



UNIVERSITI PUTRA MALAYSIA

**RADIATION – INDUCED POLYMERIZATION OF POLYMETHACRYLIC
GEL FOR DOSIMETRY**

ISKANDAR SHAHRIM MUSTAFA

FS 2006 9

**RADIATION-INDUCED POLYMERIZATION OF
POLYMETHACRYLIC GEL FOR DOSIMETRY**

By

ISKANDAR SHAHRIM MUSTAFA

**Thesis Submitted to the School of Graduate Studies, Universiti Putra Malaysia,
in Fulfilment of the Requirement for the Degree of Master of Science**

March 2006



To:

My beloved mom, Siti Ishah bt Md. Hanafi

Thanks for the encouragement, love and support in fulfilling my endeavour...



Abstract of thesis presented to the Senate of Universiti Putra Malaysia in
fulfilment of the requirement for the degree of Master of Science

**RADIATION-INDUCED POLYMERIZATION OF
POLYMETHACRYLIC GEL FOR DOSIMETRY**

By

ISKANDAR SHAHRIM MUSTAFA

March 2006

Chairman : Associate Professor Elias Saion, PhD

Faculty : Science

The aim of the present work is to investigate the effect of radiation on methacrylic acid (MAA) crosslinked with N, N'-methylene-bisacrylamide (BIS) for the purpose of developing a polymer gel dosimeter, namely, polymethacrylic gel (PMAAG). PMAAG of different concentrations of MAA and BIS were irradiated with the absorbed doses ranging from 0 Gy to 19 Gy. Using Raman spectrophotometer, five peak intensities were identified and represents the assigned bonding for C=C (2238 cm^{-1}) of MAA, C=C (1634 cm^{-1}) of BIS, C-C (2110 cm^{-1}) of PMAA, C=O (1795 cm^{-1}) of PMAA, CH₂ (2905 cm^{-1}) of PMAA, which have strong modes of vibration in the band positions. The Raman peak intensity y as a function of absorbed dose D was found to have a

monoexponential expression in the form; $y = y_0 + A \left(1 - e^{-D/D_0} \right)$. The dose sensitivity D_0 and half dose $D_{1/2}$ were obtained and found to increase with the concentrations of MAA monomer and BIS crosslinker. The dose correlation factor of BIS, k_{BIS} is always higher than k_{MAA} of MAA ($k_{\text{BIS}} > k_{\text{MAA}}$) for all C-C, C=O and CH₂ stretching modes, indicating BIS is a very significant parameter in the polymerization process. Additionally, a Nuclear magnetic resonance (NMR) spectrometer was used in order to measure the relaxation time of polymethacrylic (PMAA) gel dosimeters. The relaxation rate (ΔR_2) dose sensitivity value (12.5 ± 0.1 Gy) of MAA monomer by Lepage, et al 2001 is comparable with PMAAG experimental value gained which are 12.6 ± 0.1 Gy. A comparison between and indirect measurements of polymer formed along with consumptions of monomer and crosslinker are also obtained from this experimental work. Finally, a UV-Vis spectrophotometer was used to records PMAAG degree of absorption in order to determine the accuracy of PMAAG. The PMAAG has a mean value of absorption of 0.614 at 375 nm. The dose derived from PMAAG is comparable to Fricke dosimeter and ionization chamber readings between $4.7 \pm 0.1\%$ and $11.6 \pm 0.1\%$. The dose errors of less than $10 \pm 0.1\%$ are considered acceptable in radiation processing, an improvement of accuracy less than $5.0 \pm 0.1\%$ is acceptable in radiotherapy.

Abstrak tesis ini yang dikemukakan kepada Senat Universiti Putra Malaysia sebagai memenuhi keperluan untuk ijazah Master Sains

**RADIASI MENGARUH PEMPOLIMERAN GEL POLIMETAKRILIK
UNTUK DOSIMETRI**

Oleh

ISKANDAR SHAHRIM MUSTAFA

Mac 2006

Pengerusi : Professor Madya Elias Saion, PhD

Fakulti : Sains

Matlamat kajian pada masa ini adalah untuk menyiasat kesan terhadap asid metakrilik (MAA) yang bertaut-silang dengan “N,N'-methylene-bisacrylamide” (BIS) dengan tujuan untuk pembangunan meterdos gel polimer, gel polimetakrilik (PMAAG). PMAAG dengan berlainan kepekatan MAA dan BIS disinarkan dengan dos terserap dari 0 Gy hingga 19 Gy. Dengan menggunakan spektrofotometer Raman, keamatan lima puncak dikenalpasti dan mewakili ikatan terhadap C=C (2238 cm^{-1}) dari MAA, C=C (1634 cm^{-1}) dari BIS, C-C (2110 cm^{-1}) dari PMAA, C=O (1795 cm^{-1}) dari PMAA, CH₂ (2905 cm^{-1}) dari PMAA, yang mempunyai getaran mod kuat di dalam kedudukan ikatan. Keamatan puncak Raman γ sebagai fungsi dos terserap D didapati mempunyai ungkapan monoeksponen dalam bentuk

$y = y_0 + A \left(1 - e^{-D/D_0} \right)$. Kepekatan dos D_0 dan dos separa $D_{1/2}$ diperolehi dan didapati meningkat dengan kepekatan monomer MAA dan taut-silang BIS. Faktor sekaitan dos dari BIS, k_{BIS} sentiasa lebih tinggi daripada k_{MAA} dari MAA ($k_{BIS} > k_{MAA}$) untuk semua mod regangan C-C, C=C, C=O dan CH₂, menunjukkan BIS adalah parameter yang sangat bererti di dalam proses pempolimeran. Secara tambahan, sebuah spektrometer salunan magnet nuklear (NMR) telah digunakan, bertujuan untuk mengukur masa santaian meterdos gel polimetakrilik (PMAA). Nilai kepekaan dos (12.5 ± 0.1 Gy) kadar santaian (ΔR_2) dari monomer MAA oleh Lepage, et al 2001 adalah setanding dengan nilai yang diperoleh dari uji kaji PMAAG iaitu 12.6 ± 0.1 Gy). Perbandingan antara pengukuran langsung dengan tidak langsung pembentukan polimer dengan penggunaan monomer dan taut-silang juga diperolehi dari kerja uji kaji ini. Akhirnya, sebuah spektrofotometer UV-Vis digunakan untuk merekod darjah serapan PMAAG untuk mengenalpasti ketepatan PMAAG. PMAAG mempunyai nilai penyerapan min 0.614 pd 375 nm. Dos yang diperoleh dari PMAAG adalah setanding dengan bacaan meterdos Fricke dan kebuk pengionan di antara $4.7 \pm 0.1\%$ dan $11.6 \pm 0.1\%$. ralat dos yang kurang daripada $10.0 \pm 0.1\%$ adalah dipertimbangkan kebolehtenerimaan di dalam pemprosesan sinaran, pemberian ketepatan kurang dari $5.0 \pm 0.1\%$ adalah boleh diterima di dalam radioterapi.

ACKNOWLEDGEMENTS

All the praise and admiration for Allah, the Almighty, Beneficial and the most Merciful, who has enabled me to submit this thesis.

The authors gratefully would like to acknowledge the following people and organizations for all their advice, help and support, assists, suggestions, motivations and encouragement throughout graduate career:

My beloved mother, Siti Ishah Md. Hanafi; my supervisor and mentor, Assoc. Prof. Dr. Elias Saion from Physics Department, Faculty of Science, Universiti Putra Malaysia and staff; Dr Jumiah Hassan from Physics Department, Faculty of Science, Universiti Putra Malaysia; Assoc. Prof Dr. Mohamad Zaki Abdul Rahman from Chemistry Department, Faculty of Science, Universiti Putra Malaysia; Dr. Noriah and Mr. Taiman Kadni from Secondary Standard Dosimetry Laboratory (SSDL) Malaysian Institute for Nuclear Technology Research (MINT) and staff. Last but not least my respectful colleagues: Nordiana Sadikun, Mohd Saiful Bakhtiar Laui, Zainal Hafiz Ramly, Mohd Mohd Asri Mat Teridi, Hamezan Muhammad, Hamzah Harun, Azian Othman, Nuraihan Harun, Norazimah Yusoff, Azlina Baha, Ajis Lepit, Yousoff Mohammed, Mohammad Ahmed, Aris Doyan and Susilawati.

I certified that an Examination Committee has met on 16th March 2006 to conduct the final examination of Iskandar Shahrim bin Mustafa on his Master of Science thesis entitled "Radiation-Induced Polymerization of Polymethacrylic Gel for Dosimetry" in accordance with Universiti Pertanian Malaysia (Higher Degree) Act 1980 and Universiti Pertanian Malaysia (Higher Degree) Regulation 1981. The committee recommends that the candidate be awarded the relevant degree. Members of the Examination Committee are as follows:

ZAINAL ABIDIN TALIB, PhD

Associate Professor

Faculty of Science

Universiti Putra Malaysia

(Chairman)

ZAINAL ABIDIN SULAIMAN, PhD

Associate Professor

Faculty of Science

Universiti Putra Malaysia

(Internal Examiner)

ABDUL HALIM SHAARI, PhD

Professor

Faculty of Science

Universiti Putra Malaysia

(Internal Examiner)

AHMAD SHUKRI MUSTAFA KAMAL, PhD

Professor

Institute of Graduate Studies

Universiti Sains Malaysia

(External Examiner)

HASANAH MOHD. GHAZALI, PhD

Professor/Deputy Dean

School of Graduate Studies,

Universiti Putra Malaysia

Date:

This thesis submitted to the senate of Universiti Putra Malaysia and has been accepted as fulfilment of the requirement for the degree of Master of Science. The members of the Supervisory Committee are as follows:

ELIAS SAION, PhD

Associate Professor

Faculty of Science

Universiti Putra Malaysia

(Chairman)

MOHD ZAKI ABDUL RAHMAN, PhD

Associate Professor

Faculty of Science

Universiti Putra Malaysia

(Member)

JUMIAH HASSAN, PhD

Lecturer

Faculty of Science

Universiti Putra Malaysia

(Member)

TAIMAN KADNI, MSc

Manager

Secondary Standard Dosimetry Laboratory

Malaysian Institute for Nuclear Technology Research (MINT)

(Member)

AINI IDERIS, PhD

Professor/Dean

School of Graduate Studies

Universiti Putra Malaysia

Date:

DECLARATION

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

ISKANDAR SHAHRIM MUSTAFA

Date:

TABLE OF CONTENTS

	Page
DEDICATION	ii
ABSTRACT	iii
ABSTRAK	v
ACKNOWLEDGEMENTS	vii
APPROVAL	viii
DECLARATION	x
LIST OF TABLES	xv
LIST OF FIGURES	xvi
LIST OF ABBREVIATIONS/SYMBOL	xxiv
LIST OF GLOSSARY	xxvi
 CHAPTER	
1. INTRODUCTION	1.1
1.1. Scope of the present study	1.5
1.2. Problem statement	1.7
1.3. Objective of research	1.9
1.4. Significance of study	1.9
1.5. Outline of the thesis	1.10
2. LITERATURE REVIEW	2.1
2.1. Historical background of ionizing radiation in radiotherapy	2.1
2.2. Cobalt-60 dosimetry in radiation research and medical radiation therapy	2.2
2.3. Dosimetry	2.4
2.3.1. Ionization chamber	2.5
2.3.2. Liquid chemical system	2.6
2.3.2.1.The Fricke dosimeter	2.6
2.3.2.2.Polymer gel dosimeter	2.8
2.3.3. Related work in dosimetry	2.10
2.3.4. Chemical mechanism of polymer gel dosimeters	2.13
2.3.5. Macroscopic response of polymer gel dosimeters	2.16
2.3.6. Tissue equivalence	2.20
3. THEORY	3.1
3.1. Ionizing radiations	3.1
3.2. Interaction of gamma ray with matter	3.2
3.2.1. Rayleigh scattering	3.4
3.2.2. Photoelectric absorption	3.5
3.2.3. Compton scattering	3.7

3.2.4. Pair production	3.11
3.2.5. Total absorption coefficient	3.12
3.2.6. Absorbed dose	3.14
3.3. Radiation effects	3.14
3.3.1. Ionization and Excitation	3.14
3.3.2. Radiolysis	3.17
3.3.3. Recombination	3.17
3.3.4. Polymerization	3.18
3.3.5. Crosslinking	3.19
3.3.6. Chain Scission	3.19
3.3.7. Role of Radical Scavengers and Oxygen	3.20
3.4. UV-Visible spectroscopy	3.21
3.4.1. Electronic transitions in molecules	3.22
3.4.2. Optical absorption spectra	3.24
3.4.3. The Beer-Lambert's law	3.25
3.5. Absorbed dose of dosimetry	3.27
3.6. Raman spectroscopy	3.30
3.6.1. Inelastic scattering of light	3.30
3.6.2. The theory of Raman effect	3.33
3.7. Nuclear Magnetic Resonance	3.38
3.7.1. Magnetic properties of atomic nuclei	3.39
3.7.2. Magnetization	3.41
3.7.3. Spin-lattice relaxation time (T1)	3.44
3.7.4. Spin-spin relaxation time (T2)	3.46
4. MATERIALS AND METHODS	4.1
4.1. Sample preparation	4.1
4.1.1. Fricke dosimeter preparation	4.1
4.1.2. Polymer gel (Polymethacrylic - PMAA) Preparation	4.2
4.2. Irradiation of polymethacrylic (PMAA) Gel Dosimeters	4.7
4.3. Characterizations of PMAAG	4.12
4.3.1. UV-Visible spectroscopy	4.13
4.3.2. Raman spectroscopy	4.16
4.3.3. Nuclear Magnetic Resonance (NMR) Spectroscopy	4.20
5. RESULTS AND DISCUSSIONS	5.1
5.1. Polymerization Process of PMAAG	5.1
5.2. Characterization of PMAAG by Raman Spectroscopy	5.5
5.2.1. Identification of Raman Peaks	5.5
5.2.2. Increment of C-C stretching mode versus % MAA	5.8
5.2.3. Dose sensitivity, D_0 of C-C stretching mode versus % MAA	5.12

5.2.4. Half dose, $D_{1/2}$ of C-C stretching mode versus % MAA	5.13
5.2.5. Increment of C-C stretching mode versus % BIS	5.14
5.2.6. Dose sensitivity, D_0 of C-C stretching mode versus % BIS	5.18
5.2.7. Half dose, $D_{1/2}$ of C-C stretching mode versus % BIS	5.19
5.2.8. Increment of C=O stretching mode versus % MAA	5.20
5.2.9. Dose sensitivity, D_0 of C=O stretching versus % MAA	5.23
5.2.10. Half dose, $D_{1/2}$ of C=O stretching mode versus MAA	5.24
5.2.11. Increment of C=O stretching mode versus % BIS	5.26
5.2.12. Dose sensitivity, D_0 of C=O stretching versus % BIS	5.29
5.2.13. Half dose, $D_{1/2}$ of C=O stretching mode versus % BIS	5.30
5.2.14. Increment of CH ₂ stretching mode versus % MAA	5.31
5.2.15. Dose sensitivity, D_0 of CH ₂ stretching versus % MAA	5.34
5.2.16. Half dose, $D_{1/2}$ CH ₂ stretching mode of PMAA at constant MAA	5.35
5.2.17. Increment of CH ₂ stretching mode versus % BIS	5.36
5.2.18. Dose sensitivity, D_0 of CH ₂ stretching versus % BIS	5.39
5.2.19. Half dose, $D_{1/2}$ of CH ₂ stretching mode versus % BIS	5.40
5.3. Consumptions of MAA and BIS	5.42
5.3.1. Decrement of C=C stretching of monomer	5.42
5.3.2. Dose sensitivity, D_0 of C=C stretching of MAA	5.45
5.3.3. Half dose, $D_{1/2}$ of C=C stretching of MAA	5.47
5.3.4. Decrement of C=C stretching of BIS	5.48
5.3.5. Dose sensitivity, D_0 of C=C stretching of BIS	5.51
5.3.6. Half dose, $D_{1/2}$ of C=C stretching of BIS at constant MAA	5.52
5.4. Raman correlation factors of BIS and MAA	5.54
5.5. Characterizations of PMAAG by NMR Relaxation Method	5.56
5.5.1. Increment of Spin-lattice relaxation rate (ΔR_1) at constant MAA	5.57
5.5.2. Dose sensitivity D_0 of relaxation rate (ΔR_1) at constant MAA	5.60
5.5.3. Half dose $D_{1/2}$ of relaxation rate (ΔR_1) at constant MAA	5.61
5.5.4. Increment of Spin-lattice relaxation rate (ΔR_1) at constant BIS	5.63
5.5.5. Dose sensitivity D_0 of relaxation rate (ΔR_1) at constant BIS	5.65
5.5.6. Half Dose $D_{1/2}$ of relaxation rate (ΔR_1) at constant BIS	5.66
5.5.7. Increment of Spin-spin relaxation rate (ΔR_2) at constant MAA	5.68

5.5.8. Dose sensitivity D_0 of relaxation rate (ΔR_2) at constant MAA	5.72
5.5.9. Half Dose $D_{1/2}$ of relaxation rate (ΔR_2) at constant MAA	5.73
5.5.10. Increment of Spin-spin relaxation rate (ΔR_2) at constant BIS	5.74
5.5.11. Dose sensitivity D_0 of relaxation rate (ΔR_2) at constant BIS	5.78
5.5.12. Half Dose $D_{1/2}$ of relaxation rate (ΔR_2) at constant BIS	5.79
5.6. NMR correlation factors for BIS and MAA	5.80
5.7. Characterization PMAAG by UV-Visible Absorption Method	5.85
5.7.1. Polymer gels used as point dosimeters	5.85
5.7.2. Dose Calculation of PMAAG	5.93
6. CONCLUSIONS AND FURTHER WORKS	6.1
6.1. Raman spectra and structure characterizations	6.2
6.2. Nuclear Magnetic Resonance (NMR) relaxation time	6.4
6.3. UV-Visible absorptions and standard accuracy	6.5
6.4. Suggestion for future work	6.6
BIBLIOGRAPHY	R.1
APPENDICES	A.1
BIODATA OF THE AUTHOR	B.1
LIST OF PUBLICATIONS/SEMINARS/POSTERS	L.1

LIST OF TABLES

Table		Page
2.1	Dose sensitivity of different polymer gel dosimeters (data obtain from Lepage, <i>et al</i> 2001).	2.20
2.2	Elemental compositions of several tissues and tissue-equivalent PAG gels (6%T/50%C). Mass and electron density are also given.	2.21
4.1	Various mixture of PMAAG at 0% BIS.	4.4
4.2	Various mixture of PMAAG at 2% BIS.	4.5
4.3	Various mixture of PMAAG at 3% BIS.	4.5
4.4	Various mixture of PMAAG at 4% BIS.	4.6
4.5	Various mixture of PMAAG at 5% BIS.	4.6
5.1	Values of parameter, determine from dose absorbance equation.	5.94

LIST OF FIGURES

Figure		Page
1.1	Macromolecular configuration of polymers.	1.7
1.2	Various polymer arrangements.	1.7
2.1	Schematic drawing illustrating the creation of radiolytic products of water by ionising irradiation. The radiolytic particles are created in spurs after which they diffuse and may react with other molecules (a). As a result, the interaction radius increases with time.	2.11
3.1	The photoelectric photon absorption cross-section in lead (Leo, 1994).	3.7
3.2	Schematic diagram of Compton scattering.	3.9
3.3	Energy spectrum of electrons recoiling after Compton scattering, for various energy of the incident photon. The sharp maximum electron recoil energy is known as the <i>Compton edge</i> (Leo 1994).	3.10
3.4	Contribution of gamma-ray interactions in a medium (Perkins, 1987).	3.13
3.5	The possible transitions of π , σ and η orbital electron in molecules.	3.24
3.6	Simple model illustrating Stokes and Anti-Stokes Raman scattering.	3.34
3.7	Energy level diagrams illustrating Raman scattering, with incoming radiation on the left, scattered radiation emergent on the right. Only energy levels directly involved are depicted: (a) Stokes Raman transition and (b) Anti-Stokes Raman transition.	3.36

3.8	The nuclear will be able to absorb radiowaves in both the strong and weak magnetic field. However, the absorption occurs at a field strength dependent frequency which is higher in the strong magnetic field than in the weak magnetic field.	3.40
3.9	Stable (a) and unstable (b) position of the magnet	3.41
3.10	Nuclei submitted to an external magnetic field are precessing on two opposite cones, (a) at the equilibrium, a slight excess of spins are precessing around the external field direction B_0 (b) equivalent to have one stationary magnetic moment, M directed along B_0	3.44
3.11	Transverse relaxation phenomena induce an increasing dephasing of individual spins in order to observe a progressive decrease of the macroscopic magnetisation.	3.46
4.1	Polymer gel dosimeters containing methacrylic acid (MAA), N, N'-methylene-bisacrylamide (BIS), gelatine, and deionized water in 5 ml ampoule tubes sealed with parafilm tape.	4.4
4.2	Cobolt-60 gamma rays chamber of Eldorado type located at Secondary Standard Dosimetry Laboratory (SSDL), Malaysia Institute for Nuclear Technology (MINT) in Bangi, Selangor.	4.8
4.3	Perspex sample holder.	4.9
4.4	Sample holder positioned 5 cm from Field Surface (FS) in water phantom.	4.10
4.5	Eldorado's control panel. The electrometer and barometer were used during the procedure in calibrations of ionization chamber and Fricke dosimeter	4.11
4.6	Ionization Chamber (NE2571 #1028)	4.11

4.7	Electrometer – PTW UNIDOS	4.12
4.8	A double beam UV-VIS spectrophotometer (Camspec M350)	4.15
4.9	Quartz cuvette with a 1 cm diameter.	4.16
4.10	Raman spectrometer (RSI 2001G, Raman System, Inc.) equipped with 532 nm solid state diode.	4.18
4.11	Nuclear magnetic resonance spectroscopy which consist of magnet module and control module.	4.20
5.1	Chemical structures of (a) Methacrylic Acid (MAA); (b) N, N'-methylene-bisacrylamide (BIS); (c) Polymethacrylic (PMAA).	5.2
5.2	Initiation of chemical structure (a) methacrylic acid (MAA); (b) N, N'-methylene-bisacrylamide (BIS); (c) propagation of polymethacrylic (PMMA).	5.4
5.3	Progression in polymer structure as a function of initial crosslinker concentration. (a) A polymer solely composed of MAA alone, or BIS alone (b) PMAAG composed of low initial BIS concentration, (c) PMMAG composed of high initial BIS concentration.	5.4
5.4	Raman peak of PMAAG assigned the stretching of (a) C=C (2238 cm^{-1}) of MAA, (b) C=C (1634 cm^{-1}) of BIS, (c) C-C (2110 cm^{-1}) of PMAA, (d) C=O (1795 cm^{-1}) of PMAA, (e) CH ₂ (2905 cm^{-1}) of PMAAG.	5.8
5.5	Normalised Raman intensity of C-C stretching showing the formation of PMAAG at (a) 2%, (b) 3%, (c) 4%, and (4) 5% MAA and for different BIS concentrations.	5.11
5.6	Correlation between D_0 and the initial concentration of MAA for different BIS concentration for the formation of PMAAG due to C-C stretching at 2110 cm^{-1} .	5.13

5.7	Correlation between $D_{1/2}$ and the initial concentration of MAA for different BIS composition for the formation of PMAAG due to C-C stretching at 2110 cm^{-1} .	5.14
5.8	Normalised Raman intensity of C-C stretching showing the formation of PMAAG at (a) 2%, (b) 3%, (c) 4%, and (4) 5% BIS and for different MAA concentrations.	5.17
5.9	Dose correlation factor k_{BIS} of C-C stretching at 2110 cm^{-1} of PMAAG due to BIS crosslinking.	5.19
5.10	Dose correlation factor k'_{BIS} of C-C stretching at 2110 cm^{-1} of PMAAG due to BIS crosslinking.	5.20
5.11	Normalised Raman intensity of C=O stretching showing the formation of PMAAG at (a) 2%, (b) 3%, (c) 4%, and (4) 5% MAA and for different BIS concentrations.	5.23
5.12	Correlation between D_0 and the initial concentration of MAA for different BIS composition for the formation of PMAAG monomer due to C=O band at 1682 cm^{-1} .	5.24
5.13	Correlation between $D_{1/2}$ and the initial concentration of MAA for different BIS composition for the formation of PMAAG due to C=O band at 1682 cm^{-1} .	5.25
5.14	Normalised Raman intensity of C=O stretching showing the formation of PMAAG at (a) 2%, (b) 3%, (c) 4%, and (4) 5% BIS and for different MAA concentrations.	5.28
5.15	Dose correlation factor k_{BIS} of C=O stretching of PMAAG due to BIS crosslinking.	5.30
5.16	Dose correlation factor k'_{BIS} of C=O stretching of PMAAG due to BIS crosslinking.	5.31

5.17	Normalised Raman intensity of CH ₂ stretching showing the formation of PMAAG at (a) 2%, (b) 3%, (c) 4%, and (4) 5% MAA and for different BIS concentrations.	5.34
5.18	Correlation between D_0 and the initial concentration of MAA for different BIS composition for the formation of PMAAG at CH ₂ band (2905 cm ⁻¹).	5.35
5.19	Correlation between $D_{1/2}$ and the initial concentration of MAA for different BIS composition for the formation of PMAAG at CH ₂ band (2905 cm ⁻¹).	5.36
5.20	Normalised Raman intensity of CH ₂ stretching showing the formation of PMAAG at (a) 2%, (b) 3%, (c) 4%, and (4) 5% BIS and for different MAA concentrations.	5.39
5.21	Dose correlation factor k_{BIS} of CH ₂ stretching of PMAAG due to BIS crosslinking.	5.40
5.22	Dose correlation factor k'_{BIS} of CH ₂ stretching of PMAAG due to BIS crosslinking.	5.41
5.23	Normalised Raman intensity of C=C stretching of MAA showing the consumption of MAA at (a) 2%, (b) 3%, (c) 4%, and (4) 5% BIS and for different BIS concentrations.	5.45
5.24	Correlation between D_0 and the initial concentration of MAA for different BIS compositions for the consumption of monomer at C=C stretching (2238 cm ⁻¹).	5.46
5.25	Correlation between $D_{1/2}$ and the initial concentration of MAA for different BIS composition for the consumption of monomer at C=C stretching (2238 cm ⁻¹).	5.48

5.26	Normalised Raman intensity of C=C stretching of BIS showing the consumption of BIS at (a) 2%, (b) 3%, (c) 4%, and (4) 5% MAA and for different BIS concentrations.	5.50
5.27	Correlation between D_0 and the initial concentration of BIS for different MAA composition for the consumption of crosslinker at C=C stretching (1634 cm^{-1}).	5.52
5.28	Correlation between $D_{1/2}$ and the initial concentration of BIS for different MAA compositions for the consumption of crosslinker at C=C stretching (1634 cm^{-1}).	5.53
5.29	NMR spin-lattice relaxation rate (ΔR_1) as a function of dose D at (a) 2%, (b) 3%, (c) 4% and (d) 5% MAA and for different BIS concentrations.	5.59
5.30	Correlation between D_0 value and the initial concentration of MAA for different BIS compositions in T_1 measurement.	5.61
5.31	Correlation between $D_{1/2}$ value and the initial concentration of MAA for different BIS compositions in T_1 measurement.	5.62
5.32	NMR spin-lattice relaxation rate (ΔR_1) as a function of dose D at (a) 2%, (b) 3%, (c) 4% and (d) 5% BIS and for different MAA concentrations.	5.65
5.33	Correlation between D_0 value and the initial concentration of BIS for different MAA compositions in T_1 measurement.	5.66
5.34	Correlation between $D_{1/2}$ value and the initial concentration of BIS for different MAA compositions in T_1 measurement.	5.68
5.35	NMR spin-spin relaxation rate (ΔR_2) as a function of dose D at (a) 2%, (b) 3%, (c) 4% and (d) 5% MAA and for different BIS concentrations.	5.71

5.36	Correlation between D_0 value and the initial concentration of MAA for different BIS compositions in T_2 measurement.	5.72
5.37	Correlation between $D_{1/2}$ value and the initial concentration of MAA for different BIS compositions in T_2 measurement.	5.74
5.38	NMR spin-spin relaxation rate (ΔR_2) as a function of dose D at (a) 2%, (b) 3%, (c) 4% and (d) 5% BIS and for different MAA concentrations.	5.77
5.39	Correlation between D_0 value and the initial concentration of BIS for different MAA compositions in T_2 measurement.	5.78
5.40	Correlation between $D_{1/2}$ value and the initial concentration of BIS for different MAA compositions in T_2 measurement.	5.80
5.41	UV-visible light absorbance of PMAAG sample containing gelatine, water and 0 - 5% MAA.	5.86
5.42	Dependence of radiation response on MAA concentration. Absorbance at 375 nm due to radical formation was measured following exposure time of 3 hours (43.26 Gy) of ^{60}Co radiations.	5.87
5.43	Comparison of the absorbance of various compositions of methacrylic acid (MAA) and N, N'-methylene-bisacrylamide (BIS) at wavelength 375 nm.	5.88
5.44	Comparison between PolyMethacrylic Acid gel (various compositions of BIS and MAA) with ionization chamber and national standards.	5.89
5.45	Absorbed dose comparison of various BIS concentrations for PMAAG (MAA 2%) with Fricke dosimeter and ionization chamber.	5.90

5.46	Absorbed dose comparison of various BIS concentrations for PMAAG (MAA 3%) with Fricke dosimeter and ionization chamber.	5.91
5.47	Absorbed dose comparison of various BIS concentrations for PMAAG (MAA 4%) with Fricke dosimeter and ionization chamber.	5.91
5.48	Absorbed dose comparison of various BIS concentrations for PMAAG (MAA 5%) with Fricke dosimeter and ionization chamber.	5.92

LIST OF ABBREVIATIONS/SYMBOL

γ -ray	Gamma Ray
BIS	N, N'-methylene-bisacrylamide
MAA	methacrylic acid
PMAAG	polymethacrylic gel
Gy	Gray
MINT	Malaysian Institute for Nuclear Technology Research
IAEA	International Atomic Energy Research
NMR	Nuclear Magnetic Resonance
UV-Vis	Ultraviolet - Visible
T_1	Spin-lattice relaxation time
T_2	Spin-spin relaxation time
R_1	Spin-lattice relaxation rate
R_2	Spin-spin relaxation rate
N_A	$6.022 \times 10^{23} \text{ mol}^{-1}$ (Avogadro's number)
ΔA	the change in absorbance at peaks
ρ	the density of the dosimetry solution
ϵ	molar extinction coefficient
ζ	temperature coefficient of the molar extinction coefficient
G	G-value of the radiation yield, which is valid for electrons or photons of energy 0.5 to 16 MeV at absorbed dose rates $< 2 \times 10^7 \text{ Gy/s}$.