

Comparison of children with and without chronic disease in psychosocial terms

Kronik hastalığı olan ve olmayan çocukların psikososyal açıdan karşılaştırılması

Özden Gökdemir¹ Muhteber Colak⁴ DÜlkü Bulut Batur⁵

Seval Yaprak²

Yunus Emre Okudan³ Dilek Güldal⁶

¹⁾ IUE Faculty of Medicine, Department of Family Medicine, Asst. Prof., Balcova, İzmir. {ORCID:0000-0002-0542-5767}

²⁾ Ministry of Health, Selçuklu 70 Nolu ASM, Family Practice Specialist, Konva. {ORCID:0000-0002-1454-1612}

³⁾ Anatolian Hospital, Antalya. {ORCID:0000-0003-1330-1884}

⁴⁾ Ministry of Health, Balçova 2 Nolu ASM, Family Practice Specialist, İzmir. {ORCID:0000-0003-3684-4961}

⁵⁾ Aksaray University Faculty of Medicine, Department of Family Medicine, Asst. Prof., Aksaray. {ORCID:0000-0003-3011-0924}

⁶⁾ Dokuz Eylül University, Faculty of Medicine Department of Family Medicine, Asst. Prof., İzmir. {ORCID:0000-0002-7344-9349}

İletisim adresi: Dr. Özden Gökdemir E-mail: gokdemirozden@gmail.com Geliş tarihi: 03/08/2021 Kabul tarihi: 14/12/2021 Yayın tarihi: 25/12/2021

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Özet

Amaç: Tıptaki gelişmelere bağlı olarak kronik hastalığı olan çocuk sayısı artmıştır. Bu çalışmanın amacı, kronik hastalığı olan ve olmayan çocukların duygu durumları ve çevreleriyle olan ilişkileri açısından farklılık gösterip göstermediğini araştırmaktır.

Yöntemler: Bu olgu-kontrol çalışmasına pediatri polikliniklerine başvuran 4-11 yaş arası çocuklar dahil edildi. Herhangi bir kronik hastalığı olan çocuklar vaka grubuna dahil edilirken, tanımlanan herhangi bir kronik hastalığı olmayan, şikâyeti 10 günden az olan ya da şikâyeti olmayan çocuklar kontrol grubuna alındı. Çocukların bakım verenlerine demografik özelliklerle ilgili bir anket ile Güçler Yönler ve Güçlükler Anketi (SDQ) uygulandı. Sonuçlar SPSS 21 programı ile analiz edildi; betimleyici testler ve hipotez testleri uygulandı.

Bulgular: Hem olgu hem de kontrol grubu 125 çocuktan oluşuyordu; olgu grubundaki çocukların en uzun ikamet ettiği yerin kontrol grubundan istatistiksel açıdan anlamlı fark olarak ilçe ve/veya köy olduğu bulundu (p=0.011). SDQ ve alt grupları açısından olgu ve çalışma grupları arasındaki tek anlamlı fark duygusal zorluklardı (p=0,018). Karşılaşılan zorluklarla ilgili soruya verilen yanıtlar incelendiğinde, kronik hastalığı olan çocukların akran ilişkilerinde daha fazla sorun yaşadıkları (p=0.015) ve yaşam zorluklarının daha belirgin olduğu (p=0,038) görüldü.

Sonuç: Birinci basamakta çocuk hasta grubuna yaklaşımda, hastalığın potansiyel olarak çok yönlü etkileri akılda tutularak günlük yaşam olaylarının dikkatle ele alınması önemlidir.

Anahtar sözcükler: Birinci basamak, SDQ, akran ilişkileri, kronik hastalık, çocuk bakımı

Summary

Aim: The number of children with chronic diseases has increased due to developments in medicine. The aim of this study is to determine whether children with and without chronic disease differ in terms of their emotional states and relationships with their environment.

Methods: Children of ages between 4 and 11 who applied to pediatrics polyclinics were included in this case-control study. While children with any chronic disease were included to the case group, children without any of the identified chronic diseases, either with a complaint of less than 10 days, or no complaint, were included in the control group. A questionnaire about demographic features, and Strengths and Difficulties Questionnaire (SDQ) was applied to the caregivers of the children. The results were analyzed with SPSS 21 program; descriptive tests and hypothesis tests were applied.

Results: Both the case and control groups consisted of 125 children; and the place of the longest residence of children in the case group was found to be a district and/or village, which was statistically different than the control group (p=0.011). Regarding SDQ and its sub-groups, the only significant difference between the case and study groups was emotional difficulties (p=0,018). When we analyzed the answers to the question about challenges faced, it was observed that children with chronic diseases had more/greater problems with their peer relations (p=0.015), and life challenges were more significant (p=0,038).

Conclusion: In the approach taken to the pediatric patient group in primary care, it is important to consider carefully daily life events bearing in mind the potentially multifaceted effects of the disease.

Key words: primary care, SDQ, peer relations, chronic disease, childcare

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Introduction / Background

With advanced medical treatment methods, the number of children surviving with chronic disease has increased.^(1,2,3) Having chronic disease (mucopolysaccharidosis, diabetes mellitus, etc.) increases risk of psychological status, the psychological evaluation of this condition could also be important for family physicians. ^(3,4) Although all innovations and techniques followed in medicine are followed by their respective specialties, the subject of psychological support is not given sufficient importance to patients and/or caregivers, whose role is very important in the treatment of the patient, and whose priority and importance, particularly the treatment of the patient, and whose priority and importance, particularly in developed countries, are indisputable.⁽³⁾ The psychological state of the patient significantly affects the course of the disease. For pediatric patients, the importance of psychological support is indisputable.⁽⁵⁾

It is noteworthy that research specifically from the psychosocial perspective is insufficient. The recently accepted biopsychosocial approach emphasizes the importance of psychosocial evaluation in addition to medical treatment.⁽³⁾ Therefore, comparing the psychosocial behavior of children aged 4-11 years with chronic disease to the behavior of those without can help to identify the effects of these diseases, and to realize the challenges caused. Any significant difference will emphasize that children with chronic disease are different from children without, and that approaches should be modified accordingly. Araştırma | *Research* Yıl: 2021 Cilt: 12 Sayı: 4 / e-ISSN: 2148-550X doi: 10.15511/tjtfp.21.00471

Aim

Chronic diseases require long-term treatment, and negatively affects quality of life. Especially in childhood and adolescence, individuals face many physiological and psychological changes. The additional burden of chronic disease may hinder their social and psycholoical development or cause various other problems.^(6,7) In this research, the aim is to determine whether chronic disease affects the emotional state of children and their relationships with the environment.

The ethical permission of the study was approved by the ethical committee of Dokuz Eylul University (DEU: 2013 / 13-14 11.04.2013).

Method

Descriptive case-control study.

Population and sampling: The population of the study was 4-11 years old children selected from among the outpatient and inpatient patients admitted to DEU, Child Health and Diseases Department. Children with chronic disease were compared with children without (between January and June of 2014).

The sample size was calculated to be at least 111 children for each group with 80% power, 95 % CI, OR:3 and prevalence of 10%. The cases were selected consecutively among the patients who were hospitalized in the pediatric ward and followed up at the sub-speciality clinics.

The control group, on the other hand, were the children who applied to the general and "well-child" outpa-



tient clinic of pediatrics, either with no complaints, or with complaints of durations of less than 10 days, and who were identified in the research as not having chronic diseases. Children included in the control group were matched in terms of age and gender. "Having chronic diseases" is defined as either diseases that have existed longer than 3 months or have a high probability of continuing more than 3 months. Also, diseases having clinical symptoms 3 or more times a year were considered as chronic diseases. Seven disease groups were included according to this definition.^(8,9)

- 1. Chronic Respiratory Diseases (Asthma)
- 2. Congenital heart diseases
- 3. Hematological-oncological malignancy
- 4. Metabolic diseases (Diabetes)

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- 5. Neurometabolic disease
- 6. Chronic kidney failure

7. Rheumatologic diseases (Juvenile rheumatoid ar-thritis)

Child volunteers between the ages of 4-11 participated in the study.

Those with any psychological issues were not included in the study.

Data Collection Tools:

- Sociodemographic data form

- Strengths and Difficulties Questionnaire - SDQ: This tool was developed by Robert Godman and has been translated into different languages. This questionnaire includes the parent form and school form for ages

Table 1: Location and caregivers' status of the research group								
		Case Group		Control Group		Total		
		n	%	n	%	n	%	
Location	Province	81	64.8	99	79.2	180	72	99
	District-village	44	35.2	26	20.8	70	28	
	Total	125	100	125	100	250	100	
	Parents	120	96.0	121	96.8	241	96.4	99
Caregiver	Others	5	4.0	4	3.2	9	3.6	
	Total	125	100	125	100	250	100	
Occupation of the caregivers	Housewife	76	60.8	84	67.2	160	64	99
	Worker	9	7.2	9	7.2	18	7.2	
	Office worker	25	20.0	22	17.6	47	18.8	
	Own business	12	9.6	7	5.6	19	7.6	
	Retired	3	2.4	3	2.4	6	2.4	
	Total	125	100	125	100	250	100	
Chi-Square Tests								



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4-16, and the adolescent form for 11-16 ages. The adolescent form contains the same items as the parent form, except that for many items, the first-person singular pronoun is used instead of the third person. It contains 25 questions, which focus on either positive or negative behavior characteristics. These questions are presented in five categories: behavioral problems, attention deficit and excessive mobility, emotional problems, peer problems, social behaviors.

As each category is evaluated alone, the sum of the first four categories gives the 'total difficulty score'.⁽¹⁰⁾ The Turkish validity and reliability of the questionnaire

was performed by Güvenir et al. (Cronbach alpha values were: emotional 0.70; behavioral problems 0.50; attention deficit and excessive mobility 0.70; peer problems 0.22; social behavior 0.54; and for the total difficulty score 0.73).⁽¹¹⁾

Results

125 children were included in each group; and no differences were observed between the two groups regarding age, gender, and their caregivers. The longest place of residence for children in the case group was found to be a town or a village, which was sta-

Table 2: Compare the groups according to the cut-off value							
	C	Control Group		_			
		n	%	n	%	P	
SDQ Scores	Normal 0-14	25	20.0	36	28.8		
	Abnormal 14-40	100	80.0	89	71.2	0.105	
	Total	125	100	125	100		
	Normal 0-3	87	69.06	103	82.4	0.018*	
Emotional Difficulties	Abnormal 3-10	38	30.04	22	17.6		
	Total	125	100	125	100		
	Normal 0-2	83	66.4	90	72.0		
Behavioural Difficulties	Abnormal 3-10	42	3.6	35	28.0	0.338	
	Total	125	100	125	100		
	Normal 0-5	102	81.6	108	86.4		
Hyperactivity Problems	Abnormal 6-10	23	18.4	17	13.6	0.301	
	Total	125	100	125	100		
Peer Problems	Normal 0-2	90	72.0	98	78.4	0.241	
	Abnormal 3-10	35	28	27	21.6		
	Total	125	100	125	100		
	Normal 6-10	118	94.4	118	94.4		
Positive Social Approach	Abnormal 0-5	7	5.6	7	5.6	1.000	
	Total	125	100	125	100		
Chi-Square test							



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tistically different than the control group (p=0.011 **Table 1**). Regarding SDQ and its sub-groups, there were no significant differences between the case and study groups. Additionally, there was no significant relationship between the duration of the disease and the ratings (**Table 2**). When asked about difficulties in their lives on a Likert scale: "never, a little, some and always", 88 participants answered (56 participants with chronic disease, 32 participants from the control group). Table 3 and Table 4 give the results of these 88 participants.

Table 3: Time of diffuculties and general response							
		Case Group		Control Group			
		n	%	n	%	Р	
	<1 year	18	32.1	5	15.6		
Time of the difficulties	<1 year	38	67.9	27	84.4	0.090	
	Total	56	100	32	100		
	Normal 0	35	62.5	21	65.6		
General Response	Abnormal 1-10	21	37.5	11	34.4	0.769	
	Total	56	100	32	100		
	Never	12	21.4	2	6.3	0.038*	
Life Challengee	A bit	27	48.2	24	75.0		
	Some - Always	17	30.4	6	18.8		
	Total	56	100	32	100		

Table 4: Comparison of the total scores in case and control groups	
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	Case Group	Control Group				
	Mean ±SD	Mean ±SD	t	р		
SDQ Scores	18,21±5.131	16,77± <mark>4.881</mark>	2.273	0.024*		
Emotional Difficulties	2,41±2.024	1,90±2.209	1,881	0,061		
Behavioural Difficulties	1,72±1.543	1,70±1.636	0,080	0,937		
Hyperactivity Problems	3,37±2.535	2,77±2.332	1,947	0,053		
Peer Problems	1,86±1.763	1,34±1.530	2,452	0,015		
Positive Social Approach	8,86±1.630	9,05±1.591	-0,943	0,347		
Life Challenges (N:88)	2,77±2.036	2,59±1.073	1.006	0,317		
Independent samples T-test						

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Discussion

In our study, it was observed that children with chronic diseases were psychosocially no different from their peers according to the total scores they got from the SDQ. Rothenberger et al reported that SDQ is appropriate for screening but there is not sufficient evidence for multicultural setting.⁽¹²⁾ Richter et all revealed that SDQ may produce different results among ethnic Norwegian samples, although this tool is "internationally most frequently used screening instruments for child and adolescent mental health purposes". We did not collect information about cultural and ethnic diversity of the children included to study. The subscale scores are considered more valuable than the total. ⁽¹³⁾ When we used the cut-off scores to evaluate subscales, only emotional difficulties were significantly different between the groups.⁽¹²⁾ When children begin to attend kindergarten or elementary school, they do not gain knowledge but also the social norms and values of their communities.⁽¹⁴⁾ Emotional difficulties may hinder integration with peers in the case group.

When the responses to the question concerning difficulties / challenges faced were assessed, it was discovered that children with chronic conditions had more/ greater difficulty with peer relationships. The SDQ is a self-assessment tool, and children with chronic diseases reported having difficulty with peer relationships. Many research have led to similar findings, demonstrating that peer relationships are indeed an issue.^(15,16) The impact of peer relationship can be used to encourage healthy lifestyles, education, as well as other interests, although it varies depending on the individuals.^(15,17)

Socioeconomic disadvantage negatively affects every dimension of child development, including mental health and well-being.^(18,19) In our study, most children with chronic disease lived in small towns and/or villages. While some studies reported that low socioeconomic status was associated with greater emotional and behavioral problems, this result was not revealed in our study.^(12,18)

Academic success and satisfaction in adult life are key problems to address for the children with chronic diseases.⁽¹⁶⁾ Life challenges were more significant in the children with chronic disease group in our study. A holistic approach towards family medicine could be a key solution during the follow-up of these children. ^(20,21,22) In the approach to the pediatric patient group in primary care, daily life issues should be carefully considered, and it is important to remember that the effects of the disease may be multifaceted. "Life challenges" is a problem not only for healthcare workers but also for policymakers. Children with chronic disease may be anxious about their future lives. However, social networks, societies, and stakeholders all have responsibility for the future of these children.

Lack of probing the effect of cultural diversity may be considered as the limitation of this study.

Conclusion

Children with chronic illnesses should be supported to achieve healthy mental development not only in extraordinary times such as COVID-19 Pandemic but also in daily practice. Healthcare workers can manage this issue just to work as a team due to monitor the follow-up chronic care and produce solutions for problems that do not attract attention at first glance, especially "peer problems" which could be normalized.

The issue that most of the children with chronic diseases were located in rural areas should be investigated. Community-oriented primary care could be considered as an option to solve the vulnerable social conditions.



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