H.4 Specialist referral for psoriatic arthritis

Reference	Study type	Numbe r of patient s	Patient characteristics	Prognostic factors	Length of follow-up	Outcome measures	Source of funding
D. Kane, L. Stafford, B.	Inception cohort (prospective)	1018 Patients	Patients referred to early arthritis clinic with joint	Clinical assessment	2 years	RAI	Not stated
FitzGerald. A prospective, clinical and	Ireland/Scotland	presenti ng to Early	either active joint swelling or an elevated acute-phase	Baseline		EULAR	
radiological study of early psoriatic	August 1994 – March 2000	Synoviti s Clinic	<pre><2 years. RF factor titre < 1/80.</pre>	radiographs		HAQ	
arthritis: an early synovitis clinic	Representative: Loss to follow-up:	129 (12/7%)	Diagnosis of PsA confirmed by consultant rheumatologist			Pain	
<i>Rheumatology.</i> 42:1460-	10 patients (8%) at 1 year	ed with PsA	using Moll & Wright criteria			DMARD use	
1408.2003	SI patients (23%)at 2 years		53% male, 47% female Mean age at presentation 41.2			Swollon	
REF ID: KANE2003	119 patients followed up at 1 yr		±15.1 years Mean duration of disease at			joints	
	97 patients followed up at 2 yr		presentation 9.9 ±15.1 months Median delay from symptom			Radiologi cal assessme	

	onset to rheumatology	nt –
	referral 5.75-7 months	Sharp
		Remissio
		n.
		defined
		by
		absence
		of
		fatigue.
		stiffness
		<15 min.
		no ioint
		pain.
		complete
		absence
		ofioint
		tenderne
		ss or
		swelling
		(including
		dactylitis
		and
		enthesitis
) on
		examinati
		on and
		ESR
		<20 mm/
		h (males)
		or ESR
		<30 mm/
		h

					(females)				
Effect size	Effect size								
At presentation !	At presentation 52 (40%) had olgioarticular PsA, 77 (60%) had polyarticular disease								
ACR class at pres	entation: I: 39 (34%), II: 38 (32%), III: 33 (28%), IV 7 (6%)							
Mean HAQ score	at presentation 0.71 ±0.64, at 1	yr 0.4 ±0.6, 2 yr 0.4 ±0.6							
Overall decrease	in all clinical and lab parameters	s of inflammation at 1 and	2 yr.						
	0 yr	1 yr	2 yr						
No. of patients	n = 129	n = 119 (92%)	n = 97 (75%)						
DMARD	15 (12%)	70 (59%)	54 (56%)						
Corticosteroids	14 (11%)	6 (5%)	5 (5%)						
VAS pain	4.8 (5) ± 2.7 (<i>n</i> = 122)	3.1 (2) ± 3 (<i>n</i> = 119)	3.4 (4) ± 2.7 (<i>n</i> = 97)						
ACR class III/IV	40 (34%) (<i>n</i> = 117)	22 (19%) (<i>n</i> = 118)	16 (16%) (<i>n</i> = 97)						
HAQ score	0.7 (0.6) ± 0.6 (<i>n</i> = 74)	0.4 (0.1) ± 0.6 (<i>n</i> = 65)	0.4 (0.1) ± 0.6 (<i>n</i> = 58)						
Ritchie Index	5.6 (4) ± 6	2.4 (1) ± 3.8	1.9 (1) ± 3						

Swollen joint count	6.9 (4) ± 8	2.9 (1) ± 5.2	2.4 (1) ± 4.1	
ESR (mm/h)	24 (16) ± 27 (<i>n</i> = 124)	13 (7) ± 15 (<i>n</i> = 112)	12 (7) ± 14 (n	= 94)
CRP (mg/l)	28 (10) ± 59 (<i>n</i> = 112)	10 (5) ± 14 (<i>n</i> = 111)	8 (4) ± 12 (n =	= 94)
Enthesopathy	29 (38%)	15 (13%)	25 (26%)	
Dactylitis	37 (29%)	10 (8%)	16 (16%)	
Remission	0	31 (26%)	20 (21%)	
Remission in 26% of pat	tients at 1 yr, 21% of patien	ts at 2 yr		
Spontaneous (DMARD-	free) remission in only 11-1	2% of patients		
Radiological:				
At baseline, 32/117 (27	%) of patients had erosions	, 24 (19%) of patients had j	oint space nar	rowing and 22 (19%) of patients had periostitis
After median 24 month	s follow-up, 40/86 (47%) of	patients had erosions (des	pite early DM	ARD use), 32 (37%) had joint space narrowing and 25 (29%) of
patients had periostitis				
Baseline (<i>n</i> = 117)		Follow-up (n =	86)	
Total number of joints y	with erosions		007	
Hands		75/2510 (2.1%	()	100/2580 (3.9%)
Faat		26/1170 (2.17	וי א	100/2300 (3.3 <i>%</i>)
Feet		26/11/0 (2.2%	o)	53/860 (6.2%)
Mean no. of joints with	erosions per patient ± S.D.			

Useda	07+10	4.2 + 2.5
Hands	0.7±1.6	1.2±2.5
Feet	0.2 ± 0.8	0.6 ± 1.6
Total number of joints with joint space narrowing		
Hands	71/3510 (2.0%)	62/2580 (2.4%)
Feet	14/1170 (1.2%)	35/860 (4.1%)
Mean no. of joints with joint space narrowing per	patient ± S.D.	
Hands	0.6 ± 2.3	0.7±1.7
Feet	0.1 ± 0.5	0.4 ± 1.4
Mean sharp score at baseline 1.2 ±2.9, mean narro	owing score (hands/feet) at baseline	1.4 ±5.3
Mean Sharp score increased to 3 ±5.2 (P=0.002), m	nean narrowing score increased to 3	.2 ±7.5 (P=0.04)
Sacroileitis present in 16 (17%) of patients		
Significant functional impairment at early stage		
Author conclusion: PsA is a chronic disease with sig	gnificant functional impairment and	radiological damage at an early stage in the course of the disease

Reference	Study type	Number of patients	Patient c	haracteris	tics	Prognostic factors	Length of follow-up	Outcome measures	Source of funding
L. Punzi, M. Pianon, P. Rossini, F. Schiavon, P. F. Gambari. Clinical and laboratory manifestations of elderly onset psoriatic arthritis: a comparison with younger onset disease. <i>Ann Rheum</i> <i>Dis.</i> 58:226- 229. 1999	Prospective study Italy Elderly vs. younger onset disease	66 consecu tive PsA patients	PsA patie duration onset PsA younger of RF +ve pa M/F Mean age at onset, y	nts with d < 1 year : 1 A (>60 yrs), onset PsA d atients exc EOPsA 8/8 65.1 ±6.7	isease 6 elderly 50 (≤60 yrs). Iuded YOPsA 23/27 44.2 ±11.1	Disease duration	2 years	Clinical Laboratory Radiographic	Not stated

Effect size

DMARD (SAARD) at 2 years: 42/50 (84%) in YOPsA patients, 15/16 (94%) EOPsA patients

Mean ±SD number of radiographic erosions in hands at presentation: 2.3 ±2.1 (EOPsA), 2.2 ±2.2 (YOPsA)

Mean ±SD number of radiographic erosions in hands after 2 years: 4.4 ±3.0 (EOPsA), 2.7 ±2.0 (YOPsA)

Mean ±SD number of radiographic erosions in feet at presentation: 2.7 ±1.2 (EOPsA), 1.1 ±1.1 (YOPsA)

Mean ±SD number of radiographic erosions in feet after 2 years: 4.7 ±2.2 (EOPsA), 2.1 ±1.2 (YOPsA)

Higher number of active joints in elderly vs young onset PsA at both baseline (12.2±6.3 vs 6.7±6.6; p<0.001) and 2-year follow-up (8.1±4.2 vs 4.7±3.6; NS)

Mean ESR decreased from 64.2 ±65.3 mm/h at baseline to 38.4 ±15.2 mm/h after 2 years' follow-up in Elderly Onset PsA patients and a more modest decrease from 30.5 ±30.0 mm/h to 26.3 ±15.0 mm/h in Younger Onset PsA patients. Mean CRP levels also decreased in both groups: 3.9 ±2.0 mg/l to 2.2 ±1.0 mg/l in Elderly Onset PsA and 1.33 ±1.3 mg/l to 0.9 ±0.9 mg/l in Younger Onset PsA patients.

Reference	Study type	Numbe r of patient s	Patient characteristics	Prognostic factors	Length of follow-up	Outcome measures	Source of funding
B. J. Harrison, A. J. Silman, E. M. Barrett, D.G.I. Scottt, D.P.M. Symmons. Presence of psoriasis does not influence the presentation or short term outcome of patients with early inflammatory polyarthrtitis. <i>J</i> <i>Rheumatol.</i> 24:1744- 9.1997	Primary care inception cohort Norfolk, UK 1989	966 patient s referre d to Norfolk Psoriasi s Registry 51 patient s with psoriasi s	Patients ≥16 years old with early inflammatory polyarthritis (swelling of at least 2 joint areas that has persisted for a minimum of 4 weeks) and psoriasis in a primary care population. 49% male, 51% female Median age at psoriasis onset 52 years Median duration of arthritis at presentation 5.75 months Note: approximately 50% had RA not PsA	Clinical assessment Lab markers Radiographs	1 year	Total number of swollen joints DMARD use Remission HAQ score Radiograp hs	Not stated
REF ID: HARRISON1997							

Effect size		
	Baseline	1 year
Second line drugs/steroids		21 (41%)
Median HAQ	0.63	0.44
Median swollen joints	7 (0-32)	4 (range: 0-16)
Remission		3 (6%)
Radiological erosions		7/32 (22%)

Reference	Study type	Numbe r of patient s	Patient characteristics	Prognostic factors	Length of follow-up	Outcome measures	Source of funding
R. Queiro-Silva, J.C. Torre- Alonso, T. Tinture-	Prospective cohort – consecutive sample	71 patient s	Patients with PsA	Disease duration	10 years	ACR	Not stated
Eguren, I. Lopez-Lagunas.	Spain		44 men, 27 women			HAQ	
onset predicts erosive and deforming	1991-2001		Mean disease duration at presentation 12 ± months (without radiographical			Lab values	
disease in psoriatic arthritis. Ann			evidence of erosions at presentation)			Radiographs	
Rheum Dis. 62:68-70.2003			Mean age 47 ±12 years				
REF ID: QUEIROSILVA2 003							
		I					
Effect size							
During first 6 m	onths 5 patients (7%) had isolate	ed DIP dise	ase				
30 (42%) oligoar	thritis						

20 (28%) polyarhtritis

16 (23%) axial disease

0 arthritis mutilans

At end of study (10 years)

28 (39%) showed oligoarthritis

24 (34%) polyarthritis

17 (24%) axial disease

2 (3%) arthritis mutilans

32/71 (45%) had developed erosive and deforming arthritis

Mean time to detect erosions or narrowing of joint spaces was 20±4 months (SD)

HAQ (unclear if at baseline or follow-up): 1.2 (0.3) in those with erosive (n=32) vs 0.6 (0.4) in non erosive (n=39) (p=0.012)

NS difference in between number of months duration of arthritis (8 ±7 months versus 10±6 months) for erosive and non-erosive disease

DMARD use in 68% of patients

Author conclusion: we support the use of DMARDs as early as possible, particularly in patients with polyarticular onset

Reference	Study type	Numb er of patien ts	Patient characteristics	Prognostic factors	Length of follow-up	Outcome measures	Source of funding		
N.J. McHugh, C. Balachrishnan , S.M. Jones. Progression of peripheral joint disease in psoriatic arthritis: a 5- year prospective study. <i>Rheumatolog</i> y. 42:778- 783.2003 REF ID: MCHUGH200 3	Prospective follow-up study Bath, UK Baseline information collected between 1987 and 1990 87% available for full follow-up	87 patient s; 13 patient s with arthriti s < 1 year durati on	Patients attending a PsA clinic (established/new onset) 25% referrals from primary care, <10% from dermatology 49 females, 38 males Median age in years at follow-up (range) 53.5 (2- 85) Median disease duration at baseline: 11 years (IQR 3.5- 17 years), subgroup analysed with arthritis within 1 year of baseline	Disease duration	Median 65 months (range 39- 90)	Rates of progression of peripheral joint score (0-70) Joint score PASI HAQ Radiographs	Jules Thorn Charitable Trust Remedi UK		
Effect size 13 patients wit Median joint sc	Effect size 13 patients with <12 months duration of arthritis Median joint score at baseline: 4 (IQR 2.3-10)								

Median joint score at follow-up: 7 (IQR 4.3-13)

Rate of peripheral joint progression significantly higher in this group up to baseline assessment compared with the rate of the joint progression in the same patients over subsequent years of follow-up (4.0 vs. 0.32, P=0.003)

Highest rate of peripheral joint involvement appeared to be within 12 months of disease onset, but steady progression of peripheral joint involvement among those referred to a clinic – 0.4 joints per year)

Median rates of joint progression according to age of onset or stage of arthritis (interquartile ranges are given in parentheses)

	Total PsA group (<i>n</i> =87)	Arthritis within 1 yr of baseline (n=13)
Duration of arthritis at baseline (yr)	11 (3.5–17)	<12 months
Joint score at baseline	6 (2–15)	4 (2.3–10)
Change in joint involvement to baseline	0.88 (0.33–1.7)	4* (2.3–10)
Joint score at follow-up	11 (4.5–24)	7 (4.3–13)
Change in joint involvement to follow-up	0.76 ((0.28–1.3)	1.2 (0.6–2.4)
Change in joint involvement from baseline to follow-up	0.43 (0–1.3)	0.32 (0–1)

Full group (not just early onset; median duration at baseline = 11 years)

% of patients with erosions in hand or wrist increased from 53 to 68%, and erosive foot disease increased from 37 to 44%

% taking MTX: at baseline (referral) = 12% vs 15% at follow-up

4/87 patients in remission at follow-up

Mean HAQ score at baseline 0.375, at follow-up 0.5 (p<0.001)

Author conclusion: although a disproportionately high number of peripheral joints are involved in the first 12 months following disease onset, there is a steady progression of peripheral joint involvement in patients with PsA who are referred to a hospital clinic

Reference	Study type	Numbe r of patient s	Patient characteristics	Prognostic factors	Length of follow- up	Outcome measures	Source of funding
S.J. Bond, V.T. Farewell, C.T. Schentag, D.D. Gladman. Predictors for radiological damage in psoriatic	Prospective cohort Toronto 1978 – 2004	625 patient s	Patients referred to University of Toronto PsA Clinic Baseline characteristics:	Disease duration	26 years	Change in number of permanently damaged joints between visits (clinically/radiograp hically)	Not stated
arthritis: results from a single centre. Ann Rheum Dis. 66:370- 376. 2007	Analysis: corrected for within-patient correlation Adjusted for: Sex, age, arthritis duration, functional class, ESR, tender joint		Female/male 272/353 Median (range) age (years) 34 (9– 86) Median (range) duration of arthritis (years) 4.5 (0–47.7)			Radiological damage (Steinbrocker) hands and feet	

	count swollen joint count			
REF ID: BOND2007	and drugs (order of increasing severity: no drug,	Median (range) number of tender joints (all joints) 4 (0–43)		
	NSAID, DMARD, steroids – none were taking biologics)	Median (range) number of tender joints (hands and feet) 3 (0–35)		
		Median (range) number of swollen joints (all joints) 2 (0–33)		
		Median (range) number of swollen joints (hands and feet) 1 (0–28)		
		Median (range) ESR rate 22.5 (0– 105)		
		Functional class		
		Good (I) 29.3% (183)		
		Medium (II) 59.2% (370)		
		Poor (III, IV) 11.5% (72)		
		Damaged joints (all joints)		
		None 62.2% (389)		
		1–4 20.8% (130)		
		5–9 5.9% (37)		
		>9 11.1% (69)		
		Damaged joints (hands and feet)		
		None 68.3% (427)		
		1–4 17.3% (108)		

	5–9 5% (31) >9 9.4% (59)		
	Drugs		
	None 24.3% (152)		
	NSAIDs 30.6% (191)		
	DMARDs 40.5% (253)		
	Steroids 4.6% (29)		

Effect size

Clinical damage: presence of a limitation of range of movement of >20% of the range not related to the presence of joint effusion, the presence of joint deformities, subluxation, loosening or ankylosis.

Radiological damage: Each joint is scored as 1, normal (with possible soft tissue swelling); 2, surface or pocket erosions; 3, erosion and joint space narrowing; and 4, disorganisation (including ankylosis, pencil-in-cup change or total joint destruction) or as requiring surgery. Radiological damage is assessed only in the joints of the hands (wrists, all metacarpophalangeals, PIPs and distal interphalangeals) and feet (MTPs and interphalangeal fist toes); 42 joints in total

Strong relationships were identified between clinical damage development and swollen joints, ESR and arthritis duration

PROGRESSION OF CLINI	PROGRESSION OF CLINICAL DAMAGE (outcome = change in clinically damaged joint count):							
Factor	Single-factor analyses		All factors included					
	Relative damage rate (95% CI)	p Value	Relative damage rate (95% CI)	p Value				
Functional class		<0.001		0.1				
Good (I)	1		1					
Medium (II)	1.56 (1.24 to 1.96)		1.16 (0.89 to 1.5)					
Poor (III, IV)	1.37 (0.96 to 1.91)		0.87 (0.59 to 1.28)					
Tender joints		<0.001		0.2				
None (0)	1		1					
Low (1–4)	1.45 (1.13 to 1.86)		1.15 (0.89 to 1.51)					
Medium (5–9)	1.63 (1.19 to 2.24)		1.27 (0.91 to 1.78)					
High (>9)	2.09 (1.54 to 2.85)		1.37 (0.97 to 1.95)					
Effusions		<0.001		<0.001				
None (0)	1		1					
Low (1–4)	1.32 (1.07 to 1.63)		1.12 (0.89 to 1.42)					
Medium (5–9)	1.84 (1.33 to 2.55)		1.48 (1.02 to 2.13)					
High (>9)	2.95 (1.82 to 4.78)		2.6 (1.56 to 4.36)					

ESR		0.17		0.75		
Low (<15)	1		1			
Medium (15–30)	1.05 (0.82 to 1.39)		0.99 (0.77 to 1.28)			
High (>30)	1.27 (0.94 to 1.73)		1.09 (0.8 to 1.48)			
Arthritis duration	0.67 (0.55 to 0.8) per extra decade in clinic	<0.001	0.73 (0.6 to 0.89)	<0.001		
Drugs		0.143		0.044		
None	1		1			
NSAIDs	0.72 (0.44 to 1.18)		1.11 (0.65 to 1.91)			
DMARDs	0.89 (0.6 to 1.32)		1.32 (0.84 to 2.07)			
Steroids	1.04 (0.68 to 1.6)		1.64 (1.02 to 2.68)			
DMARD, disease-modify	ving antirheumatic drug; ESR, erythrocyte sedime	ntation rate; NS	AID, non-steroidal anti-inflammatory drug.			
Change in clinically dam	aged joint count for all joints					
Before entry to the clini the opposite: the longer	Before entry to the clinic, the longer the duration, the more damage caused by arthritis, but during duration the more damage in the clinic the effect is the opposite: the longer the follow-up, the lesser the damage					
Arthritis duration at firs	st visit is a predictor for progression in patients w	ho do not have	damage at the first visit, but once a patient h	as a damaged joint		

the predictive power of arthritis duration evaporates								
Factor	Relative damage rate	Lower 95% Cl	Upper 95% Cl	p Value				
Clinical damage								
Arthritis duration at first visit								
Damaged	1.06 per decade	0.92	1.22	0.39				
Undamaged	1.54 per decade	1.22	1.96	<0.001				
Radiological damage								
Arthritis duration at first visit								
Damaged	0.99 per decade	0.81	1.19	0.88				
Undamaged	0.84 per decade	0.63	1.12	0.23				

Reference	Study type	Numbe r of patient s	Patient characteristics	Prognostic factors	Length of follow- up	Outcome measures	Source of funding
Gladman DD, Thavaneswara n A, Chandran V, Cook RJ. Do patients with psoriatic arthritis who present early	Prospective cohort University of Toronto PsA clinic	1077 patient s (436 within 2 years of diagnos is and	Patients referred to University of Toronto PsA Clinic; divided into those first seen within 2 years of diagnosis and those first seen with more than 2 years since diagnosis	Disease duration	32 years	Change in number of permanently damaged joints between visits (clinically): defined as a limitation of movement of more than 20% of the	None

fare better than those presenting later in the disease? Ann Rheum Dis. 70: 2152 – 2154 2011	1978 – 2011 Analysis: multivariable analysis using a negative binomial model	641 with disease duratio n >2 years	Baseline characteristics: See below				range that is not related to a joint effusion, the presence of flexion contactures, fused or flail joints or evidence of surgery at a particular joint	
REF ID: GLADMAN201 1A	Adjusted for: Sex, age, arthritis duration, number of damaged joints at first visit, NSAID use at first visit; DMARD use at first visit; treatment with biologics after first visit; calendar effect (based on decade of entry into clinic)							
Effect size Demographic ar	Effect size Demographic and disease characteristics at first visit							
Variable	Ear	y PsA (n=4	36)	Late PsA	(n=641)		p-value	
Sex F/M (%)	42.4	1/57.6		44.8/55.	2		0.447	
Age at PsA diag	gnosis 40.3	3		34.2			<0.0001	
Age at first visi	t 41.:	L		45.2			<0.0001	

Duration of PsA at first visit	0.92	11.0	<0.0001
Mean number of actively inflamed joint	s 10.5	11.7	0.239
Mean number of damaged joints	3.5	9.2	<0.0001
Mean PASI	6.2	5.5	0.254
Treatment at first visit	56.4%	61.6%	0.089
NSAID	28.0%	56.8%	<0.0001
DMARD biological agents	4.1%	6.7%	0.061

Multivariate analysis of progression of clinical damage

Relative rate of joint damage progression (>2 years vs <2 years disease duration at first visit): 1.38 (1.08-1.77); p=0.01

Stratification by duration of disease at clinic entry

Duration of disease at first visit	Ν	Relative rate of joint damage progression (95% CI)	P value
1-2 years vs <1 year	212	1.53 (0.99-2.36)	0.05
2-4 years vs <1 year	248	1.70 (1.11-2.62)	0.01
5-9 years vs <1 year	201	1.83 (1.16-2.88)	0.009
10-20 years vs <1 year	204	1.83 (1.14-2.96)	0.01
>20 years vs <1 year	86	2.96 (1.64-5.34)	0.0003

Authors' conclusions:

Disease progression is more marked in patients presenting with established disease of more than 2 years' duration; there is also a clear dose/exposureresponse relationship with respect to the duration of disease. These results suggest that patients with PsA should be treated earlier in the course of their disease

Reference	Study type	Numbe r of patient s	Patient characteristics	Prognostic factors	Length of follow-up	Outco me measu res	Source of funding
J.A. Husted, B.D. Tom, V.T. Farewell, C.T. Schentag, D.D.Gladman. Description and prediction of physical functional disability in psoriatic arthritis: a longitudinal analysis using a markov model	Prospective cohort Toronto 1993 – 2003 Markov model used to characterise disability process in PsA – transitions only between no disability (1) and moderate disability (2) and between moderate	341 patient s	Patients attending University of Toronto PsA Clinic. Newly diagnosed and established PsA. 201 men, 140 women Mean age 45.9 ±12.4 years Mean duration of PsA 10.6 ±8.4 years 157 patients (46%) initial HAQ score <0.5 and thus	Disease duration	Mean ±SD follow-up 5.2 ±3.04 years	HAQ Disabil ity state	Canadia n Institute of Health Researc h and the Krembil Foundati on

approach.	disability (2) and severe	assigned an initial disability
Arthritis &	disability (3)	state of 1
Rheumatism.		
52(3):404-		
409.2005	The variables included were sex, age, duration of PsA, psoriasis severity as	134 patients (39%) had a score between 0.5 and 1.5 inclusive and were assigned
REF ID: HUSTED2005	measured by the PASI, the number of clinically deformed or damaged joints, and the number of	disability state 2
	actively inflamed joints updated at each HAQ visit.	score > 1.5 and were assigned to disability state 3
1		

Effect size

Patients with duration of PsA less than 2 years were found to have more frequent transitions to different states (either to better or worse states).

Patients with duration of PsA 2-5 years and >5 years had a reduction in transition rates of 56-70% compared with those patients with PsA duration <2 years

Multivariate model of predictors of transitions between disability states:

RR transition from 1-2 or 2-3 (worsening)

< 2 years RR = 1

2-5 years RR = 0.42 (0.16-1.09)

>5 years RR 0.33 (0.14-0.76)

i.e. significantly lower rate of transition state worsening in patients with PsA duration >5 years compared to those with duration <2 years

RR transition from 2-1 or 3-2 (improving)

<2 years RR = 1

2-5 years RR = 0.33 (0.14-0.77)

>5 years RR = 0.44 (0.21-0.90)

i.e. significantly lower rate of transition state improvement in patients with PsA duration >2 years compared to those with duration <2 years

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Mean length of follow-up with HAQ 5.2 years
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Number and type of disability transition states (HAQ)

Of 341 patients, 95 (28%) were in state 1 (no disability) throughout follow-up

42 (12%) were in state 2

20 (6%) in state 3

91 patients (26.7%) encountered a single transition to either a lower or higher disability state.

93 patients (27.3%) experienced 2 or more observed transitions

Reference	Study type	Number of patients	Patient characteristics	Prognostic factors	Length of follow-up	Outcome measures	Source of funding
U.R.C. Lindqvist, GM. Alenius, T. Husmark, E. Theander, G.	Prospective cohort	135 patients	Patients with PsA (meeting CASPAR criteria) referred to rheumatology outpatients within 2 years of onset	Disease duration	2 years	Joint count	Not stated
Holmstrom, P.T. Larsson. The Swedish early	Sweden		Accessed on inclusion and at			PASI	
psoriatic arthritis register – 2-year			follow-up after 2 years of conventional care.			Lab values	
followup: a comparison with early			58% female, 42% male			VAS	
rheumatoid arthritis. J Rheumatol.			Mean age ±SD: 47.3 ±15.2			HAQ	
REF ID: LINDQVIST2008			years			Radiographs	
			Mean duration of psoriasis ±SD: 11.4 ±6.6 months				
			DMARD on inclusion: 51 patients (38%)				

Effect size							
Radiological examination performed in 120 patients on inclusion: proliferation/destruction indicating PsA found in 24 patients (20%). 79 patients examined radiographically at 2 year follow-up, 23 (32%) of patients exhibited radiological changes consistent with PsA.							
60 patients classified as mono/oligoarthritis at inclusion, 36 of those classified as mono/olgi at 2 years, 8 as polyarticular, 1 as axial, 1 as DIP and 14 as remission							
64 polyarticular at inclusion, 26 mono/oligo at 2 years, 28 poly, 0 axial, 1 DIP, 9 remission							
Significant reduction in:							
Number of swollen joints, no of tender joints, ESR/CRP, pain (VAS), PGA							
No significant change in HAQ, PASI							
Mean ±SD HAQ score at inclusion: 0.66 ±0.56, at 2 year follow-up 0.55 ±0.79							
Outcome	Baseline	Follow-up	p-value (paired t-test)				
N swollen joints	4.4±4.5	1.8±3.4	≤0.05				
N tender joints	5.8±6.7	3.6±6.7	≤0.05				
HAQ	0.43±0.26	0.25±0.29	NS				
ESR (mm/h)	17.3±17.9	11.2±10.2	≤0.05				

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	CRP (mg/l)	41.7±21.9	7.2±7.6	≤0.05		
	N with radiological damage compatible with PsA	24	33	NS		
	Pain VAS (mm)	44±24	34±26	≤0.05		
	17% in remission, radiological damage verified on inclusion or at follow-up in 31% of PsA patients					
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Reference	Study type	Number of patients	Patient characteristics	Prognostic factors	Length of follow-up	Outcome measures	Source of
							funding
F. Cantini, L. Niccoli, C. Nannini, E. Cassara, P. Pasquetti, I. Olivieri, C. Salvarani.	Prospective case-control (comparison with RA not relevant to question; therefore cohort data used) Consecutive series	236 (6/251 lost to follow- up)	All consecutive outpatients with peripheral PsA requiring second-line drugs observed between Jan 2000 and Dec 2005 at Rheumatology Unit	Disease duration	Mean 38 months	Clinical remission DMARD/bi ologic use	None declared
Frequency and duration of clinical remission in patients with peripheral psoriatic	Italy		Mean disease duration: 13 ±7.1 months				

arthritis				
requiring				
second-line				
drugs.				
Rheumatology.				
47:872-				
876.2008				
KEF ID: 268				

Effect size

32.6% of patients were in remission after an average follow-up time of 38 months

68% were on DMARD therapy and 32% were on anti-TNF-α biologic therapy (plus methorexate) after an average follow-up time of 38 months