Education and Rehabilitation of Adult Persons with Disabilities

Thematic Collection of International Importance

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THE IMPACT OF EDUCATION AND GENDER ON THE QUALITY OF LIFE IN PERSONS WITH CEREBRAL PALSY

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SUMMARY

The concept of quality of life for people with disabilities includes all aspects of life, intrapersonal, interpersonal and extrapersonal, which are related very closely. Estimation of quality of life of persons with disabilities is a prerequisite and first step in the strategy of improving the quality of life for this population.

The aim of this study is to assess the quality of life in persons with cerebral palsy in relation to gender and educational structure. The study sample consisted of 100 people with cerebral palsy, of both sexes, aged 23 to 64 year. The questionnaire WHOQOL – BREF was used for the study.

The results show that people with cerebral palsy evaluate highly their quality of life. However, the results also show that there is statistically significant difference between male and female in the two studied variables (p = 0.036, p = 0.003) in favor of the male respondents, while in the group of respondents in relation to the level of education recorded a statistically significant difference in the three variables (p = 0.011, p = 0.009, p = 0.042) in favor of those with higher education.

It is possible that these results are in favor of subjectivity, experience and aspirations of the respondents, but also because of the restrictions in this study.

Key words: adults; education; gender; persons with disabilities; satisfaction with life

INTRODUCTION

Quality of life is a very complex term that includes a significant number of subjective and objective factors meeting the needs and self-perception of the individual. The concept of quality of life was first mentioned in 1920 (Coimbra, 1972, according to Torres et al., 2013), while the first documented use of the term in the medical literature we encounter 40 years ago in the field of transplantation medicine (Trgovčević, 2013). Depending on the needs of the profession, but also by the author, the concept of quality of life is defined in different ways. WHO (The World Health Organization – WHO) by Gojčeta et al., (2008) in the "Quality of Life / Quality of Life assessment (WHOQOL): position paper from the World Health Organization" in 1995, defines quality of life as well as the perception of the life of every individual in the context of culture and value structure to which it belongs in relation to personal goals, expectations and standards. Later, the current definitions are supplemented by a new defined quality of life as a state of complete physical, mental and social well-being, which is a multidimensional

concept and includes all of the physical and psychosocial aspects (WHO, 1998). Also, World Health Organization, dependence in daily activities, restriction of social participation and quality of life is seen as multidimensional concepts that influence each other (Karaduman et al., 2010). Bowling (1994) suggests that quality of life is defined as the optimal level of mental, physical, occupational and social functioning, including relationships with the environment, and the relationship with the perception of their own health, physical fitness, life satisfaction and well-being.

Some authors believe that the quality of life cannot be precisely defined, and generally choose studying different aspects and dimensions of quality of life.

Quality of life can be seen in two aspects, the subjective and objective aspects of quality of life. Today, there is some consensus about to combine objective and subjective aspects of quality of life, based on knowledge of the benefits and quality of each of them (Delhey et al., 2001). The subjective quality of life, where personal experience or perception of one's own life is still the main criteria is the most valid for most researchers of quality of life (Cummins, 1998). Quality of life is based on several indicators:

- 1. Medical indicators (state of organs and organ systems, functional disorders, the ability to work),
- 2. Psychological indicators (emotional status in terms of attitude to life, personal and general feelings and attitude towards ourselves and others),
- 3. Social indicators (financial status in general, employment, working conditions, living conditions, opportunities adequate nutrition, medical treatment, education, cultural needs and values) (Heller, 1978; according to Macanović, 2010).

Most people are watching the quality of life through health and over time there was a need to define the quality of life related to health. The concept of quality of life including those related to health (Health-Related Quality of Life – HRQoL), and subjective well-being (SWB-subjective well-being). HRQoL describes some difficulties due to poor health in relation to physical and mental functioning, participation in areas of life, but also the "health status". SWB includes total life satisfaction, satisfaction with life achievements, as well as the positive and negative impact on quality of life. It is believed that life satisfaction is in relation to the subjective perspective, and that quality of life should be viewed through a personal assessment of the individual and in relation to the personal aspirations and achievements (Trgovčević et al., 2011).

 $HRQoL\ is\ a\ multidimensional\ concept, which\ should\ include\ four\ basic\ dimensions:$

- 1. Physical functioning, which includes the individual: self-care (feeding, dressing), physical activity (walking, running, climbing stairs), social activities (work, home work, education),
- 2. Physical symptom associated with disease or treatment (pain, physiological needs),
- 3. Psychological functioning which includes emotional state and cognitive functioning,
- 4. Social functioning related to activities and socializing with friends, relatives and others (Testa & Simonson, 1996).

The concept of quality of life for people with disabilities includes all aspects of life, intrapersonal, interpersonal and extrapersonal, which are related very closely (Cella, 1992; by Nedovic et al., 2013). According to Jovanović (2011) the presence of physical

disability at a person can significantly affect its psychological, physical and social development. Assessing the impact of physical disability on daily life reflects what is functioning and well-being of the person from day to day and in different walks of life, which is actually a quality of life assessment. Estimate a quality of life by persons with disabilities is a prerequisite and first step in the strategy of improving the quality of life for this population.

There are those aspects of quality of life that we call universal and they are common to all people, in the sense of being, belonging and the development of a particular population. It is characteristic, that the conditions, for quality of life, are similar or even identical in humans worldwide (Power, 1999). Specific aspects related to the grade of membership and development in the domains specific to disability, or aspects that people with disabilities are considered important: a subjective well-being, satisfaction, and functioning in daily life, including self-help and social roles, external resources (material and social support), physical condition, psychological state, preserving the ability of social interaction and somatic condition (pleasure without pain, without disturbance) (Schipper, 1985). Jovanović (2011) said that the essence of the concept of quality of life for people with disabilities improve and enhance the overall life of the people, which is implemented through three levels: The first level involves the realization of basic human needs, the second level refers to the experience of satisfaction in aspects that are important for another person, and the third level is the achievement of higher levels of personal enjoyment and fulfillment.

The aim of this study is to assess the quality of life in persons with cerebral palsy, and what kind of the experience of quality of life have this person in relation to the gender structure and level of education.

METHODS

Sample

The study sample was formed by 100 people with cerebral palsy of both sexes, aged 23 to 64 years (M = 20,62, SD = 10,236). Criteria for selection of subjects were: a diagnosis of cerebral palsy (a neurological diagnosis), age over 18 years, preserved intellectual skills (finding a psychologist) and membership in the Association for Cerebral and Child Palsy in Belgrade.

Table 1 Distribution of participants by gender and level of education

		N	%
	Male	46	46
Gender	Female	54	54
	Total	100	100
	Primary school	15	15
Level of	Secondary school	77	77
education	High school/College	8	8
	Total	100	100

Legend: N = number of participants, % = percentage

Table 1 shows the distribution of participants by gender and level of education. 46% of our sample was male and 54% female. Of all participants, 77% have completed secondary school. At the same time 15% had primary education and 8% of participants have higher education.

Location and Time of Research

The research was conducted during 2013, May and June in the Association for Cerebral and Child Palsy in Belgrade.

Assessment Instrument

For the study we used a brief questionnaire about the quality of life of the World Health Organization (World Health Organization Quality of Life WHOQOL - BREF) (WHO, 1996). This questionnaire is a generic questionnaire, and it is used for selfassessment of the quality of life of adults. The questionnaire was based on a theoretical model that emphasizes the importance of individual perceptions of quality of life that a person has in relation to the culture and value system in which it is located, in relation to its goals, expectations, standards and concerns. WHOQOL-BREF contains 26 questions (two general questions of quality of life) and the rest are distributed among the domains of quality of life - Physical domain (7 questions), Psychological domain (6 questions), Social domain (3 questions), and Domain of environments (8 questions). Answers to the questions are arranged in Likert scale with a range of 1 to 5, where 1 means strongly disagree and 5 is strongly agree with a particular statement. The exception of this are question number 3, 4 and 26, where 1 means strongly agree and 5 strongly disagree with a given statement. All question from the questionnaire are related to the state of patients for the last four weeks of the date of examination. WHOQOL is psychometrically valid and reliable questionnaire which was translated into Serbian.

Statistical Analysis

For statistical analysis of the data collected, in accordance with the aim of this study, we used method of parametric and non-parametric statistics: measures of frequency and percentages, measures of central tendency (mean and standard deviation), method for determining the statistical significance between the arithmetic means for independent samples and univariate analysis of variance (ANOVA). In order to verify the reliability of the test used in this study, it was determined Coefficient of reliability Cronbach α = 0,874. The value of p<0,05 was considered statistically significant. For the purpose of data analysis it was used SPSS 21.0 for Windows.

RESULTS

Table 2 The achieved scores according to the participants' gender

Table 2 The achieved scores acc	orani						
Variables		Male	Geno		sig		
-	N	M	SD	N	M	SD	Ö
V1 How would you rate your quality of life?	46	3,89	,640	54	3,63	,784	,074
V2 How satisfied are you with your health?	46	3,93	,574	54	3,81	,729	,369
V3 How much do you feel that pain prevents you from doing what you need to do?	46	4,00	,869	54	3,69	,865	,073
V4 How much do you need medical treatment to function in your daily life?	46	3,50	1,243	54	3,15	1,071	,132
V5 How much do you enjoy life?	46	3,98	,954	54	3,72	,998	,195
V6 To what extent do you feel life to be meaningful?	46	4,72	,502	54	4,57	,792	,292
V7 How well are you able to concentrate?	46	3,67	,967	54	3,65	,872	,889,
V8 How safe do you feel in your daily life?	46	3,87	,980	54	3,80	1,035	,718
V9 How healthy is your physical environment?	46	3,57	1,128	54	3,91	1,051	,120
V10 Do you have enough energy for everyday life?	46	4,09	,812	54	3,96	,868	,465
V11 Are you able to accept your bodily appearance?	46	4,48	,836	54	4,41	,836	,674
V12 To what extent do you have enough money to meet your needs?	46	3,20	,980	54	3,06	1,280	,546
V13 How available to you is the information that you need in your day-to-day life?	46	4,00	,843	54	3,89	,965	,545
V14 To what extent do you have the opportunity for leisure activities?	46	3,70	,916	54	3,94	1,017	,205
V15 How well are you able to get around?	46	3,48	1,169	54	3,06	1,295	,092
V16 How satisfied are you with your sleep?	46	4,02	,954	54	3,72	,940	,118
V17 How satisfied are you with your ability to perform daily living activities?	46	3,83	,769	54	3,65	,935	,306
V18 How satisfied are you with your capacity for work?	46	3,83	,677	54	3,65	,828	,247
V19 How satisfied are you with yourself?	46	4,04	,759	54	3,87	,702	,239
V20 How satisfied are you with your personal relationships?	46	4,50	,658	54	4,19	,803	,036*
V21 How satisfied are you with your sex life?	46	3,50	1,111	54	2,80	1,203	,003*
V22 How satisfied are you with the support you get from your friends?	46	4,37	,826	54	4,33	,752	,819
V23 How satisfied are you with the conditions of your living place?	46	3,78	,987	54	3,65	,872	,471
V24 How satisfied are you with your access to health services?	46	3,52	,983	54	3,24	1,027	,167
V25 How satisfied are you with your transport?	46	3,46	1,242	54	3,33	1,149	,608
V26 How often do you have negative feelings, such as blue mood, despair, anxiety, depression?	46	4,07	,742	54	3,89	,816	,265

 $\label{lem:lem:mean} Legend: N=number\ of\ participants,\ M=arithmetic\ mean,\ SD=standard\ deviation,\ sig=significance,\ V=variable\ (code)$

Table 2 gives an overview of the achievements of the participants on the studied variables in relation to gender. The results indicate that in both groups saw a drop in the achievement of the maximum possible score. On two of the 26 variables, received a statistically significant difference in favor of male subjects, p < 0.05 (p = 0.36 and p = 0.03).

Table 3 Distribution of participants according to the level of education and variables in physical domain

	oloc-		Level of education								
physi		-	PS Le		SS		HS	To	tal		
doma	ain	N	%	N	%	N	%	N	%	- M	SD
	2	1	6,7	4	5,2	0	0	5	5		
V3	3	4	26,7	25	32,5	4	50	33	33	3,83	,877
٧٥	4	3	20	30	39	3	37,5	36	36	3,03	,0//
	5	7	46,7	18	23,4	1	12,5	26	26		
	1	4	26,7	5	6,5	0	0	9	9		
	2	1	6,7	10	13	0	0	11	11		
V4	3	7	46,7	27	35,1	4	50	38	38	3,31	1,161
	4	1	6,7	20	26	3	37,5	24	24		
	5	2	13,3	15	19,5	1	12,5	18	18		
	1	0	0	1	1,3	0	0	1	1		
	2	0	0	3	3,9	0	0	3	3		
V10	3	1	6,7	17	22,1	1	12,5	19	19	4,02	,841
	4	6	40	36	46,8	5	62,5	47	47		
	5	8	53,3	20	26	2	25	30	30		
	1	1	6,7	9	11,7	0	0	10	10		
	2	2	13,3	14	18,2	1	12,5	17	17		
V15	3	5	33,3	25	32,5	2	25	32	32	3,25	1,250
	4	4	26,7	12	15,6	4	50	20	20		
	5	3	20	17	22,1	1	12,5	21	21		
	1	1	6,7	1	1,3	0	0	2	2		
	2	1	6,7	3	3,9	1	12,5	5	5		
V16	3	4	26,7	21	27,3	1	12,5	26	26	3,86	,954
	4	5	33,3	34	44,2	0	0	39	39		
	5	4	26,7	18	23,4	6	75	28	28		
	1	1	6,7	0	0	0	0	1	1		
	2	0	0	10	13	0	0	10	10		
V17	3	2	13,3	14	18,2	2	25	18	18	3,73	,863,
	4	9	60	43	55,8	5	62,5	57	57		
	5	3	20	10	13	1	12,5	14	14		
	2	1	6,7	5	6,5	0	0	6	6		
V10	3	6	40	21	27,3	1	12,5	28	28	2.72	763
V18	4	5	33,3	41	53,2	7	87,5	53	53	3,73	,763
	5	3	20	10	13	0	0	13	13		
Tot	al	15	100	77	100	8	100	100	100	3,67	,958

Table 3 presents the results of the participants in relation to the level of education and variables in the physical domain. Based on the results, it can be said that the average of the results for the variable V10, V16, V17 and V18 are about 4, which corresponds to the statement "a lot". The average value of the variable V4 and V15 is around 3 ("neither good nor bad"). At the same time, the average value of V3 corresponds to the statement "a little".

Table 4 Distribution of participants according to the level of education and variables in psychological domain

psychological domain											
Varial					educati			To	tal		
psychol		PS			SS		HS	10	Lai	M	SD
dom: sco		N	N % N %		N	%	N %		1-1	35	
	1	0	0	1	1,3	0	0	1	1		
	2	1	6,7	6	7,8	0	0	7	7		
V5	3	3	20	25	32,5	2	25	30	30	3,84	,982
	4	3	20	26	33,8	2	25	31	31		
	5	8	53,3	19	24,7	4	50	31	31		
	2	0	0	2	2,6	0	0	2	2		
V6	3	1	6,7	4	5,2	0	0	5	5	1.61	4,64 ,674
VO	4	3	20	16	20,8	1	12,5	20	20	4,64	
	5	11	73,3	55	71,4	7	87,5	73	73		
	1	0	0	1	1,3	0	0	1	1		
	2	0	0	6	7,8	0	0	6	6		,913
V7	3	6	40	34	44,2	0	0	40	40	3,66	
	4	3	20	24	31,2	5	62,5	32	32		
	5	6	40	12	15,6	3	37,5	21	21		
	1	1	6,7	1	1,3	0	0	2	2		
	2	0	0	1	1,3	0	0	1	1		
V11	3	2	13,3	5	6,5	0	0	7	7	4,44	,833
	4	4	26,7	24	31,2	3	37,5	31	31		
	5	8	53,3	46	59,7	5	62,5	59	59		
	2	0	0	3	3,9	0	0	3	3		
1/10	3	3	20	17	22,1	0	0	20	20	2.05	720
V19	4	6	40	42	54,5	8	100	56	56	3,95	,730
	5	6	40	15	19,5	0	0	21	21		
	1	1	6,7	1	1,3	0	0	2	2		
	2	0	0	3	3,9	0	0	3	3		
V26	3	2	13,3	9	11,7	0	0	11	11	3,97	,784
	4	8	53,3	51	66,2	5	62,5	64	64		
	5	4	26,7	13	16,9	3	37,5	20	20		
Total	al	15	100	77	100	8	100	100	100	4,08	,819

Legend: N=number of participants, %=percentage, M=arithmetic mean, SD=standard deviation, V=variable (code), PS=primary school, SS=secondary school, HS=high school

Table 4 presents the results of the participants in relation to the level of education and variables in the psychological domain. Based on the results, recorded a very high overall average value of the variable psychological domain (M = 4,08, SD = 0,819), and this value is

in the level of claims " $_a$ lot / good". Particularly noteworthy is the variable " $_a$ what extent do you feel life to be meaningful?" in which 73% of participants saying " $_a$ lot".

Table 5 Distribution of participants according to the level of education and the variables in domain of environment

Variables-					ducatio						
environ			PS Ect		SS		HS	To	otal	3.5	an.
doma scor	iin	N	%	N	%	N	%	N	%	- M	SD
	1	1	6,7	0	0	0	0	1	1		
	2	2	13,3	9	11,7	0	0	11	11		
V8	3	2	13,3	18	23,4	1	12,5	21	21	3,83	1,006
	4	3	20	30	39	5	62,5	38	38		
	5	7	46,7	20	26	2	25	29	29		
	1	1	6,7	2	2,6	0	0	3	3		
	2	1	6,7	9	11,7	0	0	10	10		
V9	3	5	33,3	20	26	2	25	27	27	3,75	1,095
	4	3	20	23	29,9	3	37,5	29	29		
	5	5	33,3	23	29,9	3	37,5	31	31		
	1	2	13,3	9	11,7	0	0	11	11		
	2	1	6,7	18	23,4	0	0	19	19		
V12	3	1	6,7	18	23,4	6	75	25	25	3,12	1,148
	4	10	66,7	25	32,5	2	25	37	37		
	5	1	6,7	7	9,1	0	0	8	8		
	1	0	0	2	2,6	0	0	2	2		
	2	3	20	1	1,3	0	0	4	4		
V13	3	1	6,7	16	20,8	3	37,5	20	20	3,94	,908
	4	6	40	37	48,1	3	37,5	46	46		
	5	5	33,3	21	27,3	2	25	28	28		
	1	0	0	1	1,3	0	0	1	1		
	2	0	0	8	10,4	1	12,5	9	9		
V14	3	0	0	21	27,3	3	37,5	24	24	3,83	,975
	4	7	46,7	29	37,7	2	25	38	38		
	5	8	53,3	18	23,4	2	25	28	28		
	1	0	0	1	1,3	1	12,5	2	2		
	2	1	6,7	3	3,9	1	12,5	5	5		
V23	3	7	46,7	24	31,2	3	37,5	34	34	3,71	,924
	4	3	20	33	42,9	2	25	38	38		
	5	4	26,7	16	20,8	1	12,5	21	21		
	1	3	20	3	3,9	0	0	6	6		
	2	3	20	9	11,7	0	0	12	12		
V24	3	4	26,7	23	29,9	3	37,5	30	30	3,37	1,012
	4	4	26,7	35	45,5	4	50	43	43	-,	_,
	5	1	6,7	7	9,1	1	12,5	9	9		

Variab	Variables- environment		Lev	el of e	ducatio	T-4-1					
-			PS		SS		HS		- Total		SD
domain score		N	%	N	%	N	%	N	%	M	J.D
	1	1	6,7	8	10,4	1	12,5	10	10		
	2	0	0	10	13	1	12,5	11	11	3,39	1,188
V25	3	5	33,3	18	23,4	3	37,5	26	26		
	4	3	20	30	39	3	37,5	36	36	-,	_,
	5	6	40	11	14,3	0	0	17	17		
Tota	al	15	100	77	100	8	100	100	100	3,62	1,032

Legend: N=number of participants, %=percentage, M=arithmetic mean, SD=standard deviation, V=variable (code), PS=primary school, SS=secondary school, HS=high school

Table 5 shows the results of the participants in relation to the level of education and the variables in the domain of environment. The average values for the variables in this field range from M = 3,12 (SD = 1,148) to M = 3,94 (SD = 0,908), and they are located between the claims *"neither good nor bad"* and *"good"*.

Table 6 Distribution of participants according to the level of education and the variables in social domain

Variables-social domain			Lev	vel of	educati	on					
		PS			SS		HS	· To	tal	M	SD
score		N	%	N	%	N	%	N	%	-	
	1	0	0	1	1,3	0	0	1	1		
	2	0	0	6	7,8	0	0	6	6	3,75	
V1	3	1	6,7	16	20,8	1	12,5	18	18		,730
	4	11	73,3	49	63,6	7	87,5	67	67		
	5	3	20	5	6,5	0	0	8	8		
	2	2	13,3	2	2,6	0	0	4	4		
V2	3	3	20	12	15,6	2	25	17	17	3,87	,661
V Z	4	6	40	56	72,7	5	62,5	67	67		
	5	4	26,7	7	9,1	1	12,5	12	12		
	2	0	0	2	2,6	0	0	2	2		,753
V20	3	2	13,3	9	11,7	0	0	11	11	4,33	
V 20	4	7	46,7	30	39	2	25	39	39	4,33	
	5	6	40	36	46,8	6	75	48	48		
	1	5	33,3	10	13	0	0	15	15		
	2	1	6,7	10	13	1	12,5	12	12		
V21	3	6	40	19	24,7	4	50	29	29	3,12	1,208
	4	3	20	28	36,4	3	37,5	34	34		
	5	0	0	10	13	0	0	10	10		
	2	1	6,7	3	3,9	0	0	4	4		
V22	3	1	6,7	6	7,8	0	0	7	7	4,35	702
	4	7	46,7	30	39	2	25	39	39		,783
	5	6	40	38	49,4	6	75	50	50		
Total		15	100	77	100	8	100	100	100	3,88	,827

Legend: N=number of participants, %=percentage, M=arithmetic mean, SD=standard deviation, V=variable (code), PS=primary school, SS=secondary school, HS=high school

The results of the participants in relation to their level of education and variables in the social domain are shown in Table 6. Results show that the average values are between claims "neither good nor bad" and "a lot", or from M = 3,12 (SD = 1,208) to M = 4,35 (SD = 0,783). Variable "How would you rate the quality of your life?" stands out, because 73,3% of participants with primary school education, 63,6% of secondary education and even 87,5% of highly educated participants saying "good".

Table 7 Relationship between groups-ANOVA

-	itionship betwe	ANOVA	
Variable -	F	df	sig
V1	2,843	2	,063
V2	,098	2	,907
V3	,793	2	,456
V4	2,392	2	,097
V 5	2,270	2	,109
V6	,566	2	,569
V7	4,743	2	,011*
V8	,404	2	,669
V9	,524	2	,594
V10	2,801	2	,066
V11	,860	2	,426
V12	,925	2	,400
V13	,088	2	,915
V14	4,995	2	,009*
V15	,577	2	,563
V16	1,500	2	,228
V17	,386	2	,681
V18	,193	2	,825
V19	1,111	2	,333
V20	1,375	2	,258
V21	2,667	2	,075
V22	1,337	2	,268
V23	1,867	2	,160
V24	3,283	2	,042*
V25	1,738	2	,181
V26	1,163	2	,317

ANOVA analysis of the level of education and the studied variables recorded a statistically significant difference in the three studied variables (Table 7): "How well are you able to concentrate?" (F=4,743, df=2, p=0,11); "To what extent do you have the opportunity for leisure activities?" (F=4,995, df=2, p=0,009); "How satisfied are you with your access to health services?" (F=3,283, df=2, p=0,42), while in the other variables level of education is not shown as a statistically significant factor.

Table 8 Difference between groups										
	PS-SS	,066		PS-SS	1,000					
V1	PS-HS	1,000	V15	PS-HS	1,000					
	SS-HS	1,000		SS-HS	1,000					
	PS-SS	1,000		PS-SS	1,000					
V2	PS-HS	1,000	V16	PS-HS	,274					
	SS-HS	1,000		SS-HS	,406					
	PS-SS	,883		PS-SS	1,000					
V3	PS-HS	,761	V17	PS-HS	1,000					
	SS-HS	1,000		SS-HS	1,000					
	PS-SS	,135		PS-SS	1,000					
V4	PS-HS	,235	V18	PS-HS	1,000					
	SS-HS	1,000		SS-HS	1,000					
	PS-SS	,262		PS-SS	,429					
V5	PS-HS	1,000	V19	PS-HS	1,000					
	SS-HS	,450		SS-HS	1,000					
	PS-SS	1,000		PS-SS	1,000					
V6	PS-HS	1,000	V20	PS-HS	,433					
	SS-HS	,887		SS-HS	,326					
	PS-SS	,168		PS-SS	,073					
V7	PS-HS	,998	V21	PS-HS	,406					
	SS-HS	,031		SS-HS	1,000					
	PS-SS	1,000		PS-SS	1,000					
V8	PS-HS	1,000	V22	PS-HS	,332					
	SS-HS	1,000		SS-HS	,475					
	PS-SS	1,000		PS-SS	1,000					
V9	PS-HS	1,000	V23	PS-HS	,540					
	SS-HS	,999		SS-HS	,173					
	PS-SS	,065		PS-SS	,071					
V10	PS-HS	1,000	V24	PS-HS	,092					
	SS-HS	1,000		SS-HS	1,000					
	PS-SS	,775		PS-SS	,346					
V11	PS-HS	,742	V25	PS-HS	,290					
	SS-HS	1,000		SS-HS	1,000					
	PS-SS	,571		PS-SS	1,000					
V12	PS-HS	1,000	V26	PS-HS	,602					
	SS-HS	1,000		SS-HS	,401					
	PS-SS	1,000								
V13	PS-HS	1,000								
	SS-HS	1,000								
	PS-SS	,008								
V14	PS-HS	,088								
	SS-HS	1,000								

Legend: PS=primary school, SS=secondary school, HS=high school

Bonferroni post-hoc test was applied to examine between which groups the difference was statistically significant (Table 8). The results show a statistically significant difference between the groups of secondary and higher education (p = 0.31) in the variable "How well are you able to concentrate?" and between groups of primary and secondary schools in the variable "To what extent do you have the opportunity for leisure activities?" (p = 0.008).

DISCUSSION

In recent years, assessment of the quality of life of persons with disabilities became very important research problem. Accordingly, there are numerous studies dealing with quality of life of people with disabilities, but there are a few with focus of their research on adults with cerebral palsy. People with cerebral palsy are heterogeneous group in terms of their motor, cognitive, social, educational and other characteristics. In regard with this they have to deal with many physical, social and psychological consequences related to their condition (Berrin et al., 2007, according to Gojčeta, 2008). In relation to the aforementioned effects, by identifying the predictors of the quality of life of people with cerebral palsy it is possible to create a treatment in order to improve the functioning of these persons (Renk & Wiley, 2007) and to improve the conditions of daily life (Odović et al., 2012).

Cerebral palsy is not only a medical problem but also it is a social and economic, and as such it, affects the overall quality of life for these people and their environment (Križ & Prpic, 2005; according to Odović et al., 2012).

Most researches of the quality of life of people with cerebral palsy usually include a small sample of children, but not a sample based on the population, either their study was based on parental attitudes, and on their perception of quality of life (Davis et al., 2010; Fatudimu et al., 2013; Khayatzadeh 2009; Viehweger, 2008), while on the other hand they used instruments that were focused on the functions and activities, therapeutic consequences and impact on parents.

In our research, the analysis of the results enabled us to determine what the experience of quality of life is for people with cerebral palsy in relation to their level of education and in relation to gender. It was found that the male participants had higher scores on almost all variables. Between male and female, of our sample, there is a difference in perception of quality of life of the studied variables, but there are not statistically significant except for the two variables, tin favor of male respondents. Male participants their relationship with other people evaluate the claim that ranges from "good" to "very good" (M=4,50, SD=0,658), while female participants generally define this relationship by saying "good" (M=4,19, SD=0,803) (p=0,036). When it comes to satisfaction with sex life, the experience of "good or bad" to "good" are characteristics of the male participants (M=3,5, SD=1,111), while female participants satisfaction with their sex life as "a not good or bad" (M=2,80, SD=1,203) (p=0,003).

Jovanović (2011) is using the WHOQOL – BREF conducted a survey of the quality of life in 390 persons with disabilities (people with cerebral palsy, muscular dystrophy, multiple sclerosis, and paraplegia) and 105 persons without disabilities. The average

value of the age of those with disabilities was 41,86, and the highest number of respondents (60,5%) had completed high school. The results (Jovanović, 2011) were correlated with the results of the research of others, showing that people with disabilities have lower levels of subjective quality of life compared to people in the general population. However, as reported by Jovanović (2011), there are differences between people with disabilities in relation to their diagnosis. Thus, in the area of physical functioning, people with cerebral palsy evaluated its functioning like people without disabilities, as they relate to others and the support they receive from others rated significantly lower grades. The results of our study indicate that between male and female, when it comes to relations with others, there is a statistically significant difference (p=0,036) in favor of the male participants, while the support of other people did not find as statistically significant variable between the gender (p=0,819). Further analysis revealed that among the participants in relation to the level of education there is no statistically significant difference when it comes to relationships and support they receive from others, but the difference in relation to the level of education recorded in one variable of psychological domain (F=4,743, df=2, p=0,011), and in the two variables in domain of the environment (F=4,995, df=2, p=0,009), (F=3,283, df=2, p=0,042).

In a survey conducted by Jovanović (2011) it's observed a statistically significant difference in the domain of environment, but also in the social and physical domain. Participants with higher education, in the aforementioned study, had significantly higher scores in these areas compared to those with primary and secondary schools. Our findings are consistent with the results of that research, and shows that people with higher education have higher achievement on the variables in almost all domains compared to those with primary and secondary education. The difference which is recorded between the groups of those with primary schools, secondary schools and higher education levels are in statistically significant differences in the two variables. Among the group of secondary and higher education was obtained difference (p=0,31) in the variables "How well are you able to concentrate?" and between groups of ones with primary and secondary schools in the variables "To what extent do you have the opportunity for leisure activities?" (p=0,008). Persons with disabilities who have higher education have significantly better physical and social functioning and occupy a better position in their environment than people with disabilities who have lower levels of education, according to Jovanović (2011). Our results may be related to this paragraph.

The highest score on the WHOQOL-BREF is in the social domain, which refers to the overall social function and role, and the lowest in the field of environment (Jovanović, 2011), while our results, also, low achievement of participants in relation to level of education recorded in domain of environment (M=3,62, SD=1,032), but the highest attainment is recorded in the psychological domain (M=4,08, SD=0,819).

Results of research conducted by Sretenović et al., (2013), on the sample consisted of people with CP showed that respondents of both genders satisfied with the quality of their life when it comes to the social domain of quality of life.

Odović et al., (2013) conducted a study on a sample of 51 people with cerebral palsy and it was noted that the level of education of respondents represents a statistically significant factor in the field of interpersonal relationships and responsibility.

In the work of Colver (2012), we found research on outcomes for people with cerebral palsy: the expectations and quality of life. In all tested areas: standard of living, subjective well-being, participation, mental health, physical health, including pain, adults with cerebral palsy have achieved lower results compared to the representatives of the general population. These results, Colver (2012) associated with inappropriate interventions in early childhood. In addition, states that adults with cerebral palsy does not get enough attention in the research, as well as the outcomes of social participation are not properly considered in adulthood.

Trgovčević, Nedović and Kljajić (2012) investigated the determinants of quality of life and their correlation with the degree of disability. In a sample of 53 people with an injury of the cervical spinal cord, they noticed a paradox, where 54,6% of participants said that they have an excellent or good quality of life. The authors draw the conclusion that people who have suffered spinal cord injury and a high degree of disability did not possess the knowledge, the resources and the social contacts that help them to build balance and well-being. Our results showed that 75% of participants rate their quality of life as good or very good.

CONCLUSION

Our study was conducted to determine impact of level of education and gender on quality of life of people with cerebral palsy. It is notable that people with cerebral palsy have a high level of subjective quality of life, because the quality of life refers to a personal assessment of the individual in relation to his aspirations and achievements, and the essence of life satisfaction was related to the subjective perspective (Trgovčević et al., 2012). Even though quality of life is highly rated among participants with regard to gender and in relation to the level of education, and there are still a differences. The question that arise to us is: does people with cerebral palsy in addition to their limits, both personal and environmental, really believe that they have a high level of satisfaction with the quality of life? It could be argued that subjectivity, experience and aspirations of the participants have a significant stake in these results. We believe that studies which are aimed at measuring the quality of life of people with cerebral palsy is a basis for improving the quality of life of these people.

There are some limitations to this study, which may lead to difficulties in interpreting the results. The first limitation is a sample of study. The study sample is not small, but we can considered it is small without problems, if we take into account that people with cerebral palsy are very heterogeneous group of people with disabilities in relation to their motor and cognitive skills, social functioning, psychological functions, and more. The division of the sample in sub-samples, likely would give us different results. Furthermore, the lack of information on the social, financial, family status of participants, may also cause difficulties in interpreting the results. The following restriction is reflected in the absence of control group.

WHOQOL-BREF is not the only instrument that is used to assess the quality of life of people with disabilities, but given the fact that it was sufficient in empirical studies that have used this instrument to assess the quality of life of people with cerebral palsy, we believe that this design will contribute and serve as a basis for future research.

REFERENCES

- 1. Bowling, A. (1994). The concept of quality of life in relation to health. *Medicinanei secoli*, 7(3), 633-645.
- 2. Cella, F.D. (1992). Quality of life: the concept. Journal Palliat Care, 8(3), 8-13.
- 3. Colver, A. (2012). Outcomes for people with cerebral palsy: life expectancy and quality of life. *Paediatrics and Child Health*, 22(9), 384-387.
- 4. Cummins, R.A. (1998). The second approximation to an international standard for life satisfaction. *Social Indicators Research*, *43*(3), 307-334.
- 5. Davis, E., Shelly, A., Waters, E., Boyd, R., Cook, K., & Davern, M. (2010). The impact of caring for a child with cerebral palsy: quality of life for mothers and fathers. *Child: care, health and development, 36*(1), 63-73.
- 6. Delhey, J., Bohnke, P., Habich, R., & Zapf, W. (2002). Quality of life in a European Perspective: The EUROMODULE as a New Instrument for Comparative Welfare Research. *Social Indicators Research*, *58*(1), 161-175.
- 7. Fatudimu, M.B., Hamzat, T.K., & Akinyinka, O.O. (2013). Comparative quality of life of Nigerian caregivers of children with cerebral palsy. *International Journal of Therapy and Rehabilitation*, 20(3), 131-135.
- 8. Gojčeta, M., Oreb-Joković, I., & injatela, R. (2008). Neki aspekti kvalitete života adolescenata sa i bez cerebralne paralize. *Hrvatska revija za rehabilitacijska istraživanja*, 44(1), 39-47.
- 9. Jovanović, M. (2011). Invalidnost i kvalitet života. Socijalna misao, 18(2), 151-160.
- 10. Karaduman, A., Yilmaz, O., Tuzun, E., KeremGunel, M., Aras, B., Mutlu, A., Tarsuslu, T., & Aras, O. (2010). A comparasion of quality of life in children with cerebral palsy and neuromuscular diseases. *Fizyoterapi Rehabilitasyon*, 21(1), 3-10.
- 11. Khayatzadeh, M. (2009). A comparative study about quality of life in mothers of children with cerebral palsy, mental retardation and mothers of normal children. *Daneshvar Medicine*, 16(83), 214-232.
- 12. Macanović, G. (2010). Procena kvaliteta života školske dece merena medicinskim, kulturnim i socijalnim pokazateljima. *PONS medicinski časopis*, *7*(2), 60-63.
- 13. Недовић, Г., Сретеновић, И., Станисављевић, Ј. (2013). Психолошки домен квалитета живота особа са церебралном парализом. *Београдска дефектолошка школа*, вол. 19(3), бр. 57, стр. 485-496.
- 14. Odović, G., Sretenović, I., Stanisavljević, J. (2012). Ostvarivanje životnih navika osoba sa motoričkim poremećajima. *VI Međunarodni naučni skup "Specijalna edukacija i rehabilitacija danas"*, Zbornik radova, str. 258-265. Univerzitet u Beogradu, Fakultet za specijalnu edukaciju i rehabilitaciju, Beograd, ISBN: 978-86-6203-037-5.
- 15. Odović, G., Rapaić, D., Stanisavljević, J., Sretenović, I. (2012). Life habits accomplishment level of persons with cerebral palsy. *Il International Scientific Conference "Special Education and Rehabilitation Cerebral Palsy"*, Book of Proceedings and Summaries, pp. 86-96, Novi Sad, Serbia.
- 16. Odović, G., Stanisavljević, J., Sretenović, I. (2013). Ostvarivanje socijalnih uloga osoba sa cerebralnom paralizom. *VII Međunarodni naučni skup "Specijalna edukacija i rehabilitacija Danas"*, Zbornik radova, str. 277-287. Univerzitet u Beogradu, Fakultet za specijalnu edukaciju i rehabilitaciju, Beograd, ISBN 978-86-6203-045-0
- 17. Power, M., Harper, A., & Bullinger, M. (1999). The World Health Organization WHOQOL-100: Test of the University of Quality of Life in 15 Different Cultural Groups Worldwide. *Health Psychology*, 18, 495–505.
- 18. Schipper, H., & Levitt, M. (1985). Measuring quality of life. Risk and benefits. *Cancer Treat Rep*, 69, 1115–1123.

- 19. Sretenović, I., Stanisavljević, J., Milivojević, M., Šarac-Marić, G., & Kovačić, A. (2013). Socijalni domen kvaliteta života osoba sa cerebralnom paralizom. II stručno naučni skup sa međunarodnim učešćem "Aktuelnosti u edukaciji i rehabilitaciji osoba sa smetnjama u razvoju", Zbornik rezimea, str. 89, Šabac.
- 20. Testa, M.A., & Simonson, M.D. (1996). Assessment of Quality-of-Life Outcomes. *The New England Journal of medicine*, 334, 835-840.
- 21. Torres, V.M.F., Marinho, C.L.A., Oliveira, C.G.G., & Vieira, S.C.M. (2013). Quality of life in adolescents with hearing deficiencies and visual impairments. *Int. Arch. Otorhinolaryngol*, 17(2), 139-146.
- 22. Trgovčević, S., Kljajić, D., & Nedović, G. (2011). Socijalna integracija kao determinant kvaliteta života osoba sa traumatskom paraplegijom. *Godišnjak Fakulteta političkih nauka*, 5(6), 491-505.
- 23. Trgovčević, S., Nedović G., & Kljajić, D. (2012). Paradoks invaliditeta: visok kvalitet života uprkos svemu. U N. Dimić (Ed.), *Stručno-naučni seminar sa međunarodnim učešćem "Dani defektolaga Srbije"* (p.69). Zlatibor: Društvo defektologa Srbije.
- 24. Trgovčević, S. (2013). *Kvalitet života osoba sa povredom kičmene moždine.* (Doctoral disertation). Beograd: Fakultet za specijalnu edukaciju i rehabilitaciju.
- The World Health Organization Quality of Life assessment (WHOQOL). (1998).
 Development and general psychometric properties. SocSci Med, 46, 1569-1585.
- 26. Viehweger, E., Robitail, S., Rohon, M.A., Jacquemier, M., Jouve, J.L., Bollini, G., & Simeoni, M. C. (2008). Measuring quality of life in cerebral palsy children. *Annales de réadaptationet de médecine physique*, 51(2), 129-137.
- 27. Wiley, R., & Renk, K. (2007). Psychological correlates of quality of life in children with cerebral palsy. *Journal of Developmental and Physical Disabilities*, 19(5), 427-447.