EPV0586

Psychological Stressor Of End Stage Chronic Kidney Disease Patients On Dialysis. A Battle For Life

S. S. Hashmi¹ and A. Syed²*

¹Department Of Applied Psychology, National University of Modern Languages, Karachi, Pakistan and ²First Faculty of Medicine, Charles University, Prague Czechia, Prague, Czech Republic *Corresponding author. doi: 10.1192/j.eurpsy.2024.1258

Introduction: Patients with chronic kidney disease (CKD) may choose to undergo dialysis. Factors that may have led patients to prescribe psychological interventions related to dialysis are poorly understood in the literature. The purpose of this study was to explore multi-level factors surrounding dialysis modalities such as Diagnosed Mood Disorders, Existential crises, Triggering events, Social support, and Distrust towards the process of dialysis.

Objectives: The study aims to investigate the psychological battle of the client while going through the process of dialysis. The study reveals multiple mood disorders and existential crises leading to depression among chronic kidney disease patients. Therefore the study was conducted with the aim of providing a therapeutic guide line in future once the factors are investigated in detail.

Methods: Semi-structured qualitative interviews were conducted in a dialysis clinic in Karachi where 19 participants participated in this qualitative study. The age ranges from 40-76. Initiating with informed consent followed by surveys assessing demographic and clinical information were administered to participants following their interviews.

Results: Qualitative findings suggested that patients were dealing with Clinical Mood Disorders without being provided treatment. Moreover, the cohesive family support enabled them to continue with daily living activities; however, the patients with low support triggering adverse events in life lost their lives in follow-up sessions. Furthermore, nephrology care doesn't seem sufficient as they are dealing with existential crises of hopelessness, regret, condemnation, and elevated death anxiety. In CKD the misinterpretation of dialysis by cognitively substituting it as End of life increased the clinical symptoms of Mood disorders. Thus the risk factors increase disturbing the quality of life.

Conclusions: Findings point to broader factors affecting dialysis modalities with Mood disorders. The low social support and adverse triggering events precipitate the risk factors of dialysis treatment. Furthermore, distrust towards dialysis and existential crisis are recommended for therapeutic interventions

Disclosure of Interest: None Declared

EPV0587

Perceived stigma and associated factors among family caregivers of patients with severe mental disorders

R. Masmoudi, W. Abid^{*}, E. Ben salem, R. Ouali, I. Feki, I. Baati and J. Masmoudi

psychiatry A department, Hedi chaker university hospital, Sfax, Tunisia *Corresponding author. doi: 10.1192/j.eurpsy.2024.1259 **Introduction:** Giving care to someone who is experiencing mental distress is a difficult and challenging task that could be detrimental to the caregiver's overall quality of life. Stigma associated with mental illness is one of the most important mental health issues faced by these caregivers.

Objectives: Our aims were to assess perceived stigma among family caregivers of patients with severe mental disorders and to identify its associated factors.

Methods: We conducted a descriptive and analytical crosssectional study among family caregivers of patients followed at the psychiatry outpatient clinic of the Hedi Chaker University Hospital in Sfax, during the period from February 2022 to July 2022.

A structured interview questionnaire was designed to collect sociodemographic data of both patients and their caregivers. We used the Stigma Devaluation Scale (SDS) to assess stigma.

Results: A total of 90 family caregivers of severely mentally ill patients were included: 26 men and 64 women, with an average age of 50.68 ± 11.67 years.

Patients' parents accounted for 40% of family caregivers. The majority of family caregivers (83.3%) had no more than secondary education. Married people represented 70% of cases.

The median age of patients was 42 years. Schizophrenia was the diagnosis in 68.9% of cases. The mean duration of illness was 16.23 years.

Daily assistance lasted from 4 to 8 hours in 30% of cases and more than 8 hours in 66.7% of cases.

The mean score (SDS12) for family-focused stigma was 13.12 ± 2.34 with ranges from 8 to 18.

Perceived stigma scores were significantly higher among caregivers caring for non-married patients (p=0.04), with an age <50 years (p=0.04), and with a higher level of education (p=0.02).

Long duration of providing care (> 8 hours per day) (p=0.05) and insufficient information about the illness (p=0.02) were significantly associated with perceived stigma.

Conclusions: The clinicians managing patients with severe mental disorders must focus on stigma and psychological distress among the caregivers and plan intervention strategies to reduce stigma.

Disclosure of Interest: None Declared

EPV0588

Resilience and burden in family caregivers of patients with severe mental disorders

R. Masmoudi, W. Abid*, E. Ben Salem, R. Ouali, I. Feki, I. Baati and J. Masmoudi

psychiatry A department, hedi chaker university hospital, sfax, Tunisia *Corresponding author.

doi: 10.1192/j.eurpsy.2024.1260

Introduction: Family caregivers of patients suffering from severe psychiatric disorders may present with health problems, lower quality of life, and painful emotions, which can seriously compromise their well-being when they do not receive appropriate professional support.

Objectives: The aims of this study were to assess the level of burden and resilience in family caregivers of patients with severe mental disorders and to determine associated factors.

Methods: We conducted a descriptive and analytical cross-sectional study among family caregivers of patients followed at the psychiatry outpatient clinic of the Hedi Chaker University Hospital in Sfax, during the period from February 2022 to July 2022.

We used the Connor–Davidson Resilience Scale (CD-RISC) to assess resilience and the Zarit Burden Inventory to assess the level of burden. Higher scores indicate higher resilience and greater burden. **Results:** The sample included 90 family caregivers of patients with severe mental disorders. The average age was 50.68. They were the parents of patients in 40% of cases. Professionally active caregivers accounted for 57.8% of cases. Thirty family caregivers had a somatic disorder history (33.3%).

The median age of patients was 42 years. Ten patients (11.1%) were financially independent. The diagnosis was schizophrenia in 68.9% of cases. The mean duration of illness was 16.23 years. Irregular follow-up was noted in 10 patients (11.1%).

The mean scores of the Zarit Burden scale and the CD-RISC were 41.86 ± 10.33 and 58.46 ± 9.18 respectively.

Unemployed caregivers and parents experienced a higher burden (p=0.001, p=0.03 respectively). The level of burden was higher in caregivers taking care of financially dependent patients (p=0.03), with a duration of the disease greater than 15 years (p=0.04), and with irregular follow-up (p=0.008).

A low level of resilience in caregivers was correlated with spousal relationship (p=0.001), cohabitation with the patient (p=0.05), widowhood (p=0.01), low level of education (p=0.02), the presence of a somatic disorder history in the caregivers (p=0.04).

A negative correlation was observed between CD-RISC and Zarit scores (p=0.04; r=-0.21).

Conclusions: Family caregivers of mentally ill patients experienced a significant level of caregiver burden, and it was lower in caregivers with higher levels of resilience. Psycho-educational programs directed toward family caregivers are highly recommended.

Disclosure of Interest: None Declared

EPV0589

Exploring Child-Parent Relationship Therapy -CPRT-Impact on Externalised Behaviours of Foster Children Experienced Complex Trauma: A Case Study

S. Hatam¹*, S. Moss², C. Cubillo³, S. Parsafar¹ and D. Berry⁴

¹Faculty of Health; ²Charles Darwin University; ³healing circle psychology, Darwin and ⁴interplay play therapy australia, Perth, Australia

*Corresponding author. doi: 10.1192/j.eurpsy.2024.1261

Introduction: Many foster children experience traumatic events that result in a wide range of disruptive behaviours, such as temper tantrums, superficially charming, no sincere remorse, and so forth. These problematic behaviours are challenging to the implementa-

tion of holistic therapeutic interventions. **Objectives:** The purpose of the current study is to explore the

effectiveness of employing Child-Parent Relationship Therapy (CPRT) on externalised behaviours of a traumatised child at home and in social interactions.

Methods: This study used a case study to explore the influence of CPRT on externalised behaviours of traumatised foster children.

The case study focused on the externalised problematic behaviours of an Australian Aboriginal child fostered under long-term care at 18 months by a Caucasian family. The foster parent and the foster child received a 10-session structured CPRT across ten weeks. The child's externalised behaviours were evaluated through the Child Behaviour Checklist (CBCL) form.

Results: The findings describe the process of a 10-session structured CPRT with a foster child who exhibits externalised disruptive behaviours because of abandonment and complex trauma. The themes in the foster parent's role and the play therapy approach relate to traumatic events and attachment issues. The results indicated a slight reduction in these behaviours. According to the follow-up interview, the child still showed aggressive behaviours in social interactions but not at home. Interviews with the foster parent indicated both the parent and child require additional support and further sessions of CPRT.

Conclusions: This case study identified an improvement in externalised behaviours for foster children with experience of complex trauma and abandonment after a 10-session CPRT. Further research is required to explore the effectiveness of a longer-term session of CPRT alongside additional support services for foster parents.

Disclosure of Interest: None Declared

EPV0590

Improving Mental Health of Medical Students through Movement, Art, and Interpersonal Relations

E. Ajándék¹ and V. Boros²*

¹Division of Integrative Medicine, Semmelweis University and ²Collection Management, Museum of Ethnography, Budapest, Hungary

*Corresponding author.

doi: 10.1192/j.eurpsy.2024.1262

Introduction: Burnout, and symptoms of distress amongst medical students is becoming increasingly common due to the uncertainty of the Hungarian healthcare system. Change itself may be the cause of stress. Since the pandemic, the workload has been growing among health care workers. Anxiety is increasing even for beginner practitioners.

Objectives: Our examination intends to improve the mental health of the students with a variety of methods to help them develop resilience towards everyday stress, such as:

- 1. increasing body awareness
- 2. exploring the inner drivers of vocations by self-esteem, worth symbols and emblems
- 3. supporting relationships and interpersonality

Methods: We had advertised a monthly course in the mailing system of the students of the Semmelweis University (Neptun). Each occasion would go as far as 240 minutes in length. Selection criteria were: guaranteeing participation in the sessions. Any applicant suffering from mental health problems requiring medical attention, or the applicant regularly skipping occasions of the session would lead to his or her getting dismissed. Courses consists of musical aerob movement and receptive art therapy tools (exl. "Self-exhibition"). The aim was to interpret ones identity via images individually. In