

Review article:

Use of Quantitative Electroencephalography Biomarker in Characterizing Mild Cognitive Impairment in Malaysian Elderly

Ismail Samhani¹, Norhidayah Badya¹, Mohammed Faruque Reza², Nordin Simbak¹

Abstract:

The elderly population rising rapidly in Malaysia and contributes to the increasing number of cognitive problems including mild cognitive impairment (MCI). However, due to limited information regarding this problem which may progress towards severe neurologic degeneration, this problem rarely diagnosed and left untreated. Hence, the use of electroencephalography (EEG) biomarker is seen to be important with the spectral power, coherence and synchronization between the both halves of brain explain the pathophysiology underlined. This method is becoming popular for its capabilities in quantifying changes in brain electrical activity and provide early signs of brain impairment. This paper reviews the incidence of mild cognitive impairment in elderly as an early cognitive deterioration signal. It continues with the role of quantitative EEG analysis in providing the physiological meaning of their brain. This paper also provides the information about neurochemical changes which associated with MCI. In addition, this paper proposes an idea of study towards examining physiology, neurochemicals, diet and lifestyles to promote healthy lifestyle in elderly.

Keywords: Elderly, mild cognitive impairment (MCI), multivariate analysis, quantitative electroencephalography (qEEG).

*International Journal of Human and Health Sciences Vol. 04 No. 04 October '20 Page : 267-270
DOI: <http://dx.doi.org/10.31344/ijhhs.v4i4.211>*

Introduction

The pace of population aging is dramatically increasing worldwide inflicting in various social, economic and health implications. Malaysia is predicted to achieve the status of aging nation by 2030 with an estimated of 15.3% of its population comprising elderly with Terengganu and Kelantan win the race, by which is classified as one of the fastest nation to achieve aged country status within 20 years¹. The rapid growth of elderly people (65 years above) increase the incidence of aging disorders such as mild cognitive impairment (MCI).

Aging is a physiological process affecting all body tissues, hence, in relating to brain and cognition,

aging is associated with cognitive decline such as attention, memory and other cognitive domains inflicting in delay of cognitive processing such as input and coding, central processing and decision making². Areas of orientation, concentration and functioning and self-care had objectively dysfunction as well, whereas males significantly greater than females, in Indian population³.

Mild Cognitive Impairment in elderly

MCI is a condition characterized by a reduction in memory and other cognitive process. However, this cognitive decline is not sufficiently severe to be diagnosed as dementia. MCI is an intermediate state lying between normal cognition and progression towards dementia with 50% of MCI

1. Faculty of Medicine, Universiti Sultan Zainal Abidin (UniSZA) Medical Campus, 20400 Jalan Sultan Mahmud, Kuala Terengganu, Terengganu, Malaysia.
2. Department of Neurosciences, School of Medical Sciences, Universiti Sains Malaysia (Health Campus), 16150 Kubang Kerian, Kelantan.

Correspondence to: Ismail Samhani, Faculty of Medicine, UniSZA Medical Campus, Jalan Sultan Mahmud, 20400, Kuala Terengganu, Terengganu, Malaysia.

E-mail: samhanismail@unisza.edu.my

elderly will subsequently develop dementia in upcoming five years⁴. Elderly with MCI progress to Alzheimer Disease at a rate of 10-15% per year, while healthy normal elderly convert at a rate of 1-2% only^{5,6}.

MCI is a common cognitive problem but not get attention since it does not obviously exhibit clinical symptoms those interfering with daily life, subsequently it is left untreated. Roughly, 60% of people diagnosed with MCI progressed with worse neurological and cognitive degeneration inflicting in dementia and Alzheimer's disease which incur higher costs for their management. Hence, early detection and interventions are vital. But, since the information of MCI related to physiology, biology, biochemistry and such are limited, MCI rarely diagnosed, leaving them worsening.

Quantitative Electroencephalographic potential

EEG is a non-invasive measurement that generates EEG waveforms signal. It is analysed quantitatively by transforming the waveform into the digital signal data through specific algorithms such as Fast Fourier Transform, it could explain wider aspects such as cognitive and emotion which represents the underlying physiology of the elderly.

Quantitative EEG (qEEG) analysis provides information on physiologically meaningful frequency components, dynamic alterations and topography of EEG signal generators, i.e. neuronal signaling. Numerous studies have shown that qEEG measures can detect disruptions in activity, topographical distribution and synchronization of neuronal (synaptic) activity such as generalized EEG slowing, reduced global synchronization and anteriorization of neuronal generators of fast-frequency resting-state EEG activity in elderly⁷.

Recent study also found that topographic EEG revealed the increase of beta 2 power over the right anterior region in comparison with normal healthy aging which distinguished their anxiety level. Both the healthy and MCI groups exhibited a predominant distribution of theta and alpha at the frontal region. But the theta are highest at the parietal and temporal areas⁸ indicating of cognitive decline among the MCI elderly, which is a qEEG characteristic of atrophy as well as memory deficit in MCI elderly⁹. A reduction of delta power at the

prefrontal area (F3, Fz and F4) and the central regions (C3, Cz and C4) explained the cognitive decline in healthy elderly⁸.

EEG coherence measures the cortical connection functionality and quantify cortico-cortico or cortico-subcortical connection. Coherence can also be used to quantify the linear correlation and detect synchronization between two channels. A decrease in coherence is interpreted as a reduction in linear function connection and function coupling in the cortical area¹⁰. The former index recorded that the higher the coherence, the higher the synchrony while higher synchrony reflects a functional linkage between the brain regions of interest. Some studies have found that MCI exhibited a reduction in coherence¹¹.

Neurochemical changes in MCI

The causes of cognitive decline can arise through neurochemical changes. Neurotransmitters are essential neurochemicals that maintain synaptic and cognitive functions by transmitting signals across synaptic neurons. Default in cognitive function contributes to the onset of age-related dysfunctions and diseases' development, thus, measuring neurochemical messengers releases that regulate the brain function is crucial. In fact, researchers suggest that the brain generates less chemical messengers with aging, which showed a decreased level of most neurotransmitters such as acetylcholine, glutamate, dopamine, serotonin and norepinephrine activity¹²⁻¹⁵ that could play a role in declining cognition and memory.

Acetylcholine is produced in cholinergic neurons that is an essential neurotransmitter in the central and the peripheral nervous system. It also is the only neurotransmitter in the motor function of the somatic nervous system. In the peripheral nervous system, acetylcholine activates skeletal muscles as well as smooth muscle and cardiac muscle function. While within the central nervous system, it acts as a neuromodulator for the cholinergic system, which causes excitatory actions that is involved with plasticity, excitability, arousal, and reward. Acetylcholine release and signalling's disturbance can have a profound impact on neurological function¹⁶⁻¹⁸. Decreases in motor control has been postulated among a major factor leading to the decline of autonomy

capacity and quality of life in elderly populations worldwide¹⁹.

While the most plentiful neurotransmitter found in the nervous system is glutamate, where it plays a major role in cognitive functions such as memory and learning. It also appears to be involved in motor behaviour function which change with age^{20,21}. Glutamate abnormalities may disturb brain function by which dysregulation of it leads to cognitive impairment and neurodegeneration that plays a role in a number of neuropsychiatric diseases, apart from normal aging process. Hence, neurotransmitter can be a potential plasma biomarker in early diagnosis of neurodegenerative diseases, such as Alzheimer disease^{22,23}.

Diet and lifestyles

In addition, poor diets and passive lifestyle practicing and cultural engagement in the elderly also is associated with a remarkable loss of many neurological functions and cognitive decline through the ageing process. Other potential contributors to non-pathological cognitive ageing also includes cardiovascular disease, sleep,

smoking and alcohol²⁴.

It is expected that there are differences in MCI elderly's brains and healthy in terms of alpha, beta, theta, delta and gamma oscillations. Moreover, the alpha-beta ratio and alpha-theta ratio which show the cognitive decline are expected to change. Furthermore, the lifestyle, nutrition and spiritual practices would give impacts on brain activities as well as cognitive performance. By uncovering the biomarker of MCI in elderly, it gives hope to provide detail explanation of MCI in Malaysian elderly in physiological, biochemical and lifestyle perspectives.

Conclusion

Through our study, we put a hope to contribute a new knowledge about the electroencephalographic information and neurochemicals as well as lifestyle those contributes to the development of MCI in elderly. Perhaps, this research will provide insights for prevention and healthy lifestyles for elderly.

Conflict of Interest:No conflict of interest

Funding: Not related.

References:

1. Mahmud A, Mazalan M, Razak A, Rasyidee A. Ageing phenomenon: Malaysia towards 2030. 2017.
2. Ishii R, Canuet L, Aoki Y, Hata M, Iwase M, Ikeda S, et al. Healthy and Pathological Brain Aging: From the Perspective of Oscillations, Functional Connectivity, and Signal Complexity. *Neuropsychobiology*. 2017;75(4):151-61.
3. Tripathi RK, Tiwari SC. Cognitive Dysfunction in Normally Aging Urban Older Adults: A Community-based Study. *Indian J Psychol Med*. 2011;33(2):177-81.
4. Peng Z, Jiang H, Wang X, Huang K, Zuo Y, Wu X, et al. The Efficacy of Cognitive Training for Elderly Chinese Individuals with Mild Cognitive Impairment. *BioMed Research International*. 2019;2019.
5. Petersen RC. Mild cognitive impairment as a diagnostic entity. *Journal of Internal Medicine*. 2004;256(3):183-94.
6. Yin S, Zhu X, Huang X, Li J. Visuospatial characteristics of an elderly Chinese population: results from the WAIS-R block design test. *Frontiers in Aging Neuroscience*. 2015;7(17).
7. Smailovic U, Jelic V. Neurophysiological Markers of Alzheimer's Disease: Quantitative EEG Approach. *Neurology and Therapy*. 2019;8(2):37-55.
8. Fauzana N, Amrana NH. Brain Dynamics of Mild Cognitive Impairment (MCI) from EEG. *Procedia-Social and Behavioral Sciences*. 2015;165:284-90.
9. Moretti D, Zanetti O, Binetti G, Frisoni G. Quantitative EEG markers in mild cognitive impairment: degenerative versus vascular brain impairment. *International Journal of Alzheimer's Disease*. 2012;2012.
10. Al-Qazzaz NK, Ali SHB, Ahmad SA, Chellappan K, Islam M, Escudero J. Role of EEG as biomarker in the early detection and classification of dementia. *The Scientific World Journal*. 2014;2014.
11. Baker M, Akrofi K, Schiffer R, Boyle MWO. EEG Patterns in Mild Cognitive Impairment (MCI) Patients. *Open Neuroimag J*. 2008;2:52-5.
12. Lanari A, Amenta F, Silvestrelli G, Tomassoni D, Parnetti L. Neurotransmitter deficits in behavioural and psychological symptoms of Alzheimer's disease. *Mechanisms of Ageing and Development*. 2006;127(2):158-65.
13. Rodríguez JJ, Noristani HN, Verkhatsky A. The serotonergic system in ageing and Alzheimer's disease. *Progress in Neurobiology*. 2012;99(1):15-41.
14. Sarter M, Bruno JP, Parikh V. Abnormal Neurotransmitter Release Underlying Behavioral and Cognitive Disorders: Toward Concepts of Dynamic and Function-Specific Dysregulation. *Neuropsychopharmacology*. 2007;32(7):1452-61.
15. Wong D, Atiya S, Fogarty J, Montero-Odasso M, Pasternak SH, Brymer C, et al. Reduced Hippocampal Glutamate and Posterior Cingulate N-Acetyl Aspartate in Mild Cognitive Impairment and Alzheimer's Disease Is Associated with Episodic Memory Performance and White Matter Integrity in the Cingulum: A Pilot Study. *Journal of Alzheimer's Disease*. 2020(Preprint):1-21.
16. DeKosky ST, Ikonomic MD, Styren SD, Beckett L, Wisniewski S, Bennett DA, et al. Upregulation of choline acetyltransferase activity in hippocampus and frontal cortex of elderly subjects with mild cognitive impairment. *Annals of Neurology*. 2002;51(2):145-55.
17. Haense C, Kalbe E, Herholz K, Hohmann C, Neumaier B, Kraiss R, et al. Cholinergic system function and cognition in mild cognitive impairment. *Neurobiology of Aging*. 2012;33(5):867-77.
18. Rinne JO, Kaasinen V, Järvenpää T, Nägren K, Roivainen A, Yu M, et al. Brain acetylcholinesterase activity in mild cognitive impairment and early Alzheimer's disease. *Journal of Neurology, Neurosurgery & Psychiatry*. 2003;74(1):113.
19. Clark BC, Woods AJ, Clark LA, Criss CR, Shadmehr R, Grooms DR. The Aging Brain & the Dorsal Basal Ganglia: Implications for Age-Related Limitations of Mobility. *Advances in geriatric medicine and research*. 2019;1:e190008.
20. McEntee WJ, Crook TH. Glutamate: its role in learning, memory, and the aging brain. *Psychopharmacology*. 1993;111(4):391-401.
21. Segovia G, Porras A, Del Arco A, Mora F. Glutamatergic neurotransmission in aging: a critical perspective. *Mechanisms of Ageing and Development*. 2001;122(1):1-29.
22. Gruden MA, Davidova TB, Mališauskas M, Sewell RDE, Voskresenskaya NI, Wilhelm K, et al. Differential neuroimmune markers to the onset of Alzheimer's disease neurodegeneration and dementia: Autoantibodies to A β (25–35) oligomers, S100b and neurotransmitters. *Journal of Neuroimmunology*. 2007;186(1):181-92.
23. Peña-Bautista C, Flor L, López-Nogueroles M, García L, Ferrer I, Baquero M, et al. Plasma alterations in cholinergic and serotonergic systems in early Alzheimer Disease: Diagnosis utility. *Clinica Chimica Acta*. 2020;500:233-40.
24. Deary IJ, Corley J, Gow AJ, Harris SE, Houlihan LM, Marioni RE, et al. Age-associated cognitive decline. *British Medical Bulletin*. 2009;92(1):135-52.