

Neurocognitive Impairments in Major Depressive Disorder: A Clinical and Cognitive Perspective.

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Introduction

Major Depressive Disorder (MDD) is not only characterized by persistent low mood and emotional disturbances but also by notable neurocognitive impairments that significantly affect patients' daily functioning and quality of life. These impairments often persist even after the remission of mood symptoms, suggesting a distinct cognitive dimension within the disorder [1, 2, 3, 4, 5].

Clinically, patients with MDD frequently exhibit deficits in attention, memory, executive function, and processing speed. Such impairments can hinder occupational performance, interpersonal relationships, and treatment adherence. From a cognitive neuroscience perspective, these dysfunctions are linked to abnormalities in brain regions such as the prefrontal cortex, hippocampus, and anterior cingulate cortex—areas responsible for higher-order cognitive processing [6, 7, 8].

Neuroimaging studies reveal structural and functional changes in these regions, including reduced grey matter volume and hypoactivation during cognitive tasks. Furthermore, dysregulation in neurotransmitters such as serotonin, dopamine, and glutamate has been implicated in both mood dysregulation and cognitive deficits.

Early detection and targeted interventions, including cognitive remediation therapy, pharmacological strategies, and neurostimulation techniques, show promise in improving cognitive outcomes. Recognizing and addressing these cognitive impairments is essential for comprehensive treatment planning in MDD, promoting better recovery and long-term functionality [9, 10].

Conclusion

Neurocognitive impairments are a core and often overlooked feature of Major Depressive Disorder, extending beyond emotional symptoms and contributing to long-term functional disability. Understanding these deficits from both clinical and cognitive perspectives is crucial for early identification and effective management. Integrating cognitive assessments and targeted interventions into routine care can enhance treatment

outcomes, support recovery, and improve the overall quality of life for individuals affected by MDD.

References

1. Chehregosha H, Khamseh ME, Malek M, et al. A view beyond HbA1c: role of continuous glucose monitoring. *Diabetes Therapy*. 2019;10:853-63.
2. Sacks DB. Hemoglobin A1c in diabetes: panacea or pointless?. *Diabetes*. 2013;62(1):41-3.
3. Little RR, Rohlfing CL. The long and winding road to optimal HbA1c measurement. *Clinica chimica acta*. 2013;418:63-71.
4. Schnell O, Crocker JB, Weng J. Impact of HbA1c testing at point of care on diabetes management. *J Sci Technol*. 2017;11(3):611-7.
5. Gore MO, McGuire DK. A test in context: hemoglobin A1c and cardiovascular disease. *J Am Coll Cardiol*. 2016;68(22):2479-86.
6. OJun A, Beydoun G, Win KT, et al. Cultivating Expertise: Unravelling Type 2 Diabetes Associations through Incremental Knowledge-Based System Development: Ripple Down Rules or Machine Learning.
7. Gill AY, Saeed A, Rasool S, et al. Revolutionizing Healthcare: How Machine Learning is Transforming Patient Diagnoses-a Comprehensive Review of AI's Impact on Medical Diagnosis. *Sci. World J.* 2023;2(10):1638-52.
8. Baronov D. The African transformation of western medicine and the dynamics of global cultural exchange. Temple University Press; 2010.
9. Lin EC, Chiang YC, Lin HY, et al. Unraveling the Link between Periodontitis and Coronavirus Disease 2019: Exploring Pathogenic Pathways and Clinical Implications. *Biomedicine*. 2023;11(10):2789.
10. Patil N, Howe O, Cahill P, et al. Monitoring and modelling the dynamics of the cellular glycolysis pathway: A review and future perspectives. *Mol. Metab*. 2022;101635.

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