ORIGINAL ARTICLE Iran J Allergy Asthma Immunol March 2012; 11(1): 57-64.

Physicians Awareness on Primary Immunodeficiency Disorders in Iran

Keramat Nourijelyani¹, Asghar Aghamohammadi², Mohammad Salehi Sadaghiani², Nasrin Behniafard², Hassan Abolhassani^{1,3}, Sarvenaz Pourjabar², Alireza Rezvanizadeh², Joobin Khadamy², Amir Imanzaeh², Mojtaba Sedaghat³, and Nima Rezaei^{2,4}

¹ Department of Epidemiology and Biostatistics, Tehran University of Medical Sciences, Tehran, Iran ² Research Center for Immunodeficiency, Pediatrics Center of Excellence, Children's Medical Center,

Tehran University of Medical Sciences, Tehran, Iran

³ Community Medicine, Tehran University of Medical Sciences, Tehran, Iran

⁴ Department of Immunology, Molecular Immunology Research Center, School of Medicine, Tehran University of Medical Sciences, Tehran, Iran

Received: 7 July 2011; Received in revised form: 13 November 2011 ; Accepted: 11 December 2011

ABSTRACT

Primary immunodeficiency diseases (PIDs) consist of a group of genetic disorders that predispose the patients to immune-mediated complications. The aim of this study was to assess the knowledge of Iranian general practitioners and pediatricians about PIDs.

A questionnaire consisting 52 closed questions on clinical symptoms, laboratory data, associated syndromes and management of PIDs patients was made valid and reliable by a pair pilot study. Then the questionnaire was filled by pediatricians, general practitioners and pediatric residents from different regions of Iran.

Totally, 333 physicians (50 general practitioners, 52 pediatric residents, 182 pediatric specialists, and 49 pediatric sub specialists) participated in this study. The mean total score was 55.9 ± 14.3 (i.e. about 29 correct answers out of 52 questions). One hundred and five participants (31.9%) answered correctly more than two third of all questions. In order to qualitatively compare the groups a ranking system was used. Total scores was significantly different between physicians groups (p<0.01). Pediatric subspecialties gained the highest rank, which was significantly over the other participants (p<0.05).

This study showed that there is a considerable lack of awareness on PIDs in physicians. This may be one of the major reasons in late diagnosis and the delay in adequate treatment deteriorating patients' morbidity and mortality. Retraining classes and reconsidered educating schedules are needed as an efficient strategies and improving physicians' knowledge about PIDs.

Keywords: Awareness; Pediatricians; Primary immunodeficiency diseases

Corresponding Author: Asghar Aghamohammadi, MD, PhD; Children's Medical Center Hospital, 62 Qarib St., Keshavarz Blvd., Tehran 14194, Iran. Tel: (+98 21) 6642 8998, Fax: (+98 21) 6692 3054, E-mail: aghamohammadi@sina.tums.ac.ir

INTRODUCTION

Primary immunodeficiency diseases (PIDs) consist of a group of genetic disorders that affect components

Copyright© 2012, IRANIAN JOURNAL OF ALLERGY, ASTHMA AND IMMUNOLOGY. All rights reserved.

of the immune system, which predispose patients to both infectious and non-infectious immune-mediated complications. $^{\rm 1-4}$

Originally, PIDs were thought to be rare, but nowadays it has become clear that they are much more common.⁵ Among all physicians, primary care physicians and pediatricians are more likely to visit patients with PIDs in their practice; therefore, they should be familiar with these life-threatening disorders.⁶

The most significant clinical presentations in PIDs are infection,^{3,7} although the rate of autoimmune diseases and malignancies are also considerable among them.^{2,8} The consequent complications may lead to decrease in quality of life and even death in PIDs patients.⁹⁻¹³ The delay in diagnosis of PIDs patients is one of the important reason in occurrence of the permanent sequels.^{14,15}

Therefore, better quality of life, longer life saving therapy and precautions sequels establishment mainly depends on early diagnosis.¹⁵⁻²⁰

Unfortunately, the diagnosis of patients with PIDs is associated with a considerable delay.²¹ One of the responsible major problems is the lack of physicians awareness about PIDs, which was particularly pertinent to developing countries.²²⁻²⁵

The aim of this study was to evaluate the knowledge and practice of Iranian physicians about PIDs.

MATERIALS AND METHODS

Study Population

Population of this study was pediatricians (specialties and subspecialties), pediatric residents and general practitioners from different parts of Iran who participated in the 21^{st} International Pediatrics Congress, October 2009 in Tehran, Iran.

Prior to data collection the study was approved in the ethic committee of the Ministry of Health in Tehran University of Medical Sciences. Demographic data, university certificate, duration of medical practice, place of medical practice, history of previous encounter with suspected or documented primary immunodeficient patients and overall score of awareness about PIDs were evaluated for each participant. The survey was done before the initiating date of immunologic conferences.

Survey Approach

To assess a score of awareness of physician about PIDs, a prototype questionnaire was prepared based on questionnaire from a similar survey in Kuwait²⁵ which translated and modified by consulting professionals in PIDs and a professional in questionnaire making. A pilot study was performed to make the questionnaire reliable and valid (alpha koronbach= 0.7961, kappa=0.8127)

The final version of questionnaire with 52 closed questions was ready containing 26 questions on the clinical presentation of PIDs, 10 questions on associated diseases and syndromes, 14 questions were on laboratory investigations (Table 1). The last two questions were on the problems of physicians in managing PIDs patients and their needs to reeducation classes. The overall score of each participant was computed by adding the correct answers to these 52 questions. Passing the exam was defined as answering more than 2/3 of the questions.²⁵ Also in order to assess qualitatively, ten different ranks were determined, including: extremely low (score less than 12.5), very low (score from 12.5 to 25), low (score from 25 to 37.5), low-medium (score from 37.5 to 50), and high-medium (score from 50 to 62.5), high (score from 62.5 to 75), very high (score from 75 to 87.5) and extremely high (score more than 87.5).

Data Analysis

The awareness scores transformed to a common 0-100 scale and the primary analyses included 333 physicians (The non-responder rate to single items was very low in total, 0.31%) who were fully compliant with the study protocol. Correlation analyses were done product moment correlation using Pearson's coefficients; statistical tests were two-tailed intra group. Pearson chi-square from crosstab was used to compare especial category with other ranks. Moreover, to handle many observations as possible, missing data for repeated measurements were imputed using an explicit regression model (i.e., repeated measure model with unstructured covariance matrix) that included previously observed scores of the participants as well as the important covariates.

Physicians Awareness on Primary Immunodeficiency Disorders

L'Clinical features Viait panets What is the most important feature in a child with PID 1.5 Malignancy 1.5 Recurrent Infections Yes 85.9 Autoimmune disease 2.7 Growth failure 3.9 Not answered 6 6 Which of the following can be a clue to PID disease 1.5 5 Lymphoid hypoplasia Yes 7.3.6 Torticollis No 41.1 Hypophyseal failure No 28.2 Eosinophili with erythrodermia Yes 58.3 Polydactylia No 31.8 Frequent common colds No 31.8 Frequent contal candidiasis at the age of two Yes 81.1 Angioedema Yes 41.7 Simultaneous existence of two internal infections Yes 43.8 Ipomphoid hyperplasia Yes 76 No 33.9 44.3 38.9 Polomocythis proveci pneumonia Yes 76 Nomati aboutian Yes 76 No andal boutian Yes 76 <	Question	Correct answer	%
Malignancy1.5Recurrent InfectionsYes85.9Autoimmune disease2.7Growth failure3.9Not answered6Which of the following can be a clue to PID disease1.1Lymphoid hypoplasiaYes73.6TorticollisNo41.1Hypophysel failureNo28.2Eosinophilia with crythrodermiaYes58.3PolydactyliaNo31.8Frequent common coldsNo18.6Frequent common coldsNo18.6Frequent cral candidiasis at the age of twoYes81.1AngioedemaYes81.1AngioedemaYes81.1AngioedemaYes83.3JohnstomorNo33.9HypoparathyroidismYes56.2Wilms tumorNo33.9HypoparathyroidismYes76Neurand botulismNo39Poliomyelitis after receiving oral polio vaccine (OPV)Yes70.6Failure to thriveYes75.True or falseYes63.1BronchicetasiaYes75.True or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedYes75.The signs or symptoms of PID patients can emerge after the 6 monthsYes76.7True or falseThe signs or symptoms of PID patients can emerge after the 6 monthsYes76.6History of 3 otitis media during childhood	I-Clinical features		
Recurrent InfectionsYes85.9Autoimmune disease2.7Growth failure3.9Not answered6Which of the following can be a clue to PID disease73.6Lymphoid hypoplasiaYes73.6TorticollisNo41.1Hypopphyseal failureNo28.2Eosinophilia with erythrodermiaYes58.3PolydactyliaNo31.8Frequent contanto coldsNo31.8Frequent contanto coldsNo31.8Frequent contanto coldsYes81.1AngicodermaYes81.1Delay in shedding the deciduous teethYes85.2Jymboid hyperplasiaYes85.2Lymphoid hyperplasiaYes85.2Lymphoid hyperplasiaYes85.2Lymphoid hyperplasiaYes76No33.933.8Pheumocystis jiroveci pneumoniaYes76NoYes7676Nonatal botulismYes76No39.276Policomyelitis after receiving oral polio vaccine (OPV)Yes76Failure to thriveYes87.3BronchicctasiaYes76Tue or falseT75Tue or falseT75Tue or falseT75Itsory of 3 otitis media during childhoodYes83.8decade of lifeYes76.3The signs or symptoms of PID patients can emerge after the 6 monthsYes83.	What is the most important feature in a child with PID		
Recurrent InfectionsYes85.9Autoimmune disease2.7Growth failure3.9Not answered6Which of the following can be a clue to PID disease73.6Lymphoid hypoplasiaYes73.6TorticollisNo41.1Hypopphyseal failureNo28.2Eosinophilia with erythrodermiaYes58.3PolydactyliaNo31.8Frequent contanto coldsNo31.8Frequent contanto coldsNo31.8Frequent contanto coldsYes81.1AngicodermaYes81.1Delay in shedding the deciduous teethYes85.2Jymboid hyperplasiaYes85.2Lymphoid hyperplasiaYes85.2Lymphoid hyperplasiaYes85.2Lymphoid hyperplasiaYes76No33.933.8Pheumocystis jiroveci pneumoniaYes76NoYes7676Nonatal botulismYes76No39.276Policomyelitis after receiving oral polio vaccine (OPV)Yes76Failure to thriveYes87.3BronchicctasiaYes76Tue or falseT75Tue or falseT75Tue or falseT75Itsory of 3 otitis media during childhoodYes83.8decade of lifeYes76.3The signs or symptoms of PID patients can emerge after the 6 monthsYes83.	Malignancy		1.5
Growth failure3.9Not answered6Which of the following can be a clue to PID disease5Lymphoid hypoplasiaYesTorticollisNoHypophyseal failureNoEssinophilia with erythroderniaYesPolydacytliaNoPolydacytliaNoFrequent common coldsNoFrequent condication coldsNoPrequent condication coldsYesPolydacytliaYesMore than 3 weeks delay in umbilical cord separationYesAngioedemaYesDelay in shedding the deciduous teethYesSimultaneous existence of two internal infectionsYesYes85Lymphoid hyperplasiaYesWins tumorNoNo33.9Poliomyelitis after receiving oral polio vaccine (OPV)YesParial albinismYesPoliomyelitis after receiving oral polio vaccine (OPV)YesParial albinismYesTrue or falseTesThe signs or symptoms of PID patients can emerge after the 6 monthsYesPasis or symptoms of PID patients can emerge after the 6 monthsYesThe signs or symptoms of PID patients can emerge after the 6 monthsYesPistis receiving in patients can emerge after the 6 monthsYesPistis receiving so for PID patients can emerge after the 6 monthsYesPistis modia syndromeNo32.9Which of the following is associated with PIDElder-Danlos syndromeNo34.5W	Recurrent Infections	Yes	85.9
Not answered6Which of the following can be a clue to PID diseaseLymphoid hypoplasiaYes73.6TorticollisNo41.1Hypophyseal failureNo28.2Eosinophilia with erythrodermiaYes58.3PolydactyliaNo31.8Frequent common coldsNo18.6Frequent oral candidiasis at the age of twoYes91More than 3 weeks delay in umbilical cord separationYes47.1Delay in shedding the deciduous teethYes47.1Delay in shedding the deciduous teethYes47.1Delay in shedding the deciduous teethYes85Lymphoid hyperplasiaYes56.2Wilms tumorNo33.9HypoparathyroidismYes76Nonatal botulismYes76Nonatal botulismYes76.2Poliomyelitis after receiving oral polio vaccine (OPV)Yes70.6Failure to thriveYes75.True or falseT75True or falseT75The signs or symptoms of PID patients can emerge after the 6 monthsYes43.8decade of lifeT7443.5The signs or symptoms of PID patients can emerge after the 6 monthsYes43.8The signs or symptoms of PID patients can emerge after the 6 monthsYes43.8Acacade of lifeT7443.5The signs or symptoms of PID patients can emerge after the 6 monthsYes43.8Acacade of li	Autoimmune disease		2.7
Which of the following can be a clue to PID diseaseLymphoid hypoplasiaYes73.6TorticollisNo24.1Hypophyseal failureNo28.2Eosinophilia with erythrodermiaYes58.3PolydactyliaNo31.8Frequent common coldsNo18.6Frequent common coldsNo18.6Frequent oral candidiasis at the age of twoYes91More than 3 weeks delay in umbilical cord separationYes81.1AngioodemaYes41.7Delay in shedding the deciduous teethYes43.8Pheumocystis jirovcei pneumoniaYes56.2Lymphoid hyperplasiaYes56.2Uhins tumorNo33.9HypoparathyroidismYes70.6Failure to thriveYes70.6Failure to thriveYes70.6Failure to thriveYes43.8Pheumocystis jirovcei pneumoniaYes70.6Failure to thriveYes70.6Failure to thriveYes70.6 <t< td=""><td>Growth failure</td><td></td><td>3.9</td></t<>	Growth failure		3.9
Lymphoid hypoplasia Yes 73.6 Torticollis No 41.1 Hypophyseal failure No 28.2 Eosinophilia with erythrodermia Yes 58.3 Polydactylia No 31.8 Frequent common colds No 18.6 Frequent common colds No 18.6 Frequent calciditasis at the age of two Yes 91 More than 3 weeks delay in umbilical cord separation Yes 81.1 Angioedema Yes 71.1 Delay in shedding the deciduous teeth Yes 71.7 Simultaneous existence of two internal infections Yes 85 Lymphoid hyperplasia Yes 56.2 Wilms tumor No 33.9 Hypoparatyroidism Yes 43.8 Pneumocystis jiroveci pneumonia Yes 76 No 33.9 Hypoparatyroidism Yes 76 Noon 319 Pholomyeltis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 63.1 Bronchiectasia Yes 75 True or false The signs or symptoms of PID patients can emerge after the 6 months Yes 80.8 of age, when the maternal antibodies are diminished The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome Yes 72.4 Hyporelanosis of tio No 17.4 Sturge-Weber syndrome Yes 72.4 Hyporelanosis of tio No 17.4 Sturge-Weber syndrome Yes 75.7 Turter Syndr	Not answered		6
Lymphoid hypoplasia Yes 73.6 Torticollis No 41.1 Hypophyseal failure No 28.2 Eosinophilia with erythrodermia Yes 58.3 Polydactylia No 31.8 Frequent common colds No 18.6 Frequent common colds No 18.6 Frequent calciditasis at the age of two Yes 91 More than 3 weeks delay in umbilical cord separation Yes 81.1 Angioedema Yes 71.1 Delay in shedding the deciduous teeth Yes 71.7 Simultaneous existence of two internal infections Yes 85 Lymphoid hyperplasia Yes 56.2 Wilms tumor No 33.9 Hypoparatyroidism Yes 43.8 Pneumocystis jiroveci pneumonia Yes 76 No 33.9 Hypoparatyroidism Yes 76 Noon 319 Pholomyeltis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 63.1 Bronchiectasia Yes 75 True or false The signs or symptoms of PID patients can emerge after the 6 months Yes 80.8 of age, when the maternal antibodies are diminished The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome Yes 72.4 Hyporelanosis of tio No 17.4 Sturge-Weber syndrome Yes 72.4 Hyporelanosis of tio No 17.4 Sturge-Weber syndrome Yes 75.7 Turter Syndr	Which of the following can be a clue to PID disease		
Hypophyseal failureNo28.2Eosinophilia with erythrodermiaYes58.3PolydactyliaNo31.8Frequent common coldsNo31.8Frequent common coldsNo18.6Frequent oral candidiasis at the age of twoYes91More than 3 weeks delay in umbilical cord separationYes47.1Dolay in shedding the deciduous teethYes41.7Simultaneous existence of two internal infectionsYes45.2Lymphoid hyperplasiaYes56.2Wilms tumorNo33.9HypoparathyroidismYes76Neonatal botulismYes76Neonatal botulismYes70.6Failure to thriveYes70.6Failure to thriveYes70.6Failure to thriveYes70.6Failure to thriveYes73.1BronchicetasiaYes74.4Eczema and subcutaneous bleedingYes75.True or falseTTThe signs or symptoms of PID patients can emerge after the 6 monthsYes66.7birthThe signs or symptoms of PID patients can emerge from the time ofYes72.4Hypomelanosis of tioNo17.4Sturge-Weer syndrome78.4Ataxia-TelangiectasiaYes72.443.8In Associated symptoms and diseasesWich of the following is associated with PIDYes72.4Hypomelanosis of itoNo17.4Sturge-Weer syndromeYes72.4 </td <td>Lymphoid hypoplasia</td> <td>Yes</td> <td>73.6</td>	Lymphoid hypoplasia	Yes	73.6
Polydactylia with erythrodermia Yes 58.3 Polydactylia No 31.8 Frequent common colds No 18.6 Frequent common colds Pres 91 More than 3 weeks delay in umbilical cord separation Yes 91 More than 3 weeks delay in umbilical cord separation Yes 81.1 Angioedema Yes 41.7 Delay in shedding the deciduous teeth Yes 41.7 Simultaneous existence of two internal infections Yes 41.7 Simultaneous existence of two internal infections Yes 56.2 Wilms tumor No 33.9 Hypoparathyroidism Yes 76. Neonatal botulism Yes 76 Neonatal botulism No 339 Poliomyelitis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 ottis media during childhood No 29.1 Partial albinism Yes 75 True or false Yes 75 True or false Yes 75 The signs or symptoms of PID patients can emerge after the 6 months Yes 63.1 Bronchiectasia Yes 75 The signs or symptoms of PID patients can emerge during the third Yes 63.7 Briss or symptoms of PID patients can emerge during the third Yes 63.7 Directing the term of Yes 77. The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth L-Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome Yes 77.8 Ataxia-Telangiectasia Yes 77.7 Ataxia-Telangiectasia Yes 77.7 Turner syndrome Yes 77.7 Turner syndrome Yes 77.7 Turner syndrome Yes 77.7	Torticollis	No	41.1
Polydactylia No 31.8 Frequent common colds No 18.6 Frequent common colds Yes 91 More than 3 weeks delay in umbilical cord separation Yes 91 More than 3 weeks delay in umbilical cord separation Yes 41.7 Delay in shedding the deciduous teeth Yes 41.7 Simultaneous existence of two internal infections Yes 85 Lymphoid hyperplasia Yes 56.2 Wilms tumor No 33.9 Hypoparathyroidism Yes 76 Noo 33.9 Hypoparathyroidism Yes 76 Noo 39 Poliomyelitis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 63.1 Bronchiectasia Yes 75 True of false Yes 75 True of false Yes 75 True of false The signs or symptoms of PID patients can emerge after the 6 months Yes 43.8 decade of life The signs or symptoms of PID patients can emerge after the 6 months Yes 75 H - Associated symptoms and diseases Which of the following is associated with PID Eher-Danlos syndrome Yes 77.8 Ataxia-Telangiectasia Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 74.4 Sturge-Weber syndrome Yes 77.8 Ataxia-Telangiectasia Yes 77.7 Ataxia-Telangiectasia Yes 77.7 Ataxia-Telangiectasia Yes 77.8 Ataxia-Telang	Hypophyseal failure	No	28.2
Polydactylia No 31.8 Frequent common colds No 18.6 Frequent common colds Yes 91 More than 3 weeks delay in umbilical cord separation Yes 91 More than 3 weeks delay in umbilical cord separation Yes 41.7 Delay in shedding the deciduous teeth Yes 41.7 Simultaneous existence of two internal infections Yes 85 Lymphoid hyperplasia Yes 56.2 Wilms tumor No 33.9 Hypoparathyroidism Yes 76 Noo 33.9 Hypoparathyroidism Yes 76 Noo 39 Poliomyelitis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 63.1 Bronchiectasia Yes 75 True of false Yes 75 True of false Yes 75 True of false The signs or symptoms of PID patients can emerge after the 6 months Yes 43.8 decade of life The signs or symptoms of PID patients can emerge after the 6 months Yes 75 H - Associated symptoms and diseases Which of the following is associated with PID Eher-Danlos syndrome Yes 77.8 Ataxia-Telangiectasia Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 74.4 Sturge-Weber syndrome Yes 77.8 Ataxia-Telangiectasia Yes 77.7 Ataxia-Telangiectasia Yes 77.7 Ataxia-Telangiectasia Yes 77.8 Ataxia-Telang	Eosinophilia with erythrodermia	Yes	58.3
Frequent common coldsNo18.6Frequent oral candidiasis at the age of twoYes91More than 3 weeks delay in umbilical cord separationYes81.1AngioedemaYes41.7Delay in shedding the deciduous teethYes41.7Diaty in shedding the deciduous teethYes85.Lymphoid hyperplasiaYes56.2Wilms tumorNo33.9HypoparathyroidismYes76Neonatal botuismYes76Neonatal botuismYes76Neonatal botuismNo39Poliomyelitis after receiving oral polio vaccine (OPV)Yes70.6Failure to thriveYes47.4Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes75True or falseT75The signs or symptoms of PID patients can emerge after the 6 monthsYes43.8decade of lifeT77.843.8decade of lifeT77.843.8decade of lifeYes77.877.8Hastorick symptoms and diseasesYes77.877.8Mixhch Aldrich syndromeYes77.877.8Ataxia-TelangiectasiaYes72.479.1Kiskott-Aldrich syndromeYes76.877.8Ataxia-TelangiectasiaYes72.477.8Ataxia-TelangiectasiaYes72.877.8Ataxia-TelangiectasiaYes72.877.8Ataxia-Telangiectasia<		No	31.8
Frequent oral candidiasis at the age of two Yes 91 More than 3 weeks delay in umbilical cord separation Yes 81.1 Angioedema Yes 47.1 Delay in shedding the deciduous teeth Yes 41.7 Simultaneous existence of two internal infections Yes 85 Lymphoid hyperplasia Yes 56.2 Wilns tumor No 33.9 Hypoparathyroidism Yes 76 Neonatal botulism Yes 76 Neonatal botulism No 39 Poliomyelitis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 75 True or false 75 True or false 75 The signs or symptoms of PID patients can emerge after the 6 months Yes 80.8 of age, when the maternal antibodies are diminished 75 The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth H- Associated symptomes and diseases Which of the following is associated with PID Ehler-Danlos syndrome No 34.5 Wikkott-Aldrich syndrome No 34.5 Turge Veber syndrome No 34.5 Sturge-Weber syndrome Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 66.7 Dirthe Sturge Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 66.7 Dirthe Sturge Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 65.7 Turrer syndrome Yes 77.7 Turrer S		No	18.6
More than 3 weeks delay in umbilical cord separationYes81.1AngioedemaYes47.1Delay in shedding the deciduous teethYes47.1Ding in shedding the deciduous teethYes41.7Simultaneous existence of two internal infectionsYes85Lymphoid hyperplasiaYes56.2Wilms tumorNo33.9HypoparathyroidismYes43.8Pneumocystis jiroveci pneumoniaYes76Neonatal botulismNo39Poliomyelitis after receiving oral polio vaccine (OPV)Yes70.6Failure to thriveYes82.3History of 3 otitis media during childhoodNo29.1Partial albinismYes47.4Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes75True or falseThe signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedYes43.8the signs or symptoms of PID patients can emerge from the time of birthYes77.8Hacade of lifeIfe14.4The signs or symptomes and diseasesYes77.8Which of the following is associated with PID Ether-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes76.2Myenelanosis of itoNo17.4Sturge-Weber syndromeYes76.5Bardet-Biedle syndromeYes57.7Intrace syndromeYes57.7 </td <td>-</td> <td>Yes</td> <td>91</td>	-	Yes	91
AngioedemaYes47.1Delay in shedding the deciduous teethYes41.7Simultaneous existence of two internal infectionsYes85Lymphoid hyperplasiaYes56.2Wilms tumorNo33.9HypoparathyroidismYes43.8Pneumocystis jirvoeci pneumoniaYes76Neonatal botulismYes76Poliomyelitis after receiving oral polio vaccine (OPV)Yes70.6Failure to thriveYes82.3History of 3 otitis media during childhoodNo29.1Partial albinismYes47.4Eczema and subcutaneous bleedingYes75True or falseYes75The signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedYes43.8the signs or symptoms of PID patients can emerge during the thirdYes43.8decade of lifeThe signs or symptoms and diseasesWhich of the following is associated with PIDEhler-Danlos syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Bardet-Biedle syndromeYes57.7IndicationYes57.7Turger SyndromeYes57.7Turger SyndromeYes57.7Turger SyndromeYes57.7Intrace SyndromeYes57.7Turger SyndromeYes			81.1
Delay in shedding the deciduous teeth Yes 41.7 Simultaneous existence of two internal infections Yes 85 Lymphoid hyperplasia Yes 56.2 Wilms tumor No 33.9 Hypoparathyroidism Yes 43.8 Pneumocystis jiroveci pneumonia Yes 76 Neonatal botulism No 39 Poliomyelitis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 47.4 Eczema and subcutaneous bleeding Yes 63.1 Bronchiectasia Yes 75 True or false Yes 75 True or false State and subcutaneous bleeding Yes 63.1 Bronchiectasia Yes 75 True or false He maternal antibodies are diminished The signs or symptoms of PID patients can emerge after the 6 months Yes 60.8 decade of life Yes 64.7 He signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth He signs or symptoms of PID patients can emerge from the time of Yes 77.8 Ataxia-Telangiectasia Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 46.5 Bardet-Biedle syndrome Yes 46.5 Bardet-Biedle syndrome Yes 57.7 Turner syndrome Yes 57.7			
Simultaneous existence of two internal infectionsYes85Lymphoid hyperplasiaYes56.2Wilms tumorNo33.9HypoparathyroidismYes43.8Pneumocystis jiroveci pneumoniaYes76Neonatal botulismNo39Poliomyelitis after receiving oral polio vaccine (OPV)Yes70.6Failure to thriveYes82.3History of 3 otitis media during childhoodNo29.1Partial albinismYes47.4Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes75True or falseT75The signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedT43.8The signs or symptoms of PID patients can emerge during the thirdYes43.8decade of lifeII- Associated symptoms and diseasesWhich of the following is associated with PIDEhler-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Bardet-Biedle syndromeYes57.7Turrer syndromeYes57.7Turrer syndromeYes57.7Turrer syndromeYes57.7			
Lymphoid hyperplasia Yes 56.2 Wilms tumor No 33.9 Hypoparathyroidism Yes 43.8 Pneumocystis jiroveci pneumonia Yes 76 Neonatal botulism Yes 76 Neonatal botulism Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 47.4 Eczema and subcutaneous bleeding Yes 63.1 Bronchiectasia Yes 75 True or false The signs or symptoms of PID patients can emerge after the 6 months Yes 80.8 of age, when the maternal antibodies are diminished The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth IL Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome No 34.5 Wiskott-Aldrich syndrome Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 46.5 Bardet-Biedle syndrome Yes 46.5 Bardet-Biedle syndrome Yes 46.5 Bardet-Biedle syndrome Yes 57.7 Turner syndrome Yes 57.7			
Wilms tumorNo33.9HypoparathyroidismYes43.8Pneumocystis jiroveci pneumoniaYes76Neonatal botulismNo39Poliomyelitis after receiving oral polio vaccine (OPV)Yes70Failure to thriveYes82.3History of 3 otitis media during childhoodNo29.1Partial albinismYes47.4Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes75True or falseYes75The signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedYes43.8decade of lifeYes66.7The signs or symptoms of PID patients can emerge from the time ofYes66.7birthH- Associated symptoms and diseasesWhich of the following is associated with PIDFes77.8Ataxia-TelangiectasiaYes72.474.874.8Kusture-Ibanios sof fitoNo17.474.8Sturge-Weber syndromeNo29.174.874.8Kustar-ElangiectasiaYes75.874.8Hatxia-TelangiectasiaYes74.874.8Sturge-Weber syndromeNo29.174.9Kostman syndromeYes46.575.7Bardet-Biedle syndromeYes57.777.7Turner syndromeYes57.777.7Turner syndromeYes57.777.7Turner synd			
Hypoparathyroidism Yes 43.8 Pneumocystis jiroveci pneumonia Yes 76 Neonatal botulism No 39 Poliomyelitis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 63.1 Bronchiectasia Yes 63.1 Bronchiectasia Yes 75 True or false Yes 80.8 of age, when the maternal antibodies are diminished Yes 80.8 of age, when the maternal antibodies are diminished Yes 66.7 birth Signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome No 34.5 Wiskott-Aldrich syndrome Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 46.5 Bardet-Biedle syndrome Yes 77.7 Turner Syndrome Yes 77.7 Turne			
Pheumocystis jiroveci pneumonia Yes 76 Neonatal botulism No 39 Poliomyelitis after receiving oral polio vaccine (OPV) Yes 70.6 Failure to thrive Yes 82.3 History of 3 otitis media during childhood No 29.1 Partial albinism Yes 47.4 Eczema and subcutaneous bleeding Yes 63.1 Bronchiectasia Yes 75 True or false The signs or symptoms of PID patients can emerge after the 6 months Yes 80.8 of age, when the maternal antibodies are diminished The signs or symptoms of PID patients can emerge during the third Yes 43.8 decade of life The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 46.5 Bardet-Biedle syndrome Yes 57.7 Turner syndrome Yes 57.7			
Neonatal botulismNo39Poliomyelitis after receiving oral polio vaccine (OPV)Yes70.6Failure to thriveYes82.3History of 3 otitis media during childhoodNo29.1Partial albinismYes47.4Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes63.1BronchiectasiaYes75True or falseTTThe signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedYes43.8decade of lifeTYes66.7birthYes66.75birthH- Associated symptoms and diseasesYes77.8Which of the following is associated with PIDYes77.8Ehler-Danlos syndromeYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Bardet-Biedle syndromeYes57.7No20.710b's syndromeYesKostman syndromeYes57.7Turner syndromeYes57.7Turner syndromeYes57.7Turner syndromeNo48			
Poliomyelitis after receiving oral polio vaccine (OPV)Yes70.6Failure to thriveYes82.3History of 3 otitis media during childhoodNo29.1Partial albinismYes47.4Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes75True or falseYes80.8The signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal attibodies are diminishedYes43.8decade of lifeYes66.7The signs or symptoms of PID patients can emerge from the time ofYes66.7birthII- Associated symptoms and diseasesYes71.8Which of the following is associated with PIDYes72.4Hypomelanosis of itoNo17.417.4Sturge-Weber syndromeYes46.529.1Kostman syndromeYes46.520.7Job's syndromeYes57.777.7Turner syndromeYes57.777.7 <td></td> <td></td> <td></td>			
Failure to thriveYes82.3History of 3 otitis media during childhoodNo29.1Partial albinismYes47.4Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes75True or falseYes75The signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedYes43.8decade of lifeYes43.843.8decade of lifeYes66.75The signs or symptoms of PID patients can emerge from the time of birthYes66.7LI Associated symptoms and diseasesYes77.8Which of the following is associated with PIDYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Barder-Biedle syndromeYes46.5Barder-Biedle syndromeYes57.7Turner s			
History of 3 otitis media during childhood No 29.1 Partial albinism Yes 47.4 Eczema and subcutaneous bleeding Yes 63.1 Bronchiectasia Yes 75 True or false The signs or symptoms of PID patients can emerge after the 6 months Yes 80.8 of age, when the maternal antibodies are diminished The signs or symptoms of PID patients can emerge during the third Yes 43.8 decade of life The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome No 34.5 Wiskott-Aldrich syndrome Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome Yes 46.5 Bardet-Biedle syndrome Yes 57.7 Turner syndrome Yes 57.7 Turner syndrome Yes 57.7 Turner syndrome No 48			
Partial albinismYes47.4Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes75True or falseYes75The signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedYes43.8decade of lifeYes43.8The signs or symptoms of PID patients can emerge during the thirdYes43.8decade of lifeYes66.7birthII- Associated symptoms and diseasesKothich of the following is associated with PIDEhler-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Bardet-Biedle syndromeYes46.5Bardet-Biedle syndromeYes57.7Turner syndromeYes57.7Turner syndromeYes57.7Chediak-Higashi syndromeYes57.7Chediak-Higashi syndromeYes57.7			
Eczema and subcutaneous bleedingYes63.1BronchiectasiaYes75True or falseYes80.8of age, when the maternal antibodies are diminishedYes80.8of age, when the maternal antibodies are diminishedYes43.8decade of lifeYes66.7The signs or symptoms of PID patients can emerge from the time ofYes66.7birthYes66.7LI- Associated symptoms and diseasesWhich of the following is associated with PIDYes75Ehler-Danlos syndromeNo34.575Wiskott-Aldrich syndromeYes77.878Ataxia-TelangiectasiaYes72.479Hypomelanosis of itoNo17.429.1Kostman syndromeYes46.520.7Job's syndromeYes7577.7Turner syndromeNo20.720.7Cherdiak-Hizashi syndromeYes57.777.7Cherdiak-Hizashi syndromeYes57.7			
BronchiectasiaYes75True or falseYes80.8The signs or symptoms of PID patients can emerge after the 6 monthsYes80.8of age, when the maternal antibodies are diminishedYes43.8The signs or symptoms of PID patients can emerge during the thirdYes43.8decade of lifeYes66.7The signs or symptoms of PID patients can emerge from the time ofYes66.7birthII- Associated symptoms and diseasesYes77.8Hich of the following is associated with PIDYes77.8Ehler-Danlos syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Bardet-Biedle syndromeYes46.5Bardet-Biedle syndromeYes57.7Job's syndromeYes57.7Turner syndromeYes57.7Chediak–Hizashi syndromeYes57.7			
True or falseYes80.8The signs or symptoms of PID patients can emerge after the 6 months of age, when the maternal antibodies are diminishedYes80.8The signs or symptoms of PID patients can emerge during the third decade of lifeYes43.8The signs or symptoms of PID patients can emerge from the time of birthYes66.7II- Associated symptoms and diseasesWhich of the following is associated with PIDEhler-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Bardet-Biedle syndromeYes57.7Turner syndromeYes57.7Turner syndromeYes57.7No4814			
The signs or symptoms of PID patients can emerge after the 6 months of age, when the maternal antibodies are diminishedYes80.8The signs or symptoms of PID patients can emerge during the third decade of lifeYes43.8The signs or symptoms of PID patients can emerge from the time of birthYes66.7 II- Associated symptoms and diseases Which of the following is associated with PIDEhler-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Bardet-Biedle syndromeYes57.7Job's syndromeYes57.7Turner syndromeYes57.7Chediak–Hirgashi syndromeNo48		168	75
of age, when the maternal antibodies are diminished The signs or symptoms of PID patients can emerge during the third Yes 43.8 decade of life The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome No 34.5 Wiskott-Aldrich syndrome Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome No 29.1 Kostman syndrome Yes 46.5 Bardet-Biedle syndrome Yes 57.7 Turner syndrome No 48 Chediak–Higgsbi syndrome		Vac	<u>00 0</u>
The signs or symptoms of PID patients can emerge during the third decade of lifeYes43.8The signs or symptoms of PID patients can emerge from the time of birthYes66.7 II- Associated symptoms and diseasesWhich of the following is associated with PID Ehler-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeYes46.5Bardet-Biedle syndromeYes57.7Job's syndromeYes57.7Turner syndromeNo48Chediak-Higashi syndromeNo48		168	80.8
decade of life The signs or symptoms of PID patients can emerge from the time of Yes 66.7 birth II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome No 34.5 Wiskott-Aldrich syndrome Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome No 29.1 Kostman syndrome Yes 46.5 Bardet-Biedle syndrome Yes 57.7 Turner syndrome No 48 Chediak–Higgsbi syndrome		Vac	12.0
The signs or symptoms of PID patients can emerge from the time of birthYes66.7 II- Associated symptoms and diseasesWhich of the following is associated with PID VEhler-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeNo29.1Kostman syndromeYes46.5Bardet-Biedle syndromeYes57.7Turner syndromeYes57.7Chediak–Higashi syndromeNo48		res	43.8
birth II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome No 34.5 Wiskott-Aldrich syndrome Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome No 29.1 Kostman syndrome Yes 46.5 Bardet-Biedle syndrome Yes 57.7 Turner syndrome No 48 Chediak–Higashi syndrome		V	((7
II- Associated symptoms and diseases Which of the following is associated with PID Ehler-Danlos syndrome No 34.5 Wiskott-Aldrich syndrome Yes 77.8 Ataxia-Telangiectasia Yes 72.4 Hypomelanosis of ito No 17.4 Sturge-Weber syndrome No 29.1 Kostman syndrome Yes 46.5 Bardet-Biedle syndrome No 20.7 Job's syndrome Yes 57.7 Turner syndrome No 48 Chediak–Higashi syndrome No 48		res	00.7
Which of the following is associated with PIDEhler-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeNo29.1Kostman syndromeYes46.5Bardet-Biedle syndromeNo20.7Job's syndromeYes57.7Turner syndromeNo48Chediak–Higashi syndromeKostman			
Ehler-Danlos syndromeNo34.5Wiskott-Aldrich syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeNo29.1Kostman syndromeYes46.5Bardet-Biedle syndromeNo20.7Job's syndromeYes57.7Turner syndromeNo48Chediak-Higashi syndromeYes			
Wiskott-Aldrich syndromeYes77.8Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeNo29.1Kostman syndromeYes46.5Bardet-Biedle syndromeNo20.7Job's syndromeYes57.7Turner syndromeNo48Chediak-Higashi syndromeVes		N	24.5
Ataxia-TelangiectasiaYes72.4Hypomelanosis of itoNo17.4Sturge-Weber syndromeNo29.1Kostman syndromeYes46.5Bardet-Biedle syndromeNo20.7Job's syndromeYes57.7Turner syndromeNo48Chediak-Higashi syndromeYes			
Hypomelanosis of itoNo17.4Sturge-Weber syndromeNo29.1Kostman syndromeYes46.5Bardet-Biedle syndromeNo20.7Job's syndromeYes57.7Turner syndromeNo48Chediak-Higashi syndromeVes	-		
Sturge-Weber syndromeNo29.1Kostman syndromeYes46.5Bardet-Biedle syndromeNo20.7Job's syndromeYes57.7Turner syndromeNo48Chediak-Higashi syndromeVes	-		
Kostman syndromeYes46.5Bardet-Biedle syndromeNo20.7Job's syndromeYes57.7Turner syndromeNo48Chediak-Higashi syndromeKes57.7			
Bardet-Biedle syndromeNo20.7Job's syndromeYes57.7Turner syndromeNo48Chediak-Higashi syndromeNo48			
Job's syndrome Yes 57.7 Turner syndrome No 48 Chediak-Higashi syndrome	•		
Turner syndrome No 48 Chediak–Higashi syndrome			
Chediak-Higashi syndrome	Job's syndrome	Yes	
Chediak–Higashi syndrome Ves 82.6	Turner syndrome	No	48
	Chediak–Higashi syndrome	Yes	82.6

Table 1. The questions and scores*

T7	N T 1		
ĸ	Nouru	alvoni	of ol
IX.	INDUIT	jelyani,	ci ai.

III- Laboratory findings		
Which of the following directly helps us in diagnosis a PID patients		
Lymphocyte stimulation tests	Yes	77.2
Fecal occult blood test	No	42.3
Antibacterial antibody response to previous vaccines	Yes	71.2
Blood urea nitrogen, creatinine	No	48.3
Determining superficial markers of lymphocytes	Yes	82.3
Anemia panel	No	26.1
Complete blood count and differential	Yes	78.1
Serum isohemagglutinins	Yes	57.4
Hepatic function panel	No	31.8
Candida and tetanus skin test	Yes	73.3
Which of the following can be a clue in diagnosing a PID patient		
The count of blood eosinophils in a child with one and a half years of	Yes	58
age equals to 15,500		
Small platelets and thrombocytopenia	Yes	61
Serum IgG concentration in an infant with 7 months of age equals to	No	18.6
420 mg/dl		
Large granules in neutrophils	Yes	70.9
IV -Managing PID patients		
Which of the following vaccines should not be administered in a child w	vith PID	
Influenza A vaccine		
BCG	Yes	74.2
IPV		
Hepatitis B vaccine		
Which of the following medications decreases rate of infections	in child with c	common variable
immunodeficiency		
Immunoglobulin replacement therapy	Yes	63.7
Recombinant interferon		
Recurrent blood transfusion		
Plasmapheresis		
Do you have difficulties in managing patients with PID	Yes	86.2
Is retraining classes regarding the PID syndromes necessary for	Yes	95.8
general practitioners and specialists		
* The second seco		

* The score of each question is 100/52

RESULTS

A total of 333 pediatricians (50 general practitioners, 52 pediatric residents, 182 pediatric specialists, and 49 pediatric sub specialists) were included in the study which 61% of them were male. The median age of participants was 44 (range 26-88) years; the median years of practicing medicine was 16 (1-48) years.

Most of the participants (55.8%) worked in the state hospitals; 20.1% worked in their private clinics; 6.6% worked in non-state hospitals; Remaining participants (17.5%) worked in more than one center and had overlap between state, non-state hospitals and clinics. Nineteen percent of them were also academic staff in medical Universities. Most of the participants (252 persons=75.7%) had visited at least one suspected or documented PIDs case during their practice.

The mean total knowledge score was 55.9 with a standard deviation of 14.3. One hundred and five participants (31.9%) answered correctly more than 2/3 of all questions and passed the exam. The best scores were documented in management of PIDs ($68.9\pm1.32\%$), which followed by laboratory findings ($56.9\pm5.4\%$), clinical symptoms ($57.3\pm9.78\%$) and associated syndromes ($48.7\pm5.3\%$) respectively. Total scores of physicians were 46.4 ± 13.7 for general practitioners, 54.8 ± 14.8 for pediatric specialties, 61.5 ± 18.4 for pediatric residents, and 63.8 ± 14.5 for pediatric subspecialties. The scores were found to be independent of gender (p=0.54).

According to the mentioned qualitative ranking system, the performance of the groups involved in this study is demonstrated in table 2. Based upon the qualitative assessment, the rank between different groups of physicians was significantly different (p<0.01). General practitioner perceived the lowest rank as "low- medium", pediatric residents along with pediatric specialists remained in the "high-medium" group and sub-specialist gained the "high" rank. The subspecialists' rank significantly was above practitioners (p<0.01) and pediatric specialists

(*p*<0.05). Moreover the rank of residents was significantly more than general practitioners (*p*< 0.01). Furthermore, visiting 6 or more than 6 patients significantly increased the rank from low-medium to high (*p*<0.05). Moreover, working at state hospitals significantly was associated with higher rank (*p*<0.05). The period of time passed from graduation of physician had reverse association with their scores (r=-0.26, *p*<0.001) especially in the scores of associated syndromes (r=-0.75, *p*<0.001) (Table 2)."

Age group	Number (%)	Mean of scores (±SD)	Qualitative ranking	Post Hoc P value	P value
≤29 years old	32 (9.6%)	56.5 ± 10.3	High-medium	-	< 0.001
30-39 years old	97 (29.1%)	63.7 ± 8.9	High	(with more than 60<0.001)	
40-49 years old	112 (33.6%)	59.8 ± 8.8	High-medium	-	
50-59 years old	46 (13.8%)	53.3 ± 9.8	High -medium	-	
≥60 years old	46 (13%)	50.2 ± 6.5	High -medium	(with 30-39<0.001)	
Sex					
Male	204(61.3%)	56.2 ±10.0	High-medium	-	0.54
Female	129(38.7%)	55.4 ± 9.2	High-medium	-	
Place of medical practice					
A (Only in Governmental hospital)	188(55.8%)	62.4 ± 9.0	High-medium	(with G<0.001)	< 0.001
B (Only in Private hospital)	22(6.6%)	54.3±10.3	High-medium	-	
C (Only in Private office)	67(20.1%)	53.6±9.6	High-medium	-	
D (Governmental hospital and Private hospital)	8(2.4%)	64.7±5.6	High	(with G<0.001)	
E (Governmental hospital and Private office)	23(9.4%)	63.0±10.0	High	(with G<0.001)	
F (Private hospital and Private office)	14(4.2%)	50.6±8.8	High-medium	-	
G (Governmental hospital and Private hospital	5(1.5%)	42.4±3.8	Low-medium	(with A<0.001) (with	
and Private office)				D<0.001) (with E<0.001)	
University certificate					
General practitioner	50(15%)	46.4 ±13.7	Low-medium	(with resident<0.01) (with SS= 0.001)	<0.01
Pediatric specialist	182(54.7%)	54.8± 14.8	High-medium	(with SS<0.05)	
Sub-specialists	49(14.7%)	63.8±14.5	High	(with GP*=0.001) (with specialist<0.05)	
Pediatric resident	52(15.6%)	61.5±18.4	High-medium	(with GP<0.01)	
Being faculty member					
Yes	63(18.9%)	56.52±18.6	High-medium	-	0.78
No	270(81.1%)	54.1±18.1	High-medium	-	
Previous encounter with suspected or			-		
documented primary immunodeficient patients					
<6 patients	206 (61.8%)	43.1±17.5	Low-medium	-	0.01
>6 patients	46 (13.8%)	69.2±15.1	High	-	

Table 2. Comparison of awareness score in	different groups of 333 Iranian physician

GP: General Practitioner; SS: Sub-specialists

DISCUSSION

PIDs are a group of inherited primarily disorders of the immune component system.^{8,26,27} Among 180 distinct PIDs, knowledge about 20 most prevalent diseases can account for >90% of cases. The disorders vary in the severity and spectrum of symptoms, but without effective and early treatments, they can be fatal. A high index of suspicion and prompt diagnosis can lead to lifesaving treatment and substantial improvement in quality of life for persons with PIDs.

Despite advances in new molecular techniques on human genomics for identification of the responsible gene defects and in development of new therapeutic methods such as gene therapy,^{16,18,28-30} there are many lack in the public health intervention for this group of diseases.

However, appropriate defining characteristics of PIDs by common feature of increased susceptibility to chronic and recurrent infections make them candidates for a more public health attention. Prompt diagnosis and treatment of PIDs patients can be lifesaving and result in marked improvements in the quality and length of life. Therefore the foundation for a public health intervention to improve the health status of persons with PIDs is increase in accuracy of diagnostic methods; and the efficacy of early interventions. Additional obstacles include the difficulty of diagnosis in the absence of a high index of suspicion and the lack of awareness among health-care providers, which impedes the timely recognition of affected persons.

To address these impediments and improve health outcomes among patients with PIDs, Centers for Disease Control and Prevention (CDC) and associates have adapted a population-based public health framework developed as part of CDC's strategic plan for genomics and public health, for the problem of PIDs (Available at *http://www.cdc.gov/genomics/ about/strategic.htm*).

In November 2001, CDC convened a multidisciplinary panel of specialists to identify and discuss public health strategies that can be applied to PIDs (Available at *http://www.cdc.gov/genomics/ info/conference/PIsynop.htm*).

During the meeting, specialists in clinical immunology, public health, genetics, pediatrics, health communication, and ethics from state and federal agencies, academic centers, professional organizations, and advocacy foundations discussed the public health framework relating to PIDs. The framework has four components as follows: 1- Application of traditional public health methods to assess the impact of PIDs on community health; 2- Development, implementation, and evaluation of screening tests administered to newborns and clinical algorithms for early recognition of symptomatic persons to the earliest possible diagnosis and facilitate Systems;³¹ 3treatment for PIDs Surveillance Evaluation of screening and diagnostic tools to ensure their quality and appropriateness for identification of patients with PIDs; and 4- Communication with healthcare providers and the public to facilitate prompt and appropriate diagnosis and intervention.

CDC has begun to apply this framework in the context of ethical, legal, and social considerations in different conditions.³²⁻³⁴ However, educational efforts have the first priority because of the role of education on each four mentioned components. Targets of education are three major subsets of PIDs as priorities for a systematic public health assessment; include profound T-cell defects, because of their resulting high mortality in the absence of interventions; antibody deficiencies, and due to the substantial number of persons affected and the high burden of morbidity; and Chronic granulomatous disease (CGD), because of the existence of an established data set.

All these framework components need to trained and reevaluate in physicians especially pediatrics, and clinical immunologists.

Review of data obtained from National Primary Immunodeficiency Registry of Iran has shown that the mean delay in diagnosis of PIDs was almost 4 years.^{35,36}

According to basis lack of knowledge in target physicians in this study, educations of primary-care physicians group must be considered to achieve early clinical recognition by following items: lessons on the effect of early interventions on morbidity and mortality associated with PIDs, identification of a group of diseases that can benefit from using an early clinical recognition algorithm include Severe combined immunodeficiency, X-Linked Agammaglobulinemia, Common variable immunodeficiency, CGD, evaluate the usefulness and accuracy of family history, early clinical signs and symptoms and initial laboratory tests for early recognition of PIDs. Then a national system for early clinical recognition of PIDs and conduct collaborative studies among clinical centers in selected PIDs should be established.

Although these educational efforts have been ongoing for years in our country, outcomes have not been formally evaluated which led us to perform this study. Among the Iranian pediatricians who participated in this survey, awareness about the PIDs was in high-medium qualitative rank.

In this study the main cause of low knowledge in target group was general deficit in both the knowledge and practice of pediatricians in the field of PIDs which recently has also been reported in another study in Kuwait.²⁵. Although clinical manifestations of PIDs were the most important items for diagnosis; this did not appear to be well in the knowledge of Iranian pediatricians.

Most of our pediatricians did not have enough knowledge about application of para-clinical tests for their patients, but they had in desirable level of knowledge about treatment of PIDs patients.

Although those with previous PIDs patients are more likely to have high knowledge, the proportion of these physicians who had performing well interventions remains at or below 50%.

The exact limitation of this study was due to nonresponders and also who did not attend the congress which may lead to selection bias.

Amazingly, knowledge of PIDs among more experienced pediatricians with higher qualification and higher ranking was not different significantly when compared to less experienced ones. This may be related to limited availability or awareness of the pediatricians about PIDs programs. We therefore recommend implementation of strategies to improve the awareness of pediatricians about PIDs to early interventions with intravenous immunoglobulins.

These strategies may include comprehensive underand post-graduated education, organizing educational courses, and publishing educational materials. Pediatricians should also be educated about the warning signs of PIDs.

Despite rapid developments in the science of PIDs, these diseases have still a significant impact into the health system. Continuing medical education after graduation can increase the knowledge of physicians especially in younger physicians. An understanding of the reasons for lack of awareness can help us to decrease the number of mismanaged PIDs patients. With this information about pediatricians' PIDs care practices, perceptions, and beliefs; it may be possible to conduct targeted interventions to improve primary care for PIDs in Iran

REFERENCES

- Lee PP, Lau YL. Primary immunodeficiencies: "new" disease in an old country. Cell Mol Immunol 2009; 6(6):397-406.
- Cunningham-Rundles C, Bodian C. Common variable immunodeficiency: clinical and immunological features of 248 patients. Clin Immunol 1999; 92(1):34-48.
- Stiehm ER, Chin TW, Haas A, Peerless AG. Infectious complications of the primary immunodeficiencies. Clin Immunol Immunopathol 1986; 40(1):69-86.
- Aghamohammadi A, Farhoudi A, Moin M, Rezaei N, Kouhi A, Pourpak Z, et al. Clinical and immunological features of 65 Iranian patients with common variable immunodeficiency. Clin Diagn Lab Immunol 2005; 12(7):825-32.
- Casanova JL, Fieschi C, Bustamante J, Reichenbach J, Remus N, von Bernuth H, et al. From idiopathic infectious diseases to novel primary immunodeficiencies. J Allergy Clin Immunol 2005; 116(2):426-30.
- Boyle JM, Buckley RH. Population prevalence of diagnosed primary immunodeficiency diseases in the United States. J Clin Immunol 2007; 27(5):497-502.
- Eades-Perner AM, Gathmann B, Knerr V, Guzman D, Veit D, Kindle G, et al. The European internet-based patient and research database for primary immunodeficiencies: results 2004-06. Clin Exp Immunol 2007; 147(2):306-12.
- Chapel H, Geha R, Rosen F. Primary immunodeficiency diseases: an update. Clin Exp Immunol 2003; 132(1):9-15.
- Sigstad HM, Stray-Pedersen A, Froland SS. Coping, quality of life, and hope in adults with primary antibody deficiencies. Health Qual Life Outcomes 2005; 3:31.
- Zebracki K, Palermo TM, Hostoffer R, Duff K, Drotar D. Health-related quality of life of children with primary immunodeficiency disease: a comparison study. Ann Allergy Asthma Immunol 2004; 93(6):557-61.
- Nicolay U, Haag S, Eichmann F, Herget S, Spruck D, Gardulf A. Measuring treatment satisfaction in patients with primary immunodeficiency diseases receiving lifelong immunoglobulin replacement therapy. Qual Life Res 2005; 14(7):1683-91.
- Mozaffari H, Pourpak Z, Pourseyed S, Moin M, Farhoodi A, Aghamohammadi A, et al. Health-related quality of life in primary immune deficient patients. Iran J Allergy Asthma Immunol 2006; 5(1):23-7.

- Dinakar C. Alleviating disease burden in primary immunodeficiency diseases. Ann Allergy Asthma Immunol 2006; 96(2):260-2.
- Hermaszewski RA, Webster AD. Primary hypogammaglobulinaemia: a survey of clinical manifestations and complications. Q J Med 1993; 86(1):31-42.
- Champi C. Primary immunodeficiency disorders in children: prompt diagnosis can lead to lifesaving treatment. J Pediatr Health Care 2002; 16(1):16-21.
- Buckley RH. Primary immunodeficiency diseases due to defects in lymphocytes. N Engl J Med 2000; 343(18):1313-24.
- 17. Quartier P, Debre M, De Blic J, de Sauverzac R, Sayegh N, Jabado N, et al. Early and prolonged intravenous immunoglobulin replacement therapy in childhood agammaglobulinemia: a retrospective survey of 31 patients. J Pediatr 1999; 134(5):589-96.
- Myers LA, Patel DD, Puck JM, Buckley RH. Hematopoietic stem cell transplantation for severe combined immunodeficiency in the neonatal period leads to superior thymic output and improved survival. Blood 2002; 99(3):872-8.
- Antoine C, Muller S, Cant A, Cavazzana-Calvo M, Veys P, Vossen J, et al. Long-term survival and transplantation of haemopoietic stem cells for immunodeficiencies: report of the European experience 1968-99. Lancet 2003; 361(9357):553-60.
- Aghamohammadi A, Pouladi N, Parvaneh N, Yeganeh M, Movahedi M, Gharagolou M, et al. Mortality and morbidity in common variable immunodeficiency. J Trop Pediatr 2007; 53(1):32-8.
- Seymour B, Miles J, Haeney M. Primary antibody deficiency and diagnostic delay. J Clin Pathol 2005; 58(5):546-7.
- Aghamohammadi A, Moein M, Farhoudi A, Pourpak Z, Rezaei N, Abolmaali K, et al. Primary immunodeficiency in Iran: first report of the National Registry of PID in Children and Adults. J Clin Immunol 2002; 22(6):375-80.
- Matamoros Flori N, Mila Llambi J, Espanol Boren T, Raga Borja S, Fontan Casariego G. Primary immunodeficiency syndrome in Spain: first report of the National Registry in Children and Adults. J Clin Immunol 1997; 17(4):333-9.
- 24. Al-Herz W, Naguib KK, Notarangelo LD, Geha RS, Alwadaani A. Parental Consanguinity and the Risk of Primary Immunodeficiency Disorders: Report from the Kuwait National Primary Immunodeficiency Disorders Registry. Int Arch Allergy Immunol; 154(1):76-80.

- Al-Herz W, Zainal ME, Salama M, Al-Ateeqi W, Husain K, Abdul-Rasoul M, et al. Primary immunodeficiency disorders: survey of pediatricians in Kuwait. J Clin Immunol 2008; 28(4):379-83.
- Puck JM. Primary immunodeficiency diseases. Jama 1997; 278(22):1835-41.
- Primary immunodeficiency diseases. Report of an IUIS Scientific Committee. International Union of Immunological Societies. Clin Exp Immunol 1999; 118 Suppl 1:1-28.
- Aiuti A, Slavin S, Aker M, Ficara F, Deola S, Mortellaro A, et al. Correction of ADA-SCID by stem cell gene therapy combined with nonmyeloablative conditioning. Science 2002; 296(5577):2410-3.
- Buckley RH. Advances in the understanding and treatment of human severe combined immunodeficiency. Immunol Res 2000; 22(2-3):237-51.
- Cavazzana-Calvo M, Hacein-Bey S, Yates F, de Villartay JP, Le Deist F, Fischer A. Gene therapy of severe combined immunodeficiencies. J Gene Med 2001; 3(3):201-6.
- 31. Thacker SB, Stroup DF. Future directions for comprehensive public health surveillance and health information systems in the United States. Am J Epidemiol 1994; 140(5):383-97.
- 32. Brown AS, Gwinn M, Cogswell ME, Khoury MJ. Hemochromatosis-associated morbidity in the United States: an analysis of the National Hospital Discharge Survey, 1979-1997. Genet Med 2001; 3(2):109-11.
- McDonnell SM, Witte DL, Cogswell ME, McIntyre R. Strategies to increase detection of hemochromatosis. Ann Intern Med 1998; 129(11):987-92.
- Wetterhall SF, Cogswell ME, Kowdley KV. Public health surveillance for hereditary hemochromatosis. Ann Intern Med 1998; 129(11):980-6.
- 35. Mir Saeid Ghazi B, Aghamohammadi A, Kouhi A, Farhoudi A, Moin M, Rezaei N, et al. Mortality in Primary Immunodeficient Patients, Registered in Iranian Primary Immunodeficiency Registry. Iran J Allergy Asthma Immunol 2004; 3(1):31-6.
- 36. Rezaei N, Aghamohammadi A, Moin M, Pourpak Z, Movahedi M, Gharagozlou M, et al. Frequency and clinical manifestations of patients with primary immunodeficiency disorders in Iran: update from the Iranian Primary Immunodeficiency Registry. J Clin Immunol 2006; 26(6):519-32.