REVIEW ARTICLE

Correlation Between the Dopamine and ROS Levels in Depression Disorders

Zuhair I. Al Mashhadani¹, Muneam H. Ali¹, Mona N. Al-Terehi^{2*}

¹Al-Nisour University College, Baghdad, Iraq ²College of Science, University of Babylon, Hillah, Iraq

Received: 15th August, 2021; Revised: 03rd September, 2021; Accepted: 02nd November, 2021; Available Online: 25th December, 2021

ABSTRACT

Depression is one of the mode disorder diseases which recorded in high percentage in last years in the Iraqi population. The current study suggested to determine the correlation between dopamine level with reactive oxygen species (ROS) in depressed patients, results exhibit that ROS was higher in depressed patients than control in significant differences (p 0.000), dopamine decreased in patients in comparing with control in non-significant differences (p 0.273). The correlations among study variables in patient group were non-significant weak positive correlation between age and dopamine (r 0.033, p 0.894), non-significant inverse relation between BMI and dopamine (r -0.297, p 0.217), non-significant inverse weak correlation observed between Dopamine and ROS (r-0.069, p 0.780). The correlation between ROS and age was an inverse correlation (r -0.060, p 0.808) and with BMI weak positive correlation (r 0.190, P 0.790) in non-significant differences in both variables. In the control group the correlation between age and dopamine non-significant weak positive relation was observed (r 0.098, p 0.691). Non-significant inverse relation between BMI and dopamine (r -0.062, P 0.801), non-significant inverse weak correlation observed between Dopamine and ROS (r-0.064, p 0.794). The correlation between ROS and age was an inverse correlation (r -0.018, p 0.940) and with BMI inverse correlation (r -0.417, p 0.075) in non-significant differences in both variables. The current results concluded that the dopamine was slightly affected by ROS, age and BMI in depression patients.

Keywords: Dopamine, Depression disorder, Correlation, ROS.

International Journal of Pharmaceutical Quality Assurance (2021); DOI: 10.25258/ijpga.12.4.18

How to cite this article: Al Mashhadani ZI, Ali MH, Al-Terehi MN. Correlation Between the Dopamine and ROS Levels in

Depression Disorders. International Journal of Pharmaceutical Quality Assurance. 2021;12(4):328-330.

Source of support: Nil. **Conflict of interest:** None

INTRODUCTION

Oxidative stress is a complex phenomenon characterized by elevation in free radicals productions or decreased in degeneration, ROS or reactive oxygen species is one of the important types of free radicals groups consist of superoxide anion (O_2^-) , hydroxyl anion (O_1^-) and hydrogen peroxide (H_2O_2) . 1-3

Dopamine is one of the neurotransmitter has a major role in the brain regulation, behavior modulation, voluntary movement, motivation, prolactin production inhibition, sleeping, and other functions,⁴ and it precursor of other related catecholamines such as norepinephrine and epinephrine.⁵ The association between dopamine and oxidative stress have been described.^{6,7} It is oxidize to form quinones and H₂O₂ which react with iron to form 'OH,⁸ thus dopamine contributed in elevation oxidative stress when antioxidant system decline, and this involvement if some disease like Parkinson's disease and depression disorders.^{9,10}

The association of ROS with dopamine levels in depression disorder was suggested to study at present work in Iraqi patients as a pilot study.

MATERIALS AND METHODS

A pilot study included 19 depressed patients and 19 healthy individuals as control group were enrolled in the present study, ethical approval of the environment and health ministry of Iraq was depended in sample and data collection. The ROS detected by colorimetric method, dopamine detection by ELIZA technique. Data represented as mean \pm SE, independent T test and correlation coefficients used to significantly analyze at p < 0.05.

RESULTS AND DISCUSSION

The current results show that non-significant differences in age (p = 0.950) and BMI (p = 0.144) between groups, ROS was higher in depressed patients than control in significant differences (p > 0.000), dopamine decreased in patients in comparing with control in non-significant differences (p < 0.273) (Table 1). The association of dopamine in the pathophysiology of depression have been studied. 11,12 Numerous studies focusing on the role of the dopamine system in depression converge on a down regulation of this system. 13

Table 1: mean differences of study variables for both groups

	•		-
Variables	Depression Patients	Control	Sig
Age	38.10 ± 2.92	38.36 ± 2.94	0.950
BMI	24.98 ± 0.910	27.02 ± 1.01	0.144
ROS	105.20 ± 12.162	25.53 ± 1.93	0.00
Dopamine pg/mL	13.35 ± 0.915	21.91 ± 7.05	0.273

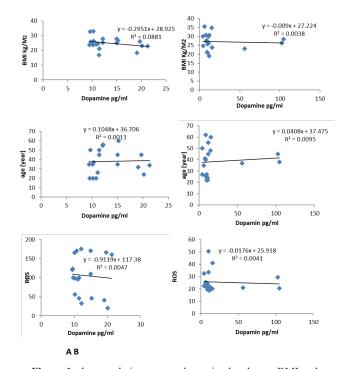


Figure 1: the correlation among dopamine level, age, BMI and ROS, A patient, B control

The correlation between dopamine and study variables were detected, in patient group non-significant weak positive correlation between age and dopamine (r 0.033, p > 0.894), non-significant inverse relation between BMI and dopamine (r -0.297, p > 0.217), non-significant inverse weak correlation observed between dopamine and ROS (r-0.069, p > 0.780) (Figure 1A). Also, the correlation between ROS and age was inverse correlation (r -0.060, p > 0.808) and with BMI weak positive correlation (r 0.190, p > 0.790) in non-significant differences for both variables (Figure 2 A).

In the control group the correlation between age and dopamine non-significant weak positive relation was observed (r 0.098, p 0.691). Non-significant inverse relation between BMI and dopamine (r -0.062, p >0.801). Non-significant inverse weak correlation observed between Dopamine and ROS (r-0.064, p >0.794) (Figure 1, B). However, the correlation between ROS and age was an inverse correlation (r -0.018, p 0.940) and with BMI inverse correlation (r -0.417, p 0.075) in non-significant differences for both variables (Figure 2B).

Studies found that the level of dopamine in the human striatum declines up to 50% with age.¹⁴ the elevated level of dopamine was documented in young than older individuals.^{15,16} Others found that the effect age in dopamine was occurred

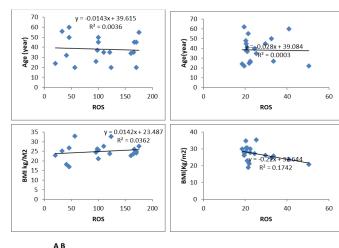


Figure 2: The correlation between ROS, Age and BMI in study groups, A Patients, B control group

by D2 dopamine receptor binding sites which mainly post-synaptically located was decrease with normal aging consequence with impaired of the presynaptic nigrostriatal dopaminergic neuronal system.¹⁷

The cellular and molecular mechanisms linking oxidative stress with dopaminergic neuron death are complex and not well characterized.¹³ Decreased dopamine receptor function increases ROS activity and vice versa working in concert or independently of each other. 18 Different factor contributed in the relation of ROS with age or body mass index (BMI). The lifestyle is an important factor to enhance ROS regulation in the body which may be controlled in the harmful effect of ROS. 19 Nevertheless, the avoiding of chronic ROS exposure contributed in the lowering of oxidative stress effected, the association ROS with age observed by the aging of cell which loosed ability to repair damage and impaired antioxidant molecules to scavenge ROS. ²⁰⁻²² The current study conclude that dopamine was slightly affected by age, BMI and ROS in depressed patients, ROS was also slightly affected by age and BMI in the patients group, for further investigations the present results need a wide spectrum of sample number of different Iraqi populations.

REFERENCES

- 1. Paravicini TM, Touyz RM. Redox signaling in hypertension. Cardiovasc Res. 2006;71:247-258.
- Wilcox CS. Oxidative stress and nitric oxide deficiency in the kidney: A critical link to hypertension? Am J Physiol Regul Integr Comp Physiol. 2005;289:R913

 –R935.
- Dikalov S, Griendling KK, Harrison DG. Measurement of reactive oxygen species in cardiovascular studies. Hypertension. 2007;49:717–727. [PMC free article]
- Calabresi P, Picconi B, Tozzi A, Di Filippo M, "Dopaminemediated regulation of corticostriatal synaptic plasticity," Trends in Neurosciences, 2007;30(5):211-219
- An JH, Oh BK, Choi JW. Detection of tyrosine hydroxylase in dopaminergic neuron cell using gold nanoparticles-based barcode DNA. Journal of biomedical nanotechnology. 2013 Apr 1:9(4):639-643.

- Juarez Olguin H, Calderon Guzman D, Hernandez Garcia E, Barragan Mejia G. The role of dopamine and its dysfunction as a consequence of oxidative stress. Oxidative medicine and cellular longevity. 2016 Oct;2016. https://doi.org/10.1155/2016/9730467.
- Grima G, Benz B, Parpura V, Cuénod M, Do KQ. Dopamineinduced oxidative stress in neurons with glutathione deficit: implication for schizophrenia. Schizophrenia research. 2003 Aug 1;62(3):213-224.
- Dusek P, Schneider SA, Aaseth J. Iron chelation in the treatment of neurodegenerative diseases. Journal of Trace Elements in Medicine and Biology. 2016 Dec 1;38:81-92.
- Weng M, Xie X, Liu C, Lim KL, Zhang CW, Li L. The sources of reactive oxygen species and its possible role in the pathogenesis of Parkinson's disease. Parkinson's Disease. 2018 Sep 2;2018.
- Dhir A, Kulkarni SK. Involvement of nitric oxide (NO) signaling pathway in the antidepressant action of bupropion, a dopamine reuptake inhibitor. European journal of pharmacology. 2007 Jul 30;568(1-3):177-185.
- Brown AS, Gershon S. Dopamine and depression. J Neural Transm Gen Sect. 1993;91(2-3):75-109. doi: 10.1007/BF01245227. PMID: 8099801.
- Belujon P, Grace AA. Dopamine system dysregulation in major depressive disorders. International Journal of Neuropsychopharmacology. 2017 Dec;20(12):1036-1046.
- Guo JD, Zhao X, Li Y, Li GR, Liu XL. Damage to dopaminergic neurons by oxidative stress in Parkinson's disease. International journal of molecular medicine. 2018 Apr 1;41(4):1817-25.
- Morgan DG. The dopamine and serotonin systems during aging in human and rodent brain. A brief review. Progress in Neuro-psychopharmacology and Biological Psychiatry. 1987 Jan 1;11(2-3):153-157.

- Knutson B, Fong GW, Adams CM, Varner JL, Hommer D (2001)
 Dissociation of reward anticipation and outcome with event-related fMRI. Neuroreport 12:3683-3687.
- Knutson B, Adams CM, Fong GW, Hommer D. Anticipation of increasing monetary reward selectively recruits nucleus accumbens. Journal of Neuroscience. 2001 Aug 15;21(16):RC159.
- Antonini A, Leenders KL, Reist H, Thomann R, Beer HF, Locher J. Effect of age on D2 dopamine receptors in normal human brain measured by positron emission tomography and 11C-raclopride. Archives of Neurology. 1993 May 1;50(5):474-480.
- Zeng C, Villar VA, Yu P, Zhou L, Jose, PA. Reactive oxygen species and dopamine receptor function in essential hypertension. Clinical and experimental hypertension (New York, N.Y.: 1993), 2009;31(2):156-178. https://doi.org/10.1080/10641960802621283.
- Fukui T, Yamauchi K, Maruyama M, Yasuda T, Kohno M, Abe Y. Significance of measuring oxidative stress in lifestyle-related diseases from the viewpoint of correlation between d-ROMs and BAP in Japanese subjects. Hypertens Res. 2011 Sep;34(9):1041-1045. doi: 10.1038/hr.2011.76. Epub 2011 Jun 16. PMID: 21677660.
- Lam YT, "Critical Roles of Reactive Oxygen Species in Age-Related Impairment in Ischemia-Induced Neovascularization by Regulating Stem and Progenitor Cell Function", Oxidative Medicine and Cellular Longevity, 2016;;14. https://doi. org/10.1155/2016/7095901.
- Kadhum AM, Abd Al-Kadim BM, Kareem TC, Nayyef AA, Al-Terehi MN. Associations of Tumor Necrosis Factor-[alpha] G-308A (rs1800629) Gene Polymorphisms with Major Depressive Disorder. NeuroQuantology. 2021 Jul 1;19(7):133-137.
- 22. Kadhum AM, Abd ES, Taleb RA, Hussain KN, Al-Terehi MN. Genetic Polymorphisms of Mitochondrial Genes (ND1, ATP6c) in Depression Patients. NeuroQuantology. 2021 Jul 1;19(7):128-133.