

# Chronic pain among Lebanese individuals with spinal cord injury: Pain interference and impact on quality of life

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## Keywords

Chronic Pain; Spinal Cord Injuries; Quality of Life; Lebanon

## Abstract

**Background:** Chronic pain is one of the most disabling consequences of spinal cord injury (SCI). Although studies have identified a link between chronic pain and decreased quality of life (QOL) among this population, few studies have looked into the experience of chronic pain in Lebanese individuals with SCI and the impact of pain characteristics on QOL. Thus, the present study evaluated the chronic pain experience and its associated factors among Lebanese individuals with SCI in order to determine the impact of pain on QOL.

**Methods:** A cross-sectional study was conducted on 81 Lebanese individual with SCI between August 1<sup>st</sup> and October 31, 2022. The collected information included sociodemographic characteristics, SCI-related information, pain-related variables, and the 12-item Short Form Health Survey (SF-12). Factors associated with pain interference were evaluated using a linear

regression model. One-way ANOVA and independent sample t-test were used to evaluate the association of different baseline and pain characteristics with QOL.

**Results:** In the present study, 81.5% of participants reported chronic pain with the majority of them having neuropathic pain type. Employment status ( $P = 0.034$ ), type of pain ( $P = 0.009$ ), and pain severity ( $P = 0.028$ ) were significantly associated with pain interference. Unemployed participants and those with severe chronic pain, particularly neuropathic pain, had lower QOL.

**Conclusion:** Chronic pain was found to be highly prevalent among Lebanese patients with SCI. Pain interference and QOL were significantly affected by employment status and pain type. Therefore, targeting chronic pain and its associated factors in rehabilitation practice is warranted.

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## Introduction

Spinal cord injury (SCI) is a serious medical condition that leads to a significant disability.<sup>1</sup> Various health complications result from SCI including sensory and motor function impairments, neurogenic bladder and bowel function problems,<sup>2,3</sup> cardiovascular and respiratory complications,<sup>4</sup> sexual disorders,<sup>5</sup> spasticity,<sup>6</sup> chronic pain,<sup>7</sup> and others.<sup>8</sup> These underlying problems affect the patient's physical and psychological functioning, which significantly impacts their quality of life (QOL).<sup>9-11</sup>

QOL,<sup>12</sup> commonly reduced in the SCI population,<sup>13-15</sup> has been a key outcome measure in rehabilitation.<sup>16,17</sup> QOL is considered multidimensional, and several studies have emphasized various factors as producing negative impacts on QOL.<sup>18-20</sup> Age, gender, employment, education, as well as injury-related factors such as injury level and duration have been studied as factors that lower QOL levels in SCI.<sup>21-23</sup> Therefore, chronic pain has been proposed to have an eventual major impact on QOL due to its effect on mood, sleep, recreational and vocational activities, and different rehabilitation outcomes.<sup>24-28</sup>

Chronic pain is one of the most debilitating consequences of SCI.<sup>29,30</sup> It has been reported in approximately 68% of the SCI population;<sup>7</sup> it is classified into the two types of nociceptive and neuropathic pain with the latter being considered the most disabling.<sup>31,32</sup> The experience of chronic pain after SCI is infrequently resolved<sup>33</sup> and it is influenced frequently with various factors.<sup>34-36</sup> Previous studies found that pain experience is associated with sociodemographic factors, injury characteristics and complications, and other psychological and environmental factors.<sup>36-38</sup>

The experience of chronic pain in SCI encompassing its characteristics, types, and associations has been an area of interest for researchers in the last decade with the aim to improve pain management procedures.<sup>32,39-41</sup> Although various studies have examined the relationship between chronic pain experience and different clinical and sociodemographic factors,<sup>38</sup> it remains unclear and inconclusive in the literature,<sup>42,43</sup> and a need for studies on the characteristics of pain severity and interference with daily life and their predictive factors has emerged.<sup>44</sup> In addition, studies have reported the association of chronic pain with decreased QOL,<sup>9,32,45,46</sup> thus, limited studies have investigated the impact of pain characteristics including severity, interference, and

specific types on QOL.<sup>46,47</sup>

Accordingly, it has been recently recommended that clinical research focus on pain interference and its influencing factors rather than only focusing on pain severity, which is frequently studied in pain research.<sup>48</sup> Its precise influence on QOL has yet failed to be conclusive specifically in individuals with SCI in developing countries.<sup>49</sup> To the best of our knowledge, no studies have been performed to evaluate the prevalence of chronic pain experience, its characteristics among individuals with SCI, and its impact on their QOL particularly in Lebanon. In this regard, proceeding from the necessity of understanding this relationship as a great concern to update efficient rehabilitative protocols for chronic pain management in individuals with SCI, we tried initially to assess this concept in the Lebanese population with SCI. The primary aim of the present study was to evaluate the experience of chronic pain and to identify the factors associated with pain interference in Lebanese individuals with SCI. The second aim was to determine the impact of different factors, particularly pain, on their QOL.

## Materials and Methods

**Participants:** The participants included individuals with SCI who were recruited mainly from the Health, Rehabilitation, Integration, and Research Center (HRIR), a tertiary outpatient rehabilitation center for people with disabilities located in three main geographic areas (Beirut, South, and Bekaa) in Lebanon, and through contacting healthcare professionals from different inpatient and outpatient rehabilitation centers nationwide to randomly include Lebanese individuals with SCI. Individuals with chronic SCI (time since injury > 1 year) who were 18 years of age or older, lived in Lebanon, and were able to read and comprehend the questionnaire were considered eligible for inclusion. Participants were excluded if they had significant psychological or cognitive impairment, congenital disorders, neurodegenerative illnesses, infections, or unstable medical conditions.

**Study design and procedure:** This cross-sectional study was carried out on 81 Lebanese individuals with SCI over a period of 3 months extending from August 1<sup>st</sup> until the end of October 2022. This study was conducted according to the research ethics guidelines of the Declaration of Helsinki<sup>50</sup> and was approved by the Institutional Review Board of HRIR, Beirut, Lebanon. A

detailed explanation of the study aims and procedures were presented to the participants during a telephone call interview. Throughout the conversation, further information was provided as needed. Involvement in the study was requested with a clear indication of voluntary participation and a focus on respect for confidentiality in data collection and storage. Following their oral consent, written informed consent was obtained via an electronic system from participants who agreed to participate. Participants were invited to complete a self-report questionnaire as an online survey utilizing a link to "Google Forms" software via email or any other social media platform of their choice, or through a telephone interview. It should be highlighted that all information in the survey was gathered anonymously and treated with strict confidentiality.

**Instrument:** The questionnaire was presented with a definition of chronic pain experienced after SCI, and a description of the purpose of the study and its procedure, followed by a reassurance question on the consent to participate. The first part of the questionnaire included questions on socio-demographic and injury attributes related to age, gender, educational level, occupational and marital status, time since SCI, level of SCI (cervical, thoracic, or lumbar), and degree of impairment (tetraplegia or paraplegia). The second section comprised questions on the experience of chronic pain after SCI. Participants were asked if they have experienced and are experiencing chronic pain after an injury. The questionnaire also included questions on the characteristics of pain including its pattern, duration, triggering factors, and relieving factors. In addition to a question that asked about the use of pain medication in the past week. Evaluation of pain severity and interference with daily activities was assessed using the Arabic version of the Brief Pain Inventory (BPI).<sup>51</sup> The type of pain (neuropathic or non-neuropathic pain) experienced was concluded from the 7 self-report items of the Arabic Version of the Neuropathic Pain Questionnaire [Douleur Neuropathique en 4 (DN4)].<sup>52</sup> The third section of the questionnaire encompasses the 12-item Short Form Health Survey (SF-12), a QOL measure.<sup>53</sup>

### Measures

**Arabic version of the BPI:** We used the first screening item of the BPI which is related to the presence of pains other than everyday pains, and the BPI pain intensity and interference scores. The BPI is a self-administered questionnaire that was

initially intended to evaluate cancer pain<sup>54</sup> and is currently used as a common pain measure for different chronic pain conditions.<sup>55</sup> The BPI intensity and interference scores are regularly used in SCI pain research studies.<sup>37,56,57</sup> The BPI has been translated into Lebanese and validated among Lebanese cancer pain patients.<sup>51</sup>

The pain severity score is obtained as the sum of the scores of the 4 questions on pain intensity divided by 4. These 4 questions measure pain severity on a 0-10 numeric rating scale (NRS), with 0 denoting no pain and 10 denoting the worst possible pain, throughout the course of the previous week ((worst pain, least pain, pain at average, present).

The pain interference score is calculated as the sum of the scores of 7 items divided by 7. The 7 questions are rated on a 0-10 NRS (does not interfere-completely interfere), which inquire about the extent of pain-related impairment in general activity, mood, walking ability, normal work, relations with other people, sleep, and enjoyment of life. Greater interference is reflected by higher scores.

**Arabic version of the DN4 questionnaire:** The 7 symptom questions of the Arabic DN4 questionnaire, which include characteristics (burning, painful coldness, and electric shocks) and symptoms (tingling, pins and needles, numbness, and itching), were utilized to diagnose the presence of neuropathic pain symptoms.<sup>52</sup> These verbal descriptions are scored 1 to each positive item and 0 to each negative item (total score range 0-7). We adopted a cut-off score of 3 reflecting neuropathic pain, which had previously been shown to be accurate and reliable.<sup>58</sup> The DN4 was originally designed in French as a 10-item diagnostic test for neuropathic pain with 3 clinical questions and 7 self-reported questions. It has been translated into other languages and validated in Arabic among Lebanese populations.<sup>52</sup> Its sensitivity and specificity for identifying neuropathic pain are 70% and 67%, respectively.<sup>59</sup>

**Arabic version of the SF-12:** To measure the QOL of participants, we used the Arabic version of the SF-12. It is a health-related QOL measure derived from the Medical Outcomes Study 36-Item Short-Form Health Survey (SF-36).<sup>60</sup> The SF-12 evaluates QOL in the 8 domains of physical functioning (PF), role limitations due to physical problems (RP), bodily pain (BP), general health (GH), vitality (VT), social functioning (SF), role limitations due to emotional problems (RE), and

mental health (MH), and generates the physical component score (PCS) and mental component score (MCS). It has been used widely and extensively established as reliable and valid in different health clinical conditions.<sup>61-63</sup> An Arabic version of the SF-12 has been recently validated in the Lebanese population.<sup>53</sup>

The generated data from "Google Forms" on an excel spreadsheet were transported to the SPSS statistical software (version 26, IBM Corp., Armonk, NY, USA) for analysis. Given that the response to all questions was required, there was no missing data to substitute. Descriptive statistics were used to describe the baseline characteristics of participants. Categorical variables were presented as frequency and percentage, and mean  $\pm$  standard deviation (SD) was used to present continuous variables. To identify potential predictors of pain interference, a univariate analysis was performed for each variable using a non-parametric test. All variables in the univariate analysis with a p-value  $< 0.1$  were entered into a multiple linear regression model.

To evaluate the association of different baseline and pain characteristics with QOL, mean scores of the SF-12 domains and its subscales (PCS and MCS) were compared between groups using one-way analysis of variance (ANOVA) and independent samples t-test. Two-tailed P-values  $< 0.05$  were considered significant.

## Results

### *Baseline characteristics of individuals with SCI:*

The population included 81 individuals with SCI. Men constituted the majority of the total sample (86.4%). The mean age of the participants was  $38.15 \pm 11.75$  years with 40.7% aged between 26 and 35 years. Out of the total sample, 54.3% were married, 67.9% did not have a college degree, and 66.7% were unemployed. Regarding the injury characteristics, 59.3% had a lumbosacral injury, and 77.8% were paraplegic. Finally, concerning the pain experience characteristics, 18.5% did not experience chronic pain, and 81.5% are experiencing chronic pain with the majority of them ( $n = 48$ ) presenting a DN4 symptoms score of above 3, reflecting neuropathic pain. As for pain severity in participants with chronic pain, the average score of pain severity was  $5.48 \pm 2.27$  with 56.1% of them describing their pain as moderate and 31.8% as severe. The average score of interference of pain with daily activities was  $4.29 \pm 2.39$ . A detailed

description of the baseline characteristics of participants is presented in table 1.

**Multiple linear regression evaluating associations with pain interference:** Results of the univariate analysis illustrated 6 possible predictors of high pain interference ( $P = 0.10$ ) to be included in the multivariate analysis, age range, marital status, occupational status, SCI level, type of pain (Neuropathic/Non-neuropathic), and pain severity scores. Table 2 depicts the associations between the resulting variables from the univariate analysis and pain interference scores. The regression model showed a significant correlation between pain interference and occupational status (Wald-Chi square = 5.38;  $P = 0.02$ ), SCI level (Wald-Chi square = 5.65;  $P = 0.05$ ), type of pain (Wald-Chi square = 4.18;  $P = 0.04$ ), and pain severity (Wald-Chi square = 3.69;  $P = 0.05$ ). Participants with thoracic injuries are more prone to higher pain interference, 4.118 times (95% CI: 0.05-0.95) that in those with lumbosacral injury. Unemployed participants were 4.268 times (95% CI: 1.25-14.55) more likely to have high pain interference scores compared with employed participants. The risk of higher pain interference among individuals with neuropathic pain (DN4  $\geq 3$ ) was 3.430 times (95% CI: 1.05-11.17) higher than those with non-neuropathic pain. Moreover, participants with pain severity  $\geq 4$  were more likely to have higher pain interference, 3.699 times (95% CI: 0.97-14.05) those with pain severity scores of less than 4.

**Comparison of QOL domains in individuals with SCI according to baseline and pain characteristics:** The SF-12 domains, PCS, and MCS scores for different baseline characteristics are presented in table 3.

The mean GH domain score was significantly lower in individuals older than 35 years compared to those below 35 years of age ( $P = 0.004$ ). The mean PF domain score was significantly lower in individuals with tetraplegia than in those with paraplegia ( $P = 0.007$ ), and lower in individuals who experience neuropathic pain as compared with those with non-neuropathic pain and those who do not experience chronic pain ( $P = 0.030$ ). Moreover, it was lower in individuals with severe and moderate pain as compared with those with mild pain ( $P = 0.032$ ). The role limitation due to physical problems domain presents a statistically significant difference between different pain severity scores ( $P = 0.030$ ).

**Table 1.** Baseline characteristics of individuals with spinal cord injury (SCI)

Baseline characteristic	n (%)	Baseline characteristic	n (%)
Age range (year)		Level of injury	
18-25	9 (11.1)	Paraplegia	63 (77.8)
26-35	33 (40.7)	Quadriplegia	18 (22.2)
36-45	15 (18.5)	Locomotion	
46-55	15 (18.5)	Wheelchair dependent	61 (75.3)
56-65	9 (11.1)	Walking ability	20 (24.7)
Gender		Pain pattern	
Male	70 (86.4)	Always	23 (34.6)
Female	11 (13.6)	No pattern	28 (28.4)
Marital status		1-3 days a month	2 (2.5)
Single	37 (45.7)	1-2 days a week	6 (7.4)
Married	44 (54.3)	3-6 days a week	7 (8.6)
Educational level		No pain	15 (18.5)
< 12 years of education	55 (67.9)	Duration of pain	
> 12 years of education	26 (32.1)	Not specified	34 (42.0)
Employment status		Few days	7 (8.6)
Unemployed	54 (66.7)	Few times a day	16 (19.8)
Employed	27 (33.3)	5 min-1 hour	5 (6.2)
Time since SCI (year)		< 5 min	4 (4.9)
1-2	8 (9.9)	No pain	15 (18.5)
2-5	3 (3.7)	Pain type	
> 5	70 (86.4)	No pain	15 (18.5)
SCI level		Neuropathic pain (DN4 > 3)	48 (59.3)
Cervical	21 (25.9)	Non-neuropathic pain (DN4 < 3)	18 (22.2)
Thoracic	12 (14.8)	Pain severity	
Lumbosacral	48 (59.3)	Mild	8 (12.1)
Hospital visits to take pain medication (during the past month)		Moderate	37 (56.1)
0 hospital visits	42 (64.6)	Severe	21 (31.8)
1-2 times	19 (29.2)	Pain interference	
> 3 times	4 (6.2)	Mild	19 (28.8)
Age (mean ± SD)	38.15 ± 11.75	Moderate	34 (51.5)
Pain severity score (Pain subjects) (mean ± SD)	5.48 ± 2.27	Severe	13 (19.7)
Pain interference score (Pain subjects) (mean ± SD)	4.29 ± 2.39		
PCS (mean ± SD)	36.26 ± 21.36		
MCS (mean ± SD)	52.16 ± 22.94		

SCI: Spinal cord injury; DN4: Douleur Neuropathique en 4; PCS: Physical component score; MCS: Mental component score; SD: Standard deviation

However, the BP domain varied significantly with occupational status ( $P = 0.050$ ), presence of pain and its type ( $P = 0.010$ ), pain severity ( $P = 0.005$ ), and pain interference ( $P = 0.006$ ). A statistically significant difference was found in the role limitation due to emotional problems domain depending on the educational level ( $P = 0.019$ ), and in the SF domain according to the type of paralysis ( $P = 0.050$ ). Mean PCS scores were significantly higher in employed individuals, those not experiencing pain or with non-neuropathic pain type, and those with mild pain severity and interference compared to their counterparts. The mean score of MCS differed only between employed and non-employed individuals ( $P = 0.041$ ).

## Discussion

The present study examined a sample of Lebanese individuals with SCI in terms of their experiences with chronic pain and factors that contribute to pain interference, and assessed the impact of several factors, most notably pain characteristics, on their QOL. This study demonstrated that 81% of the participants experienced chronic pain, and more than 70% of them experienced neuropathic pain symptoms. These results are consistent with the prevalence rate of any kind of pain after SCI reported in earlier studies,<sup>38,64</sup> and the high proportion is justified by the time since the injury being more than 5 years in most individuals, as previous studies have reported that a longer time since the injury presented high pain prevalence.<sup>65</sup>

**Table 2.** Factors associated with pain interference

	Factors	OR	95% CI	Wald-Chi square	P	Effect size (required sample size)
Dependent variable: Pain interference	Age (year)			0.98	0.32	0.12 (68)
	< 35	0.55	0.17-1.77			
	> 35		Reference			
	Marital status			1.13	0.28	0.15 (55)
	Single	0.49	0.13-1.82			
	Married		Reference			
	SCI level			5.65	0.06	0.25 (34)
	Cervical	0.30	0.08-1.12			
	Thoracic	0.21	0.04-0.94			
	Lumbar		Reference			
	Occupational status			5.38	0.02	0.26 (33)
	Unemployed	4.26*	1.25-14.58			
	Employed		Reference			
	Type of pain			4.18	0.04	0.22 (39)
	Neuropathic	3.43*	1.05-11.17			
Non-neuropathic		Reference				
Pain severity			3.69	0.05	0.25 (34)	
> 4	3.69*	0.97-14.04				
< 4		Reference				

SCI: Spinal cord injury; OR: Odds ratio; CI: Confidence interval

\*P-value < 0.05 was considered significant.

The required sample size was estimated assuming 80% of statistical power.

Thus, most of the individuals experiencing pain also had pain location below the level of the SCI, which is similar to the finding that neuropathic pain location below the level of SCI develops gradually over months to years after an injury.<sup>66</sup> Overall, the study sample presented an average pain intensity of 5.48 and pain interference of 4.29, reflecting moderate ranges according to pain severity and interference categories.<sup>67</sup> These levels are comparable to those reported in the literature in chronic SCI participants.<sup>28,64,68</sup>

Demographics and injury-related attributes proved not to be associated with pain interference ratings; this is consistent with the findings of prior studies.<sup>28,43</sup> Our study highlighted a significant association with employment status, type of pain, and pain intensity levels. The results showed that unemployed individuals with SCI are more susceptible to higher pain interference when compared to employed individuals; these findings are in line with those of previous studies.<sup>28,69,70</sup> In addition, the magnitude of higher levels of pain interference is predicted by pain intensity and type of pain. In terms of pain intensity, our findings are consistent with previous research outlining its relationship with interference, as well as the role of pain intensity in mediating the interference with activities.<sup>44</sup> Moreover, we found that the participants with neuropathic pain were more

likely to experience higher levels of interference with daily activities, which is in accordance with a recent study by Felix et al. that compared pain interference between neuropathic pain and non-neuropathic pain.<sup>64</sup>

The findings of the present study revealed different determinants of QOL in individuals with SCI. In terms of sociodemographic characteristics, individuals over the age of 35 years had lower levels of GH than younger individuals, which is in line with previous findings showing that age is not closely associated with global QOL.<sup>71</sup> Moreover, unemployed individuals exhibited lower levels of QOL in the BP domain as well as in the PCS and MCS subscales which is in line with the results of various studies that have shown a significant association between employment status and QOL.<sup>72,73</sup>

Individuals who had not completed a university education had lower scores on the RE domain, which is supported by the findings of Halvorsen et al. indicating the significant impact of education on QOL and social participation,<sup>20</sup> as well as the proven relationship between educational level and different emotional problems and the MH component of QOL.<sup>74,75</sup> Furthermore, we found that individuals with tetraplegia had lower scores on the PF and SF domains than individuals with paraplegia in terms of injury characteristics.

**Table 3.** Comparison of quality of life (QOL) domains for individuals with spinal cord injury (SCI) (according to baseline and pain characteristics)

Variable	GH	PF	RP	BP	VT	RE	MH	SF	PCS	MCS
Age (year)										
< 35	48.81 ± 28.12	27.98 ± 28.79	20.24 ± 38.34	63.69 ± 31.81	57.74 ± 24.37	40.48 ± 48.43	63.99 ± 25.92	51.19 ± 34.01	40.17 ± 21.34	53.34 ± 19.72
> 35	31.41 ± 24.81	32.05 ± 29.77	11.54 ± 31.33	53.21 ± 34.01	52.56 ± 0.23	46.15 ± 49.18	55.45 ± 27.92	49.36 ± 29.51	32.05 ± 20.83	50.88 ± 26.17
P	0.004	0.530	0.260	0.150	0.390	0.620	0.150	0.790	0.080	0.630
Educational level										
< 12 years of education	42.27 ± 27.58	31.82 ± 31.71	14.55 ± 34.25	55.00 ± 33.47	55.00 ± 27.38	34.55 ± 47.01	58.86 ± 26.96	53.18 ± 31.56	35.90 ± 21.32	50.39 ± 22.07
> 12 years of education	36.54 ± 28.48	25.96 ± 22.89	19.23 ± 37.62	66.35 ± 31.57	55.77 ± 27.66	61.54 ± 47.55	62.02 ± 27.72	44.23 ± 31.86	37.01 ± 21.86	55.88 ± 24.70
P	0.390	0.400	0.570	0.150	0.900	0.010	0.620	0.230	0.280	0.820
Occupational status										
Unemployed	37.50 ± 26.04	28.24 ± 30.73	12.96 ± 32.48	68.52 ± 28.24	52.31 ± 28.01	37.96 ± 47.51	56.94 ± 26.67	46.76 ± 57.41	33.10 ± 19.52	48.49 ± 22.45
Employed	46.30 ± 30.77	33.33 ± 25.94	22.22 ± 40.03	53.70 ± 34.48	61.11 ± 25.31	53.70 ± 49.85	65.74 ± 27.42	57.41 ± 29.26	42.59 ± 23.77	59.49 ± 22.52
P	0.180	0.460	0.260	0.050	0.170	0.170	0.170	0.150	0.050	0.041
Paralysis										
Paraplegic	41.27 ± 28.45	34.52 ± 26.72	15.87 ± 34.577	56.75 ± 33.05	57.94 ± 27.97	38.10 ± 47.27	60.91 ± 28.44	53.97 ± 32.13	37.10 ± 21.53	52.72 ± 23.66
Quadriplegic	37.50 ± 26.080	13.89 ± 32.33	16.67 ± 38.348	65.28 ± 33.36	45.83 ± 23.08	61.11 ± 50.16	56.25 ± 21.96	37.50 ± 27.45	33.33 ± 21.11	50.17 ± 20.68
P	0.610	0.007	0.930	0.330	0.090	0.070	0.500	0.050	0.510	0.680
Pain type										
No pain	36.67 ± 22.88	39.18 ± 10.11	30.00 ± 45.51	80.00 ± 30.17	45.00 ± 33.00	50.00 ± 50.00	54.17 ± 31.57	51.67 ± 29.07	22.24 ± 5.74	50.20 ± 26.76
Non-neuropathic pain	34.72 ± 25.92	33.36 ± 7.86	22.22 ± 39.19	61.11 ± 29.97	66.67 ± 25.72	50.00 ± 48.50	64.58 ± 23.96	41.67 ± 24.25	21.10 ± 4.97	55.72 ± 21.25
Neuropathic pain	43.75 ± 29.84	21.51 ± 3.10	9.38 ± 28.53	51.04 ± 32.59	54.17 ± 24.91	38.54 ± 48.64	59.90 ± 26.91	53.13 ± 34.83	20.10 ± 2.90	51.43 ± 22.63
P	0.430	0.030	0.090	0.010	0.060	0.580	0.550	0.420	0.035	0.740
Pain severity										
Mild	46.74 ± 28.50	42.39 ± 38.01	28.26 ± 44.78	71.74 ± 34.79	53.26 ± 31.35	45.65 ± 49.80	56.52 ± 28.17	50.00 ± 32.85	47.28 ± 24.33	51.36 ± 23.43
Moderate	41.22 ± 28.39	29.76 ± 30.22	23.81 ± 40.67	60.81 ± 24.67	59.46 ± 23.82	37.84 ± 49.16	65.20 ± 23.59	53.38 ± 32.89	32.09 ± 14.14	53.97 ± 22.67
Severe	32.14 ± 25.17	22.30 ± 18.43	4.05 ± 18.17	40.48 ± 37.48	50.00 ± 28.50	50.00 ± 47.43	54.17 ± 30.95	45.24 ± 29.17	31.54 ± 24.88	49.85 ± 23.72
P	0.210	0.030	0.030	0.005	0.410	0.630	0.260	0.640	0.010	0.790
Pain interference										
Mild	47.50 ± 28.12	30.83 ± 24.28	21.67 ± 40.86	69.17 ± 31.95	50.83 ± 28.22	53.33 ± 49.01	60.83 ± 30.39	56.67 ± 37.67	42.29 ± 22.18	55.41 ± 26.79
Moderate	36.84 ± 25.16	34.87 ± 32.64	17.11 ± 35.40	58.55 ± 31.44	57.24 ± 23.90	36.84 ± 47.48	59.87 ± 25.02	48.03 ± 26.24	36.84 ± 20.99	50.49 ± 18.34
Severe	34.62 ± 33.13	13.46 ± 24.19	00.00 ± 00.00	34.62 ± 29.82	59.62 ± 34.66	38.46 ± 50.63	57.69 ± 26.78	42.31 ± 31.26	20.67 ± 11.79	49.51 ± 26.32
P	0.210	0.070	0.170	0.006	0.520	0.350	0.940	0.330	0.008	0.610

GH: General health; PF: Physical functioning; RP: Role limitations due to physical problems; BP: Bodily pain; VT: Vitality; SF: Social functioning; RE: Role limitations due to emotional problems; MH: Mental health; PCS: Physical component score; MCS: Mental component score

This finding is consistent with the findings of several research studies that attribute the difference to higher levels of dependency in more severe injuries.<sup>49,76,77</sup>

Regarding the experience of chronic pain and its effect on QOL, the current study emphasizes that individuals with higher pain intensity have lower levels of QOL in the PF, RP, and BP domains, and the PCS subscale. These findings comply with the research by Ataoglu et al. demonstrating lower levels of QOL in all domains in individuals with higher pain intensity.<sup>78</sup> The specificity of the relationship between pain intensity and the physical dimensions of QOL reflects the demonstrated association between pain intensity and physical function in people living with pain.<sup>79,80</sup> Similarly for pain interference, higher levels are associated with lower BP domain and PCS subscale scores, which is in accordance with numerous studies that have reported the impact of pain on PF,<sup>25,81,82</sup> as well as studies that have reported that pain interference is more significant for the conceptualization of the link between pain and QOL.<sup>83</sup>

The current study found that having chronic pain reduced the BP, PF, and PCS subscale scores of the SF-12. In particular, neuropathic pain has been demonstrated to exacerbate the decline in QOL levels in SCI. These results are consistent with recent studies that have demonstrated the highest reductions in QOL among individuals with SCI reporting neuropathic pain (NP).<sup>46,84</sup>

To the best of our knowledge, this is the first study to evaluate the chronic pain experience of Lebanese individuals with SCI. This cross-sectional study assessed the impact of neuropathic pain type on QOL, since the majority of previous research has examined pain in general, without distinguishing between types. Thus, this study emphasized the significance of the employment of individuals with SCI on the improvement of their chronic pain experience, pain interference, and QOL. The findings of this study will add to the literature concerning the Lebanese individuals with SCI on the Lebanese SCI population since data targeting this population remains scarce in Lebanon.<sup>85</sup> Furthermore, since the application of legislative actions taken by the Lebanese government such as regulations for schools, workplaces, and government programs remains lacking<sup>86</sup> and 80% of Lebanese people with disabilities stay unemployed,<sup>87</sup> this study can help

to support the importance of including people with disabilities, particularly individuals with SCI, in the labor force in order to enhance their QOL.

Our study does have certain limitations. Although several variables evaluating pain experience were gathered, it is possible that some important variables such as coping and pain adaptation, along with the perceived effect of adopted pain management procedures, were underestimated. It is also worth noting that the sample size is inadequately similar to SCI-pain studies which might be related to the lack of population-based records and registers on individuals with physical abilities.<sup>88</sup> The findings of this study were based on self-reported data, thus increasing the possibility that the responses were influenced by personal and psychological factors since the psychological state of participants was not evaluated.

The current study raises intriguing questions and suggests possible future directions. Hence, further research with a greater sample size is required, with an emphasis on understanding the effective pain-relieving methods utilized by Lebanese individuals with SCI.

## Conclusion

A high proportion of Lebanese individuals with SCI experience chronic pain. This study highlights the substantial and negative influence of chronic pain, particularly neuropathic pain, on QOL. Pain interference is a better effective method to understand the experience of pain, and employment status is the main factor affecting interference and QOL. It is critical to broaden the current study's findings in order to promote QOL by warranting chronic pain management and incorporating employment and social reintegration as primary rehabilitation goals for individuals with SCI in Lebanon.

## Conflict of Interests

The authors declare no conflict of interest in this study.

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## References

- Bennett J, Emmady PD. Spinal cord injuries. In: StatPearls [Internet]. Treasure Island, FL: StatPearls Publishing; 2022.
- Hamid R, Averbeck MA, Chiang H, Garcia A, Al Mousa RT, Oh SJ, et al. Epidemiology and pathophysiology of neurogenic bladder after spinal cord injury. *World J Urol* 2018; 36(10): 1517-27.
- Tate DG, Wheeler T, Lane GI, Forchheimer M, Anderson KD, Biering-Sorensen F, et al. Recommendations for evaluation of neurogenic bladder and bowel dysfunction after spinal cord injury and/or disease. *J Spinal Cord Med* 2020; 43(2): 141-64.
- Wecht JM, Harel NY, Guest J, Kirshblum SC, Forrest GF, Bloom O, et al. Cardiovascular autonomic dysfunction in spinal cord injury: Epidemiology, diagnosis, and management. *Semin Neurol* 2020; 40(5): 550-9.
- Courtois F, Charvier K. Sexual dysfunction in patients with spinal cord lesions. *Handb Clin Neurol* 2015; 130: 225-45.
- Tamburin S, Filippetti M, Mantovani E, Smania N, Picelli A. Spasticity following brain and spinal cord injury: Assessment and treatment. *Curr Opin Neurol* 2022; 35(6): 728-40.
- Hunt C, Moman R, Peterson A, Wilson R, Covington S, Mustafa R, et al. Prevalence of chronic pain after spinal cord injury: A systematic review and meta-analysis. *Reg Anesth Pain Med* 2021; 46(4): 328-36.
- Perrouin-Verbe B, Lefevre C, Kieny P, Gross R, Reiss B, Le FM. Spinal cord injury: A multisystem physiological impairment/dysfunction. *Rev Neurol (Paris)* 2021; 177(5): 594-605.
- Andresen SR, Biering-Sorensen F, Hagen EM, Nielsen JF, Bach FW, Finnerup NB. Pain, spasticity and quality of life in individuals with traumatic spinal cord injury in Denmark. *Spinal Cord* 2016; 54(11): 973-9.
- Tulsky DS, Kisala PA, Victorson D, Tate DG, Heinemann AW, Charlifue S, et al. Overview of the Spinal Cord Injury-Quality of Life (SCI-QOL) measurement system. *J Spinal Cord Med* 2015; 38(3): 257-69.
- Kreuter M, Siosteen A, Erholm B, Bystrom U, Brown DJ. Health and quality of life of persons with spinal cord lesion in Australia and Sweden. *Spinal Cord* 2005; 43(2): 123-9.
- Post MW. Definitions of quality of life: What has happened and how to move on. *Top Spinal Cord Inj Rehabil* 2014; 20(3): 167-80.
- Trgovcevic S, Milicevic M, Nedovic G, Jovanic G. Health condition and quality of life in persons with spinal cord injury. *Iran J Public Health* 2014; 43(9): 1229-38.
- Yazdanshenas Ghazwin M, Chaibakhsh S, Latifi S, Tavakoli A H, Koushki D. Quality of life in Iranian men with spinal cord injury in comparison with general population. *Arch Neurosci* 2015; 2(2): e21529.
- Lude P, Kennedy P, Elfstrom ML, Ballert CS. Quality of life in and after spinal cord injury rehabilitation: A longitudinal multicenter study. *Top Spinal Cord Inj Rehabil* 2014; 20(3): 197-207.
- Duan R, Qu M, Yuan Y, Lin M, Liu T, Huang W, et al. Clinical benefit of rehabilitation training in spinal cord injury: A systematic review and meta-analysis. *Spine (Phila Pa 1976)* 2021; 46(6): E398-E410.
- Quel de Oliveira C, Refshauge K, Middleton J, de Jong L, Davis GM. Effects of activity-based therapy interventions on mobility, independence, and quality of life for people with spinal cord injuries: A systematic review and meta-analysis. *J Neurotrauma* 2017; 34(9): 1726-43.
- Sturm C, Gutenbrunner CM, Egen C, Geng V, Lemhofer C, Kalke YB, et al. Which factors have an association to the Quality of Life (QoL) of people with acquired Spinal Cord Injury (SCI)? A cross-sectional explorative observational study. *Spinal Cord* 2021; 59(8): 925-32.
- Lee JS, Kim SW, Jee SH, Kim JC, Choi JB, Cho SY, et al. Factors affecting quality of life among spinal cord injury patients in Korea. *Int Neurolog J* 2016; 20(4): 316-20.
- Halvorsen A, Pape K, Post MWM, Biering-Sorensen F, Mikalsen S, Hansen AN, et al. Participation and quality of life in persons living with spinal cord injury in Norway. *J Rehabil Med* 2021; 53(7): jrm00217.
- Zwecker M, Heled E, Bluvstein V, Catz A, Bloch A, Zeilig G. Assessment of the unmediated relationship between neurological impairment and health-related quality of life following spinal cord injury. *J Spinal Cord Med* 2022; 45(2): 293-300.
- Ebrahimzadeh MH, Soltani-Moghaddas SH, Birjandinejad A, Omidi-Kashani F, Bozorgnia S. Quality of life among veterans with chronic spinal cord injury and related variables. *Arch Trauma Res* 2014; 3(2): e17917.
- Tavakoli SA, Kaviani M, Bakhsh SC, Ghajarzadeh M, Hamedan MS, Ghazwin MY, et al. Is Level of injury a determinant of quality of life among individuals with spinal cord injury? A Tertiary Rehabilitation Center Report. *Oman Med J* 2016; 31(2): 112-6.
- Norrbrink BC, Hultling C, Lundeberg T. Quality of sleep in individuals with spinal cord injury: A comparison between patients with and without pain. *Spinal Cord* 2005; 43(2): 85-95.
- Nicholson PK, Nicholas MK, Middleton J. Spinal cord injury-related pain in rehabilitation: A cross-sectional study of relationships with cognitions, mood and physical function. *Eur J Pain* 2009; 13(5): 511-7.
- Craig A, Tran Y, Siddall P, Wijesuriya N, Lovas J, Bartrop R, et al. Developing a model of associations between chronic pain, depressive mood, chronic fatigue, and self-efficacy in people with spinal cord injury. *J Pain* 2013; 14(9): 911-20.
- Taran S, Conti J, Routhier F, Latimer-Cheung AE, Noreau L, Sweet SN. Leisure time physical activity, perception of impact of pain and life satisfaction after spinal cord injury. *Ann Phys Rehabil Med* 2018; 61(4): 273-5.
- Ullrich PM, Jensen MP, Loeser JD, Cardenas DD. Pain intensity, pain interference and characteristics of spinal cord injury. *Spinal Cord* 2008; 46(6): 451-5.
- Widerstrom-Noga EG, Finnerup NB, Siddall PJ. Biopsychosocial perspective on a mechanisms-based approach to assessment and treatment of pain following spinal cord injury. *J Rehabil Res Dev* 2009; 46(1): 1-12.
- Masri R, Keller A. Chronic pain following spinal cord injury. *Adv Exp Med Biol* 2012; 760: 74-88.
- Scholz J, Finnerup NB, Attal N, Aziz Q, Baron R, Bennett MI, et al. The IASP classification of chronic pain for ICD-11: Chronic neuropathic pain. *Pain* 2019; 160(1): 53-9.
- Burke D, Fullen BM, Stokes D, Lennon O. Neuropathic pain prevalence following spinal cord injury: A systematic review and meta-analysis. *Eur J Pain* 2017; 21(1): 29-44.
- Jensen MP, Hoffman AJ, Cardenas DD. Chronic pain in individuals with spinal cord injury: A survey and longitudinal study. *Spinal Cord* 2005; 43(12): 704-12.
- Rintala DH, Hart KA, Priebe MM. Predicting consistency of pain over a 10-year period in persons with spinal cord injury. *J Rehabil Res Dev* 2004; 41(1): 75-88.
- Norrbrink BC, Lund I, Hultling C, Levi R, Werhagen L, Ertzgaard P, et al. Gender related differences in pain in spinal cord injured individuals. *Spinal Cord* 2003; 41(2): 122-8.
- Nicholson PK, Nicholas MK, Middleton J, Siddall P. Psychological characteristics of people with spinal cord injury-related persisting pain referred to a tertiary pain management center. *J Rehabil Res Dev* 2009; 46(1): 57-67.
- Khazaeipour Z, Ahmadipour E, Rahimi-Movaghar V, Ahmadipour F, Vaccaro AR, Babakhani B. Association of pain, social support and socioeconomic indicators in patients with spinal cord injury in Iran. *Spinal Cord* 2017; 55(2): 180-6.
- Muller R, Brinkhoff MW, Arnet U, Hinrichs T, Landmann G, Jordan X, et al. Prevalence and associated factors of pain in the Swiss spinal cord injury population. *Spinal Cord* 2017; 55(4): 346-54.
- Rekand T, Hagen EM, Gronning M. Chronic pain following spinal cord injury.

- Tidsskr Nor Laegeforen 2012; 132(8): 974-9.
40. Jang JY, Lee SH, Kim M, Ryu JS. Characteristics of neuropathic pain in patients with spinal cord injury. *Ann Rehabil Med* 2014; 38(3): 327-34.
  41. Turner JA, Cardenas DD, Warms CA, McClellan CB. Chronic pain associated with spinal cord injuries: A community survey. *Arch Phys Med Rehabil* 2001; 82(4): 501-9.
  42. Hagen EM, Rekan T. Management of neuropathic pain associated with spinal cord injury. *Pain Ther* 2015; 4(1): 51-65.
  43. Dijkers M, Bryce T, Zanca J. Prevalence of chronic pain after traumatic spinal cord injury: A systematic review. *J Rehabil Res Dev* 2009; 46(1): 13-29.
  44. Li C, DiPiro ND, Clark JMR, Krause JS. Mediating effects of pain interference on the relationships between pain intensity and probable major depression among participants with spinal cord injury. *Arch Phys Med Rehabil* 2022; 103(4): 747-54.
  45. Finnerup NB. Pain in patients with spinal cord injury. *Pain* 2013; 154(Suppl 1): S71-S76.
  46. Burke D, Lennon O, Fullen BM. Quality of life after spinal cord injury: The impact of pain. *Eur J Pain* 2018; 22(9): 1662-72.
  47. Nagoshi N, Kaneko S, Fujiyoshi K, Takemitsu M, Yagi M, Iizuka S, et al. Characteristics of neuropathic pain and its relationship with quality of life in 72 patients with spinal cord injury. *Spinal Cord* 2016; 54(9): 656-61.
  48. Ord AS, Lad SS, Shura RD, Rowland JA, Taber KH, Martindale SL. Pain interference and quality of life in combat veterans: Examining the roles of posttraumatic stress disorder, traumatic brain injury, and sleep quality. *Rehabil Psychol* 2021; 66(1): 31-8.
  49. Nasidi MA, Akindele MO, Ibrahim AA, Ahmad AA, Musa A. Health-related quality of life and related characteristics of persons with spinal cord injury in Nigeria. *Iran J Neurol* 2019; 18(2): 50-6.
  50. Williams JR. The Declaration of Helsinki and public health. *Bull World Health Organ* 2008; 86(8): 650-2.
  51. Ballout S, Nouredine S, Huijjer HA, Kanazi G. Psychometric evaluation of the arabic brief pain inventory in a sample of Lebanese cancer patients. *J Pain Symptom Manage* 2011; 42(1): 147-54.
  52. Chatila N, Pereira B, Maarrawi J, Dallel R. Validation of a new Arabic version of the Neuropathic Pain Diagnostic Questionnaire (DN4). *Pain Pract* 2017; 17(1): 78-87.
  53. Haddad C, Sacre H, Obeid S, Salameh P, Hallit S. Validation of the Arabic version of the "12-item short-form health survey" (SF-12) in a sample of Lebanese adults. *Arch Public Health* 2021; 79(1): 56.
  54. Cleeland CS, Ryan KM. Pain assessment: Global use of the Brief Pain Inventory. *Ann Acad Med Singap* 1994; 23(2): 129-38.
  55. Keller S, Bann CM, Dodd SL, Schein J, Mendoza TR, Cleeland CS. Validity of the brief pain inventory for use in documenting the outcomes of patients with noncancer pain. *Clin J Pain* 2004; 20(5): 309-18.
  56. Finnerup NB, Jensen MP, Norrbrink C, Trok K, Johannesen IL, Jensen TS, et al. A prospective study of pain and psychological functioning following traumatic spinal cord injury. *Spinal Cord* 2016; 54(10): 816-21.
  57. Hand BN, Velozo CA, Krause JS. Measuring the interference of pain on daily life in persons with spinal cord injury: A Rasch-validated subset of items from the Brief Pain Inventory interference scale. *Aust Occup Ther J* 2018; 65(5): 405-11.
  58. Bouhassira D, Attal N, Alchaar H, Boureau F, Brochet B, Bruxelle J, et al. Comparison of pain syndromes associated with nervous or somatic lesions and development of a new neuropathic pain diagnostic questionnaire (DN4). *Pain* 2005; 114(1-2): 29-36.
  59. Timmerman H, Steegers MAH, Huygen FJPM, Goeman JJ, van Dasselaaar NT, Schenkels MJ, et al. Investigating the validity of the DN4 in a consecutive population of patients with chronic pain. *PLoS One* 2017; 12(11): e0187961.
  60. Ware J, Kosinski M, Keller SD. A 12-Item Short-Form Health Survey: construction of scales and preliminary tests of reliability and validity. *Med Care* 1996; 34(3): 220-33.
  61. Gandek B, Ware JE, Aaronson NK, Apolone G, Bjorner JB, Brazier JE, et al. Cross-validation of item selection and scoring for the SF-12 Health Survey in nine countries: Results from the IQOLA Project. *International Quality of Life Assessment. J Clin Epidemiol* 1998; 51(11): 1171-8.
  62. Haggell P, Westergren A. Measurement properties of the SF-12 health survey in Parkinson's disease. *J Parkinsons Dis* 2011; 1(2): 185-96.
  63. Arovah NI, Heesch KC. Assessment of the validity and reliability of the Indonesian version of Short Form 12 (SF-12). *J Prev Med Hyg* 2021; 62(2): E421-9.
  64. Felix ER, Cardenas DD, Bryce TN, Charlifue S, Lee TK, MacIntyre B, et al. Prevalence and impact of neuropathic and nonneuropathic pain in chronic spinal cord injury. *Arch Phys Med Rehabil* 2022; 103(4): 729-37.
  65. van Gorp S, Kessels AG, Joosten EA, van Kleef M, Patijn J. Pain prevalence and its determinants after spinal cord injury: A systematic review. *Eur J Pain* 2015; 19(1): 5-14.
  66. Siddall PJ, McClelland JM, Rutkowski SB, Cousins MJ. A longitudinal study of the prevalence and characteristics of pain in the first 5 years following spinal cord injury. *Pain* 2003; 103(3): 249-57.
  67. Jensen MP, Tome-Pires C, de la V, Galan S, Sole E, Miro J. What determines whether a pain is rated as mild, moderate, or severe? the importance of pain beliefs and pain interference. *Clin J Pain* 2017; 33(5): 414-21.
  68. Burke D, Fullen BM, Lennon O. Pain profiles in a community dwelling population following spinal cord injury: a national survey. *J Spinal Cord Med* 2019; 42(2): 201-11.
  69. Mugoya GC, Hooper LM, Tomek S, George DS, Bolland A, Ufomadu J, et al. The interrelationships among pain interference, depressive symptoms, loneliness, and employment status: A moderated mediation study. *Clin Rehabil* 2018; 32(7): 967-79.
  70. Teasell RW, Bombardier C. Employment-related factors in chronic pain and chronic pain disability. *Clin J Pain* 2001; 17(4 Suppl): S39-S45.
  71. Hu Y, Mak JN, Wong YW, Leong JC, Luk KD. Quality of life of traumatic spinal cord injured patients in Hong Kong. *J Rehabil Med* 2008; 40(2): 126-31.
  72. Moghimian M, Kashani F, Cheraghi MA, Mohammadnejad E. Quality of Life and Related Factors Among People With Spinal Cord Injuries in Tehran, Iran. *Arch Trauma Res* 2015; 4(3): e19280.
  73. Geyh S, Ballert C, Sinnott A, Charlifue S, Catz A, D'Andrea Greve JM, et al. Quality of life after spinal cord injury: A comparison across six countries. *Spinal Cord* 2013; 51(4): 322-6.
  74. Khazaeipour Z, Taheri-Otaghsara SM, Naghdi M. Depression following spinal cord injury: Its relationship to demographic and socioeconomic indicators. *Top Spinal Cord Inj Rehabil* 2015; 21(2): 149-55.
  75. Saadat S, Javadi M, Divshali BS, Tavakoli AH, Ghodsi SM, Montazeri A, et al. Health-related quality of life among individuals with long-standing spinal cord injury: A comparative study of veterans and non-veterans. *BMC Public Health* 2010; 10: 6.
  76. Sabour H, Soltani Z, Latifi S, Norouzi-Javidan A, Arman F, Emami-Razavi SH, et al. Injury-related characteristics and quality-of-life among Iranian individuals with spinal cord injury. *Iran J Neurol* 2015; 14(3): 136-41.
  77. Gurcay E, Bal A, Eksioglu E, Cakci A. Quality of life in patients with spinal cord injury. *Int J Rehabil Res* 2010; 33(4): 356-8.
  78. Ataoglu E, Tiftik T, Kara M, Tunc H, Ersoz M, Akkus S. Effects of chronic pain on quality of life and depression in patients with spinal cord injury. *Spinal Cord* 2013; 51(1): 23-6.
  79. Karayannis NV, Sturgeon JA, Chih-Kao M, Cooley C, Mackey SC. Pain interference and physical function demonstrate poor longitudinal association in people living with pain: A PROMIS investigation. *Pain* 2017; 158(6): 1063-8.
  80. Suso-Ribera C, Camacho-Guerrero L, Osma J, Suso-Vergara S, Gallardo-Pujol D. A Reduction in pain intensity is more strongly associated with improved physical functioning in frustration tolerant individuals: A longitudinal moderation study in chronic pain patients. *Front Psychol* 2019; 10: 907.
  81. Alschuler KN, Jensen MP, Sullivan-Singh

- SJ, Borson S, Smith AE, Molton IR. The association of age, pain, and fatigue with physical functioning and depressive symptoms in persons with spinal cord injury. *J Spinal Cord Med* 2013; 36(5): 483-91.
82. Modirian E, Pirouzi P, Soroush M, Karbalaeei-Esmaeili S, Shojaei H, Zamani H. Chronic pain after spinal cord injury: Results of a long-term study. *Pain Med* 2010; 11(7): 1037-43.
83. Cuff L, Fann JR, Bombardier CH, Graves DE, Kalpakjian CZ. Depression, pain intensity, and interference in acute spinal cord injury. *Top Spinal Cord Inj Rehabil* 2014; 20(1): 32-9.
84. Richardson EJ, Brooks LG, Richards JS, Bombardier CH, Barber J, Tate D, et al. Changes in pain and quality of life in depressed individuals with spinal cord injury: Does type of pain matter? *J Spinal Cord Med* 2016; 39(5): 535-43.
85. Sunna T, Elias E, Summaka M, Zein H, Elias C, Nasser Z. Quality of life among men with spinal cord injury in Lebanon: A case control study. *NeuroRehabilitation* 2019; 45(4): 547-53.
86. International Labour Organization. Emerging good practices related to training and job placement of persons with disabilities in Lebanon. Beirut, Lebanon: International Labour Organization, Regional Office for Arab States; 2013.
87. Combaz E. Situation of persons with disabilities in Lebanon. Brighton, UK: Institute of Development Studies; 2018.
88. Summaka M, Zein H, Naim I, Fneish S. Assessing the psychological impact of COVID-19 outbreak and its related factors on Lebanese individuals with physical disabilities. *Disabil Health J* 2021; 14(3): 101073.