



**UNIVERSITI PUTRA MALAYSIA**

**QUALITY OF LIFE OF STROKE SURVIVORS AND THEIR FAMILY  
CAREGIVERS AT SELECTED REFERRAL REHABILITATION CARE  
CENTERS**

**KHAW WAN-FEI**

**FPSK(p) 2018 3**



**QUALITY OF LIFE OF STROKE SURVIVORS AND  
THEIR FAMILY CAREGIVERS AT SELECTED  
REFERRAL REHABILITATION CARE CENTERS**

**By**

**KHAW WAN-FEI**

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

**November 2017**

All material contained within the thesis, including without limitation text, logos, icons, photographs and all other artwork, is copyright material of Universiti Putra Malaysia unless otherwise stated. Use may be made of any material contained within the thesis for non-commercial purposes from the copyright holder. Commercial use of material may only be made with the express, prior, written permission of Universiti Putra Malaysia.

Copyright © Universiti Putra Malaysia



© COPYRIGHT UPM

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

**QUALITY OF LIFE OF STROKE SURVIVORS AND THEIR FAMILY  
CAREGIVERS AT SELECTED REFERRAL REHABILITATION CARE  
CENTERS**

By

**KHAW WAN-FEI**

**November 2017**

**Chair: Syed Tajuddin Syed Hassan, PhD**  
**Faculty: Medicine and Health Sciences**

Stroke disability results in hardship to both patients and families. Care for stroke patients can be stressful and lead to deterioration of quality of life (QOL). Despite the large number of stroke cases, limited local data are available on the impact of stroke on QOL of stroke survivors and caregivers, and little is known regarding the interdependence of QOL within the dyad. Thus, this study was developed to evaluate overall QOL of stroke survivors and caregivers, and determined factors predicting QOL. Dyadic analysis using Actor-Partner Interdependence Model (APIM) examined whether stroke survivor's and caregiver's QOL at baseline, predicts his or her own QOL at six months follow-up (actor effect) and partner's QOL at follow-up (partner effect).

This was a prospective study that involved 160 stroke survivors and 160 caregivers recruited from neurology clinics in Hospital Kuala Lumpur and Hospital Rehabilitasi Cheras in Klang Valley, Malaysia. The six months follow-up included 38 stroke survivors and 38 caregivers. Socio-demographic and medical conditions, caregiving appraisal, coping strategies and QOL were obtained from stroke survivors and caregivers. The data were entered and analysed using the IBM SPSS for Windows, Version 22.0 and IBM SPSS AMOS, Version 22.0. Univariate analysis was used to clean and check the quality of data. Bivariate analysis was performed to examine the bivariate relationships among predictor variables and stroke survivor's and caregiver's QOL. Confirmatory factor analysis (CFA) was used to test dimensionality of the QOL measure of five indicators: physical health, mental health, general health, peace, and faith. Path analysis was used to determine the direct and indirect effects of predictors on QOL in both stroke survivors and caregivers, and to assess the APIM model for survivor-caregiver dyads.

CFA results indicated acceptable goodness-of-fit of one-factor QOL model ( $\chi^2(12) = 42.265, p < 0.001$ ; RMSEA = 0.063; SRMR = 0.038; CFI = 0.970), which included five indicators: physical health, mental health, general health, peace, and faith. Stroke survivors rated lower QOL scores compared to caregivers ( $M = 57.96, SD = 15.06$  versus  $M = 45.50, SD = 16.65$ ). Six determinants with both a direct and indirect effect on the QOL of stroke survivors were age, duration of stroke, stroke severity, cognitive function, physical function, and psychological distress. Seven determinants with both a direct and indirect effect on the QOL of stroke caregivers were caregiver's age, stroke severity, social support, adaptive coping, maladaptive coping, caregiving burden, and caregiving satisfaction. Actor effects between stroke survivor's QOL ( $\beta = 0.79, p < 0.001$ ) and caregiver's QOL ( $\beta = 0.76, p < 0.001$ ) at baseline and follow-up were significant. This indicates that stroke survivor's and caregiver's QOL at baseline were positively related to their own QOL at 6-months-follow-up.

These findings highlight the overall QOL score integrates all the measures of health status of stroke survivors and their caregivers. The APIM dyadic analysis provides insight into the relationships between stroke survivor's and caregiver's QOL. Thus, this study definitively indicates the necessity for arranging interventions targeting both stroke survivors and family caregivers, to enhance their QOL in the rehabilitation process.

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

**KUALITI HIDUP PESAKIT STROK DAN PENJAGA MEREKA DI  
PUSAT RUJUKAN RAWATAN DAN PEMULIHAN YANG TERPILIH**

Oleh

**KHAW WAN-FEI**

**November 2017**

**Pengerusi: Syed Tajuddin Syed Hassan, PhD**  
**Fakulti: Perubatan dan Sains Kesihatan**

Ketidakupayaan pesakit akibat strok menyebabkan kesukaran kepada kedua-dua pesakit dan keluarga. Menjaga pesakit strok boleh menyebabkan stres dan kemerosotan kualiti hidup (“QOL”). Walaupun bilangan kes strok yang tinggi, namun hasil kajian tempatan mengenai kesan stroke terhadap kualiti hidup pesakit and penjaga keluarga adalah terhad. Tambahan pula, pengetahuan mengenai kualiti hidup bagi pasangan pesakit dan penjaga dalam konteks saling bergantung masih kekurangan. Oleh itu, kajian ini bertujuan menilai “Overall QOL” bagi pesakit strok dan penjaga mereka. Kajian ini juga menentukan faktor-faktor yang mempengaruhi kualiti hidup. Analisis diad dijalankan dengan menggunakan “Actor-partner interdependence model” (“APIM”) untuk mengenalpasti pengaruh kualiti hidup terhadap sendiri, kesan actor dan kualiti hidup terhadap pasangan mereka, kesan partner, di peringkat awal dan susulan.

Kajian dijalankan secara prospektif dengan melibatkan 160 pesakit strok dan 160 penjaganya, dari klinik neurologi Hospital Kuala Lumpur dan Hospital Rehabilitasi Cheras di Lembah Klang, Malaysia. Seramai 38 pesakit strok dan 38 penjaga mereka terlibat dalam kajian susulan enam bulan kemudian. Ciri sosio-demografi dan perubatan, penilaian penjagaan, strategi menghadapi stress, dan kualiti hidup diperolehi daripada pesakit strok dan penjaga mereka. Data yang diperolehi dianalisis dengan menggunakan perisian IBM SPSS for Windows, Version 22.0 dan IBM SPSS AMOS, Version 22.0. Analisis univariat digunakan untuk meneliti dan memeriksa kualiti data. Analisis bivariat dilakukan untuk meneliti hubungan bivariat antara pembolehubah ramalan dan QOL bagi pesakit dan penjaga strok. Analisis faktor konfirmatori (“CFA”) digunakan untuk menguji dimensi skor QOL bagi lima indikator: kesihatan fizikal, kesihatan mental, kesihatan umum, keamanan, dan iman. Analisis jalur (“Path analysis”) juga dijalankan untuk menentukan kesan langsung dan tidak

langsung terhadap QOL, bagi kedua-dua pesakit dan penjaga strok, serta menilai model diad “APIM” bagi pesakit strok dan penjaga mereka.

Keputusan analisis “CFA” menunjukkan “One-factor QOL model” adalah bersesuaian dengan data ( $\chi^2(12) = 42.265, p < 0.001$ ; RMSEA = 0.063; SRMR = 0.038; CFI = 0.970). Model ini merangkumi lima indikator: kesihatan fizikal, kesihatan mental, kesihatan umum, keamanan, dan iman. Nilai min skor “QOL” bagi pesakit adalah lebih rendah berbanding dengan penjaga ( $M = 57.96, SD = 15.06$  berbanding  $M = 45.50, SD = 16.65$ ). Enam penentu yang mempengaruhi “QOL” bagi pesakit strok secara langsung dan tidak langsung ialah umur pesakit, tempoh strok, tahap strok, fungsi kognitif, fungsi fizikal, dan tekanan psikologi. Tujuh penentu yang mempengaruhi “QOL” bagi penjaga stroke secara langsung dan tidak langsung ialah umur penjaga, tahap strok, sokongan sosial, “adaptive coping”, “maladaptive coping”, beban penjagaan, dan kepuasan penjagaan. Pada kedua-dua peringkat pengukuran, awalan dan enam bulan kemudian kesan aktor adalah signifikan, bagi kedua-dua pesakit “QOL” ( $\beta = 0.79, p < 0.001$ ) dan penjaga “QOL” ( $\beta = 0.76, p < 0.001$ ). Dengan menggunakan model “APIM” bagi pesakit dan penjaga, nilai “QOL” pada peringkat awal mempengaruhi nilai “QOL” pada peringkat susulan secara positif.

Hasil kajian ini jelas menunjukkan skor “QOL” adalah penting untuk menilai status kesihatan pesakit strok dan penjaga mereka. Model “APIM” pula dapat memberikan gambaran tentang hubungan antara “QOL” bagi pesakit dan penjaganya. Oleh itu, kajian ini menekankan kepentingan untuk merancang intervensi bagi kedua-dua pesakit dan penjaga strok supaya QOL mereka dapat ditingkatkan bersama dalam proses pemulihan.

## ACKNOWLEDGEMENTS

First of all, I am truly indebted to Prof. Dr Syed Tajuddin Syed Hassan, my supervisor, for his consistent coaching, guidance, experience-sharing and funding. His thoughtfulness towards the educational welfare of his students has inspired me tremendously. Moreover, his patience and countless contribution in finishing this project were greatly appreciated.

Secondly, my heartfelt thanks to my co-supervisors, Prof. Dato' Dr Lye Munn Sann and Dr Siti Irma Fadhilah Ismail, for their assistance, expertise and suggestions to improve in this research. Next, I would like to acknowledge the Directors, Heads of departments and staff of Hospital Kuala Lumpur and Hospital Rehabilitasi Cheras for their invaluable cooperation and assistance.

My utmost gratitude to my colleagues and seniors for their knowledge and assistance which has been very helpful in completing this research. Not to be forgotten, I deeply acknowledge a bunch of my best friends, Lai Kuan, Heng Yaw, Hui Ceng and others for their kind understanding, encouragements and assistance throughout my study. We had shared so many good and bad times; being there to support, comfort and cheer each other at times we were really stressed up throughout the study.

And last but not least, there are no words to fully express my deepest gratitude to my beloved parents and family members, because of you all I am here today. Your endless supports, contributions and sacrifices would never be forgotten.



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:

**Syed Tajuddin Syed Hassan, PhD**

Professor  
Faculty of Medicine and Health Sciences  
Universiti Putra Malaysia  
(Chairman)

**Lye Munn Sann, PhD**

Professor  
Faculty of Medicine and Health Sciences  
Universiti Putra Malaysia  
(Member)

**Siti Irma Fadhilah Ismail, PhD**

Senior Lecturer  
Faculty of Medicine and Health Sciences  
Universiti Putra Malaysia  
(Member)

---

**ROBIAH BINTI YUNUS, PhD**

Professor and Dean  
School of Graduate Studies  
Universiti Putra Malaysia

Date:

## Declaration by graduate student

I hereby confirm that:

- this thesis is my original work;
- quotations, illustrations and citations have been duly referenced;
- this thesis has not been submitted previously or concurrently for any other degree at any other institutions;
- intellectual property from the thesis and copyright of thesis are fully-owned by Universiti Putra Malaysia, as according to the Universiti Putra Malaysia (Research) Rules 2012;
- written permission must be obtained from supervisor and the office of Deputy Vice-Chancellor (Research and Innovation) before thesis is published (in the form of written, printed or in electronic form) including books, journals, modules, proceedings, popular writings, seminar papers, manuscripts, posters, reports, lecture notes, learning modules or any other materials as stated in the Universiti Putra Malaysia (Research) Rules 2012;
- there is no plagiarism or data falsification/fabrication in the thesis, and scholarly integrity is upheld as according to the Universiti Putra Malaysia (Graduate Studies) Rules 2003 (Revision 2012-2013) and the Universiti Putra Malaysia (Research) Rules 2012. The thesis has undergone plagiarism detection software.

Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Name and Matric No.: Khaw Wan-Fei, GS30303

## Declaration by Members of Supervisory Committee

This is to confirm that:

- the research conducted and the writing of this thesis was under our supervision;
- supervision responsibilities as stated in the Universiti Putra Malaysia (Graduate Studies) Rules 2003 (Revision 2012-2013) are adhered to.

Signature: \_\_\_\_\_

Name of Chairman  
of Supervisory  
Committee: \_\_\_\_\_

Signature: \_\_\_\_\_

Name of Member  
of Supervisory  
Committee: \_\_\_\_\_

Signature: \_\_\_\_\_

Name of Member  
of Supervisory  
Committee: \_\_\_\_\_

## TABLE OF CONTENTS

	<b>Page</b>
<b>ABSTRACT</b>	i
<b>ABSTRAK</b>	iii
<b>ACKNOWLEDGEMENTS</b>	v
<b>APPROVAL</b>	vi
<b>DECLARATION</b>	viii
<b>LIST OF TABLES</b>	xiii
<b>LIST OF FIGURES</b>	xiv
<b>LIST OF ABBREVIATIONS</b>	xv
<b>CHAPTER</b>	
<b>1 INTRODUCTION</b>	<b>1</b>
1.1 Background	1
1.2 Statement of problem	2
1.3 Research questions	4
1.4 Objectives of study	4
1.5 Research hypotheses	5
1.6 Research conceptual framework	5
1.7 Significance of study	8
<b>2 LITERATURE REVIEW</b>	<b>10</b>
2.1 Introduction	10
2.2 Definition of stroke	10
2.3 Epidemiology of stroke	11
2.3.1 Global epidemiology of stroke	11
2.3.2 Epidemiology of stroke in Malaysia	12
2.3.3 Risk factors of stroke	13
2.3.4 Burden of stroke	14
2.3.5 Rehabilitation of stroke	15
2.4 Quality of life after stroke	17
2.4.1 Definition of quality of life	17
2.4.2 Measurement of quality of life	19
2.4.3 Quality of life after stroke	20
2.4.4 Factors associated post-stroke quality of life	21
2.5 Family caregiving for stroke patients	28
2.5.1 Quality of life of family caregivers of stroke survivors	31
2.5.2 Factors associated with caregiver quality of life	32
2.6 Interrelation between person with stroke and caregivers	36
2.6.1 Actor-partner interdependence model (APIM)	38
2.6.2 Applications of the APIM in the stroke literature	40

<b>3</b>	<b>METHODOLOGY</b>	43
3.1	Study location	43
3.2	Study design	44
3.3	Study population	44
3.4	Sampling	44
3.4.1	Inclusion criteria	45
3.4.2	Exclusion criteria	45
3.5	Sample size estimation	46
3.6	Sampling method	47
3.7	Study variables and operational definition	47
3.7.1	Dependent variables	47
3.7.2	Independent variables	47
3.7.3	Operational definition	48
3.8	Instruments	49
3.8.1	Socio-demographic and medical conditions	50
3.8.2	Quality of life	51
3.8.3	Caregiving appraisal	52
3.8.4	Coping strategies	52
3.8.5	Social support	53
3.9	Validity and reliability of the study instrument	53
3.9.1	Validity of questionnaire	54
3.9.2	Reliability of questionnaire	55
3.10	Data collection	55
3.11	Ethical approval	56
3.12	Data analysis	56
3.12.1	Preliminary statistical analysis	56
3.12.2	Confirmatory factor analysis	57
3.12.3	Path analysis	58
3.12.4	Dyadic data analysis	60
<b>4</b>	<b>RESULTS</b>	61
4.1	Description of sample	61
4.1.1	Demographic characteristics of respondents	61
4.1.2	Mean scores of QOL measures for respondents	63
4.1.3	Clinical characteristics of stroke survivors	63
4.1.4	Descriptive statistics for caregiving characteristics of stroke caregivers	65
4.2	Testing for a one-factor QOL model	67
4.3	Testing relationships between predictors and QOL in stroke survivors	70
4.3.1	Correlation analysis of the variables	70
4.3.2	Significant relationship between observed variables	72
4.3.3	Mediating factors	75
4.4	Testing relationships between predictors and QOL in stroke caregivers	77
4.4.1	Correlation analysis of the variables	77
4.4.2	Significant relationship between observed variables	80
4.4.3	Mediating factors	83

4.5	Testing relationship between QOL among stroke survivors and their caregivers over two time points	85
4.5.1	Changes in QOL over two time points	85
4.5.2	Dyadic relationship between physical function and QOL over time	86
<b>5</b>	<b>DISCUSSION</b>	<b>89</b>
5.1	Introduction	89
5.2	QOL scores for stroke survivors and caregivers	89
5.3	Relationship between predictors and QOL in stroke survivors	91
5.4	Relationship between predictors and QOL in caregivers	93
5.5	Dyadic relationship between QOL over two time points	96
<b>6</b>	<b>CONCLUSION AND RECOMMENDATIONS</b>	<b>98</b>
6.1	Conclusion	98
6.2	Recommendations for future research	99
6.3	Limitations of study	101
	<b>REFERENCES</b>	<b>103</b>
	<b>APPENDICES</b>	<b>116</b>
	<b>BIODATA OF STUDENT</b>	<b>132</b>
	<b>LIST OF PUBLICATIONS</b>	<b>133</b>

## LIST OF TABLES

Table		Page
1	Study instruments	49
2	Demographic characteristics of respondents ( $N = 320$ )	62
3	Mean score of QOL ( $N = 320$ )	63
4	Clinical characteristics of stroke survivors ( $N = 160$ )	64
5	Caregiving characteristics of stroke caregivers ( $N = 160$ )	66
6	Mean score of overall QOL ( $N = 320$ )	69
7	Pearson correlation coefficient matrix of the measured variables	71
8	Relationship between independent and dependent variables	74
9	Standardized indirect, direct, and total effects of variables on QOL in stroke survivors	76
10	Pearson correlation coefficient matrix of the measured variables	79
11	Relationship between independent and dependent variables	82
12	Standardized indirect, direct, and total effects of variables on QOL in stroke caregivers	84
13	Comparisons between stroke survivors and caregivers on QOL at baseline and 6 months follow-up ( $N = 38$ dyads)	86
14	Pearson correlation coefficient matrix of the study variables in stroke survivor-caregiver dyads	86
15	Relationship between QOL for stroke survivors and caregivers over two time points	88

## LIST OF FIGURES

Figure	Page
1 Conceptual framework of the QOL model of stroke survivors and caregivers	7
2 The Actor Partner Interdependence Model (APIM)	39
3 Multiple mediation model	59
4 Initial one-factor model for the QOL score	68
5 Final one-factor model for the QOL score	69
6 Significant pathways of the final model of factors influencing QOL in stroke survivors	73
7 Significant pathways of the final model of factors influencing QOL in stroke caregivers	81
8 The APIM demonstrating actor and partner effects of QOL over two time points	87



## LIST OF ABBREVIATIONS

ADL	Activities of daily living
APIM	Actor-Partner Interdependence Model
BI	Barthel Index
CFA	Confirmatory factor analysis
CFI	Comparative Fit Index
CI	Confidence interval
CRA	Caregiver Reaction Assessment Scale
DALYs	Disability-adjusted life years
FACIT-Sp	Functional Assessment of Chronic Illness Therapy-Spiritual Wellbeing Scale
HADS	Hospital Anxiety and Depression Scale
HRQOL	Health-related quality of life
ICF	International Classification of Functioning, Disability, and Health
MCS	Mental component summary
MMSE	Mini-Mental State Examination
MOS-SSS	Medical outcomes study social support survey
OR	Odds ratio
PA	Path analysis
PAC	Positive Aspects of Caregiving Scale
PCS	Physical component summary
QOL	Quality of life
REGARDS	REason for Geographical and Racial Differences in Stroke study
RMSEA	Root mean square error of approximation
SEM	Structural equation modelling
SF-12	12-Item Short-Form Health Survey
SF-36	36-Item Short-Form Health Survey
SRMR	Standardized root mean square residual
WHOQOL	World Health Organization Quality of Life Group

# CHAPTER 1

## INTRODUCTION

### 1.1 Background

Stroke is the second leading cause of death globally and common cause of permanent disability. Worldwide, an estimated 16.9 million people are affected by stroke and 5.9 million deaths annually (Feigin et al., 2014). These numbers are expected to rise to 70 million stroke survivors and 12 million stroke deaths by 2030. For the year 2009, 52,000 cases of hospitalisation for stroke and 2,300 deaths were estimated to have occurred in government hospitals in Malaysia. The Malaysian National Burden of Disease Study showed stroke ranked second among top 10 leading cause of burden of disease. Stroke incidence in Malaysia has been estimated at 96/ 100,000 for ischemic stroke and 21/ 100,000 for hemorrhagic stroke in 2014 (Aziz et al., 2015).

Stroke is the main cause of disability in adult life and this has placed a tremendous burden on healthcare systems, families, and economics. About 15% to 30% of stroke patients with permanent impairments and 20% need institutional care after a stroke. Among those who survived, more than 60% stroke patients had incomplete recovery (Scherbakov, Von Haehling, Anker, Dirnagl, & Doehner, 2013). Stroke-related disability includes physical, social and cognitive functions. Approximately 70-80% of survivors manifest paralysis, speech disabilities or emotional problems. Apart from functional disability, the long-term effects of stroke include fatigue, cognitive and psychological problems (Mohd Zulkifly, Ghazali, Che Din, Singh, & Subramaniam, 2016). These impairments were linked to lower levels of quality of life (QOL) (Algurén, Fridlund, Cieza, Sunnerhagen, & Christensson, 2012; Azlin & Rizal, 2009).

Cross-sectional data suggest that QOL after stroke is significantly impaired (Bach et al., 2011; Cerniauskaite et al., 2012). However, data from longitudinal studies are limited and they showed similar trend to the cross-sectional studies (Algurén et al., 2012; Haley, Roth, Kissela, Perkins, & Howard, 2011; Hamza, Al-Sadat, Loh, & Jahan, 2014). A research that examined 85 stroke patients during rehabilitation and 6 months after discharge, noted that the patients' QOL scores improved significantly during inpatient rehabilitation, followed by a decline in the 6 months after discharge (Hopman & Verner, 2003). In addition, patients reported poorer QOL were associated with reduced functional recovery (Nichols-Larsen, Clark, Zeringue, Greenspan, & Blanton, 2005).

Stroke is a leading cause of persistent disability and handicap, affects all aspects of patient life including physical, behavioural, psychological, and social functioning. These resulting impairments after stroke can affect both stroke survivors and their caregivers (Algurén et al., 2012). Around 80% of stroke survivors return to community living after stroke, they rely on emotional and physical support from family members, such as spouses, adult children and siblings, close friends and sitters to aid in their recovery. In addition, stroke is a sudden and unpredictable event and caregivers often feel unprepared for their caregiving role, and these changes can be very stressful for both stroke patients and caregivers (Ostwald, Bernal, Cron, & Godwin, 2009).

Stroke caregivers have lower QOL than norms and caregivers' QOL are lower than those of the stroke patients (Akosile, Okoye, Nwankwo, Akosile, & Mbada, 2011; McPherson, Wilson, Chyurlia, & Leclerc, 2011; Parag et al., 2008). Caregivers of stroke survivors have been associated with higher levels of depression and reduced QOL (Godwin, Swank, Vaeth, & Ostwald, 2013b). The QOLs of stroke patients and their caregivers were influenced by various factors, such as ages and gender of patients and caregivers, patients' functional ability and social involvement (Jönsson, Lindgren, Hallström, Norrving, & Lindgren, 2005).

Although caregiving burden has been shown to decrease caregivers' physical and psychological QOL, caregivers have expressed increased in life satisfaction, enjoyments in caregiver role, and better problem solving and coping ability (Haley et al., 2009). Some caregiver studies draw conclusions that suggest caregivers' experiences are more positive than negative (Cameron, Stewart, Streiner, Coyte, & Cheung, 2014; Haley et al., 2009). For example, stroke family caregivers reported that caregiving enabled to appreciate life more, feeling needed and appreciated and developing a more positive attitude toward life. Positive caregiving experiences are associated to better physical and mental health, as well as better QOL. In a systematic review, it has been suggested that the need for more research examining both the positive and negative experiences of caregiving in stroke (Mackenzie & Greenwood, 2012). The significance of positive caregiving experiences is pronounced, and future work is needed to determine factors associated with positive impacts of caregiving and to establish strategies to improve positive caregiving experiences.

## **1.2 Statement of problem**

Quality of life (QOL) is a useful indicator of overall health. It captures data on the physical, psychological, social and spiritual health of individuals. QOL of stroke survivors and caregivers are usually assessed by using multiple indicators of self-rated health status, physical and emotional functioning, social and spiritual well-being (Algurén et al., 2012; Hamza et al., 2014; Zhang, Sun, Wu, & Xia, 2013). However, these measures have been used separately and a cohesive picture of overall QOL has not been published. There is a need to identify a measure of QOL score, including general health, physical, mental, faith, and peace in stroke survivors and caregivers.

Much of the stroke research identifying possible socio-demographic, functional status, and behavioural factors contributing to QOL have used linear regression to identify predictor variables significantly associated with QOL (Saha, Harries, & Gilhooly, 2016). Despite a number of studies examining the factors that predict QOL, little is known about the complex inter-relationships between numerous predictors (Algurén et al., 2012; Ellis, Grubaugh, & Egede, 2013; Haley et al., 2011). Path analysis, an alternative statistical technique, is considered as an appropriate statistical method for such an explorative study to better understand the complex interrelation between the variables. Path analysis identifies direct and indirect pathways in which a set of predictor variables influences outcome variable. However, these factors have not been extensively studied using path analysis and available studies show inconsistent findings (Chen et al., 2015; Howitt et al., 2011; Morris, van Wijck, Joice, & Donaghy, 2013).

Care for stroke patients has been reported as a complex caregiving process. The relationship between caregiving stressors and caregiver outcome is influenced by caregiving appraisal and coping variables. Some studies have documented that caregivers adapt to challenges and problems of caregiving over time, whereas others have shown caregiving strain and burden are continuing to increase over time (Godwin et al., 2013b; McLennon, Bakas, Jessup, Habermann, & Weaver, 2014). However, the benefits of caregiving have been reported (Haley et al., 2009). Nevertheless, research on identifying of both positive and negative impacts of caregiving is underdeveloped. Information on factors that improve or deteriorate QOL is lacking. Also, the data on how survivors and family caregivers influence each other's QOL after a stroke are not available.

Despite the large number of stroke cases, there are limited local data on the impact of stroke on the family caregivers. There are many interpersonal and contextual variables influencing caregiver quality of life, many of which interact with one another, thus creating a distinctively unique experience for each caregiver. Caregiving stressors, appraisal, coping and social supports, and care-recipient conditions may influence caregiver QOL, but few studies considered the combined impact of all the variables (Ostwald et al., 2009; Yu, Hu, Efirid, & McCoy, 2013). Existing studies examining predictors of caregiver burden and QOL are limited in terms of scope because they do not consider all important variables in a single comprehensive model.

Research on the impact of stroke on QOL tends to focus on the patients. However, there is evidence that the outcomes of caregivers and patients are interdependence (McCarthy, Lyons, & Powers, 2011). Significant positive correlations have been found between the outcomes of caregivers and care-recipients on QOL (Cramm, Strating, & Nieboer, 2012), distress, and depression (Ostwald et al., 2009). Since stroke survivors and their caregivers health outcomes are affected by patient's health situation, interactions in patient and caregiver dyads should be considerate in stroke studies. There is interdependence between stroke survivor and caregiver, because they share thoughts and feelings, extend mutual support, and encouragement from each other.

Ostwald et. al. (2009) have shown the mutual influence both stroke patients' and family members' stress and QOL. Thus, there is a need to study the impact of stroke on health outcomes that considers dyads as the unit of analysis.

Even though many QOL studies have reported experiences of stroke patients and caregivers by focusing on the individual level of analysis (Algurén et al., 2012; Haley et al., 2009; Saha et al., 2016), there has been very little research reported using the dyad level of analysis. The need of dyadic research in post-stroke QOL has been emphasized, and studies have identified relational factors in QOL following stroke for patients and caregivers, such as coping patterns, social supports, and burden (Opara & Jaracz, 2010; Visser et al., 2015). Moreover, dyadic studies have linked patients' QOL with their caregivers' QOL (Bergström, Eriksson, von Koch, & Tham, 2011; Cramm et al., 2012), but no studies have longitudinally studied the dyadic relations between patients' and their caregivers' QOL to understand the process of patient's recovery and caregiving experiences. It was necessary to study how a stroke impacts both persons individually as well as together over time.

Previous findings have shown in a dyad, mutual impacts on QOL occurred (Bergström et al., 2011; Cramm et al., 2012). Information on QOL of stroke survivors and caregivers are limited from developed countries. Since there are ethnic and cultural differences in QOL measures, there may also be differences in the patterns of QOL scores of stroke patients and caregivers in Malaysia. However, there are no local baseline data on the interrelationship of stroke survivors and caregivers QOL during the rehabilitation process. Thus, there is a need to investigate a dyadic relationship of stroke survivors and caregivers QOL across six months of stroke rehabilitation.

### **1.3 Research questions**

The following research questions guided this study:

1. What are the overall QOL scores for stroke survivors and caregivers?
2. What are the factors associated with QOL scores of stroke survivors and caregivers?
3. Is there an interrelationship between QOL scores stroke survivors and caregivers over time?

### **1.4 Objectives of study**

General objective:

To determine QOL scores in stroke survivors and caregivers and dyadic relations of stroke survivors' and their caregivers' QOL scores over two time points.



Specific objectives:

1. To identify a one-factor QOL model for stroke survivors and caregivers.
2. To determine the relationships between stroke survivor socio-demographic and medical conditions, functioning and disability, and QOL in stroke survivors.
3. To determine the relationships between caregiver variables, caregiving appraisal, coping strategies, and QOL in stroke caregivers
4. To determine the dyadic relationships between stroke survivors' and caregivers' QOL scores over two time points.

## **1.5 Research hypotheses**

1. There is a one-factor QOL model that underlies scores on general health, physical health, mental health, peace, and faith for stroke survivors and caregivers.
2. There are direct and indirect effects of stroke survivor socio-demographic and medical conditions, functioning and disability, and QOL in stroke survivors.
3. There are direct and indirect effects of caregiver variables, caregiving appraisal, coping strategies, and QOL in stroke caregivers.
4. Stroke survivors' and caregivers' QOL scores would be positively related to their own and others' QOL over two time points.

## **1.6 Research conceptual framework**

The research framework for this study is presented in Figure 1. This research framework displays the factors that influenced QOL scores of stroke survivors and caregivers. Next, the dyadic relationships among stroke survivors' and caregivers' QOL over time were examined.

In this study, QOL in stroke survivors and caregivers consisted of physical, mental, social, and spiritual domains. The first goal of this study was to identify overall QOL scores for stroke survivors and their caregivers. It hypothesized that there was one-factor QOL model that underlies the indicators on general health, physical health, mental health, peace, and faith. CFA was used to test dimensionality of the measure for five QOL indicators: physical health, mental health, general health, peace, and faith. The overall QOL score values were generated using weighted factor score values from one-factor QOL model.

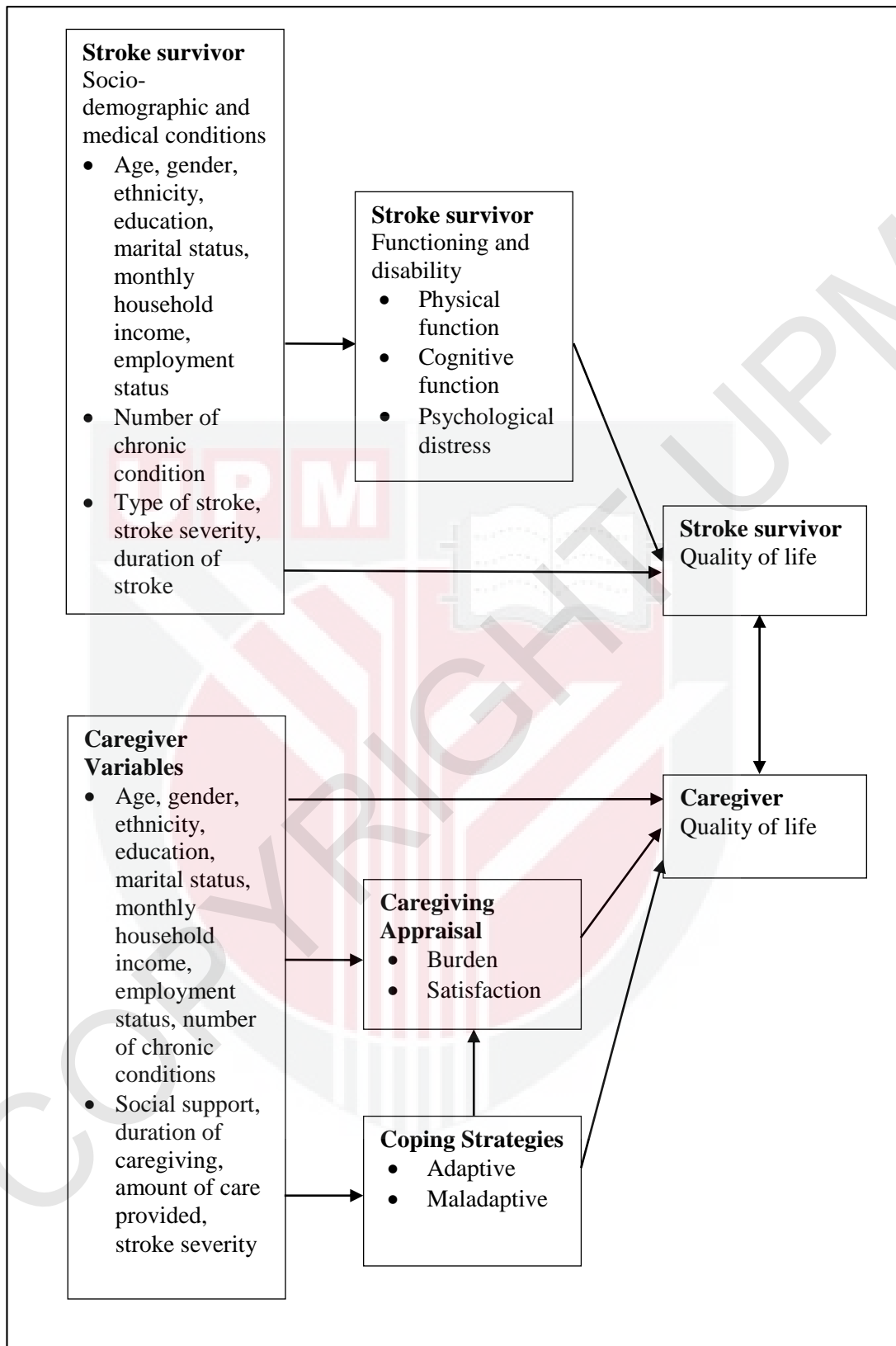
QOL concepts and integrative biopsychosocial model of functioning, disability and health had been applied in the stroke rehabilitation and outcomes (Geyh, Cieza, Kollerits, Grimby, & Stucki, 2007). This model conceptualizes a person's functioning and disability as a dynamic interaction between health conditions and contextual factors, such as individual, social, and environmental factors. In this connection, and

to address the second goal of this study related to identify the factors that influenced QOL scores of stroke survivors were analysed by using path analysis. This research included the following categories of variables: socio-demographic and medical conditions, and functioning and disability (e.g., physical function, cognitive function, and psychological distress). Socio-demographic and medical conditions of stroke survivors were presumed to affect stroke survivor's QOL directly as well as indirectly through levels of functioning and disability.

The review of literature had examined the impact of numerous variables on caregiver's QOL, all of which were examined using structural equation modelling. The caregiver model for studying determinants of QOL in stroke caregivers was developed by identifying relevant factors from literature and mapping with coping and stress models (Chronister & Chan, 2006). Using this framework as a guide and theoretical and empirical evidence that supports the role of coping strategies and social support in adjustment to stress in stroke caregiving (Nir, Greenberger, & Bachner, 2009; Ostwald et al., 2009; Visser-Meily et al., 2009; Yu et al., 2013). In this research framework for caregiver's QOL, factors that affected caregivers' QOL include: caregiver variables, caregiving appraisal (e.g., satisfaction, burden), and coping strategies (e.g., adaptive, maladaptive coping). The framework was designed to provide a comprehensive depiction of determinants of caregiver QOL.

The purpose of this study was to determine caregiver's QOL, caregiving satisfaction, and caregiving burden during caregiving period. It was hypothesized that caregiver variables influenced caregiver QOL directly as well as indirectly through measures of caregiving appraisal and coping strategies. Next, the role of caregiving satisfaction, burden, and coping on caregivers' overall QOL scores were examined. Coping strategies were hypothesized to have direct and indirect effects on caregivers' QOL. In addition, coping strategies and caregiving appraisal were mediated the relationships between the caregiver variables and caregiver QOL.

Existing studies examined the presence of interdependence relationship between QOL among stroke survivors and caregivers (Kim, Reed, Hayward, Kang, & Koenig, 2011). The illness of a family member causes physical, emotional, and financial stresses, which influences the quality of life of other family members. Family members have a strong influence on a patient's psychological adjustment and management of illness. Further, the Actor-Partner Interdependence Model (APIM) provided both a conceptual and methodological view of understanding interdependent influences within dyadic relationship (Kenny, Kashy, & Cook, 2006). The APIM dyadic analysis uses the dyad as the unit of analysis and not individuals and provides estimates of actor and partner effects. Based on the APIM, it is possible to explore the actor and partner effects of QOL among stroke survivors and their caregivers at two time points. The study examined whether stroke survivor's and caregiver's QOL at baseline predicts his or her own QOL at six months of rehabilitation (e.g., an actor effect) and his or her partner's QOL at six months of rehabilitation (e.g., a partner effect).



**Figure 1: Conceptual framework of the QOL model of stroke survivors and caregivers**



## 1.7 Significance of study

In order to view QOL comprehensively, researchers have sought a way to summarize these QOL measures. An overall QOL score, including general health, physical, mental, faith, and peace indicators, was assessed in this study among stroke survivors and caregivers. The resulting overall QOL score may be applied to determine stroke population health status, assess health disparities among stroke survivors and caregivers, and identify risk factors of overall QOL. Measuring overall QOL score of stroke survivors and caregivers could be used to assist health professionals in assessing the effects of illness and treatment or rehabilitation.

This study makes an important contribution to the literature by filling a void that addresses a public health issue and provides preliminary research of family caregiving following stroke. Since caregiving has been shown to have an important and increasing impact on stroke patients, this first study using local samples has been carried out. The study provided a comprehensive model in explaining caregiving appraisal and coping strategies and its relation to QOL. Every caregiver adapts to burden differently during caregiving process. Knowledge on coping strategies and social support will guide healthcare providers to intervene using effective strategies to improve supportive care and psychological adjustment for caregivers to enhance their overall QOL. Hence, interventions designed to meet family caregivers' needs need to be implemented.

In addition, family caregiving of stroke patients not only have negative experiences, but also positive feelings in their caregiver role. This information is important as to determine whether stroke caregivers find benefits in their role and activities or caregivers experience high levels of stress and burden. This information will enrich caregiving knowledge within the Malaysian culture and thus help healthcare providers view comprehensive picture of family caregiving in stroke rehabilitation. Different coping strategies can be introduced to caregivers with different situation. Rehabilitation programs highlighting coping resources, including problem-focused coping styles and positive relationship coping strategies for caregivers, and psychoeducation programs to inform caregivers about stroke-related information, and readjustment of caregiver's life (Cheng, Chair, & Chau, 2014) can be used to minimize negative caregiving experiences.

The effects of stroke on daily living and well-being are interdependence between two persons in the dyad. Better understanding of dyad situation may serve as a baseline data for identifying vulnerable dyads in order to provide targeted support. Furthermore, a better understanding of the dyadic relations between stroke survivors and caregivers QOL over time can provide additional evidence and implications to stroke rehabilitation. QOL scores of stroke survivors and caregivers have important clinical implications for exploration of risk factors for stroke survivors and caregivers at risk of poor health outcomes.

This study assesses important caregiver and survivor interdependent relationship that could add to the existing literature on QOL studies after a stroke. Stroke disability extracts a toll on the QOL of patients and caregivers, these caregivers have an impact on the patient's adjustment to illness and recovery. Since there are interdependence relationship of QOL among dyads, there may also be changes in QOL among stroke patients and caregivers over time. Therefore, this study is the first kind in Malaysia to determine the dyadic relationships in QOL of stroke patients and caregivers over two time points. However, the APIM model in longitudinal dyadic study to examine the change in QOL are virtually unknown. The information can also contribute to the existing literature and prompt other researchers to investigate further on the mechanisms that contribute to dyadic relations of QOL of stroke survivors and caregivers. Consequently, the information can be used to promote the important of family interventions in successful stroke rehabilitation, not only direct rehabilitation for stroke survivors but also caregivers are an integral part of stroke rehabilitation process.



## REFERENCES

- Akosile, C. O., Okoye, E. C., Nwankwo, M. J., Akosile, C. O., & Mbada, C. E. (2011). Quality of life and its correlates in caregivers of stroke survivors from a Nigerian population. *Quality of Life Research*, 20(9), 1379-1384.
- Akyüz, M., Ünalın, H., Palamar, D., Demirdalı, A., Kutlu, A., Misirlioglu, T. Ö., . . . Akarirmak, Ü. (2015). Correlation of Upper Extremity Function to Quality of Life of Primary Caregivers of Ambulatory Stroke Survivors Living in the Community. *Neurosurgery Quarterly*, 25(1), 29-33.
- Algurén, B., Fridlund, B., Cieza, A., Sunnerhagen, K. S., & Christensson, L. (2012). Factors Associated With Health-Related Quality of Life After Stroke A 1-Year Prospective Cohort Study. *Neurorehabilitation and neural repair*, 26(3), 266-274.
- Appelros, P., Nydevik, I., & Viitanen, M. (2003). Poor outcome after first-ever stroke. *Stroke*, 34(1), 122-126.
- Aziz, A. F. A., Nordin, N. A. M., Aziz, N. A., Abdullah, S., Sulong, S., & Aljunid, S. M. (2014). Care for post-stroke patients at Malaysian public health centres: self-reported practices of family medicine specialists. *BMC family practice*, 15(1), 40.
- Aziz, Z. A., Lee, Y. Y., Ngah, B. A., Sidek, N. N., Looi, I., Hanip, M. R., & Basri, H. B. (2015). Acute Stroke Registry Malaysia, 2010-2014: Results from the National Neurology Registry. *Journal of Stroke and Cerebrovascular Diseases*, 24(12), 2701-2709.
- Azlin, N., & Rizal, A. (2009). Health related quality of life (HRQOL) among stroke survivors attending rehabilitation centres in Selangor. *Jurnal Kesihatan Masyarakat*, 15(2), 83-90.
- Bach, J. P., Riedel, O., Pieper, L., Klotsche, J., Dodel, R., & Wittchen, H. U. (2011). Health-related quality of life in patients with a history of myocardial infarction and stroke. *Cerebrovasc Dis*, 31(1), 68-76. doi:10.1159/000319027
- Bai, M., & Lazenby, M. (2015). A systematic review of associations between spiritual well-being and quality of life at the scale and factor levels in studies among patients with cancer. *Journal of palliative medicine*, 18(3), 286-298.
- Balhara, Y. P., Verma, R., Sharma, S., & Mathur, S. (2012). A study of predictors of anxiety and depression among stroke patient-caregivers. *J Midlife Health*, 3(1), 31-35. doi:10.4103/0976-7800.98815
- Baumann, M., Couffignal, S., Le Bihan, E., & Chau, N. (2012). Life satisfaction two-years after stroke onset: the effects of gender, sex occupational status, memory

- function and quality of life among stroke patients (Newsqol) and their family caregivers (Whoqol-bref) in Luxembourg. *BMC neurology*, 12(1), 1.
- Bergström, A. L., Eriksson, G., von Koch, L., & Tham, K. (2011). Combined life satisfaction of persons with stroke and their caregivers: associations with caregiver burden and the impact of stroke. *Health and quality of life outcomes*, 9(1), 1.
- Bertrand, R. M., Saczynski, J. S., Mezzacappa, C., Hulse, M., Ensrud, K., & Fredman, L. (2012). Caregiving and cognitive function in older women: Evidence for the healthy caregiver hypothesis. *Journal of Aging and Health*, 24(1), 48-66.
- Bluvol, A., & Ford-Gilboe, M. (2004). Hope, health work and quality of life in families of stroke survivors. *J Adv Nurs*, 48(4), 322-332. doi:10.1111/j.1365-2648.2004.03004.x
- Boden-Albala, B., & Sacco, R. L. (2000). Lifestyle factors and stroke risk: exercise, alcohol, diet, obesity, smoking, drug use, and stress. *Curr Atheroscler Rep*, 2(2), 160-166.
- Bowling, A. (1995). *Measuring disease : a review of disease specific quality of life measurement scales*. Buckingham ; Philadelphia: Open University Press.
- Bredle, J. M., Salsman, J. M., Debb, S. M., Arnold, B. J., & Cella, D. (2011). Spiritual well-being as a component of health-related quality of life: the functional assessment of chronic illness therapy—spiritual well-being scale (FACIT-Sp). *Religions*, 2(1), 77-94.
- Cameron, J. I., Stewart, D. E., Streiner, D. L., Coyte, P. C., & Cheung, A. M. (2014). What makes family caregivers happy during the first 2 years post stroke? *Stroke*, 45(4), 1084-1089.
- Carod-Artal, F. J., & Egido, J. A. (2009). Quality of life after stroke: the importance of a good recovery. *Cerebrovasc Dis*, 27 Suppl 1, 204-214. doi:10.1159/000200461
- Carter, J. H., Lyons, K. S., Stewart, B. J., Archbold, P. G., & Scobee, R. (2010). Does age make a difference in caregiver strain? Comparison of young versus older caregivers in early - stage Parkinson's disease. *Movement Disorders*, 25(6), 724-730.
- Cerniauskaite, M., Quintas, R., Koutsogeorgou, E., Meucci, P., Sattin, D., Leonardi, M., & Raggi, A. (2012). Quality-of-life and disability in patients with stroke. *Am J Phys Med Rehabil*, 91(13 Suppl 1), S39-47. doi:10.1097/PHM.0b013e31823d4df7
- Chen, C.-M., Tsai, C.-C., Chung, C.-Y., Chen, C.-L., Wu, K. P., & Chen, H.-C. (2015). Potential predictors for health-related quality of life in stroke patients undergoing inpatient rehabilitation. *Health and quality of life outcomes*, 13(1), 1.

- Chen, Y., Lu, J., Wong, K. S., Mok, V. C., Ungvari, G. S., & Tang, W. K. (2010). Health-related quality of life in the family caregivers of stroke survivors. *International Journal of Rehabilitation Research*, 33(3), 232-237.
- Cheng, H. Y., Chair, S. Y., & Chau, J. P.-C. (2014). The effectiveness of psychosocial interventions for stroke family caregivers and stroke survivors: A systematic review and meta-analysis. *Patient education and counseling*, 95(1), 30-44.
- Chronister, J., & Chan, F. (2006). A stress process model of caregiving for individuals with traumatic brain injury. *Rehabilitation Psychology*, 51(3), 190.
- Chung, M. L., Bakas, T., Plue, L. D., & Williams, L. S. (2016). Effects of self-esteem, optimism, and perceived control on depressive symptoms in stroke survivor-spouse dyads. *Journal of Cardiovascular Nursing*, 31(2), E8-E16.
- Clark, A. N., Sander, A. M., Pappadis, M. R., L Evans, G., & Chiou-Tan, F. Y. (2010). Caregiver characteristics and their relationship to health service utilization in minority patients with first episode stroke. *NeuroRehabilitation*, 27(1), 95-104.
- Coull, A. J., Lovett, J. K., Rothwell, P. M., & Oxford Vascular, S. (2004). Population based study of early risk of stroke after transient ischaemic attack or minor stroke: implications for public education and organisation of services. *BMJ*, 328(7435), 326. doi:10.1136/bmj.37991.635266.44
- Cramm, J. M., Strating, M. M., & Nieboer, A. P. (2012). Satisfaction with care as a quality-of-life predictor for stroke patients and their caregivers. *Quality of Life Research*, 21(10), 1719-1725.
- Davis, L. C., Sander, A. M., Struchen, M. A., Sherer, M., Nakase-Richardson, R., & Malec, J. F. (2009). Medical and psychosocial predictors of caregiver distress and perceived burden following traumatic brain injury. *The Journal of head trauma rehabilitation*, 24(3), 145-154.
- Dhamoon, M., Moon, Y., Paik, M., Boden-Albala, B., Rundek, T., Sacco, R. L., & Elkind, M. (2010). Quality of life declines after first ischemic stroke The Northern Manhattan Study. *Neurology*, 75(4), 328-334.
- Di Legge, S., Koch, G., Diomedi, M., Stanzione, P., & Sallustio, F. (2012). Stroke Prevention: Managing Modifiable Risk Factors. *Stroke Res Treat*, 2012. doi:10.1155/2012/391538
- Dimitrov, D. M., Raykov, T., & AL-Qataee, A. A. (2015). Developing a Measure of General Academic Ability: An Application of Maximal Reliability and Optimal Linear Combination to High School Students' Scores. *Educational and Psychological Measurement*, 75(3), 475-490.
- Ellis, C., Grubaugh, A. L., & Egede, L. E. (2013). Factors associated with SF-12 physical and mental health quality of life scores in adults with stroke. *J Stroke Cerebrovasc Dis*, 22(4), 309-317. doi:10.1016/j.jstrokecerebrovasdis.2011.09.007



- Epstein-Lubow, G. P., Beevers, C. G., Bishop, D. S., & Miller, I. W. (2009). Family functioning is associated with depressive symptoms in caregivers of acute stroke survivors. *Archives of physical medicine and rehabilitation*, 90(6), 947-955.
- Fatimang, L., & Rahmah, M. (2011). Penjagaan pesakit strok: adakah ia satu bebanan? apa yang penjaga persepikan? *Jurnal Kesihatan Masyarakat*, 17(1), 32-41.
- Fatoye, F. O., Komolafe, M. A., Adewuya, A. O., & Fatoye, G. K. (2006). Emotional distress and self-reported quality of life among primary caregivers of stroke survivors in Nigeria. *East Afr Med J*, 83(5), 271-279. doi:10.4314/eamj.v83i5.9433
- Feigin, V. L., Forouzanfar, M. H., Krishnamurthi, R., Mensah, G. A., Connor, M., Bennett, D. A., . . . Truelsen, T. (2014). Global and regional burden of stroke during 1990–2010: findings from the Global Burden of Disease Study 2010. *The Lancet*, 383(9913), 245-255.
- Feigin, V. L., Krishnamurthi, R. V., Parmar, P., Norrving, B., Mensah, G. A., Bennett, D. A., . . . Group, G. B. D. S. P. E. (2015). Update on the Global Burden of Ischemic and Hemorrhagic Stroke in 1990-2013: The GBD 2013 Study. *Neuroepidemiology*, 45(3), 161-176. doi:10.1159/000441085
- Ferrell, B. R. (1995). The impact of pain on quality of life. A decade of research. *Nurs Clin North Am*, 30(4), 609-624.
- Franzén - Dahlin, Å., & Laska, A. C. (2012). Gender differences in quality of life after stroke and TIA: a cross - sectional survey of out - patients. *Journal of clinical nursing*, 21(15 - 16), 2386-2391.
- Fredman, L., Lyons, J. G., Cauley, J. A., Hochberg, M., & Applebaum, K. M. (2015). The Relationship Between Caregiving and Mortality After Accounting for Time-Varying Caregiver Status and Addressing the Healthy Caregiver Hypothesis. *J Gerontol A Biol Sci Med Sci*, 70(9), 1163-1168. doi:10.1093/gerona/qlv009
- Fróes, K. S. d. S. O., Valdés, M. T. M., Lopes, D., & Silva, C. (2011). Factors associated with health-related quality of life for adults with stroke sequelae. *Arquivos de neuro-psiquiatria*, 69(2B), 371-376.
- Fung, T. T., Stampfer, M. J., Manson, J. E., Rexrode, K. M., Willett, W. C., & Hu, F. B. (2004). Prospective study of major dietary patterns and stroke risk in women. *Stroke*, 35(9), 2014-2019. doi:10.1161/01.STR.0000135762.89154.92
- Geyh, S., Cieza, A., Kollerits, B., Grimby, G., & Stucki, G. (2007). Content comparison of health-related quality of life measures used in stroke based on the international classification of functioning, disability and health (ICF): a systematic review. *Quality of Life Research*, 16(5), 833-851.

- Giaquinto, S., Spiridigliozzi, C., & Caracciolo, B. (2007). Can faith protect from emotional distress after stroke? *Stroke*, 38(3), 993-997. doi:10.1161/01.str.0000257996.26950.59
- Godwin, K. M., Ostwald, S. K., Cron, S. G., & Wasserman, J. (2013a). Long-term health-related quality of life of stroke survivors and their spousal caregivers. *J Neurosci Nurs*, 45(3), 147-154. doi:10.1097/JNN.0b013e31828a410b
- Godwin, K. M., Swank, P. R., Vaeth, P., & Ostwald, S. K. (2013b). The longitudinal and dyadic effects of mutuality on perceived stress for stroke survivors and their spousal caregivers. *Aging & mental health*, 17(4), 423-431.
- Guyatt, G. H., Feeny, D. H., & Patrick, D. L. (1993). Measuring health-related quality of life. *Ann Intern Med*, 118(8), 622-629.
- Haas, B. K. (1999). Clarification and integration of similar quality of life concepts. *Image: The Journal of Nursing Scholarship*, 31(3), 215-220.
- Haghgoo, H. A., Pazuki, E. S., Hosseini, A. S., & Rassafiani, M. (2013). Depression, activities of daily living and quality of life in patients with stroke. *Journal of the neurological sciences*, 328(1), 87-91.
- Haley, W. E., Allen, J. Y., Grant, J. S., Clay, O. J., Perkins, M., & Roth, D. L. (2009). Problems and benefits reported by stroke family caregivers results from a prospective epidemiological study. *Stroke*, 40(6), 2129-2133.
- Haley, W. E., Roth, D. L., Howard, G., & Safford, M. M. (2010). Caregiving strain and estimated risk for stroke and coronary heart disease among spouse caregivers differential effects by race and sex. *Stroke*, 41(2), 331-336.
- Haley, W. E., Roth, D. L., Kissela, B., Perkins, M., & Howard, G. (2011). Quality of life after stroke: a prospective longitudinal study. *Qual Life Res*, 20(6), 799-806. doi:10.1007/s11136-010-9810-6
- Hamza, A. M., Al-Sadat, N., Loh, S. Y., & Jahan, N. K. (2014). Predictors of poststroke health-related quality of life in Nigerian stroke survivors: a 1-year follow-up study. *BioMed research international*, 2014.
- Hayes, J., Chapman, P., Young, L. J., & Rittman, M. (2015). The prevalence of injury for stroke caregivers and associated risk factors. *Topics in stroke rehabilitation*.
- Hejazi, S. M. A., Mazlan, M., Abdullah, S. J. F., & Engkasan, J. P. (2015). Cost of post-stroke outpatient care in Malaysia. *Singapore medical journal*, 56(2), 116.
- Hopman, W. M., & Verner, J. (2003). Quality of life during and after inpatient stroke rehabilitation. *Stroke*, 34(3), 801-805.
- Howitt, S. C., Jones, M. P., Jusabani, A., Gray, W. K., Aris, E., Mugusi, F., . . . Walker, R. W. (2011). A cross-sectional study of quality of life in incident stroke

survivors in rural northern Tanzania. *J Neurol*, 258(8), 1422-1430. doi:10.1007/s00415-011-5948-6

Hulley, S. B., Cummings, S. R., Browner, W. S., Grady, D. G., & Newman, T. B. (2013). *Designing clinical research*: Lippincott Williams & Wilkins.

Hussain, N. A., Abdullah, M. R., Esa, A. R., Mustapha, M., & Yusoff, N. (2014). Predictors Of Life Satisfaction Among Family Caregivers Of Hospitalized First-Ever Stroke Patients In Kelantan. *ASEAN Journal of Psychiatry*, 15(2), 164-175.

Ibrahim, N. M., Shohaimi, S., Chong, H.-T., Rahman, A. H. A., Razali, R., Esther, E., & Basri, H. B. (2009). Validation study of the Mini-Mental State Examination in a Malay-speaking elderly population in Malaysia. *Dementia and geriatric cognitive disorders*, 27(3), 247-253.

Jeong, B.-O., Kang, H.-J., Bae, K.-Y., Kim, S.-W., Kim, J.-M., Shin, I.-S., . . . Yoon, J.-S. (2012). Determinants of quality of life in the acute stage following stroke. *Psychiatry investigation*, 9(2), 127-133.

Jeong, Y.-G., Jeong, Y.-J., Kim, W.-C., & Kim, J.-S. (2015). The mediating effect of caregiver burden on the caregivers' quality of life. *Journal of physical therapy science*, 27(5), 1543.

Johnstone, B., Franklin, K. L., Yoon, D. P., Burris, J., & Shigaki, C. (2008). Relationships among religiousness, spirituality, and health for individuals with stroke. *J Clin Psychol Med Settings*, 15(4), 308-313. doi:10.1007/s10880-008-9128-5

Jönsson, A.-C., Lindgren, I., Hallström, B., Norrving, B., & Lindgren, A. (2005). Determinants of quality of life in stroke survivors and their informal caregivers. *Stroke*, 36(4), 803-808.

Joo, H., Dunet, D. O., Fang, J., & Wang, G. (2014). Cost of informal caregiving associated with stroke among the elderly in the United States. *Neurology*, 83(20), 1831-1837.

Keller, S. D., Ware, J. E., Bentler, P. M., Aaronson, N. K., Alonso, J., Apolone, G., . . . Kaasa, S. (1998). Use of structural equation modeling to test the construct validity of the SF-36 health survey in ten countries: Results from the IQOLA project. *Journal of clinical epidemiology*, 51(11), 1179-1188.

Kendall, E., & Terry, D. (2009). Predicting emotional well-being following traumatic brain injury: a test of mediated and moderated models. *Soc Sci Med*, 69(6), 947-954. doi:S0277-9536(09)00376-1 [pii]10.1016/j.socscimed.2009.06.021

Kenny, D., Kashy, D., & Cook, W. (2006). *Dyadic Data Analysis*: New York: Guilford.



- Kim, S. S., Reed, P. G., Hayward, R. D., Kang, Y., & Koenig, H. G. (2011). Spirituality and psychological well - being: Testing a theory of family interdependence among family caregivers and their elders. *Research in nursing & health*, 34(2), 103-115.
- Kissela, B. M., Khoury, J. C., Alwell, K., Moomaw, C. J., Woo, D., Adeoye, O., . . . La Rosa, F. D. L. R. (2012). Age at stroke temporal trends in stroke incidence in a large, biracial population. *Neurology*, 79(17), 1781-1787.
- Kline, R. B. (2015). *Principles and practice of structural equation modeling*: Guilford publications.
- Krishnamoorthy, M. (2007, 2007 Apr 24 ). Killer stroke: Six Malaysians hit every hour. *The Star*. Retrieved from <http://www.thestar.com.my/story/?file=%2F2007%2F4%2F24%2Fnation%2F17524877>
- Krishnamurthi, R. V., Deveber, G., Feigin, V. L., Barker-Collo, S., Fullerton, H., Mackay, M. T., . . . Lo, W. (2015). Stroke prevalence, mortality and disability-adjusted life years in children and youth aged 0-19 years: data from the global and regional burden of stroke 2013. *Neuroepidemiology*, 45(3), 177-189.
- Kruithof, W. J., van Mierlo, M. L., Visser-Meily, J. M., van Heugten, C. M., & Post, M. W. (2013). Associations between social support and stroke survivors' health-related quality of life—a systematic review. *Patient education and counseling*, 93(2), 169-176.
- Lazarus, R. S., & Folkman, S. (1984). Coping and adaptation. *The handbook of behavioral medicine*, 282-325.
- Lee, C. D., Folsom, A. R., & Blair, S. N. (2003). Physical activity and stroke risk: a meta-analysis. *Stroke*, 34(10), 2475-2481. doi:10.1161/01.STR.0000091843.02517.9D
- Lee, M., Wu, Y. L., & Ovbiagele, B. (2016). Trends in Incident and Recurrent Rates of First-Ever Ischemic Stroke in Taiwan between 2000 and 2011. *J Stroke*, 18(1), 60-65. doi:10.5853/jos.2015.01326
- Lee, S.-H., Bae, S.-H., Hwang, J.-A., & Kim, K.-Y. (2017). The Effects of Stroke Patients' Physical Functions and Emotional Characteristics on their Quality of Life. *International Journal of Bio-Science and Bio-Technology*, 9, 53-64. doi:10.14257/ijbsbt.2017.9.1.04.
- Louie, D. R., & Eng, J. J. (2016). Powered robotic exoskeletons in post-stroke rehabilitation of gait: a scoping review. *Journal of neuroengineering and rehabilitation*, 13(1), 53.
- Lua, P. L., Mustapha, N., Abdullah, R., & Rahman, A. K. A. (2014). The Experiences And Challenges In Caring For HIV/AIDS Patients: A Qualitative Exploration Among. *ASEAN Journal of Psychiatry*, 120-130.

- Luengo-Fernandez, R., Gray, A. M., Bull, L., Welch, S., Cuthbertson, F., & Rothwell, P. M. (2013). Quality of life after TIA and stroke Ten-year results of the Oxford Vascular Study. *Neurology*, *81*(18), 1588-1595.
- Lui, M. H. L., Lee, D. T., Greenwood, N., & Ross, F. M. (2012). Informal stroke caregivers' self-appraised problem-solving abilities as a predictor of well-being and perceived social support. *Journal of clinical nursing*, *21*(1 - 2), 232-242.
- MacIsaac, L., Harrison, M. B., & Godfrey, C. (2010). Supportive care needs of caregivers of individuals following stroke: a synopsis of research. *Canadian journal of neuroscience nursing*, *32*(1), 39-46.
- Mackenzie, A., & Greenwood, N. (2012). Positive experiences of caregiving in stroke: a systematic review. *Disability and rehabilitation*, *34*(17), 1413-1422.
- Mahmud, W. M. R. W., Awang, A., & Mohamed, M. N. (2004). Psychometric evaluation of the medical outcome study (MOS) Social Support Survey among Malay postpartum women in kedah, north west of peninsular Malaysia. *The Malaysian journal of medical sciences: MJMS*, *11*(2), 26.
- Mayo, N. E., Scott, S. C., Bayley, M., Cheung, A., Garland, J., Jutai, J., & Wood-Dauphinee, S. (2015). Modeling health-related quality of life in people recovering from stroke. *Quality of Life Research*, *24*(1), 41-53.
- McCarthy, M., Lyons, K., & Powers, L. (2011). Expanding poststroke depression research: movement toward a dyadic perspective. *Topics in stroke rehabilitation*, *18*(5), 450.
- McLennon, S. M., Bakas, T., Jessup, N. M., Habermann, B., & Weaver, M. T. (2014). Task difficulty and life changes among stroke family caregivers: Relationship to depressive symptoms. *Archives of physical medicine and rehabilitation*, *95*(12), 2484-2490.
- McPherson, C. J., Wilson, K. G., Chyurlia, L., & Leclerc, C. (2011). The caregiving relationship and quality of life among partners of stroke survivors: a cross-sectional study. *Health and quality of life outcomes*, *9*(1), 1.
- Meyer, B. (2001). Coping with severe mental illness: Relations of the Brief COPE with symptoms, functioning, and well-being. *Journal of Psychopathology and Behavioral Assessment*, *23*(4), 265-277.
- Ministry of Health Malaysia, H. I. C. (2010). *Health Facts 2009*. Retrieved from [http://www.moh.gov.my/images/gallery/publications/md/hi/hi\\_2009.pdf](http://www.moh.gov.my/images/gallery/publications/md/hi/hi_2009.pdf).
- Mohd Zulkifly, M. F., Ghazali, S. E., Che Din, N., Singh, D. K. A., & Subramaniam, P. (2016). A Review of Risk Factors for Cognitive Impairment in Stroke Survivors. *The Scientific World Journal*, 2016.

- Morimoto, T., Schreiner, A. S., & Asano, H. (2003). Caregiver burden and health-related quality of life among Japanese stroke caregivers. *Age Ageing*, 32(2), 218-223. doi:10.1093/ageing/32.2.218
- Morris, J. H., van Wijck, F., Joice, S., & Donaghy, M. (2013). Predicting health related quality of life 6 months after stroke: the role of anxiety and upper limb dysfunction. *Disability and rehabilitation*, 35(4), 291-299.
- Moser, A., Stuck, A. E., Silliman, R. A., Ganz, P. A., & Clough-Gorr, K. M. (2012). The eight-item modified Medical Outcomes Study Social Support Survey: psychometric evaluation showed excellent performance. *Journal of clinical epidemiology*, 65(10), 1107-1116.
- Neelamegam, M., Looi, I., Cheah, W. K., Narayanan, P., Hamid, A. M. A., & Ong, L. M. (2013). Stroke incidence in the South West District of the Penang Island, Malaysia: PEARLS: Penang Acute stroke research longitudinal study. *Preventive medicine*, 57, S77-S79.
- Nichols-Larsen, D. S., Clark, P., Zeringue, A., Greenspan, A., & Blanton, S. (2005). Factors influencing stroke survivors' quality of life during subacute recovery. *Stroke*, 36(7), 1480-1484.
- Nir, Z., Greenberger, C., & Bachner, Y. G. (2009). Profile, burden, and quality of life of Israeli stroke survivor caregivers: a longitudinal study. *Journal of Neuroscience Nursing*, 41(2), 92-105.
- Nur, M., & Sulong, S. (2012). Direct medical cost of stroke: findings from a tertiary hospital in Malaysia. *Med J Malaysia*, 67(5), 473.
- O'Donnell, M. J., Xavier, D., Liu, L., Zhang, H., Chin, S. L., Rao-Melacini, P., . . . investigators, I. (2010). Risk factors for ischaemic and intracerebral haemorrhagic stroke in 22 countries (the INTERSTROKE study): a case-control study. *Lancet*, 376(9735), 112-123. doi:10.1016/S0140-6736(10)60834-3
- Opara, J. A., & Jaracz, K. (2010). Quality of life of post-stroke patients and their caregivers. *Journal of medicine and life*, 3(3), 216.
- Ostwald, S. K., Bernal, M. P., Cron, S. G., & Godwin, K. M. (2009). Stress experienced by stroke survivors and spousal caregivers during the first year after discharge from inpatient rehabilitation. *Topics in stroke rehabilitation*, 16(2), 93-104.
- Othman, Z., Wong, S. T., Drahman, I., & Zakaria, R. (2014). Caregiver Burden is Associated with Cognitive Decline and Physical Disability of Elderly Post-Stroke Patients. *Middle-East Journal of Scientific Research*, 22(9), 1265-1271.
- Owens, P. L., Bradley, E. H., Horwitz, S. M., Viscoli, C. M., Kernan, W. N., Brass, L. M., . . . Horwitz, R. I. (2002). Clinical assessment of function among women

- with a recent cerebrovascular event: a self-reported versus performance-based measure. *Annals of internal medicine*, 136(11), 802-811.
- Owolabi, M. (2013). Consistent determinants of post - stroke health - related quality of life across diverse cultures: Berlin–Ibadan study. *Acta Neurol Scand*, 128(5), 311-320.
- Padilla, G., Grant, M., Ferrell, B., & Presant, C. (1996). Quality of life-cancer. *Quality of life and pharmacoeconomics in clinical trials*, 301-308.
- Parag, V., Hackett, M. L., Yapa, C. M., Kerse, N., McNaughton, H., Feigin, V. L., & Anderson, C. S. (2008). The impact of stroke on unpaid caregivers: results from The Auckland Regional Community Stroke study, 2002–2003. *Cerebrovasc Dis*, 25(6), 548-554.
- Perrin, P. B., Norup, A., Caracuel, A., Bateman, A., Tjørnlund, M., & Arango - Lasprilla, J. C. (2016). An Actor - Partner Interdependence Model of Acquired Brain Injury Patient Impairments and Caregiver Psychosocial Functioning: A Dyadic - Report, Multinational Study. *Journal of Clinical Psychology*.
- Persson, J., Holmegaard, L., Karlberg, I., Redfors, P., Jood, K., Jern, C., . . . Forsberg-Wärleby, G. (2015). Spouses of Stroke Survivors Report Reduced Health-Related Quality of Life Even in Long-Term Follow-Up Results From Sahlgrenska Academy Study on Ischemic Stroke. *Stroke*, 46(9), 2584-2590.
- Prasomsri, J., Bovonsunthonchai, S., Khemthong, S., & Jalayondeja, C. (2014). *The effect of stress, anxiety and depression on functional recovery in individuals with stroke*. Paper presented at the 2nd ASEAN Plus Three Graduate Research Congress (2ndAGRC), Bangkok.
- Pucciarelli, G., Vellone, E., Savini, S., Simeone, S., Ausili, D., Alvaro, R., . . . Lyons, K. S. (2017). Roles of Changing Physical Function and Caregiver Burden on Quality of Life in Stroke: A Longitudinal Dyadic Analysis. *Stroke*, 48(3), 733-739. doi:10.1161/strokeaha.116.014989
- Pusat Informatik Kesehatan, & Perancangan, B. *Petunjuk Kesehatan 2014*. (1551-4589 MOH/S/RAN/74.14(TR)). Retrieved from [http://www.moh.gov.my/index.php/file\\_manager/dl\\_item/554756755a584a6961585268626939515a57356c636d4a706447467549465630595731684c304a31613356665547563064573571645774664d6a41784e4639496558426c636d7870626d73756347526d](http://www.moh.gov.my/index.php/file_manager/dl_item/554756755a584a6961585268626939515a57356c636d4a706447467549465630595731684c304a31613356665547563064573571645774664d6a41784e4639496558426c636d7870626d73756347526d).
- Rigby, H., Gubitz, G., & Phillips, S. (2009). A systematic review of caregiver burden following stroke. *International Journal of Stroke*, 4(4), 285-292.
- Romero, J. R., & Wolf, P. A. (2013). Epidemiology of Stroke: Legacy of the Framingham Heart Study. *Glob Heart*, 8(1), 67-75. doi:10.1016/j.gheart.2012.12.007



- Saha, B. K., Harries, P., & Gilhooly, K. J. (2016). Living with stroke in Bangladesh: Factors influencing quality of life (QoL). *World Federation of Occupational Therapists Bulletin*, 1-9.
- Salaycik, K. J., Kelly-Hayes, M., Beiser, A., Nguyen, A. H., Brady, S. M., Kase, C. S., & Wolf, P. A. (2007). Depressive symptoms and risk of stroke: the Framingham Study. *Stroke*, 38(1), 16-21. doi:10.1161/01.STR.0000251695.39877.ca
- Salter, K., Zettler, L., Foley, N., & Teasell, R. (2010). Impact of caring for individuals with stroke on perceived physical health of informal caregivers. *Disability and rehabilitation*, 32(4), 273-281.
- Saposnik, G., Cohen, L. G., Mamdani, M., Pooyania, S., Ploughman, M., Cheung, D., . . . Stroke Outcomes Research, C. (2016). Efficacy and safety of non-immersive virtual reality exercising in stroke rehabilitation (EVREST): a randomised, multicentre, single-blind, controlled trial. *Lancet Neurol*. doi:10.1016/S1474-4422(16)30121-1
- Saposnik, G., Levin, M., & Outcome Research Canada Working, G. (2011). Virtual reality in stroke rehabilitation: a meta-analysis and implications for clinicians. *Stroke*, 42(5), 1380-1386. doi:10.1161/STROKEAHA.110.605451
- Scherbakov, N., Von Haehling, S., Anker, S. D., Dirnagl, U., & Doehner, W. (2013). Stroke induced Sarcopenia: muscle wasting and disability after stroke. *International journal of cardiology*, 170(2), 89-94.
- Seshadri, S., Beiser, A., Pikula, A., Himali, J. J., Kelly-Hayes, M., Debette, S., . . . Wolf, P. A. (2010). Parental occurrence of stroke and risk of stroke in their children: the Framingham study. *Circulation*, 121(11), 1304-1312. doi:10.1161/CIRCULATIONAHA.109.854240
- Simon, C., Kumar, S., & Kendrick, T. (2009). Cohort study of informal carers of first-time stroke survivors: profile of health and social changes in the first year of caregiving. *Social science & medicine*, 69(3), 404-410.
- Sousa, K. H., & Kwok, O. M. (2006). Putting Wilson and Cleary to the test: analysis of a HRQOL conceptual model using structural equation modeling. *Qual Life Res*, 15(4), 725-737. doi:10.1007/s11136-005-3975-4
- Spaccavento, S., Craca, A., Del Prete, M., Falcone, R., Colucci, A., Di Palma, A., & Loverre, A. (2014). Quality of life measurement and outcome in aphasia. *Neuropsychiatr Dis Treat*, 10, 27-37. doi:10.2147/NDT.S52357
- Tarlow, B. J., Wisniewski, S. R., Belle, S. H., Rubert, M., Ory, M. G., & Gallagher-Thompson, D. (2004). Positive aspects of caregiving: Contributions of the REACH project to the development of new measures for Alzheimer's caregiving. *Research on aging*, 26(4), 429-453.

- Testa, M. A., & Simonson, D. C. (1996). Assessment of quality-of-life outcomes. *The New England Journal of Medicine*, 334(13), 835-840.
- Tramonti, F., Fanciullacci, C., Giunti, G., Rossi, B., & Chisari, C. (2014). Functional status and quality of life of stroke survivors undergoing rehabilitation programmes in a hospital setting. *NeuroRehabilitation*, 35(1), 1-7.
- van Mierlo, M. L., Schroder, C., van Heugten, C. M., Post, M. W., de Kort, P. L., & Visser-Meily, J. M. (2014). The influence of psychological factors on health-related quality of life after stroke: a systematic review. *Int J Stroke*, 9(3), 341-348. doi:10.1111/ijss.12149
- Van Peppen, R. P., Kwakkel, G., Wood-Dauphinee, S., Hendriks, H. J., Van der Wees, P. J., & Dekker, J. (2004). The impact of physical therapy on functional outcomes after stroke: what's the evidence? *Clinical rehabilitation*, 18(8), 833-862.
- Vincent - Onabajo, G., Ali, A., & Hamzat, T. (2013). Quality of life of Nigerian informal caregivers of community - dwelling stroke survivors. *Scandinavian journal of caring sciences*, 27(4), 977-982.
- Visser-Meily, A., Post, M., van de Port, I., Maas, C., Forstberg-Wärleby, G., & Lindeman, E. (2009). Psychosocial functioning of spouses of patients with stroke from initial inpatient rehabilitation to 3 years poststroke course and relations with coping strategies. *Stroke*, 40(4), 1399-1404.
- Visser, M. M., Heijenbrok-Kal, M. H., van't Spijker, A., Oostra, K. M., Busschbach, J. J., & Ribbers, G. M. (2015). Coping, problem solving, depression, and health-related quality of life in patients receiving outpatient stroke rehabilitation. *Archives of physical medicine and rehabilitation*, 96(8), 1492-1498.
- Ware, J. E., Jr. (1991). Conceptualizing and measuring generic health outcomes. *Cancer*, 67(3 Suppl), 774-779.
- WHOQOL Group. (1993). Study protocol for the World Health Organization project to develop a Quality of Life assessment instrument (WHOQOL). *Qual Life Res*, 2(2), 153-159.
- Wilson, I. B., & Cleary, P. D. (1995). Linking clinical variables with health-related quality of life. A conceptual model of patient outcomes. *JAMA*, 273(1), 59-65.
- World Health Organization. (1978). Cerebrovascular disorders (offset publication). Geneva: World Health Organization.
- Yin, S., Njai, R., Barker, L., Siegel, P. Z., & Liao, Y. (2016). Summarizing health-related quality of life (HRQOL): development and testing of a one-factor model. *Population health metrics*, 14(1), 22.

- Yu, Y., Hu, J., Eford, J. T., & McCoy, T. P. (2013). Social support, coping strategies and health - related quality of life among primary caregivers of stroke survivors in China. *Journal of clinical nursing*, 22(15-16), 2160-2171.
- Yusoff, M. S. B. (2011). The validity of the Malay Brief COPE in identifying coping strategies among adolescents in secondary school. *Int Med J*, 18, 29-33.
- Yusoff, N., Low, W., & Yip, C. (2009). Reliability and validity of the Malay version of Brief COPE scale: A study on Malaysian women treated with adjuvant chemotherapy for breast cancer. *Malaysian Journal of Psychiatry*, 18(1).
- Yusoff, N., Low, W. Y., & Yip, C.-H. (2011). Psychometric properties of the Malay Version of the hospital anxiety and depression scale: a study of husbands of breast cancer patients in Kuala Lumpur, Malaysia. *Asian Pac J Cancer Prev*, 12(4), 915-917.
- Yusoff, U. N., Mahat, D., Omar, A., Teh, C. H., Yoep, N., & Saari, R. (2013). Burden of premature mortality in Malaysia. *International Journal of Public Health Research*, 3(1), 249-258.
- Zhang, X., Sun, Q., Wu, M., & Xia, G. (2013). Health-related quality of life after stroke: a 2-year prospective cohort study in Wuhan, China. *Int J Neurosci*, 123(2), 138-141. doi:10.3109/00207454.2012.746336

## BIODATA OF STUDENT

This is Khaw Wan-Fei, who was born in Penang. She obtained her primary education at S.R.J.K. (c) Jit Sin (B) in year 1996 and then secondary education at S.M.J.K. Jit Sin, Bukit Mertajam in 2003. Following this in year 2004, she continued her tertiary education in Universiti Putra Malaysia and graduated with Bachelor of Science (Biomedical Sciences) in 2007. She completed her Master of Science in Community Health in 2010. Currently, she pursues a PhD in Epidemiology and Biostatistics at Faculty of Medicine and Health Sciences, UPM.





## LIST OF PUBLICATIONS

- Khaw WF, Hassan STS, Lye MS & Siti Irma FI. 2017. Comparing Spirituality and Quality of Life between Stroke Survivors and Their Family Caregivers. *International Journal of Public Health and Clinical Sciences*, 4(6), 41-51.
- Khaw WF, Hassan STS, Lye MS, Siti Irma FI, Rosna AR, Faisal I. 2017. Depression, anxiety and quality of life in stroke survivors and their family caregivers: A pilot study using an actor/partner interdependence model. *Electronic Physician* 9 (8), 4924-4933.





**UNIVERSITI PUTRA MALAYSIA**

**STATUS CONFIRMATION FOR THESIS / PROJECT REPORT AND COPYRIGHT**

**ACADEMIC SESSION :** \_\_\_\_\_

**TITLE OF THESIS / PROJECT REPORT :**

---

---

---

**NAME OF STUDENT :** \_\_\_\_\_

I acknowledge that the copyright and other intellectual property in the thesis/project report belonged to Universiti Putra Malaysia and I agree to allow this thesis/project report to be placed at the library under the following terms:

1. This thesis/project report is the property of Universiti Putra Malaysia.
2. The library of Universiti Putra Malaysia has the right to make copies for educational purposes only.
3. The library of Universiti Putra Malaysia is allowed to make copies of this thesis for academic exchange.

I declare that this thesis is classified as :

\*Please tick (v )

**CONFIDENTIAL**

(Contain confidential information under Official Secret Act 1972).

**RESTRICTED**

(Contains restricted information as specified by the organization/institution where research was done).

**OPEN ACCESS**

I agree that my thesis/project report to be published as hard copy or online open access.

This thesis is submitted for :

**PATENT**

Embargo from \_\_\_\_\_ until \_\_\_\_\_  
(date) (date)

**Approved by:**

\_\_\_\_\_  
(Signature of Student)  
New IC No/ Passport No.:

\_\_\_\_\_  
(Signature of Chairman of Supervisory Committee)  
Name:

Date :

Date :

**[Note : If the thesis is CONFIDENTIAL or RESTRICTED, please attach with the letter from the organization/institution with period and reasons for confidentially or restricted. ]**