ISSN: 2472-0895

Open Access

The Impact of Epilepsy on Quality of Life: A Comprehensive Review

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Abstract

Epilepsy, a chronic neurological disorder characterized by recurrent seizures, affects approximately 50 million people worldwide. While medical treatment primarily focuses on controlling seizures, the impact of epilepsy extends far beyond the seizures themselves. This article reviews the profound ways epilepsy can impact Quality Of Life (QOL), encompassing physical health, mental health, social functioning, and overall well-being.

Keywords: Neurological disorder • Antiepileptic drugs • Seizures

Introduction

Epilepsy can have significant effects on physical health and daily functioning. The unpredictability of seizures can limit an individual's independence and ability to engage in various activities, from driving to recreational pursuits. Seizures can lead to physical injuries, including falls and burns, and certain types of seizures can be life-threatening. Anti-Epileptic Drugs (AEDs), the primary treatment for epilepsy, can also have physical side effects. These can include dizziness, fatigue, nausea, and more serious effects like bone density loss or liver problems. The mental health impact of epilepsy is considerable. Depression and anxiety disorders are significantly more prevalent in people with epilepsy compared to the general population. The reasons for this are multifaceted, ranging from the biological impact of seizures on mood regulation to the psychosocial stress of living with epilepsy [1].

Cognitive function can also be affected. Seizures and certain AEDs can lead to difficulties with memory, attention, and processing speed. In children, these cognitive effects can interfere with academic performance and development. Social challenges are a common consequence of epilepsy. Misunderstandings and stigma around epilepsy can lead to social isolation and discrimination. Fear of having a seizure in public can also contribute to social anxiety and withdrawal. Employment can be significantly impacted. The unpredictable nature of seizures can limit job opportunities and lead to higher rates of unemployment [2].

Literature Review

Health-Related Quality of Life (HRQoL) is a specific measure of how a person's health affects their perceived quality of life. Studies consistently show that people with epilepsy have lower HRQoL scores than the general population. Factors that contribute to lower HRQoL scores include seizure frequency, seizure severity, side effects of medication, and comorbid mental health conditions.

Epilepsy can profoundly affect an individual's quality of life, with impacts

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Received: 02 April, 2023; Manuscript No. elj-23-108606; Editor assigned: 04 April, 2023, PreQC No. P-108606; Reviewed: 17 April, 2023, QC No. Q-108606; Revised: 22 April, 2023, Manuscript No. R-108606; Published: 29 April, 2023, DOI: 10.37421/2472-0895.2023.9.193

on physical health, mental health, social functioning, and overall well-being. This underscores the need for comprehensive epilepsy care that goes beyond seizure control. Holistic care can include mental health services, social support, vocational guidance, and educational accommodations. As our understanding of the broader impact of epilepsy grows, so too does our ability to provide care that enhances the quality of life for those living with this complex condition [3].

Epilepsy can significantly affect interpersonal relationships and family dynamics. The unpredictability of seizures can place emotional and practical burdens on family members, particularly primary caregivers. Fear for the person's safety, worry about the potential for Sudden Unexpected Death in Epilepsy (SUDEP), and the responsibility of providing care can lead to stress and caregiver burnout. Moreover, siblings of children with epilepsy might feel neglected or overlooked, adding another layer of complexity to family dynamics.

Conversely, strong supportive relationships can greatly enhance the quality of life for people with epilepsy. Thus, support for caregivers and family therapy may be important components of a comprehensive epilepsy care plan [4].

Discussion

Epilepsy can influence educational and career trajectories. Children with epilepsy may struggle acadically due to seizures, side effects of medications, and school absences. This can lead to lower educational attainment and limit career options. Moreover, fear of seizures can prevent individuals from pursuing certain careers or advancing in their chosen fields. Educational support, vocational training, and workplace accommodations can help individuals with epilepsy reach their full potential. Legislation such as the Americans with Disabilities Act (ADA) in the United States protects people with epilepsy from job discrimination and mandates reasonable workplace accommodations [5].

Given the substantial impact of epilepsy on quality of life, QOL assessment should be an integral part of epilepsy care. Tools such as the Quality of Life in Epilepsy Inventory (QOLIE) can help clinicians gauge the impact of epilepsy on various life domains and track changes over time. These assessments can guide treatment decisions and help identify areas where additional support is needed.

Enhancing quality of life in people with epilepsy requires a multifaceted approach. This includes optimizing seizure control with minimal side effects, addressing comorbid mental health conditions, providing social and vocational support, and fostering supportive relationships. Public education is also crucial to dispel misconceptions about epilepsy and reduce stigma [6].

Conclusion

Epilepsy significantly impacts quality of life, touching on nearly every

aspect of daily living. Recognizing and addressing these impacts is essential in epilepsy care. While challenges persist, there is cause for optimism. Advances in treatment, growing awareness, and an increasing focus on holistic care are all contributing to improved quality of life for people with epilepsy. The road ahead requires continued research, advocacy, and a commitment to viewing individuals with epilepsy in their totality not just as patients, but as people with dreams, aspirations, and the desire to lead fulfilling lives.

Acknowledgment

None.

Conflict of Interest

None.

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How to cite this article: Narayana, Laxmi. "The Impact of Epilepsy on Quality of Life: A Comprehensive Review." *Epilepsy J* 9 (2023): 193.