Scintigraphic evaluation of disease activity in juvenile rheumatoid arthritis

Ву

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List of abbreviation

% percent

⁰C Degree Centigrade

99Tc-MDP Technitium Methelene Di Phosphonate ACR American College of Rheumatology

ANA Antinuclear Antibody

Anti-ds-DNA Anti double stranded antibody

b,i,d twice per day

C2 Complement factor 2
C3 Complement factor 3
CBC Complete Blood Count
CD4 Cluster of differentiation 4

CRP C reactive protein

CT Computerized tomography
DIP Distal Interphalangeal

dl deciliter

DMARDs Disease Modifying Anti Rheumatic Drugs

ECG Electrocardiography

ESR Erythrocyte sedimentation rate

Ga Gallium

GH Growth hormone

HDD Hydroxymethelene Di Phosphonate

HLA Human leukocytic antigens

IgA Immunoglobulin A
IGF1 Insulin Growth factor 1
IGF2 Insulin Growth factor 2
IgG Immunoglobulin G
IgM Immunoglobulin M

In Indium

JRA Juvenile rheumatoid arthritis.

Kg Kilogram L Litre

LN Lymph Nodes

M² Square meter

mCi milli curi

MCP Metacarbophalangeal
MDP Methelene diphosphate

mg milligram
mm millimeter
mm³ Cubic millimeter

MRI Magnetic resonance imaging

MTP Metatarsophalangeal

NC Nanocolloid

List of abbreviations Cont.

NSAIDs Non Steroidal Anti-Inflammatory Drugs

OD Once daily

PIP Proximal Interphalangeal PPD Protein Purified Derivative

q,i,d Four times per day
RA Rheumatoid Arthritis
RF Rheumatoid factor

Rh +ve Rheumatoid factor positive Rh -ve Rheumatoid factor negative

SAARDs Slow Acting Anti Rheumatic Drugs SLE Systemic Lupus Erythematosus

TB Tuberculosis

t,I,d Three times per day

Y Year(s)

Introduction & Aim of work

Rheumatoid arthritis is characterized by an inflammatory process of the synovial membrane with increased capillary permeability, exudation of plasma proteins into the synovial stroma and the joint cavity, and infiltration of cellular element (Arnett et al., 1988).

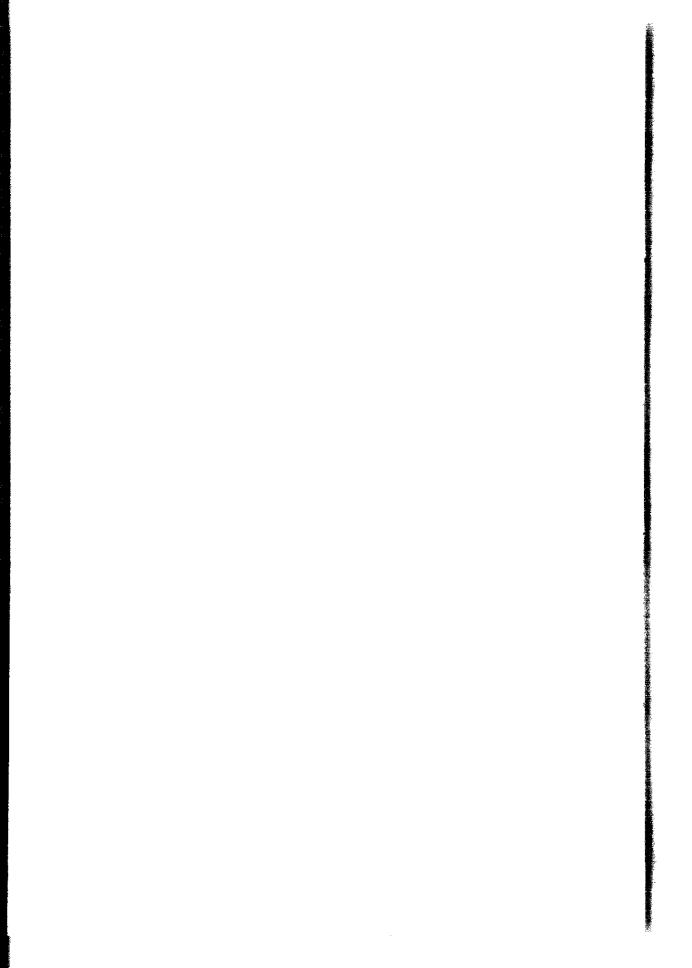
The clinical assessment of this condition is normally based on the evaluation of pain and swelling experienced by the patient, physical examination of the joints and laboratory investigations. This type of assessment is obviously rather subjective and is not always able to evaluate the real state of the disease. Objective, non-invasive test, aimed at measuring the synovitis, is of great interest in clinical practice (*Breedveld et al 1989*).

Nuclear medicine has been currently applied to evaluate disease activity (Breedveld et al 1989). Technetium-99m-labeled, non-specific, polyclonal, human Immunoglobulin G (99mTc-HIG) has been used to quantify synovial inflammation in rheumatoid arthritis, as it can be detected in inflamed joints as a result of the increased vascular permeability of the synovial membrane (Fishman et al 1988).

Aim of the work:

To evaluate this scintigraphic imaging with ^{99m}Tc-HIG as an objective test for assessing synovitis and compare the results of this technique with clinical examination and laboratory indices of disease activity.

Review of literature



Review of literature

Chapter 1

Juvenile Rheumatoid Arthritis

Juvenile rheumatoid arthritis (JRA) is a disease or a group of diseases characterized by chronic synovitis and associated with a number of extra-articular manifestations. A confusing number of names have been applied, including juvenile arthritis, Still disease, juvenile chronic polyarthritis, and chronic childhood arthritis. JRA encompass several broad clinical subgroups (table 1).

- 1- Rheumatoid factor-positive (RF +ve) polyarticular disease.
- 2- Rheumatoid factor-negative (RF -ve) polyarthritis.
- 3- Pauciarticular disease type I.
- 4- Pauciarticular disease type II.
- 5-Systemic-onset disease.

Recognition of these subgroups is useful in the diagnosis, follow-up, and appropriate care of children with chronic arthritis (Schaller, 1996).

Table 1: Subgroups of JRA

	Poly-articular	Poly-articular	* ***		
Characteristics	rheumatoid factor-	rheumatoid factor-	Pauci-articular	Pauci-articular type	Systemic onset
	negative	positive	type I	II	
Percentage of JRA	20-30	5-10	30-40	10-15	10-20
patients					
Sex	90 % girls	80 % girls	80 % girls	90 % boys	60 % boys
Age at onset	Any	Late childhood	Early Childhood	Late childhood	Any
Joints	Any, multiple	Any, multiple	Few large joints:	Few large joints: hip	Any, multiple
			Knee, ankle, elbow	girdle	
Sacroiliitis	No	Rare	No	Common	No
I ridocy clitis	Rare	No	30 % chronic	10-20 % acute	No
			iridocyclitis	iridocyclitis	
Rheumatoid Factor	Negative	100 %	Negative	Negative	Negative
Antinuclear	25 %	75 %	90 %	Negative	Negative
antibodies					
HLA studies	?	HLA DR4	HLA DR5, DR6,	HLA B27	?
			and DR8		
Ultimate morbidity	Severe arthritis, 10-15	Severe arthritis, >	Polyarthritis,	Subsequent	Severe arthritis,
	%	50 %	20 %	spondyloarthropathy,?	25 %

(Cassidy & Petty, 1995).