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# Prevalence of the Metabolic Syndrome (MS) in Patients with Psoriasis: Moroccan Experience

Ramli I, Rachadi H, Amarouch H, Bouhllab J, Senouci K and Hassam B

Departement of Dermatology and Venereology, University hospital Avicenna, University Mohammed V, Rabat-Morocco.

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#### ABSTRACT

Several studies have objectified a high prevalence of metabolic syndrom(MS) in psoriasis patients. Objective of the study is to determine the prevalence of metabolic syndrom and its components in psoriasis patients compared with controls, also the factors determinants the occurrence of MS in our patients. This was a prospective study including 100 psoriasis patients versus 100 controls seen in consultation or in hospitals. Conducted for a period of 12 months. The prevalence of MS was higher in patients with psoriasis compared with controls (OR= 2.89, p=0,0002). with a significant increase in obesity (OR = 3.25, p = 0.001), hypertriglyceridemia (OR = 3.9, p = 0.013) and glucose intolerance (OR = 2.13, p = 0.046). Risk factors determining the appearance of SM in psoriatic were age, family history of psoriasis, seniority of psoriasis, but we did not find a correlation with sex, severity of psoriasis and the clinical form. The management of psoriasis will integrate research risk factors for a possible metabolic syndrome and its treatment through a multidisciplinary approach.

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### Introduction

Psoriasis is a chronic, immune-mediated inflammatory skin disease, associated with metabolic and cardiovascular disease. Psoriasis is estimated to affect about 1- 3% of the population in Maghreb. A number of risk factors have been recognized in the etiology and pathogenesis of psoriasis, including family history and environmental risk factors, such as diet, obesity, smoking, stress, and alcohol consumption. Several studies have demonstrated a high prevalence of metabolic syndrome (MS) in psoriasis patients. It is responsible for a significant morbidity and mortality and amending prognosis and management of this disease. Many studies suggest that psoriasis is often associated with the MS to the point that some have suggested that psoriasis could be one of the elements of MS.

The objective of our study was to determine the prevalence of MS and its components in patients with psoriasis compared to controls, as well as the factors that determine the occurrence of MS in our patients and to compare our results with the literature data.

#### Material and methods

This is a prospective case control studycarried out at the Department of dermatology, universityhospitalAvicenna, Rabat-Morocco in the periodfromFebruary 2013 to February 2014 for psoriasis patients and control patients consultant in the sameperiod, receiving no treatmentthatcouldinterferewith the MS parameters. Weexcludedfrom the study nonconsenting patients and psoriatic patients whoreceivedsystemictreatment of psoriasis, includingacitretin, methotrexate or PUVA for at least one monthbefore the start of the study. The statistical data wereentered and

Tele: +212 6 61377771 E-mail address: inssaf.ramli@gmail.com © 2016 Elixir All rights reserved

analyzedusing	SPSS	13.0.,	And	р	< 0.05
wasconsideredsi	gnificant.				

 
 Table I. The prevalence of MS and its components in the psoriatic population and case controls

	Psoriaticgroup n (%)		Group control	OR	IC (95%)	Р			
	`	<i>,</i>	n (%)		× /				
MS	32	(32%)	14	2,891	1,430-	0.002			
			(14%)		5,84				
Glucose	22	(22%)	13	2,134	1,003-	0.046			
intolerance			(13%)		4,54				
Hypertension	36	(36%)	26	1,312	0,72-	0.36			
•••			(26%)		2,37				
abdominal	31	(31%)	13	3,255	1,57-	0.001			
obesity			(13%)		6,71				
Elevatedtriglyce	14	(14%)	4	3,907	1,23-	0.013			
rides			(4%)		12,32				
DecreaseHDL	37	(37%)	28	1,510	0,832-	0.17			
			(28%)		2,741				

Table II.	Factorsassociatedwit	h the development	of MS in
	patients with	n psoriasis	

MS	OR	Р	IC(96%)				
Age	1,033	0,016	[1,006-1,061]				
Sex	1.30	0.36	[0.702 -2.637]				
Family history of psoriasis	2,816	0,072	[0,913-8,682]				
PASI	0,988	0,383	[0,962-1,015]				
Evolution of psoriasis	1,061	0,031	[1,005-1,120]				
	• .	1					

OR : odds Ratio, IC confidence interval

#### Results

169 psoriasis patients werecollected of which 100 met the inclusion criteria. These patients wereseen in consultation in 22% of cases and in 78% are hospitalized cases. The averageage of our patients was 40.3 years +/- 19.5 years, rangingfrom 6-89 years. A male predominancewasnoted with a percentage of 59%. The sex ratio male / femalewas 1.44.

	Psoriasis	controls	OR	IC	Р
Italy[1]	30.1%	20.6%	1.65	1.16-2.35	
USA[2]	34%	26%	1.5	1.40-1.61	
Tunisia [6]	35.5%	30.8%	1.39	0.88-2.18	0.095
JAPAN [7]			1.82	1.12-3.21	
Germany [10]	4.3%	1.1%	5.92	2.78-12.8	0.001
Our series	32%	14%	2,891	1,430-5,845	0.002

Table III. Comparative table between prevalence of MS among psoriatic patients vs controls in the different series published

Table IV. The prevalence of different components of MS in psoriatic patients in differentseries published

SMparameters	USA[2] Tu		Tunisi	sia [4] Gern		any [6]	Our series	
	OR	р	OR	Р	OR	р	OR	Р
Glucose intolerance		0.58	1.2	0.3	2.48	< 0.001	2,13	0.04
hypertension		0.02	1.01	0.3	3.2	< 0.001	1,31	0.36
Abdominal obesity		0.003	2.2	0.002	-	-	3,25	0.001
Elevatedtrigly cerides		0.12	0.79	0.2	2.09	0.05	3,90	0.013
DecreaseHDL	2	0.004		0.64	-	-	1,51	0.17

The seniority of psoriasis was  $7.86 \pm 9$  years (2 months to 50 years). The mode of evolutionwas progressive in 85.4% of cases. A family psoriasis wasnoted in 19% of patients and 24% of psoriasis patients were smoking. Regarding control cases, the averageagewas 41.15 +/- 18.12 years. A male wasfoundwith a percentage of 57%. A chronic smoking wasnoted in 26% of cases. We found a higherprevalence of MS in patients with psoriasis compared to controlswith a statistically significant difference (OR = 2.89, p = 0.0002). By studying each parameter SM, we found a high prevalence of all components of the SM but this increase was significant for obesity (OR = 3.25, p = 0.001) hypertrigly ceridemia (OR = 3.9, p = 0.013) and glucose intolerance (OR = 2.13, p = 0.046) (Table II).

After an analytical survey statistics, risk factors determining the onset of MS in patients with psoriasis wereage (significantly for patients aged over 40 with p <0.001), family history of psoriasis (p = 0.05), the seniority of psoriasis (disease duration 10 years, p = 0.07), but we have not found a correlation with sex, psoriasis severity (p = 0.46) and clinical presentation (p = 0.18) (Table II).

#### Discussion

In the literatureseveral recent studies have shown an increase of MS and its components in patients with psoriasis. This relationship has not been adequately described in our Moroccancontext. The increase in the prevalence of MS in patients with psoriasis was observed in several countries including Italy, Israel, India, Japan, China, Tunisia and the United States with a prevalence ranging from 4.3 % to 35.5% and an OR of 1.39 to 5.92 (Table III) [1, 2].

Finally, new data indicatethat psoriasis association withMSoccursearly in the course of the diseasewhen psoriasis isassociated with obesity and hyperlipidemia [3]. In ourstudy, the prevalence of metabolic syndrome washigher in psoriatic with patients compared to controls cases а statisticallysignificantdifference 0.002). (p evenafterlogistic regression for age and sexwhichjoined data literature [1, 2]. The Tunisian and Japanesestudiesobjectified a highprevalence of MS in patients with psoriasis compared to controls but not statistically significant (p = 0.095) [4] [5]. In ourseries, the separateanalysis of eachparameter of MS showed a highprevalence of all parameters in psoriaticpatients butonly glucose intolerance, abdominal

obesity and hypertriglyceridemia were significant. Otherseriesfoundvaryingresults [2, 4, 6] (Table IV).

This demonstrates the epidemiological variability of psoriasis, probably related to genetic variations and the influence of socio-economic and environmental factors specific to each population [7]. This association has clinical implications for the treatment of patients with psoriasis, as regards screening MS in these patients, taking steps to reduce cardiovascular risk in these patients. So the recognition of the MS can have an impact on the safety and efficacy of the therapy of psoriasis. Thus our results and literature data may partly explain the increased risk of cardiovascular disease, metabolic and mortality in people with psoriasis reported in previous studies [8]. In the literature several factors influence the onset of MS in psoriasis [9].

In ourstudy, the factors associated with MS were first, the age and evolution of psoriasis with high prevalence in patients with psoriasis after 40 years and when the disease progresses over 10 years. This was consistent with published studies [1, 4, 6]. And secondly, the family history of psoriasis was significantly associated with MS in our patients (OR = 2.816 and p = 0.05). This could be explained by a common genetic predisposition between psoriasis and MS, in particular the existence of genetic locipleiotropic (PSORS2-4, CDKAL1 and ApoE4) [7]. In a Tunisian study [4], femalegender and plaque psoriasis were factors influencing the onset of MS in the psoriasis have no impact on the occurrence of MS [10].

#### Conclusion

After a review of the literature and our knowledge we report the first study evaluating the association of psoriasis with metabolic syndrome in Morocco. Our studys howedthatmetabolic syndrome, obesity, impaired glucose tolerance and hypertriglyceridemia were frequently associated in psoriasis patients compared to controls. Thus the metabolic syndrome should be routinely screened in the presence of these risk factors to ensuretaking adequate and comprehensive care to reduce cardiovascularrisk. **Conflict of interest** 

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None

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