reduction through washing treatment” in accordance with Universiti Pertanian Malaysia (higher Degree) Act 1980 and Universiti Pertanian Malaysia (Higher Degree) Regulations 1981. The committee recommends that the candidate be awarded the relevant degree. Members of the Examination Committee are as follows:

AZIZAH ABDUL-HAMID, PhD

Associate Professor
Faculty of Food Science and Technology
Universiti Putra Malaysia
(Chairman)

SON RADO, PhD

Professor
Faculty of Food Science and Technology
Universiti Putra Malaysia
(Internal Examiner)

MD JELAS HARON, PhD

Professor
Faculty of Science
Universiti Putra Malaysia
(Internal Examiner)

SUKIMAN SARMANI, PhD

Professor
Faculty Science and Technology
Universiti Kebangsaan Malaysia
(External Examiner)

HASANAH MOHD. GHAZALI, Ph.D.

Professor/ Deputy Dean
School of Graduate Studies
Universiti Putra Malaysia

Date:



This thesis was submitted to the Senate of Universiti Putra Malaysia and has been accepted as fulfilment of the requirement for the degree of Doctor of Philosophy. The members of the Supervisory Committee were as follows:

Jinap Selamat, PhD

Professor
Faculty of Food Science and Technology
Universiti Putra Malaysia
(Chairman)

Jamilah Bakar, PhD

Professor
Faculty of Food Science and Technology
Universiti Putra Malaysia
(Member)

Fatimah Abu Bakar, PhD

Associate Professor
Faculty of Food Science and Technology
Universiti Putra Malaysia
(Member)

Ahmad Ismail, PhD

Associate Professor
Faculty Science
Universiti Putra Malaysia
(Member)

HASANAH MOHD. GHAZALI, PhD

Professor and Dean
School of Graduate Studies
Universiti Putra Malaysia

Date: 17 July 2009



DECLARATION

I declare that the thesis is my original work except for quotations and citations, which have been duly acknowledged. I also declare that it has not been previously and is not concurrently submitted for any other degree at UPM or at any other institutions.

PARVANEH HAJEB

Date:



TABLE OF CONTENTS

	Page
DEDICATION	ii
ABSTRACT	iii
ABSTRAK	v
ACKNOWLEDGEMENTS	viii
APPROVAL	xii
DECLARATION	xii
LIST OF TABLES	xvii
LIST OF FIGURES	xix
LIST OF APPENDICES	xxi
LIST OF ABBREVIATIONS	xxii
CHAPTER	
1 INTRODUCTION	1
Background of study	1
Importance of study	3
Objectives	4
2 LITERATURE REVIEW	5
Consumption of seafood in Malaysia	5
The nutritional considerations of fish	8
Fish proteins and its metal binding properties	10
Heavy metal in foods/ fish	14
Mercury and methylmercury	17
Physical and chemical properties of mercury	18
Mercury exposure and risk evaluation for humans	20
Mercury poisoning	22
Mercury species and their transformation in aquatic environment	27
Mercury studies on seafood in Malaysia	29
Methods of mercury removal in food	34
Mercury chelating agents and their mechanisms	35
Cysteine	39
Acids	42
Citric acid	44
Ethylenediaminetetraacetic acid (EDTA)	47
Methods of chemical analysis of mercury in food/ fish	50
Speciation of mercury	51
Chromatographic separation of mercury species	55
High performance liquid chromatography (HPLC)	55
Gas chromatography (GC)	58
Capillary electrophoresis (CE)	60



	Spectrometric detection	61
	Cold vapor atomic absorption spectrophotometry (CV-AAS)	61
	Cold vapor atomic Fluorescence spectrophotometry (CV-AFS)	62
	Atomic Emission Spectrometry (AES)	63
	Microwave induced plasma-atomic emission spectrometry (MIP-AES)	63
	Inductively coupled plasma-mass spectrometry (ICP-MS)	63
	Other detectors used for mercury determination	65
	UV	65
	Mass spectrometry (MS)	65
	Electron capture detector (ECD)	66
	Response surface methodology (RSM) and product optimization	67
3	METHOD OPTIMIZATION FOR METHYLMERCURY DETERMINATION IN FISH SAMPLES USING GC-μECD	70
	Introduction	70
	Materials and methods	72
	Chemicals and materials	72
	Instrumentation	73
	Calibration	75
	Experimental design	75
	Methylmercury extraction in fish samples	77
	Limit of detection (LOD) and limit of quantification (LOQ)	78
	Statistical analysis	78
	Model validation	80
	Results and discussion	80
	Fitting the models	80
	Interpretation of response surface model	84
	Validation of the procedure and application to real samples	89
	Conclusions	90
4	CORRELATION BETWEEN TOTAL MERCURY AND METHYLMERCURY CONCENTRATIONS IN COMMONLY CONSUMED MARINE FISH SPECIES IN MALAYSIA	91
	Introduction	91
	Materials and methods	94
	Chemicals and materials	94
	Fish samples	95
	Instrumentation	96
	Calibration	97
	Total mercury extraction in fish samples	97



	Methylmercury extraction in fish samples	98
	Recovery of total mercury and methylmercury in fish samples	98
	Limit of detection (LOD) and limit of quantification (LOQ)	99
	Estimated weekly intake of total mercury and methylmercury	99
	Statistical analysis	100
Results and discussion		100
	Recovery, Limit of detection (LOD) and limit of quantification (LOQ)	100
	Total mercury and methylmercury concentration in muscle and liver of fish species	101
	Correlation between total mercury and methylmercury in fish species	107
	Total mercury and methylmercury concentration in muscle and liver of short-bodied mackerel and long tail tuna from the east and west coasts of Peninsular Malaysia	112
	Correlation between total mercury and methylmercury in two fish species from the east and west coasts of Peninsular Malaysia	120
	Correlation between total mercury level and fishes weight	124
	Assessment of total mercury and methylmercury exposure in Malaysians	127
Conclusions		129
5	EFFECTS OF WASHING PRETREATMENTS ON THE MERCURY CONCENTRATION (INDUSTRIAL PROTOCOL)	131
	Introduction	131
	Materials and methods	134
	Chemicals and materials	134
	Instrumentation	135
	Calibration	135
	Experimental design	136
	Sample preparation	137
	Mercury extraction in fish samples	139
	EDTA extraction in fish samples	139
	Verification of the model	139
	Limit of detection (LOD) and limit of quantification (LOQ)	140
	Recovery of total mercury in fish samples	140
	Statistical analysis	140
Results and discussion		142
	Recovery, Limit of detection (LOD) and limit of	142

	quantification (LOQ)	
	Fitting the response surface models	142
	Interpretation of response surface model	146
	The main effect of independent variables	148
	The interaction effect of independent variables	152
	Optimization of mercury reduction in fish samples	152
	Conclusions	153
6	EFFECTS OF WASHING PRETREATMENTS ON THE MERCURY CONCENTRATION (HOME-USED PROTOCOL)	155
	Introduction	155
	Materials and methods	156
	Chemicals and materials	156
	Instrumentation	156
	Calibration	157
	Experimental design	157
	Sample preparation	158
	Mercury extraction in fish samples	159
	Verification of the model	159
	Limit of detection (LOD) and limit of quantification (LOQ)	160
	Recovery of total mercury in fish samples	160
	Statistical analysis	160
	Results and discussion	162
	Recovery, Limit of detection (LOD) and limit of quantification (LOQ)	162
	Fitting the response surface models	162
	Interpretation of response surface model	164
	Optimization of mercury reduction in fish samples	168
	Conclusions	168
7	SUMMARY, GENERAL CONCLUSION AND RECOMMENDATION FOR FUTURE RESEARCH	170
	REFERNCES	176
	APPENDICES	210
	BIODATA OF THE STUDENT	220
	LIST OF PUBLICATIONS	221



LIST OF TABLES

Table		Page
2.1	Fish Consumption and Trade, 2000.	5
2.2	Top 10 seafood consumption by country.	6
2.3	Mercury level in fish/biota muscle, liver and heart from Malaysia reported by other studies.	32
2.4	Stability constants ($\log k_1$ (mol l^{-1}) at 25 °C) for methylmercury (MeHg^+) with inorganic and organic ligands.	38
3.1	GC-micro ECD operating condition for methylmercury determination in fish.	74
3.2	GC-MS operating condition for methylmercury determination in fish.	74
3.3	Uncoded and coded independent variables used in RSM design for recovery of methylmercury in BCR-463.	76
3.4	Experimental points of the Central Composite Design for recovery of methylmercury in BCR-463.	77
3.5	Design matrix, experimental values and predicted values in the screening design for recovery of methylmercury in BCR-463.	81
3.6	Analysis of variance of the regression coefficients of the fitted quadratic equations for recovery of methylmercury in BCR-463.	83
3.7	The calculated optimum points for recovery of methylmercury in BCR-463.	89
3.8	Methylmercury concentration in muscle tissue of different fish sample.	90
4.1	Recovery of total mercury and methylmercury using CRM (BCR-463).	101
4.2	Characteristics of the fish samples collected from local market in Peninsular Malaysia.	103
4.3	Total mercury and methylmercury concentrations in liver and muscle of fish species collected from local market in Peninsular Malaysia.	104

4.4	Total mercury and methylmercury level ($\mu\text{g/g}$ wet wt.) in long tail tuna and short-bodied mackerel from the east and west coast of the Peninsular Malaysia.	113
4.5	Total mercury and methylmercury levels ($\mu\text{g/g}$ wet wt) in tuna and mackerel species from other studies.	118
5.1	Operating condition for HPLC-PDA for EDTA determination.	135
5.2	Uncoded and coded independent variables used in RSM design for mercury reduction.	137
5.3	Experimental points of the Central Composite Design for mercury reduction.	138
5.4	pH adjustment using different concentration of HCl and NaOH.	138
5.5	Design matrix, experimental values and predicted values in the screening design for mercury reduction.	144
5.6	Analysis of variance of the regression coefficients of the fitted quadratic equations for mercury reduction.	145
5.7	Optimum conditions, predicted and experimental value of mercury removal at that condition.	153
6.1	Uncoded and coded independent variables used in RSM design for mercury removal.	157
6.2	Design matrix, experimental values and predicted values in the screening design for mercury removal.	158
6.3	pH adjustment using different concentration of citric acid and NaOH.	159
6.4	Design matrix, experimental values and predicted values in the screening design for mercury removal.	163
6.5	Analysis of variance of the regression coefficients of the fitted quadratic equations for mercury removal..	164
6.6	Optimum conditions, predicted and experimental value of mercury removal at that condition.	168

LIST OF FIGURES

Figure		Page
2.1	Domain nature of metallothionein. Four and three atoms of cadmium (Cd) are coordinated in α - and β -domains of metallothionein, respectively.	13
2.2	Chelation of metals by citric acid.	46
2.3	Structural formula of EDTA.	48
2.4	Metal-EDTA chelate.	49
3.1	Calibration curve for methylmercury standard solutions detected by GC- μ ECD.	75
3.2	Three-dimensional response surface showing the effect of the extraction time, sulfuric acid, cysteine concentration and toluene volume on recovery of methylmercury in certified reference material BCR-463.	88
4.1	Map of fish sampling locations in Peninsular Malaysia.	96
4.2	Calibration curve for total mercury standard solutions using CV-AAS.	97
4.3	Correlation between total mercury and methylmercury in different fish species.	109
4.4	Correlation between total mercury and methylmercury levels in the muscle of short-bodied mackerel from (A) Kuantan, (B) Kuala Perlis, (C) Chendring.	122
4.5	Correlation between total mercury and methylmercury levels in the muscle of long tail tuna from (A) Kuantan, (B) Kuala Perlis, (C) Chendring.	123
4.6	Relationship between body weight and mercury level in the muscle of long tail tuna from (A) Kuantan, (B) Kuala Perlis, (C) Chendring. (○) Total mercury and (●) methylmercury levels.	125
4.7	Relationship between body weight and mercury level in the muscle of short-bodied mackerel from (A) Kuantan, (B) Kuala Perlis, (C) Chendring. (○) Total mercury and (●) methylmercury levels.	126

4.8	Estimated weekly intake (EWI) of total mercury and methylmercury ($\mu\text{g}/\text{kg}$ body wt.) of different fish species.	129
5.1	Calibration curve for EDTA standard solutions using HPLC-PDA.	136
5.2	Three-dimensional response surface showing the effect of the pH, cysteine concentration, salt concentration, EDTA concentration and exposure time on removal of mercury in fish fillet.	147
6.1	Three-dimensional response surface plots showing the effect of the pH, salt concentration and exposure time on removal of mercury in fish fillet.	166



LIST OF APPENDICES

APPENDIX		Page
A1	A: Typical GC- μ ECD chromatogram of methylmercury standard calibrant solution (1 ng ml ⁻¹); B: GC- μ ECD chromatogram of methylmercury in BCR-463 (3.04 μ g g ⁻¹).	210
A2	Total ion chromatogram for methylmercury chloride in BCR-463 (3.04 μ g g ⁻¹), obtained by GC-MS.	211
A3	Contour plots showing the effect of the extraction time, sulfuric acid and cysteine concentration and toluene volume on recovery of methylmercury in certified reference material BCR-463.	212
B1	A: Typical HPLC-PDA chromatogram of EDTA standard calibrant solution (100 μ g/mL); B: HPLC-PDA chromatogram of EDTA in fish tissue.	213
B2	Fish samples (Short-bodied mackerel) used for mercury reduction treatments in chapter 5 & 6.	214
B3	Fish fillets (Short-bodied mackerel) used for mercury reduction treatments in chapter 5 & 6.	215
B4	Mercury reduction treatments (chapter 6 & 7)	216
B5	Contour plots showing the effect of the pH, cysteine concentration, salt concentration, EDTA concentration and exposure time on removal of mercury in fish fillet.	217
C1	Contour plots showing the effect of the pH, salt concentration and exposure time on removal of mercury in fish fillet.	219



LIST OF ABBREVIATIONS

Ag	Silver
AES	Atomic Emission Spectrometry
ANOVA	Analysis of variance
AOAC	Association of Official Analytical Chemists
Bi	Bismuth
Br⁻	Bromide
CaNa₂EDTA	Calcium disodium ethylenediaminetetraacetate
CH₃Hg⁺	Methylmercury
CH₃HgCl	Methylmercury chloride
(CH₃)₂Hg	Dimethylmercury
C₆H₅Hg⁺	Phenylmercury
CCD	Central composite design
CCFAC	Codex Committee on Foods Additives and Contaminants
Cd	Cadmium
CE	Capillary electrophoresis
CP	Center point
Cl	Calcium
Cr	Chromium
CRM	Certified reference materials
CV-AAS	Cold vapor atomic absorption spectrophotometry
CV-AFS	Cold vapor atomic Fluorescence spectrophotometry



DMPS	2,3 dimercaptopropane-1-sulfonate
DMSA	Dimercaptosuccinic acid
DPA	D-penicillamine
ECD	Electron capture detector
EDTA	Ethylenediaminetetraacetic acid
EPA	Environmental Protection Agency
Eq	Equation
EWI	Estimated weekly intake
F⁻	Fluoride
FAO	Food and Agricultural Organization
FDA	Food and drug analysis
g	Gram
GC	Gas chromatography
GC-AFS	Gas chromatography-atomic fluorescence spectrometry
GC-ICP-MS	Gas chromatography-inductively coupled plasma-mass spectrometry
GC-MS	gas chromatography-mass spectrometry
g/s	Gram/ second
h	Hour
HCl	Hydrochloric acid
Hg	Mercury
HNO₃	Nitric acid
H₂SO₄	Sulfuric acid
HPLC	High performance liquid chromatography



HPLC-MS	High performance liquid chromatography-mass spectrometry
I	Iodide
ICP-MS	Inductively coupled plasma-mass spectrometry
IUPAC	International Union for Pure and Applied Chemistry
JECFA	Joint Expert Committee on Food Additives
KBr	Potassium bromide
Kg	Kilogram
L	Liter
LC	liquid chromatography
LKIM	Lembaga Kemajuan Ikan Malaysia
LOD	Limit of detection
LOQ	limit of quantification
M	Molar
mg	Milligram
MeHg	methylmercury
MeOH	Methanol
min	Minute
MIP-AES	Microwave induced plasma-atomic emission spectrometry
mL	Milliliter
MOH	Ministry of Health Malaysia
MS	Mass spectrometry
MT	Metallothionein