





Gender differences in quality of life among Bell's palsy patients: a cross-sectional study

Diferenças entre gêneros na qualidade de vida entre pacientes com paralisia de Bell: um estudo transversal

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ABSTRACT | BACKGROUND: Functional and psychosocial consequences are reported to be associated with Bell's palsy. Quality outcomes in bell's palsy could be influenced by the sociodemographic factors in males and females, but their predicting factors are scantly reported. This study is designed to compare the quality outcomes in males and females with Bell's palsy. METHODS: Nonprobability sampling and a convenient method were used to recruit 196 participants in this cross-sectional study. The study protocol followed the STROBE guidelines and the institutional review board approval for the study was obtained. Participants are screened for eligibility prior to enrollment. The data collection form is used to get the participant characteristics. The facial disability index and World Health Organization Quality of life Brief version (WHO QOL BREF) are used to assess the quality outcomes. RESULTS: Chi-square statistics for the sociodemographic data in males and females were significant for the type of employment (χ < 0.001). Unpairedt-testt analysis showed significant differences (P < 0.05) in the mean scores of the physical and socialcomponentst of the facial disability index and physiological, environmenta, I and overall components of WHOQOL BREF indicating that females are affected more than males. Further regression analysis revealeda significant association for a higher level of education (β = 2.77, CI = 0.52 to 5.02, P = 0.046) and private employment (β = 4.30, CI = 0.06 to 8.54, P = 0.047) with poor quality outcomes in females. CONCLUSION: Quality of life is affected more in females with Bell's palsy and the predicting factors are higher levels of education and employment in private organizations. Longer duration of symptoms predicted poor quality in both males and females.

KEYWORDS: Facial Paralysis. Quality of Life. Gender Differences.

RESUMO | INTRODUÇÃO: Relata-se que as consequências funcionais e psicossociais estão associadas à paralisia de Bell. Os resultados de qualidade na paralisia de Bell podem ser influenciados pelos fatores sociodemográficos em homens e mulheres, mas seus fatores preditivos são pouco relatados. Este estudo foi desenvolvido para comparar os resultados de qualidade em homens e mulheres com paralisia de Bell. MÉTODOS: Amostragem não probabilística e método conveniente foram utilizados para recrutar 196 participantes neste estudo transversal. O protocolo do estudo seguiu as diretrizes STROBE e a aprovação do conselho de revisão institucional para o estudo foi obtida. Os participantes são selecionados quanto à sua elegibilidade antes da inscrição. O formulário de coleta de dados é utilizado para obter as características dos participantes. O índice de incapacidade facial e a versão resumida da qualidade de vida da Organização Mundial da Saúde (WHO QOL BREF) são usados para avaliar os resultados de qualidade. RESULTA-DOS: As estatísticas do qui-quadrado para os dados sociodemográficos em homens e mulheres foram significativas para o tipo de emprego (x < 0.001). A análise do teste t não pareado mostrou diferenças significativas (P < 0.05) nos escores médios dos componentes físico e social do índice de incapacidade facial e dos componentes fisiológicos, ambientais e gerais do WHOQOL BREF, indicando que as mulheres são mais afetadas que os homens. Análises de regressão adicionais revelaram associação significativa para maior nível de escolaridade (β = 2,77, CI = 0,52 a 5,02, P = 0,046) e emprego privado (β = 4,30, CI = 0,06 a 8,54, P = 0,047) com resultados de baixa qualidade em mulheres. CONCLUSÃO: A qualidade de vida é mais afetada em mulheres com paralisia de Bell e os fatores preditivos são maior nível de escolaridade e emprego em organizações privadas. A maior duração do sintoma previu má qualidade em homens e mulheres.

PALAVRAS-CHAVE: Paralisia Facial. Qualidade de Vida. Diferenças de Gênero.

Submitted Aug. 27th, 2024, Accepted Nov. 1st, 2024, Published Dec. 12th, 2024

J. Physiother. Res., Salvador, 2024;14:e5898

http://dx.doi.org/10.17267/2238-2704rpf.2024.e5898 | ISSN: 2238-2704

Assigned editors: Elen Beatriz Pinto, Marina Makhoul

How to cite this article: Ravichandran H, Shetty KS, Mhabemo N, Shetty A, Bangera N, Janakiraman B. Gender differences in quality of life among Bell's palsy patients: a cross-sectional study. J Physiother Res. 2024;14:e5898. http://dx.doi.org/10.17267/2238-2704rpf.2024.e5898



1. Introduction

More than identity, the face plays a crucial role in communication and it also reflects the individual's sense of oneself. Facial expressions are fundamental for the extraction and transmission of information's during a conversation. The concern for physical appearance and aesthetics is increasingly common nowadays.¹

Bell's palsy is a non-progressive neurological condition characterized by seventh cranial nerve paralysis and facial muscle paralysis on one side of the face. 1.2 Bell's palsy impairments occur due to the lesion in the facial nerve at any one of its branches at intracranial, infratemporal and extratemporal levels.2 Facial nerve has motor, sensory and parasympathetic functions over the face. The functional problems associated with facial impairments are difficulty in chewing, drinking, swallowing, speaking, smiling, blinking eyes, and other facial expression activities that could distress the individual resulting in anxiety, irritability and affect their psychological status.3 Although the spontaneous recovery occurs within 3 weeks in Bell's palsy patients, there is always a risk of residual paresis and associated facial impairment following recovery.4 Residual facial impairment is quite disfiguring and it is estimated that approximately 29% of Bell's palsy patients develop residual hemifacial impairments. 5 Synkinesia, a form of abnormal movement occurs with Bell's palsy patients with poor prognosis and facial impairments. Despite the type of prognosis, facial impairments associated with Bell's palsy could negatively impact the patient's quality of life throughout its course.1

Individuals with facial impairments perceive their body image negatively, added with their poor prognosis, and they tend to withdraw themselves from social / community participation activities.^{2,5} It is evident that body image dissatisfaction leads to several negative health outcomes and psychopathological states such as depression and low mood status in these individuals.⁴ Thus the facial deficits in Bell's palsy could potentially influence the biopsychosocial aspects of an individual.

Although the condition is self-limiting and spontaneously recovers in the majority of the

independent factors individuals, several associated with the biopsychosocial adverse effect of facial impairment in Bell's palsy. Age, level of education, type of occupation, location of residence, and marital status are some of the factors that potentially could affect the biopsychosocial wellness in Bell's palsy individuals. In a recent study by Bylund et al.6 it is reported that facial impairments of Bell's palsy are associated with psychosocial functions in the majority of the participants. In a cross-sectional study^z the authors reported significant levels of psychological distress in Bell's palsy patients. In a case-control study from China, the authors concluded a strong association of psychological distress in Bell's palsy patients.8

In patients with poor prognosis and with chronicity of the condition, the mechanism disturbing their biopsychosocial wellness is described by the illness intrusiveness phenomenon.⁹ Illness intrusiveness are illness-induced disruptions to performing routine lifestyle activities that can compromise psychosocial well-being and contribute to emotional stress.^{10,11}

In Bell's palsy, the biopsychosocial disturbances could hamper the quality of life of individuals. Quality of life is characterized by four domains such as physical, psychological, social and environmental. The quality of life of an individual with Bell's palsy is determined by his sociodemographic, clinical and psychosocial factors. ¹² In the clinical practice guidelines published by the American Academy of Neurology and the American Academy of Otolaryngology – Head and Neck Surgery Foundation (AAO-HNSF) ¹³ the strategies to improve quality of life in Bell's palsy patients are not clearly reported.

There is a need for identifying gender differences in biopsychosocial and wellness aspects in individuals with Bell's palsy. The results of the study could identify the most affected quality of life component in Bell's palsy to help the clinicians and rehabilitation professionals for screening and to provide interventions accordingly. The present study aims to analyze the association of sociodemographic variables in quality of life among males and females with the diagnosis of Bell's palsy. The study is also hypothesized to test the gender differences in quality of life among Bell's palsy individuals.

2. Methods

This cross-sectional study is conducted according to STROBE guidelines for cross-sectional design. The study protocol was approved (ACP/OP/2023/OL42) by the institutional review board of Alva's College of Physiotherapy and Research Centre, Moodbidri, DK, Karnataka, India. The study was registered in the Clinical Trial Registry of India (CTRI/2024/04/066270). G*power statistical application is used to estimate the sample size. Based on a previous study¹⁴ findings and power analysis using a priori computed with two tails, ratio of variances (0.749), alpha error probability (0.05) and power (0.80) parameters, the estimated sample size for the study is 196 participants. Through non-probability and convenience sampling methods, the eligible participants were recruited from 14 hospitals in Dakshina Kannada district, Karnataka. The diagnosis of Bell's palsy is confirmed by a qualified neurologist or a general physician. Qualified Physiotherapy professionals in those hospitals were trained for screening and data collection processes. The data-collecting personnel were supervised by the authors of the study.

Both genders, aged from 21 to 60 years, with idiopathic Bell's palsy, lower etiology from motor neuron lesion, and one-sided facial weakness with no sparing of forehead muscles are considered for inclusion. Participant presenting with bilateral facial weakness, Bell's palsy secondary to other pathologies, bell's palsy associated with any other comorbid conditions, recurrent Bell's palsy, with history of any form of disabilities or deformities, with history of previous psychiatric illness and not willing to participate were excluded from the study. Participants fulfilling the selection criteria were explained about the study protocol and their consent was obtained. Participants' characteristics and quality of life were the data collected from the participants (Details enclosed in Appendix I: Data collection form).

2.1 Participant characteristics

A standard data collection form is used to collect data from all the respondents. The characteristics of respondents collected were gender, age, marital status, educational qualification, employment status, location of residence, date of diagnosis of Bell's palsy, duration of days from the onset of Bell's palsy symptom, comorbid illness and its duration.

2.2 Quality of life

The facial disability index and WHO QOL BREF questionnaires were the quality of life scales administered to all the enrolled participants.

2.2.1 WHO QOL BREF measure

It is cross-culturally valid, reliable, and available in over 40 languages. It measures the quality of life through individual domains such as physiological, psychological, environmental, social, and overall. The mean score is calculated from each domain and multiplied by 4 to make it comparable to the WHO QOL 100 version. It measures quality outcomes in the general population and individuals with disabling conditions. It is considered an alternative to WHO QOL 100, which is lengthy and time-consuming to administer.¹⁵

2.2.2 Facial disability index measure

It is a facial neuromuscular system disorder-specific, self-report instrument measuring disabilities through physical, social, and overall components. The score for each component in the facial disability index was calculated according to the number of questions answered and the raw score for each question. The overall scores ranged from 0 (worst) to 100 (better). It is Psychometrically proven and easy to administer in Bell's palsy individuals.¹⁶

Trained data collectors ensured the completeness of both the self-reported measures from all the participants.

2.3 Statistical analysis

The collected data were analyzed using Statistical Package for Social Sciences software (version 26.0. Armonk, NY: IBM Corp). Descriptive statistical analysis was performed for participant's characteristics such as age, marital status, education status, employment status, location of residence and duration from the onset of Bell's palsy symptom. An Independent t-test is performed to compare the mean and standard deviation of all the components of the facial disability index and WHO QOL BREF outcomes in males and females. Multiple logistic regression is performed to analyze the influence of demographic characteristics on quality of life outcome.

3. Results

3.1 Participant characteristics

A total of 196 Bell's palsy individuals enrolled in this study from multi-centers in Dakshina Kannada. Both male (n = 91) and female (n = 105) subjects were included in this study. The age criteria for participants included in this study varied from 21 to 75 years and the majority of the participants, 67.34% (n = 132) belong to 41 to 60 years of age. The majority of the participants (Males = 78, Females = 84) with Bell's palsy were married. Male Bell's palsy individuals had a similar proportion for educational categories such as no education, elementary, higher secondary and college level education, compared to the majority of females with Bell's palsy with higher education qualifications. The majority of the males 48.35% (n = 44) were employed in the government sector whereas in females 45.71% (n = 48) were unemployed. Most of the participants are from rural (males = 55, females = 64) place and in the duration of Bell's palsy at the time of data collection was 1 – 2 weeks in both genders (males = 77, females = 93). The chi-square statistical analysis for the demographic characteristics of Bell's palsy participants were significant in education (P = 0.015) and employment (P = 0.000) characteristics (table 1).

Table 1. Gender-wise categorization of sociodemographic characteristics of participants with Bell's palsy

		Male	Female	
Demog	raphic	Frequency (Percentage)	Frequency (Percentage)	χ*
	21-40	24 (24.1%)	28 (25%)	
Age	41-60	60 (69.1%)	72 (65.7%)	0.692
	> 61	7 (6.6%)	5 (9.2%)	0.092
	Single	7 (16.6%)	17 (7.8%)	
Marital status	Married	78 (80.0%)	84 (84.2%)	0.149
	Widow	6 (3.3%)	4 (7.8%)	0.149
	Uneducated	22 (25.8%)	26 (24.76%)	
Education	Elementary	25 (9.1%)	8 (7.6%)	•
status	Higher sec.	27 (30%)	32 (30.47%)	0.015
	College	17 (35%)	39 (37.14%)	•
	Unemployed	17 (18.68%)	48 (45.71%)	
Employment status	Self employed	25 (27.47%)	53 (50.47%)	0.000*
Status	Private	5 (5.49%)	4 (3.80%)	
	Government	44 (48.35%)	0 (0%)	
Location	Rural	55 (60.43%)	64 (60.95%)	
20044011	Urban	36 (39.56%)	41 (39.04%)	0.942
Duration	1-2 weeks	77 (84.61%)	93 (88.57%)	0.415
2	>2 weeks	14 (15.38%)	12 (11.42%)	

3.2 The gender-wise difference in Facial disability index and WHOQOL - BREF outcomes

Facial disability index and WHOQOL – BREF mean scores were compared between genders using independent t-tests. The mean of physical (56.76 \pm 26.1) and social component (45.02 \pm 8.5) of facial disability index were significantly (P < 0.05) lower in females compared to males. The mean of WHOQOL – BREF outcome in environmental (25.31 \pm 3.8) and overall components (6.99 \pm 2.0) were lower in females indicating a significant difference between the genders (table 2).

Table 2. Gender wise comparison of FDI and WHOQOL-BREF

		N	Male		F	emale		Significance
Ou	tcome	M	SD	N	M	SD	N	Significance
	Physical	64.67	25.19	91	56.76	26.14	105	0.033*
FDI	Social	48.48	10.61	91	45.02	8.54	105	0.012*
	Overall	113.15	28.42	91	117.39	22.90	105	0.250
	Physical	21.70	3.32	91	20.61	4.12	105	0.046*
WHOOM	Psychological	16.04	2.80	91	15.00	3.51	105	0.025
WHOQOL- BREF	Social	9.85	1.88	91	9.22	2.36	105	0.043
	Environmental	26.50	3.27	91	25.31	3.84	105	0.022*
	Overall	7.75	1.75	91	6.99	2.02	105	0.005*

Source: the authors (2024).

3.3 Association between gender and quality of life

After controlling sociodemographic variables, the multiple regression analysis was conducted between gender and quality of life (facial disability index and WHOQOL – BREF outcomes). The physical component outcomes of facial disability index (P = 0.012, β = 8.486, CI = 1.856 to 15.117) and WHOQOL – BREF (P = 0.000, β = 2.639, CI = 1.631 to 3.647) were significantly correlated with female gender (table 3 and 4).

Table 3. Gender-wise difference in quality of life (FDI)

						FI	I						
	Physical				Social				Overall				
	В	SE	p	CI	В	SE	P	CI	В	CI			
Female	8.486	3.362	0.012	1.856 to 15.117	-3.391	1.355	0.013	-6.023 to -0.720	5.095 3.751 0.176 -2.303 12.49				
Male		R	ef	•		I	Ref			Re	f	•	

Table 4. Gender-wise difference in quality of life (WHOQOL-BREF)

									WHO	OQOL-B	REF									
	Physica	Physical Psychological						Enviro	ıment			Social				Overall				
	В	SE	P	CI	В	SE	p	CI	В	SE	p	CI	В	SE	p	CI	В	SE	p	CI
Female	2.639	0.511	.000	1.631 to 3.647	0.297	297 0.404 0.491 -0.518 to 1.076				0.945 0.491 0.056 .024 to 1.914			0.594 0.273 2.177 0.056 to 1.133				-0.637	0.281	0.025	-1.191 to -0.082
Male		R	ef			R	ef		Ref			Ref				Ref				

3.4 Factors predicting and protecting poor quality of life outcomes in both genders

Bell's palsy males in the age group from 21 to 40 years (β = - 7.85,CI = -13.281 to -2.430, P = 0.005) and from 41 to 60 years (β = - 6.54, CI = -11.562 to - 1.529, P = 0.011) are protective against poor quality outcomes in physiological component of WHOQOL BREF.

In elementary-level education males are protected against higher scores (β = - 14.71, CI = -29.19 to -0.24, P = 0.046) in the overall component of facial disability index. College-level education in females, significantly predicted (β = 2.77, CI = 0.52 to 5.02, P = 0.046) the higher scores in environmental component of WHOQOL BREF scale.

Females working in the private sector are associated with higher odds of scoring poor outcomes in the physiological component (β = 4.30, CI = 0.06 to 8.54, P = 0.047) of WHOQOL BREF. Whereas males working in private sectors are protected from low scores in physical (β = -26.77, CI = -52.20 to -1.35, P = 0.039) and overall (β = -32.40, CI = -60.96 to -3.84, P = 0.027) component in facial disability index. Similarly, the self-employed males had an inverse association (β = -2.47, CI = -4.73 to -0.22, P = 0.031) with the psychological component of WHOQOL BREF outcome (table 5,6,7,8).

Table 5. Quality of life in females (FDI)

				Physica	l			Socia	ıl			Overa	11
FDI		В	SE	P	CI	В	SE	P	CI	В	SE	P	CI
	21 – 40 years	- 7.58	16.3	0.644	- 40.06 to 24.89	-1.46	6.0	0.809	- 13.457 to 10.527	2.359	14.5	0.872	-26.564 to 31.283
AGE	41 – 60 years	- 14.41	15.1	0.343	- 44.45 to 15.62	2.28	5.5	0.684	- 8.808 to 13.369	- 2.90	13.4	0.830	-29.648 to 23.841
	Above 61 years			1			<u> </u>	1				1	
	Married	19.64	16.5	0.239	- 13.26 to 52.56	- 1.69	6.1	0.783	- 13.846 to 10.458	- 5.29	14.7	0.721	- 34.606 to 24.013
MARITAL STATUS	Widow			1				1			l	1	
	Single	25.73	18.5	0.169	- 11.16 to 62.62	3.58	6.8	0.602	- 10.032 to 17.210	0.63	16.5	0.970	-32.220 to 33.485
	Uneducated			1				1	I			1	
EDUCATIONAL	Elementary	- 1.60	9.7	0.869	- 20.92 to 17.71	- 1.66	3.5	0.645	- 8.794 to 5.471	10.71	8.6	0.219	- 6.493 to 27.914
STATUS	Higher sec.	- 0.29	10.0	0.977	- 20.22 to 19.64	0.93	3.7	0.801	- 6.424 to 8.294	12.35	8.9	0.17	- 5.398 to 30.10
	College	- 1.34	9.9	0.893	- 21.01 to 18.33	0.18	3.6	0.959	- 7.075 to 7.451	16.81	8.8	0.06	- 0.700 to 34.33
	Unemployed			1	ı			1				1	
EMPLOYMENT	Self employed	-3.64	5.7	0.529	- 15.11 to 7.82	1.38	2.1	0.518	- 2.850 to 5.620	0.58	5.1	0.909	-9.629 to 10.803
STATUS	Private	-0.07	12.7	0.996	- 25.43 to 25.29	2.54	4.7	0.590	- 6.818 to 11.911	-8.48	11.3	0.458	-31.071 to 14.102
	Urban			1				1			<u> </u>	1	
LOCATION STATUS	Rural	6.880	5.4	0.213	- 4.020 to 17.780	1.87	2.0	0.357	- 2.150 to 5.899	4.49	4.8	0.314	- 4.760 to 14.652
	< 2 weeks		-	Ref				1	I		1	1	
DURATION STATUS	>2 weeks	37.447	7.5	0.000	22.470 to 52.424	- 0.270	2.7	0.923	- 5.799 to 5.259	29.14	6.7	0.000	15.810 to 42.482

 Table 6. Quality of life in females (WHOQOL-BREF)

WHOOO	RRFF	Physi	ologic	al		Psych	ologic	al		Envi	onme	ntal		Social	l,			Overa	11		
Whogo	L-DKL1	В	SE	P	CI	В	SE	P	CI	В	SE	P	CI	В	SE	P	CI	В	SE	P	CI
	21 – 40 years	2.96	1.7	0.097	-6.47 to	0.03	1.4	0.983	-2.95 to 2.88	- 1.14	1.6	0.493	-4.446 to 2.161	1.00	0.9	0.317	-0.980 to 2.985	-0.85	0.9	0.370	-2.728 to 1.026
AGE	41 – 60 years	1.80	1.6	0.268	-5.03 to 1.41	0.23	1.3	0.864	-2.44 to 2.91	0.64	1.5	0.675	-3.673to 2.392	0.48	0.9	0.601	-1.339 to 2.300	-0.76	0.8	0.378	-2.490 to 0.956
	Above 61 years	1				1		'		1	1	1	1	1	1	1	1	1	1	1	1
	Single	4.95	3.1	0.114	-1.211 to	- 1.68	2.8	0.556	-7.335 to 3.967	- 2.27	3.0	0.463	-8.395 to 3.853	3.57	1.8	0.053	-0.047 to	-0.26	1.6	0.872	-3.477 to 2.956
MARITAL STATUS	Married	1.95	2.7	0.482	-3.545 to 7.450	2.18	2.5	0.392	-7.224 to 2.860	- 1.54	2.7	0.575	-7.012 to 3.915	2.50	1.6	0.127	-0.725 to 5.739	0.488	1.4	0.736	-2.382 to 3.357
	Widow			1	7.430	2.10		1	2.000	1.57	I	1	5.715			1	0.137			1	W 3.331
	Uneducated			1				1				1				1				1	
EDUCATIONAL	Elementary	1.46	1.6	0.370	-4.692 to 1.762	1.29	1.4	0.388	-1.667 to 4.252	0.73	1.6	0.651	-2.474 to 3.940	0.457	0.9	0.634	-1.440 to 2.354	-0.69	0.8	0.415	-2.378 to 0.990
STATUS	Higher sec.	1.89	1.6	0.262	-5.219 to 1.439	2.00	1.5	0.196	-1.050 to 5.057	0.86	1.6	0.60	-2.443 to 4.174	0.016	0.9	0.987	-1.941 to	-1.26	0.8	0.151	-3.006 to 0.470
	College	0.22	1.2	0.850	-2.61 to	1.23	0.9	0.219	-0.75 to 3.22	2.77	1.1	0.016	0.52 to 5.02	-0.17	0.6	0.797	-1.523 to 1.173	0.14	0.6	0.828	-1.137 to 1.417
EMPLOYMENT	Unemployed			1				1			l	1				1				1	
STATUS	Self employed	1.49	0.9	0.126	-0.426 to 3.406	0.70	0.8	0.425	-2.466 to 1.048	0.78	0.9	0.415	-1.119 to 2.689	-0.36	0.5	0.525	-1.489 to 0.764	-0.14	0.5	0.782	-1.140 to 0.860
	Private	4.30	2.1	0.047	0.067 to 8.540	0.45	1.9	0.818	-4.337 to 3.434	0.77	2.1	0.716	-3.435 to 4.985	-0.81	1.2	0.520	-3.301 to 1.680	0.59	1.1	0.597	-1.620 to 2.803
	Rural			1								1				1				1	
LOCATION STATUS	Urban	0.05	0.9	0.951	-1.765 to	0.31	0.8	0.709	-1.984 to	1.02	0.9	0.265	-2.330 to 2.641	-0.22	0.5	0.678	- 1.295 to 0.846	0.28	0.4	0.559	-0.670 to 1.231
	< 2 weeks			1	1			1				1	1		'	1				1	
DURATION STATUS	>2 weeks	2.69	1.2	0.035	0.194 to 5.197	0.98	1.1	0.397	-1.310 to 3.278	0.15	1.2	0.901	-2.330 to 2.641	1.87	0.7	0.013	0.402 to 3.343	-0.62	0.6	0.345	-1.930 to 0.681

 Table 7. Quality of life in males (FDI)

				Physical				Social				Overall	
FDI		В	SE	P	CI	В	SE	P	CI	В	SE	P	CI
	21 – 40 years	6.10	11.1	0.584	- 16.00 to 28.212	- 10.10	5.4	0.068	- 20.98 to 0.77	- 3.99	12.4	0.749	- 28.83 to 20.83
AGE	41 – 60 years	8.50	10.1	0.407	- 11.78 to 28.80	- 8.74	5.0	0.085	- 18.72 to 1.24	- 0.23	11.4	0.984	- 23.03 to 22.56
	Above 61 years			1				1				1	
	Married	4.91	10.9	0.656	- 16.95 to 26.77	4.40	5.4	0.457	- 6.71 to 14.80	8.95	12.3	0.470	- 15.60 to 33.51
MARITAL STATUS	Widow			1				1				1	
	Single	16.42	13.2	0.217	- 9.86 to 42.70	8.40	6.4	0.199	- 4.52 to 21.34	24.83	14.8	0.098	- 4.69 to 54.35
	Uneducated			1				1				1	
EDUCATIONAL	Elementary	- 12.27	6.4	0.062	- 25.16 to 0.61	- 2.44	3.1	0.446	- 8.78 to 3.90	- 14.71	7.2	0.046	- 29.19 to - 0.24
STATUS	Higher sec.	- 2.80	7.3	0.704	- 17.42 to 11.82	1.41	3.6	0.697	- 5.78 to 8.61	- 1.38	8.2	0.867	- 17.81 to 15.04
	College	- 2.01	7.5	0.791	- 17.05 to 13.02	5.81	3.7	0.122	- 1.58 to 13.21	3.79	8.4	0.656	- 13.09 to 20.69
	Unemployed			1				1				1	
EMPLOYMENT	Self employed	6.66	8.5	0.439	-10.39 to 23.71	4.01	4.2	0.344	- 4.37 to 12.40	10.67	9.6	0.271	- 8.48 to 29.83
STATUS	Private	- 26.77	12.7	0.039	-52.20 to -1.35	- 5.62	6.2	0.373	- 18.13 to 6.88	- 32.40	14.3	0.027	- 60.96 to - 3.84
	Government	-1.69	7.8	0.830	-17.35 to 13.95	1.74	3.8	0.654	-5.96 to 9.44	0.04	8.8	0.996	- 17.53 to 17.62
LOCATION STATELS	Urban			1				1				1	
LOCATION STATUS	Rural	4.86	5.1	0.345	- 5.32 to 15.06	1.62	2.5	0.520	- 3.38 to 6.64	6.49	5.7	0.262	- 4.95 to 17.94
DVD 1570316515	< 2 weeks			1				1				1	
DURATION STATUS	>2 weeks	34.66	6.4	0.000	21.74 to 47.57	1.75	3.1	0.584	- 4.59 to 8.11	36.41	7.2	0.000	21.91 to 50.92

 Table 8. Quality of life in males (WHOQOL-BREF)

WHOOO	DDEE		Ph	ysiologi	cal		Psyc	chologic	al		Envi	ronmen	ıtal			Social			C	verall	
WHOQOI	-BKEF	В	SE	P	CI	В	SE	P	CI	В	SE	P	CI	В	SE	P	CI	В	SE	P	CI
	21 – 40 years	- 7.85	2.7	0.005	-13.281 to - 2.430	1.71	2.5	0.495	-3.259 to 6.692	2.49	2.7	0.361	-7.882 to 2.901	-1.19	1.6	0.459	-4.384 to 1.994	-0.10	1.4	0.942	-2.936 to 2.728
AGE	41 – 60 years	- 6.54	2.5	0.011	-11.562 to – 1.529	2.36	2.3	0.310	-2.234 to 6.967	3.41	2.5	0.177	-8.404 to 1.567	-1.24	1.4	0.405	-4.191 to	-0.44	1.3	0.738	-3.062 to 2.176
	Above 61 years			1				1				1	l			1	ı			1	
	Single	1.71	2.0	0.417	-2.46 to 5.88	2.97	1.7	0.092	-0.49 to 6.44	0.75	1.9	0.704	-4.680 to 3.175	0.70	1.1	0.555	-1.655 to 3.059	0.07	1.1	0.950	-2.160 to 2.303
MARITAL STATUS	Married	0.58	1.7	0.738	-2.88 to 4.06	2.35	1.4	0.108	-0.52 to 5.24	1.30	1.6	0.427	-4.576 to 1.958	0.02	0.9	0.981	-1.938 to 1.984	0.58	0.9	0.536	-1.277 to 2.436
	Widow			1				1				1				1				1	
	Uneducated			1				1				1				1				1	
	Elementary	1.07	1.0	0.301	-3.11 to 0.97	0.94	0.8	0.271	-0.75 to 2.65	0.68	0.9	0.479	-1.237 to 2.614	-0.94	0.5	0.106	-2.105 to 0.207	0.41	0.5	0.451	-0.678 to 1.510
EDUCATIONAL	Higher sec.	0.56	1.1	0.627	-1.75 to 2.89	-0.03	0.9	0.974	-1.96 to 1.90	0.22	1.0	0.841	-1.965 to 2.405	-0.77	0.6	0.241	-2.090 to 0.533	0.078	0.6	0.901	-1.164 to 1.319
STATUS	College	0.85	1.6	0.608	-4.138 to 2.433	0.96	1.5	0.528	-2.051 to 3.975	1.15	1.6	0.484	-2.111 to 4.419	0.068	0.9	0.944	-1.863 to 2.000	-1.10	0.86	0.204	-2.819 to 0.611
	Unemployed			1				1				1	•			1				1	
EMPLOYMENT	Self employed	0.67	1.3	0.624	-2.04 to 3.38	-2.47	1.1	0.031	-4.73 to - 0.22	1.50	1.2	0.243	-1.042 to 4.055	0.91	0.7	0.238	-0.616 to 2.443	-0.57	0.7	0.433	-2.022 to 0.875
STATUS	Private	1.74	2.0	0.394	-2.30 to 5.78	-3.25	1.6	0.057	-6.61 to 0.10	0.67	1.1	0.567	-1.664 to 3.014	0.73	1.1	0.523	-1.546 to 3.015	-0.25	1.0	0.819	-2.408 to 1.909
	Government	1.27	1.2	0.309	-1.20 to 3.76	- 2.492	1.0	0.019	-4.55 to 0.42	3.24	1.9	0.093	-0.553 to 7.045	0.03	0.7	0.960	-1.368 to 1.439	-0.19	0.6	0.774	-1.522 to 1.136
	Rural			1				1				1				1				1	
LOCATION STATUS	Urban	0.30	0.8	0.711	-1.922 to 1.317	0.73	0.6	0.280	-0.61 to 2.08	- 0.64	0.7	0.404	-2.165 to 0.881	-0.25	0.4	0.576	-1.172 to 0.656	0.775	0.4	0.079	-1.640 to 0.091
	< 2 weeks			1				1				1				1				1	
DURATION STATUS	>2 weeks	1.85	1.0	0.076	-0.19 to 3.90	0.445	0.8	0.605	-1.26 to 2.15	- 0.90	0.9	0.354	-2.833 to 1.026	0.74	0.5	0.202	-0.410 to 1.906	0.245	0.5	0.658	-1.341 to 0.852

The poor outcome in the physical and overall component of facial disability index is significantly predicted by symptoms lasting more than 2 weeks in males and females. Whereas the poor quality in the physiological and social component of WHOQOL BREF is significantly predicted by the persistence of symptoms for more than 2 weeks in females compared to males.

4. Discussion

The objective of the present study is to compare the quality of life outcome scores in males and females with Bell's palsy and the statistical analysis results indicated that females had poor quality outcomes compared to males and the poor outcomes are significantly reflected in physical, physiological, environmental and social contexts. In the same vein, age, lower educational qualification, and private and self-employment status in males protected them from lower quality scores. Despite the chances for a higher recovery rate, factors such as severity of nerve injury, extent of muscles paralyzed, age and comorbidities challenge and delays the recovery in individuals with Bell's palsy. The functional and aesthetic dysfunction impairs the quality of life in Bell's palsy individuals with poor prognosis.

Similar to the present study findings, Bylund et al.6 reported that females with Bell's palsy suffer from psychosocial dysfunction compared to males. Contrastingly in the present study, there is no significant association identified in psychological outcomes for Bell's palsy patients. At the same time, the predicting factors associated with Bell's palsy females for physical, physiological, social and environmental components varied inconsistently. In this study Bell's palsy individuals with less than 2 weeks and more than 2 weeks were included and hence it is probably challenging to predict the influence of demographic characteristics in psychological component. This could be a contributing factor that the present study finding showed insignificant changes in the psychological component of the WHOQOL BREF scale.

The study findings revealed that graduated females lack quality in the environmental component of WHOQOL BREF. Females with facial deviations in Bell's palsy had low satisfactory scores for

traveling and participating in leisure or recreational activities. This could be due to the perception of low self-esteem and the stigmatization effect following Bell's palsy. According to Fugere et al. 12 and Crerand et al.18 females are more concerned about their physical appearance and they consider as the most desirable trait. This could be a factor, hindering their satisfaction with public transportation and social participation. Low satisfaction with family members and spouse is also reported in Bell's palsy females. The inability to cope with the stress related to facial impairment might interfere their family relationships. Lack of communication, conflicts, avoidant actions, and disengagement from family hours could predict home environmental quality outcomes in females with Bell's palsy.

In this study, the physiological capacity of the WHOQOL BREF scale is significantly correlated with females employed in private sectors. Facial appearance is one of the components of body image perception. 19,20 Appearance-related dissatisfaction is reported more commonly among females than males in the workplace.²¹⁻²³ Appearance culture dominates in most workplaces in which the employees are insisted on achieving for a perfect look.²⁴⁻²⁶ Inability to smile, drooping of the eyelid, drooling, and difficulty in speaking (pronunciation) negatively impact the personality and self-confidence during work and affect their performances. Additionally, incomplete closure of eyelids in Bell's palsy results in eye dryness and sleep deprivation leading to fatigue and lack of concentration during work. This could be attributed to significant findings associated with poor satisfaction for the physical component of WHOQOL BREF in the private sector.

The longer duration of facial impairment is a predicting factor for physical and overall components of facial disability index in males and females. In females, the longer duration of symptoms is associated with the physiological and social component of WHOQOL BREF. This indicates that a poor prognosis increases the risk of poor quality of life in Bell's palsy.

The first limitation of the present study is that the outcome measures lack components analyzing the prognosis with quality of life. Such outcome analysis could reflect the impact of disease prognosis or

therapy outcomes on the quality of life among Bell's palsy individuals. Secondly, the cross-sectional design methodology could not address the causal inferences. Several confounding factors such as participant's knowledge about the condition, family/ spouse/healthcare provider support, insurance/ reimbursement coverage benefits, treatments undergone, and adherence might affected the causality which is not addressed in this study. Additionally, the individuals included in this study were not categorized according to the stages of Bell's palsy.

The strength of the present study is the objective to identify the factors predicting quality outcomes in males and females with Bell's palsy. Additionally, the predicting factors and associated poor-quality outcomes could hamper the recovery in the form of a vicious cycle. The finding of the present study providing knowledge on the potential predicting factors could help to design strategies to improve quality of life.

The clinical application of the present study's interpretation could benefit the participants, clinicians/therapists, and stakeholders. As the quality of life outcome is affected more in females, clinicians/therapist-provided education about their diagnosis could reflect changes in their quality of life and wellbeing. Access to healthcare, assurance, and financial assistance from stakeholders could reduce the poor quality of life outcome-predicting factors in females such as pursuing higher education, private employees, and the presence of symptoms for more than 2 weeks.

Further studies examining the association of facial disability outcomes using the FaCE scale, Sunny Brook scale, or House Brackmann scale will add more insights to the present study.

5. Conclusion

The study results suggest that females are more susceptible to poor quality of life outcomes in Bell's palsy than males. In females, higher education and private sector employment are the predicting factors for poor outcomes in quality of life. Longer duration of symptoms predicted poor quality outcomes in both genders.

Authors contributions

The authors declared that they have made substancial contributions to the work in terms of the conception or design of the research; the acquisition, analysis or interpretation of data for the work; and the writing or critical review for relevant intellectual content. All authors approved the final version to be published and agreed to take public responsability for all aspects of the study.

Conflicts of interest

No financial, legal, or political conflicts involving third parties (government, private companies, and foundations, etc.) were declared for any aspect of the submitted work (including but not limited to grants and funding, advisory board participation, study design, manuscript preparation, statistical analysis, etc.).

Indexers

The Journal of Physiotherapy Research is indexed by <u>DOAJ</u>, <u>EBSCO</u>, <u>LILACS</u> and <u>Scopus</u>.









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Appendix

Appendix I. Data collection form

S.no	Information	A – Characteristics of the participants Response	Code
5.110	intormation	Response	Code
1	Participant ID code		
2	Assessors ID code		
3	Date of screening and place		
4	Date and time of enrollment		
5	Gender	Male G1 Female G2	
5	How old are you?		
7	Marital status	Single Married Widow/Divorced	
8	Educational qualification	Not educated Elementary School Higher secondary school College level higher education	
9	Employment status	Unemployed Self employed Private employee Government employee House wife (if females)	
10	Location of residence	Rural R1 Urban R2	
11	Date of diagnosis of Bell's Palsy		
12	Duration of days from the onset of bells palsy symptom		
13	Comorbid illness	No Yes (if No skip further questions and go to Section B)	
14	Answer the type of comorbid condition you are suffering from	Cardiovascular disorder C1 Neurological disorder C2 Metabolic disorder C3 Oncological disorder C4 Musculoskeletal disorder C5 Respiratory disease C6 Gastrointestinal disease C7 Hematologic disease C8	
15	Mention the duration of the comorbid condition that you are suffering from	Cardiovascular disorder D1 duration Neurological disorder D2 duration Metabolic disorder D3 duration Oncological disorder D4 duration Musculoskeletal disorder D5 duration Respiratory disease D6 duration Gastrointestinal disease D7 duration Hematologic disease D8 duration	
	SE	CCTION B – Quality of Life Outcome	
	Qulity of life outcomes	G1 G2	
A	Facial disability index Total Score Social Physical		
В	WHO QOL BREF Total score Physical Psychological Social Environmental		