# Study of Brain Morphology in Healthy Aging and Alzheimer Disease in Sudan using MRI

# Inaam Eltoum<sup>1</sup>, Mohamed Yousef<sup>1, 2</sup>, Shazaly Khojaly<sup>1, 4</sup> Ahmed Abukonna<sup>1</sup>, Mohammed Salih<sup>3</sup>

<sup>1</sup> Sudan University for Sciences and Technology, College of Radiological Sciences, Khartoum, Sudan

<sup>2</sup>Batterjee Medical College, Department of Radiological Science, Jeddah, Saudi Arabia

<sup>3</sup>Royal Care International Hospital, Khartoum, Sudan

<sup>4</sup>Al-Ghad International College for Applied Medical Science, Medical Imaging Technology Department, Abha, KSA

Abstract: The present study fills a gap in the Study of Brain Morphology in Healthy Aging and Alzheimer disease in Sudan using MRI and the main objective of this study is to evaluate the Morphology of the Brain using MRI and CT in Healthy Aging and Alzheimer's in Sudan. This study was done in Sudan University of science and technology and Department of Diagnostic Radiology, Khartoum state Hospitals, Khartoum, Sudan, during the period from April 2015 to June 2017. This study included two groups. Group (A) were 100 healthy individuals, (66 males (66%), 33 females (44%). Group (B) 300 patients, 198 male (66. %) and 102 females (44.0%) with high probability of AD. The mean age of all patients is 45years ranging (15-75) years additional to control group. MRS studies were performed on 1.5 Tesla Toshiba Exclelart Vantage whole body MR systems using standard imaging head coil. The results of this study revealed that the most affected patients by Alzheimer's disease (AD) were male 202 (67%), their ages above 7th decay; the most common sign is Apraxia (77%) and Amnesia (76%). And this study revealed that Predemntia is most type affected by patients (35%) (72 male, 35 female). The majority of patients diagnosed in moderate stage of AD. Early stage of Alzheimer's disease (AD) is rarely affected by patients or rarely diagnosed because with hidden signs and symptoms. The causes the Alzheimer's disease (AD) in this study as the following the most cause of AD is Genetic factor 44% (80male, 52 females), Amyloid Hypothesis factor is 43 % (78 male, 52 females) and rare cause factor is Tau Hypothesis 07 %(15 male, 06 females). Diagnostic Imaging Factors the Alzheimer's disease revealed that the Hypocampal Atrophy imaging factor is most predictive factors in diagnosis by Magnetic Resonance Imaging (MRI) (71%), Temporal lobe atrophy (MTA) factor (71%), Diffuse cerebral Atrophy factor is (68%) and Increase size of Ventricles factor(66%). This study concluded that the majority of patients were male, the most common sign is Apraxia, the majority of patients diagnosed in moderate stage of AD, the most cause of is Genetic factor and the Hypocampal Atrophy imaging factor is most predictive factors in diagnosis by Magnetic Resonance Imaging (MRI.

Keywords: Alzheimer disease, neuropathology, brain atrophy, MRI

### 1. Introduction

Alzheimer's disease (AD) is a multifaceted disease in which cumulative pathological brain insults result in progressive cognitive decline that ultimately leads to dementia. Amyloid plaques, neurofibrillary tangles (NFTs), neurodegeneration, and inflammation are the well-established pathological hallmarks of AD. A plausible model for the development of AD posits that amyloid deposition occurs early in the process but by itself does not directly cause clinical symptoms [1, 2]. Neuronal and synaptic losses appear to be key determinants of cognitive impairment in AD [3, 4]. If neuronal loss leads to cerebral atrophy (as is likely), then it can be expected that cognitive decline and atrophy will be closely associated.

On the basis of this evidence, it has been hypothesized that AD pathological cascade is a two-stage process in which amyloidosis and neuronal pathology are largely sequential rather than simultaneous processes [5, 6]. There is also sufficient literature to support the fact that atrophy of the brain structures or neurodegeneration is the most proximate substrate of cognitive impairment in AD [7-9]. Structural magnetic resonance imaging (sMRI) measures brain morphometry and therefore can capture gray matter atrophy related to the loss of neurons, synapses, and dendritic dearborization that occurs on a microscopic level in AD; white matter atrophy related to the loss of structural integrity of white matter tracts, presumably resulting from demyelination and dying back of axonal processes; and exvacuo expansion

of cerebrospinal fluid (CSF) spaces. It has been shown that neuronal loss correlates with but exceeds NFT density in AD and is related directly to impaired cognitive function [10]. Neuronal loss also correlates with Braak NFT stage and quantitative NFT burden, validating sMRI as an AD biomarker [11-13]. According to our knowledge, information on dementia prevalence in Sudan is very limited.

The present study fills a gap in the Study of Brain Morphology in Healthy Aging and Alzheimer disease in Sudan using MRI and the main objective of this study is to evaluate the Morphology of the Brain using MRI and CT in Healthy Aging and Alzheimer's in Sudan.

### 2. Material and Method

The study is performed In Sudan. In Khartoum state hospitals (Omdurman Military Hospital, Khartoum Teaching hospital, Khartoum Public teaching Hospital, Khartoum North, Imperial Private Hospital and polyclinics privates).

This study include two groups.Group (A) were 100 healthy individuals, (66 males (66%), 33 females (44%).Group (B) 300 patients, 198 male (66. %) And 102 females (44.0%). with high probability of AD. The mean age of all patients is 45 years ranging (15-75) years additional to control group. Standardized forms were used to collect data on mores variables divided into main categories, namely age, sex,

DOI: 10.21275/23081715

#### International Journal of Science and Research (IJSR) ISSN (Online): 2319-7064 Index Copernicus Value (2017): 78.96 | Impact Factor (2015): 6.391

Signs the (AD), Types the (AD), and causes (AD) and Diagnostic Imaging Factors the (AD). Data collection according to work sheet (Appendix) includes all above variables data. (Attached sheet). Data analysis by using SPSS. 16 MRS studies were performed on 1.5 Tesla Toshiba whole body MR systems using standard imaging head coil. Routine brain MRI was performed in 3 orthogonal planes, including at least T1, T2, and fluid-attenuated inversion recovery (FLAIR) weighted images.

Coronal-oblique T1-weighted images are used for the assessment of medial temporal lobe and hippocampal atrophy. They are obtained in a plane orthogonal to the long axis of the hippocampus; this plane is orientated parallel to the brainstem. These should be thin-section images and are ideally obtained by reformatting a sagittal 3D T1 sequence through the entire brain. Additional sagittal reconstructions will enable the assessment of midline structures as well as parietal atrophy, which may be involved in certain neurodegenerative disorders. FLAIR images are used to assess global cortical atrophy (GCA), vascular white matter hyperintensities and infarctions. T2-weighted images are used to assess infarctions, in particular lacunar infarctions in the thalamus and basal ganglia, which can be missed on FLAIR images.

T2\*-weighted images are necessary to detect microbleeds in amyloid angiopathy. These images can also depict calcifications and iron deposition.

DWI should be considered as a supplemental sequence in young patients or in rapidly progressive neurodegenerative disorders (DD - vasculitis, CJD).

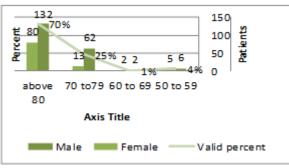
## 3. Result Presentation:

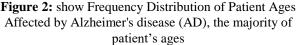
Table 1: Shows Frequency Distribution of Diagnosti	ic
Imaging Factors the Alzheimer's disease	

Diagnostic Imaging Factors	Male	Female	Total	Valid
Of AD				percent
Hypocampal Atrophy	130	86	216	71%
Temporal lobe atrophy	152	63	215	71%
(MTA)				
Atrophy (Koedam score)	126	39	165	55%
Posterior cortical atrophy	99	76	175	58%
Diffuse cerebral Atrophy	132	73	205	68%
Enlarge Sulci	84	49	133	44%
Increase size of Ventricles	121	79	200	66%
white matter lesions	141	48	189	63%
(Fazekas Scales)				

	Frequency	Percent	Valid Percent
Male	202	67	67
Female	100	33	33
Total	302	100	100

Figure 1: Show frequency Distribution of Patient Gender Affected by Alzheimer's disease (AD)





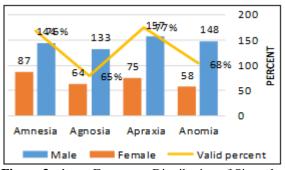


Figure 3: shows Frequency Distribution of Signs the Alzheimer's disease (AD)

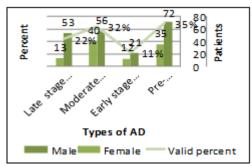


Figure 4: shows Frequency Distribution of Types the Alzheimer's disease

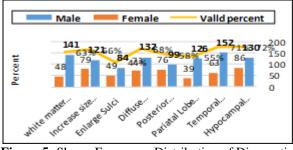


Figure 5: Shows Frequency Distribution of Diagnostic Imaging

## 4. Discussion

This study was done at Sudan University of science and technology and Department of Diagnostic Radiology, Khartoum state Hospitals, - Khartoum, Sudan, during the period from April 2015 to June 2017. The results of this study that the majority of patients were male 202 (67%) more than female 100(33%).and the Frequency Distribution of Patient Ages Affected by Alzheimer's disease (AD), the majority of

# Volume 6 Issue 10, October 2017

<u>www.ijsr.net</u>

Licensed Under Creative Commons Attribution CC BY

#### International Journal of Science and Research (IJSR) ISSN (Online): 2319-7064 Index Copernicus Value (2017): 78.96 | Impact Factor (2015): 6.391

patients ages above 7th decay approximately male more female. The results of this study revealed that the most affected patients by Alzheimer's disease (AD) were male 202 (67%), their ages above 7th decay; the most common sign is Apraxia (77%) and Amnesia (76%). And this study revealed that Predemntia is most type affected by patients (35%) (72 male, 35 female). The majority of patients diagnosed in moderate stage of AD. Early stage of Alzheimer's disease (AD) is rarely affected by patients or rarely diagnosed because with hidden signs and symptoms. The causes the Alzheimer's disease (AD) in this study as the following the most cause of AD is Genetic factor 44% (80male, 52 females), Amyloid Hypothesis factor is 43 %( 78male, 52 females) and rare cause factor is Tau Hypothesis 07 %(15 male, 06 females). Diagnostic Imaging Factors the Alzheimer's disease revealed that the Hypocampal Atrophy imaging factor is most predictive factors in diagnosis by Magnetic Resonance Imaging (MRI) (71%), Temporal lobe atrophy (MTA) factor (71%), diffuse cerebral Atrophy factor is (68%) and Increase size of Ventricles factor (66%). This study concluded that the majority of patients were male, the most common sign is Apraxia, the majority of patients diagnosed in moderate stage of AD, the most cause of is Genetic factor and the Hypocampal Atrophy imaging factor is most predictive factors in diagnosis by Magnetic Resonance Imaging (MRI).

## 5. Conclusion

This study concluded that the majority of patients were male, the most common sign is Apraxia, the majority of patients diagnosed in moderate stage of AD, the most cause of is Genetic factor and the Hypocampal Atrophy imaging factor is most predictive factors in diagnosis by Magnetic Resonance Imaging (MRI)

## References

- [1] Jack CR Jr., Lowe VJ, Weigand SD, Wiste HJ, Senjem ML, Knopman DS, Shiung MM, Gunter JL, Boeve BF, Kemp BJ, Weiner M, Petersen RC; Alzheimer's Disease Neuroimaging Initiative: Serial PIB and MRI in normal, mild cognitive impairment and Alzheimer's disease: implications for sequence of pathological events in Alzheimer's disease. Brain 2009, 132:1355-1365.
- [2] Mormino EC, Kluth JT, Madison CM, Rabinovici GD, Baker SL, Miller BL, Koeppe RA, Mathis CA, Weiner MW, Jagust WJ; Alzheimer 's disease Neuroimaging Initiative: Episodic memory loss is related to hippocampalmediated beta-amyloid deposition in elderly subjects. Brain 2009, 132:1310-1323.
- [3] DeKosky ST, Scheff SW: Synapse loss in frontal cortex biopsies in Alzheimer's disease: correlation with cognitive severity. Ann Neurol 1990, 27:457-464.
- [4] Terry RD, Masliah E, Salmon DP, Butters N, DeTeresa R, Hill R, Hansen LA, Katzman R: Physical basis of cognitive alterations in Alzheimer's disease: synapse loss is the major correlate of cognitive impairment. Ann Neurol 1991, 30:572-580.
- [5] Ingelsson M, Fukumoto H, Newell KL, Growdon JH, Hedley-Whyte ET, Frosch MP, Albert MS, Hyman BT,

Irizarry MC: Early Abeta accumulation and progressive synaptic loss, gliosis, and tangle formation in AD brain. Neurology 2004, 62:925-931.

- [6] Jack CR Jr., Knopman DS, Jagust WJ, Shaw LM, Aisen PS, Weiner MW, Petersen RC, Trojanowski JQ: Hypothetical model of dynamic biomarkers of the Alzheimer's pathological cascade. Lancet Neurol 2010, 9:119-128.
- [7] Savva GM, Wharton SB, Ince PG, Forster G, Matthews FE, Brayne C; Medical Research Council Cognitive Function and Ageing Study: Age, neuropathology, and dementia. N Engl J Med 2009, 360:2302-2309.
- [8] Fox NC, Scahill RI, Crum WR, Rossor MN: Correlation between rates of brain atrophy and cognitive decline in AD. Neurology 1999, 52:1687-1689.
- [9] Frisoni GB, Fox NC, Jack CR Jr., Scheltens P, Thompson PM: The clinical use of structural MRI in Alzheimer disease. Nat Rev Neurol 2010, 6:67-77.
- [10] Gómez-Isla T, Hollister R, West H, Mui S, Growdon JH, Petersen RC, Parisi JE, Hyman BT: Neuronal loss correlates with but exceeds neurofi brillary tangles in Alzheimer's disease. Ann Neurol 1997, 41:17-24.
- [11] Gosche KM, Mortimer JA, Smith CD, Markesbery WR, Snowdon DA: Hippocampal volume as an index of Alzheimer neuropathology: findings from the Nun Study. Neurology 2002, 58:1476-1482.
- [12] Jack CR Jr., Dickson DW, Parisi JE, Xu YC, Cha RH, O'Brien PC, Edland SD, Smith GE, Boeve BF, Tangalos EG, Kokmen E, Petersen RC: Antemortem MRI findings correlate with hippocampal neuropathology in typical aging and dementia. Neurology 2002, 58:750-757.
- [13] Silbert LC, Quinn JF, Moore MM, Corbridge E, Ball MJ, Murdoch G, Sexton G, Kaye JA: Changes in premorbid brain volume predict Alzheimer's disease pathology. Neurology 2003, 61:487-492.

# **Author Profile**



Ms. Inaam Eltoom AbdAlla Mohamed (Sudan) received the B.Sc. and MSc degree in diagnostic radiology technology, SUST 2000, and 2004 respectively, she was working as radiology technologist, in radiology department in Zaied military

hospital (EAU), and she has been active in MRI and diagnostic radiology researches Inc.



Mr. Shazaly Nader Khojaly Mansour (Sudan) received (B.Sc.) in diagnostic radiology technology and (M.Sc.) in medical ultrasound imaging from College of Medical radiological Science, Sudan University of Science and Technology in 2011 and 2015 respectively. Now as lecturer at AlGhad international college for medical sciences, Medical imaging technology department, KSA, Abha from 2016 up to now.



branch.

Dr. Mohammed Abdelwahab Idris Salih, PhD in diagnostic radiology(SUST).Radiology manager at Royal care international hospital -Khartoum/Sudan, local operator for Toshiba company CT scan), Sudan

Volume 6 Issue 10, October 2017 www.ijsr.net

## Licensed Under Creative Commons Attribution CC BY DOI: 10.21275/23081715