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Clinical Possibilities of Manifestation of Juvenile Idiopathic Arthritis

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Introduction

Juvenile Idiopathic Arthritis is the most common rheumatologic disease in children and one of the most common chronic diseases of childhood. The term JIA encompasses a heterogeneous group of disorders that all share the clinical manifestation of chronic joint inflammation. The etiology is unknown, but both genetic and environmental factors are believed to be involved. Management of the disease has improved in recent years due to advances of pharmacologic treatment options. The prognosis of patients with juvenile idiopathic arthritis is better.

Clinical

Chronic joint inflammation must be present for at least 6 weeks in the same joint before the diagnosis of juvenile idiopathic arthritis be made. Disease onset is insidious or sudden and it is characterized by morning stiffness and arthralgia during the day. Usually children with juvenile idiopathic arthritis were absent from school and activities during physical education classes is limited reflecting disease severity. Affecting the lower extremities causing limping in children, so a morning limp that improves during the day can be found [1-3].

Arthritis can be present in many conditions. A preceding illness must bring into question the possibility of infectious trigger for post infectious arthritis. Reactive arthritis should be called into question in any child with gastroenteritis and arthritis of large joints of the lower extremities. Lyme disease caused by Borrelia burgdorferi, is a major health problem in endemic areas. Arthritis is a late manifestation of the disease. So a history of travel to endemic areas exposed to ticks raises the possibility of Lyme disease. If the patient complains of severe joint pain, diagnostic alternatives include acute rheumatic fever, acute lymphocytic leukemia, septic arthritis and osteomyelitis [4,5]. Gastrointestinal symptoms, microcytic anemia, and elevated inflammatory markers raise the possibility of inflammatory bowel disease. Weight loss in the absence of diarrhea may be observed in patients with active juvenile idiopathic arthritis. This sign can also be seen in patients with acute lymphocytic leukemia and in patients with inflammatory bowel disease. Differential diagnoses include systemic lupus erythematosus. Pericarditis with orthopnea can be observed both in juvenile idiopathic arthritis and systemic lupus erythematosus [6-8].

Classification of Juvenile Idiopathic Arthritis

Systemic Arthritis

Definition: Arthritis in one or more joints with or proceeded by fever of at least 2 weeks' duration that is documented to be daily for at least 3 days [9-12].

inclusion criteria					
1.	Evanescent erytematous rash				
2.	Generalized lymph node				
	enlargement				
3.	Hepatosplenomegaly				
4.	Serositis				

Signs and symptoms

- Arthralgia
- Chest pain and shortness of breath are signs of pericarditis or pleuritis Muscle tenderness
- Myalgia

Exclusion criteria

- Psoriasis or a history of psoriasis in the patient or first-degree relative Arthritis in an HLA-B27 positive male begin-
- ning after the 6th birthday
- Ankylosing spondylitis, enthesitis related arthritis, sacroiliitis with inflammatory bowel disease, Reiter's syndrome, or acute anterior uveitis, or a history of one of these disorders in a first-degree relative
- The presence of IgM rheumatoid factor on at least 2 occasions at least 3 months apart

Oligoarthritis

Definition: Arthritis affecting one to 4 joints during the first 6 months of disease [9,13].

Inclusion criteria	Exclusion criteria		
Persistent oligoarthritis: affecting not more than 4 joints throughout the disease course. Extended ologoarthritis: affecting a total of more than 4 joints after the first 6 months of disease. Signs and symptoms Larger joints such as the knees, ankles, wrists are affected Limp Extensor muscle atrophy Flexion contractures in the knees and the wrists Anterior uveitis	1. Psoriasis or a history of psoriasis in the patient or first-degree relative 2. Arthritis in an HLA-B27 positive male beginning after the 6th birthday 3. Ankylosing spondylitis, enthesitis related arthritis, sacroiliitis with inflammatory bowel disease, Reiter's syndrome, or acute anterior uveitis, or a history of one of these disorders in a first-degree relative 4. The presence of IgM rheumatoid factor on at least 2 occasions at least 3 months apart 5. The presence of systemic JIA in the patient		

Polyarthritis

Definition: Arthritis affecting 5 or more joints during the first 6 months of disease [9,14,15].

months of disease [5,14,15].					
Inclusion criteria	Exclusion criteria				
Rheumatoid factor negative: a test for RF is negative. Signs and symptoms	Psoriasis or a history of psoriasis in the patient or first-degree relative Arthritis in an HLA-B27 positive male beginning after the 6 th birthday Ankylosing spondylitis, enthesitis related arthritis, sacroiliitis with in-				
 ✓ Both large and small joints can be affected, with symmetrical distribution ✓ Fever ✓ Severe limitations in motion, muscle weakness, decreased physical function 	flammatory bowel disease, Reiter's syndrome, or acute anterior uveitis, or a history of one of these disorders in a first-degree relative 4. The presence of IgM rheumatoid factor on at least 2 occasions at least 3 months apart 5. The presence of systemic JIA in the patient				
Rheumatoid factor positive: 2 or more tests for RF at least 3 months apart during the first 6 months of disease are positive. Signs and symptoms Both large and small joints can be	Psoriasis or a history of psoriasis in the patient or first-degree relative Arthritis in an HLA-B27 positive male beginning after the 6 th birthday Ankylosing spondylitis, enthesitis related arthritis, sacrolliitis with in-				
affected, with symmetrical distribution ✓ Fever ✓ Severe limitations in motion, muscle weakness, decreased physical function ✓ Rheumatoid nodules can be found	flammatory bowel disease, Reiter's syndrome, or acute anterior uveitis, or a history of one of these disorders in a first-degree relative 4. The presence of systemic JIA in the patient				

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Psoriatic Arthritis

Definition: Arthritis and psoriasis, or arthritis and at least 2 of inclusion criteria [9,16].

Inclusion criteria			Exclusion criteria		
2.	Dactylitis Nail pitting or onycholysis		Arthritis in an HLA-B27 positive male beginning after the 6th birthday		
3.	3. Psoriasis in a first-degree relative	2.	Ankylosing spondylitis, enthesitis related arthritis, sacroiliitis with inflammatory		
	Signs and symptoms		bowel disease, Reiter's syndrome, or		
/	Monoarticular arthritis		acute anterior uveitis, or a history of one		
/	Tenosynovitis		of these disorders in a first-degree relative		
/	Sacroiliitis	3.	The presence of IgM rheumatoid factor		
√	Disordered bone growth with resultant shortening		on at least 2 occasions at least 3 months apart		
		4.	The presence of systemic JIA in the patient		

Enthesitis Related Arthritis

Definition: Arthritis and enthesitis, or arthritis or enthesitis with at least 2 of inclusion criteria [9,15,17].

Inclusion criteria			Exclusion criteria		
1.	The presence of or a history of sacro-	1.	Psoriasis or a history of psoria-		
	iliac joint tenderness and/or inflamma-		sis in the patient or first-degree		
	tory lumbosacral pain The presence of		relative		
	HLA B-27 antigen	2.	The presence of IgM rheuma-		
2.	Onset of arthritis in a male over 6 years		toid factor on at least 2 occa-		
	of age		sions at least 3 months apart		
3.	Acute anterior uveitis	3.	The presence of systemic JIA in		
4.	History of ankylosing spondylitis, en-		the patient		
	thesitis related arthritis, sacroilliitis with				
	inflammatory bowel disease, Reiter's				
	syndrome, or acute anterior uveitis in a				
	first-degree relative				
	Signs and symptoms				
/	Pain and tenderness at the enthesis				
/	Swelling				

Undifferentiated Arthritis

Definition: Arthritis that fulfills criteria in no category or in 2 or more of the above categories [9,18].

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