

Ramírez Hinojosa Ma *

Research Article

Comparison of Family Functioning Between Women with and Without Motor Disabilities

Ramírez Hinojosa Ma

Lucina Family Therapy Institute CENCALLI Mexico City

*Corresponding Author: Delaram Dehnashi, Ramírez Hinojosa Ma, Lucina Family Therapy Institute CENCALLI Mexico City.

Received date: December 22, 2023; Accepted date: March 04, 2024; Published date: March 14, 2024

Citation: Ramírez Hinojosa Ma, (2024), Comparison of Family Functioning Between Women with and Without Motor Disabilities, *Psychology and Mental Health Care*, 8(3): DOI:10.31579/2637-8892/248

Copyright: © 2024, Ramírez Hinojosa Ma. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Multiple Sclerosis, an autoimmune disease, can significantly impact a person's daily functioning and overall well-being. Patients often experience emotional disorders and find it challenging to regulate their cognitive-emotional responses during the course of their illness. The objective of this current study was to evaluate the impact of MBSR training on quality of life of patients with Multiple Sclerosis (MS). The study consisted of 40 participants with multiple sclerosis, all of whom were enrolled in the MS Society of Shiraz in Fars Province, Iran. The participants were evenly divided into two groups: an experimental group of 20 individuals and a control group of 20 individuals. The experimental group underwent a series of eight MBSR training sessions, held once a week for 60 minutes over a period of 60 days. The control group did not receive any interventions. The findings of the present study demonstrate a notable disparity between the experimental and control groups in terms of quality of life, specifically in areas such as physical function, pain management, mental well-being, energy levels, health perception, and cognitive function in relation to health changes (P<0.05). However, while these differences were statistically significant, no significant differences were noted among the dimensions of role limitation caused by physical problems, mental challenges, and social functioning (P>0.05). The results of the present study showed that MBSR training improves some dimensions of the quality of life in patients with Multiple Sclerosis and reduced negative emotion regulation strategies. **Keywords:** mbsr; emotion; quality of life; multiple sclerosis

Introduction

For the World Health Organization (WHO), the family is understood as a group of people who make up the household and have some degree of kinship by blood, adoption or marriage^{1.}

The Structural Systemic School defines the family as a natural group that over time has developed patterns of interaction; these constitute the family structure, which in turn governs the functioning of its members, defines their range of behaviors and facilitates their interaction with each other²; It is also known as a powerful primary system, to which a person belongs; it is a living system, in constant transformation that is modified over time in order to ensure continuity and psychosocial growth of its members and also involves multiple factors that modify its conformation, structure and family dynamics^{3.}

The family structure functions as an interactive and interdependent unit; and when an event or experience occurs that affects one member of the group in particular, it affects the rest of the members, i.e. all members are affected and involved in the disability process, this impact may be at different levels and dimensions of their daily life⁴.

At this point, a reorganization within the family is necessary, which on the one hand leads to the individuation of each of its members and on the other hand provides a feeling of belonging, security and warmth³. For this reason, the family plays an important role in the process of assimilation and acceptance of disability and its different challenges. Parents tend to react in different ways to the disability of one of their members, which may depend on their degree of severity, vulnerability, their strengths, the stage of the life cycle they are going through, the medical, economic, social, educational aspects and the social support network they have ^{5,6.}

It has been documented that illness, disability and death are universal experiences that place families before one of life's greatest challenges and involve different reactions in each member of the family Interacting with a member with a disability involves adapting to its different phases and demands ^{5,6.}

The family impact of disability is reflected in the appearance of various reactions and affects such as: feelings of helplessness, aggressiveness,

guilt, anger, ambivalence and fear of the future, and even emotional discomfort (anxiety, depression) physical discomfort (illness) or even abandonment, neglect and violence for the person with disability, which are characteristics of unresolved grief $^{7,8.}$

In this way, the presence of a member with a disability generates the need to modify family roles in order to adapt, which will become more complex as they demand high levels of availability and tolerance for the care and attention of the person, so that flexibility and adjustment to new demands will allow the construction of opportunities and alternatives for the entire close family nucleus ^{6,7}.

As mentioned above, the presence of a child with a disability becomes a stressful event within the family and can trigger the breakdown of marital and family relationships. Parents often become directly and exclusively responsible for the education and rehabilitation of their child with a disability. In a study of parents of children with Down's syndrome, it was observed that they use the same parenting skills as parents of children without disabilities, but the differences lie in the high degree of parental stress due to the cognitive disability, which generates strong emotional stress^{4,8}.

exclusion are more evident in women with disabilities in any of their manifestations, as they are the group who live longer with the family of origin, have higher illiteracy rates, have fewer years of schooling, are less employed in companies, and have fewer opportunities for development within the company⁸.

These differences corroborate what several studies have pointed out: women with disabilities experience double discrimination they are often excluded from different family and social processes because they are women, and not considered competent because they have a disability or physical limitation ^{9,10,11}.

The present study aimed to investigate the impact of motor disability on the family functioning of women with disabilities and to compare them with women without disabilities.

Family functioning instrument

The family functions scale that was applied, is made up of items with factor loadings greater than or equal to .40, was standardized in the Mexican population, is made up of 22 items, divided into 4 factors, and its response style is on a Likert scale¹².

Various studies point out that the processes of overprotection, care and

Factor	Positive family environment	Cohesion and rules	Hostility and conflict avoidance	Command Problems in the expression of feelings
Items	(7)	(5)	(5)	(5)
	12,14,15,17,18,21,22	1,6,10,9,19	2,3,5,16,20	4,7, 8,11 ,13

Note. Overall Cronbach's Alpha of .8533

Table 1: Family functions scale factors

Procedure, materials and subjects

A standardized family functions scale was administered in a Mexican population to adult women with and without motor disabilities, all of them of legal age, who agreed to participate in the study.

A sample of 203 women, who agreed to participate in the study on a voluntary basis, was formed, the participants were informed of the objective of the study and their personal data were treated with confidentiality.

The family functions scale was applied in different hospitals in Mexico City¹² where they received medical and rehabilitation treatment. A total

sample of 203 women was recruited, interviewed and given a semistructured interview on socio-demographic aspects.

A group of 100 women with motor disabilities and a second group of 103 people without disabilities were formed. This group was collected in hospitals, schools and recreational centers.

The study was a quantitative, comparative and correlational onemeasurement study, with a significance level of $p \le 0.05$.

3.Results

The results obtained from the socio-demographic data are presented in the following table:

	Tota	WwD	_	WsD	
	n	n	%	n	%
Education					
College	46	35	35	11	10.7
High school	28	13	13	15	14.6
Technical career	21	9	9	12	11.7
Incomplete career	12	9	9	3	2.9
Bachelor's degree	63	26	26	37	35.9
Master's degree	31	7	7	24	23.3
Doctorate	2	1	1	1	1
With whom she lives					
Father/mother	47	30	30	17	16.5
Couple	90	34	34	56	54.4
Brothers	10	6	6	4	3.9
Sons	44	22	22	22	21.4
Alone	10	8	8	2	1.9
Friends	2	0	0	2	1.9
Marital Status					
Single	71	46	46	25	24.3
Married	85	32	32	53	51.5
Separated/divorced	21	14	14	7	6.8
Free Married	20	5	5	15	14.5
Widow	6	3	3	3	2.9

Table 2: Description data

A comparison of means between the groups of Women with disability (WwD) and Women without disability (WsD) was performed and no significant differences were found with respect to family functioning.

Scala	WwD		WsD				
	М	DE	М	DE	gl	t	р
Family functions	57.1	6.4	56.9	5.4	200	.294	.769

Note. N=203. WwD n=100 WsD n=103. Difference in average

*p≤.05. **p≤.01.

Table 3: Analysis comparison of total mean

The following table shows the results of the comparison of means between the factors of the scale for both study groups, carried out by means of Studen' t-test, with the following results: differences were observed in the factors positive family environment in WwD (M=21. 8, SD=4.6), in WwD (M=23.2, SD=3.8), cohesion and norms WwD (M=15.9, SD=3.5), in WwD (M=16.4, SD=3.2), hostility and conflict

avoidance WwD (M=9. 8, SD=2.9), in WsD (M=8.9, SD=2.3), and in the factor command problems in expressing feelings WwD (M=9.5, SD=3.5), in WsD (M=8.5, SD=2.8).

The significant differences detected between the two groups on the family functioning scale were in the factors: positive family environment, hostility and conflict avoidance and problems in expressing feelings.

Variable	WwD		WsD				
	М	DE	М	DE	gl	t	р
1.Positive family environment	21.8	4.6	23.2	3.8	201	-2.44	.015*
2.Cohesion and rules	15.9	3.5	16.4	3.2	197.9	428	.669
3.Hostility and conflict avoidance	9.8	2.9	8.9	2.3	201	2.44	.016*
4. Control problems in expression	9.5	3.5	8.5	2.8	200	2.23	.026*
of feelings							

Note. N=203. *WwD n*=100 *WsD n*=103.

p*≤.05. *p*≤.01.

Table 4: Analysis of differences of means by factor scale family functions

Discussion

The socio-demographic data show important differences between the two groups: the first group of WwD has a lower educational level, lives longer with the family of origin, lives less in a couple and has fewer years of vocational training, which shows that the group of women with disabilities experiences different levels of exclusion and backwardness compared to the WsD group [10].

To determine whether the differences between the two groups were significant, a comparison of means per factor of the scale applied was carried out, in this analysis it was observed that the WwD group presented significant differences in positive family environment, where they scored low, but not in the factors hostility and conflict avoidance, and problems in the expression of feelings, where the difference was significant, a high mean was obtained in relation to the WsD.

The results indicate that the family environment is perceived by WwD less positive, they have a perception of greater hostility, they determine for conflict avoidance and identify greater problems in the expression of feelings, which implies less expression of their feelings, discomfort or discomfort, disagreements, as well as states of wellbeing ^{7,9}.

It was also observed that the group of women with disabilities perceive their family environment with more negative and excluding aspects compared to the group without disabilities, which may hinder their personal development and their adaptation to the social environment ^{7,11}.

Parents are important figures who must adapt to each of the stages of disability (diagnosis, treatment, rehabilitation, education, employment) and where changes in family functioning will be determinant in achieving family cohesion, building harmonious relationships, facilitating the adjustment of roles and the redistribution of tasks within the family ^{4,5,6}.

Therefore, in the care process, the family must be considered, addressing the various conflict dynamics that can be generated within the family, which can become sources of conflict and stress among its members. Influencing family functioning will facilitate the process of acceptance and adaptation to disability ^{4,5,7} as several studies have pointed out, the attitude of acceptance of disability is strongly related to the adaptation of the family group ^{7,8}.

Family functioning is important in the construction of strategies and the development of adaptive psychological resources. Some of these structural elements within the family are: the autonomy of each of the

members, the existence of clear limits, flexibility in the face of changes, the expression of affection in a clear and direct way, openness to information and the creation of new relationships outside the family group^{5,6}. A positive family environment in which conflict resolution and the expression of emotions, as well as disagreement, are favored are determining factors in the development of problem-solving strategies ^{2,3}.

It is common for parents to focus more on the member with a disability, which can lead to abandoning or neglecting the other members (children, partner), which has an impact on the quality of affective relationships, coming to perceive the member with a disability in a negative way, sick, and promoting hostile and competitive relationships within the family ^{5,7}.

Several studies point out that it is common for parents to focus on the disabled member, which can lead to abandoning or neglecting the other members of the family (children, partner), which has repercussions on the quality of affective and fraternal relationships, leading to a negative perception of the disabled member, or perceiving him or her as ill, which fosters hostile and competitive relationships between family members ^{5,7}.

Knowing the characteristics of families in the face of the challenge that disability implies is of utmost importance, as a series of links and experiences are developed in the family that will be fundamental for the person in the process of adaptation, this will allow them to achieve greater adjustment and adaptation in various spheres of their development ^{4,5,7}.

Conclusion

Disability is an experience that brings families into strong and constant states of tension and involves facing challenges, deconstructing the family system where all members can express their emotions. In the study, WwD rated themselves as having lower family functioning compared to the group of non-disabled women. The differences detected in the study were located in the factors less positive family environment, greater hostility and conflict avoidance and greater problems in the expression of feelings. Clinical work with the family and with the person with disabilities in the different facets of their evolutionary cycle is recommended in order to achieve greater integration of the experience and its intrinsic challenges.

Acknowledgments to women who agreed to participate in the study. Funding None.

J. Psychology and Mental Health Care

Conflicts of Interest Author declare that there is no conflict of interest.

References

- 1. Informe mundial sobre discapacidad, (2011). OMS, Banco Mundial.
- Minuchin, S., & Fishman, H.Ch. (2013). Técnicas de terapia familiar. Paidós.
- Velasco, M.L., & Sinibaldi, J.J.F. (2001). Manejo del enfermo crónico y su familia (sistemas, historias y creencias). Manual moderno.
- Rea, A. A. C., Acle T. G., Ampudia, R. A., & García, M. M. (2014). Caracterización de los conocimientos de las madres sobre la discapacidad de sus hijos y su vínculo con la dinámica familiar. Acta Colombiana de Psicología, 17(1), 91–103.
- Grau, C., & Fernández, M. (2010) Familia y enfermedad crónica pediátrica. Anales del sistema sanitario de Navarra, 33(2), 203-212.
- 6. Rolland, J.S. (2000). Familias enfermedad y discapacidad.

Gedisa.

- Quintero, F.J.; Amaris, M. del C. & Pacheco, R.A. (2020). Afrontamiento y funcionamiento en familias en situación de discapacidad. Revista Espacios, 41(17), 1-10. http://es.revistaespacios.com/a20v41n17/a20v41n17p21.pdf
- Villavicencio, A. C., Romero, M.M., Criollo, A. M., & Peñaloza, P.W. (2018). Discapacidad y familia: Desgaste emocional. Revista de investigación en Ciencias Sociales y Humanidades. 5 (1), 1-10.
- Prieto, de la R. A. (2013). Discriminación múltiple: mujeres con discapacidad en México. Género y salud en cifras, 11 (2), 21-30.
- 10. Asamblea General de la ONU (2016). Las mujeres y las niñas con discapacidad necesitan empoderamiento, no piedad. O.N
- 11. Encuesta Nacional sobre Discriminación, ENADIS.(2017).
- García, M.M., Rivera, A.S., Reyes, L. I., & Díaz, L. R. (2006). Construcción de una escala de funcionamiento familiar. Revista iberamericana de diagnóstico y evaluación psicológica, 2 (22), 91-110.



This work is licensed under Creative Commons Attribution 4.0 License

To Submit Your Article Click Here:

Here: Submit Manuscript

DOI: 10.31579/2637-8892/248

Ready to submit your research? Choose Auctores and benefit from:

- fast, convenient online submission
- rigorous peer review by experienced research in your field
- rapid publication on acceptance
- > authors retain copyrights
- > unique DOI for all articles
- immediate, unrestricted online access

At Auctores, research is always in progress.

Learn more https://auctoresonline.org/journals/psychology-and-mental-healthcare