



Archives • 2021 • vol.2 • 1-6

BIOCHEMICAL MARKERS TO EVALUATE THE RISK OF DEVELOPMENT CARDIOVASCULAR DISEASE IN PATIENTS WITH PSORIASIS

¹Iqbal Ghalib Farhood, MBChB, FICMS, ²Rawaa Ghalib Farhood, MBChB, MSc, PhD, ³Afraa <u>Mamoori</u>, MBChB, MSc, PhD

¹Section of Dermatology & Venereology, Department of Medicine, College of Medicine, Al-Nahrain University. Baghdad, Iraq

²Department of Pathology and Forensic Medicine, College of medicine, University of Babylon, Babylon, Iraq.

*³Department of Pathology and Forensic Medicine, College of medicine, University of Babylon, Babylon, Iraq.

med.afraa.mamoori@uobabylon.edu.ig

Abstract

Psoriasis is common chronic recurrent immune-mediated inflammatory genetically determined disorders in which outcome affected by environmental factors. It is considered as a systemic disease associated with increased cardiovascular abnormalities, hypertension, dyslipidemia, atherosclerosis, diabetes mellitus, obesity, stroke, osteoporosis.

To evaluate the risk factors of cardiovascular disease in patients with psoriasis.

This study is case-controlled was carried out in AL-Imamein AL-Kadhemein Medical City, in outpatient's clinic at Dermatology department, during the period of September 2018 to September 2019.

The study was done on 60 patients of psoriasis who attended Dermatology outpatient's clinic, a control group of healthy persons of 60 age and sex matched.

We need to calculate the body mass index which is the weight in kilograms (kg) which was divided by the square of the height in meters (m²). Psoriasis severity was assessed by psoriatic area and severity index (PASI) score.

Under aseptic precaution; 5ml of fasting blood samples was taken from all subjects and was tested for Fasting blood sugar. High density lipoprotein (HDL); Triglycerides (TG) and serum total cholesterol (TC). Sixty patients were included in this study, 33 (55%) male and 27(45%) female patients. Serum HDL in psoriatic patient was low in compare to controls which is statistically highly significant. Serum TG, LDL levels and serum glucose were high in psoriatic patient in compare to controls which is statistically highly significant.

Psoriasis severity assessed using the Psoriasis Area Severity Index; It was higher in male psoriatic patient (14.97±1.81) than in females (13.61+0.98) which is statistically highly significant. There were statistical differences in TG level and duration of disease which was high in males than in female's psoriatic patients. Between male and female's psoriatic patients; there were no statistical differences concerning age, body mass index, and fasting serum glucose level.

There was positive correlation between the Psoriasis Area Severity Index and age, duration of psoriasis and with serum LDL and serum TG in patients with psoriasis. Negative correlation was between the Psoriasis Area Severity Index with serum HDL level.

PhOL Farhood, et al. 2 (pag 1-6)

We conclude that psoriatic patients considered at high risk for development of cardiovascular disease. Treatment of Dyslipidemias and dietary antioxidants supplementation should be considered in the management of psoriasis to reduce the morbidity from cardiovascular events.

Keywords: Psoriasis, Cardiovascular Diseases, Risk Factors

Introduction

Psoriasis is one of the most common chronic recurrent genetically determined, immune-mediated inflammatory disorders in which environmental factors affecting its outcome (1). Clinically; it varies from a few scaly plaques at extensor sites, scalp and nail involvement, Psoriasis vulgaris most common type affects 85 to 90% of all patients (2).

Psoriasis is considered as a systemic disease associated with increased cardiovascular abnormalities, hypertension, dyslipidemia, atherosclerosis, diabetes mellitus, obesity, stroke, osteoporosis, cancer and depression which suggest that systemic inflammation reflects a causal relationship (3). Systemic treatment for psoriasis with methotrexate, cyclosporine, acitretin, and biological drugs increase the cardiovascular disease (CVD) risks as hypertension, dyslipidemia (4).

oxidative stress and abnormal lipid metabolism were proposed the suitable explanation for increasing risk of cardiovascular disease (5). Pathogenesis of psoriasis is thought to be due to deficient antioxidant system and Increased reactive oxygen species (ROS) (6), increasing the atherosclerotic risk with subsequent cardiovascular events (7).

Adipose tissue overproduces multiple proinflammatory cytokines in response to obesity. Tumor necrosis factor (TNF-a), interleukin (IL)-6, IL-8 and reactive C protein involve in pathogenesis of psoriasis (8). This study aims to assess of risk factors of cardiovascular disease in patients with psoriasis.

Methods

This study is case-controlled was carried out in Al-Imamein AL-Kadhemein Medical City, in outpatient's clinic at Dermatology department, during the period of September 2018 to September 2019. The study was done on 60 patients of psoriasis who attended Dermatology outpatient's clinic, a control group of healthy persons of 60 age and sex matched.

Information includes demographics, medical history, lifestyle habits, drugs use and laboratory findings.

The inclusion criteria: Patients with severe plaque psoriasis of 1-year duration, PASI of more than 10, no history for at least 6 months prior to recruitment of any systemic anti-psoriatic therapy.

The exclusion criteria: significant cardiovascular disease, diabetics, hypertension, dyslipidemia, and any patients with other types of psoriasis.

The severity of psoriasis was assessed by PASI score (8). Scoring of the clinical signs in each area are summed and are finally weighted according to the area's proportion of the body, which ranged from 0 to a theoretical maximum of 72.

- Patients with a PASI > 12 have severe psoriasis.
- Clinically significant plaques covering less than 10% of the integument is moderate plaque psoriasis and result in a PASI = 7.
- If a patient with PASI < 7 as mild chronic plaquetype psoriasis (9, 10,11).

We need to calculate the body mass index which is the weight in kilograms (kg) which was divided by the square of the height in meters (m²). Psoriasis severity was assessed by PASI score (12).

Under aseptic precaution; 5ml of fasting blood samples was taken from all subjects and was tested for Fasting blood sugar. High density lipoprotein (HDL); Triglycerides (TG) and serum total cholesterol (TC).

Statistical Analysis:

Statistical analysis was done using unpaired student "t" test and probability value (p) of <0.05 was considered as statistically significant.

PhOL Farhood, et al. 3 (pag 1-6)

Results

Sixty patients were included in this study, 33 (55%) male and 27(45%) female patients. Age were ranged from 30 – 60 years with 43.58+10.13, Psoriasis duration ranged from 2 - 25 years with mean 8.78+6.58, Mean Body mass index was 28.29+3.78 statistically significant in psoriatic patients in comparison to controls (26.4+4.83). Serum HDL in psoriatic patient was low in compare to controls which is statistically highly significant, Serum TG was high in psoriatic patient in compare to controls which is statistically highly significant, Serum LDL was high in psoriatic patient in compare to controls which is statistically highly significant, Serum glucose level was was high in psoriatic patient in compare to controls which is statistically highly significant as in table (1).

Severity of psoriasis assessed using the Psoriasis Area Severity Index; in male psoriatic patient was 14.97±1.81 higher than in females was 13.61+0.98which is statistically highly significant. There were statistical differences in TG level and duration of disease which was high in males than in females. There were no statistical differences between male and female's psoriatic patients concerning age, body mass index, and fasting serum glucose level as in table (2).

There was positive correlation between the Psoriasis Area Severity Index with age, duration and with serum LDL and serum TG in patients with psoriasis. Correlation was negative between the Psoriasis Area Severity Index with serum HDL level as in table (3).

Correlation was positive between the duration of psoriasis and age, the Psoriasis Area Severity Index, serum TG, serum LDL, and negative correlation between the duration of disease with serum HDL level as in table (4).

There were 20 (33.3) psoriatic patients with hypertension show highly significant statistical differences to those who were non hypertensive psoriatic patients in age, serum TG level, serum LDL level, duration of disease and in the Psoriasis Area Severity Index as in table (5,6)

Discussion

Psoriasis is common chronic inflammatory skin disorders. Many organs not only the skin affected by inflammatory process including cardiovascular system (4). Psoriatic disease has clear association with obesity and its related metabolic abnormalities. Clinical cardiovascular provoked by dyslipidaemia, glucose intolerance and hypertension (13). In our study, I evaluate obesity by using BMI which was significantly increased in patients with psoriasis in comparison with controls. Abnormal lipid profile is a well-known cardiovascular risk factor. In our study I found highly significant elevated triglycerides and highly significant lower HDL levels which is inconsistent with other study which found increased serum HDL (14). Decreased antioxidant activity with the process of chronic inflammation resulting from elevated cholesterol level. All these together with effects of drugs used in the treatment of psoriasis as cyclosporine and acitretin.

In this study20 (33.3%) patients had hypertension which is statistically significant as in other studies ⁽¹⁵⁾. There were highly significant statistical differences to those who were non hypertensive psoriatic patients in age, serum TG level, serum LDL level, duration of disease and in the Psoriasis Area Severity Index. Many studies showed that the chronic inflammatory process responsible for hypertension as Huskic et al. found that psoriatic patients had increased concentration of tissue angiotensin-converting enzyme ⁽¹⁶⁾.

the Fasting glucose level was elevated and show highly significant differences between psoriatic patient and the control groups but there was negative correlation between the Psoriasis Area Severity Index and serum glucose level.

We conclude that psoriatic patients considered at high risk for development of cardiovascular disease. Treatment of Dyslipidemias and dietary antioxidants supplementation should be considered in the management of psoriasis to reduce the morbidity from cardiovascular events.

Acknowledgments

We appreciate the role of all persons who participate in this study.

PhOL Farhood, et al. 4 (pag 1-6)

References

- 1-Pavithra K, Karunakaran, Apama P, Ragunath S. (2008). Disorders of keratinisation. In: Valia RG, Ameet RV, editors. IADVL Textbook of Dermatology, vol 1. 3rd ed. Mumbai, Bhalani publishing house,1021-1056.
- 2. Griffiths CE, Barker JN. (2007). Pathogenesis and clinical features of psoriasis. Lancet, 370:263-71
- 3. Aldona Pietrzak, Anna Michalak-Stoma, Grazyna Chodorowska, Jacek Szepietowski C. (2010). Lipid disturbances in psoriasis: An update. Mediators of Inflammation 2010, Article ID 535612.
- 4. Pietrzak A, Bartosińska J, Chodorowska G, Szepietowski J. C, Paluszkiewicz P, and. Schwartz R. A. (2013). "Cardiovascular aspects of psoriasis: an updated review," International Journal of Dermatology, vol. 52, no. 2, pp. 153–162.
- 5. Rocha-Pereira P, Santos-Silva A, Rebelo I, Figueiredo A, Quintanilha A, Teixeira F. (2001). Dyslipidemia and oxidative stress in mild and in severe psoriasis as a risk for cardiovascular disease. Clin Chim Acta, 303 (12):33-9.
- 6. Kiymet Baz, Burak Cimen MY, Aysin Kokturak, Ayca CordanY, Gulcin Eskandari, Guliz Ikizoglu et al. (2003). Oxidant/Antioxidant status in patients with psoriasis. Yonsei Med J, 44(6):987-90.
- 7. Tripti Sexena, Agarwal BK, Pawan Kare. (2011). Serum paraoxonase activity and oxidative stress in acute myocardial infarction patients. Biomedical Research, 22(2):215-9.
- 8. Hamminga EA, van der Lely AJ, Neumann HA, Thio HB. (2006). Chronic inflammation in psoriasis and obesity: implications for therapy. Med Hypotheses 67: 768-773.
- 9. Parameswaran N, Patial S. (2010) . Tumor necrosis factor- α signaling in macrophages. Crit Rev Eukaryot Gene Expr 20: 87-103.

- 10. Fredriksson T, Pettersson U. (1978). Severe psoriasis: oral therapy with a new retinoid. Dermatologica,157(4):238–44
- 11. Sampogna F, Sera F, Abeni A. (2004). Measures of clinical severity, quality of life, and psychological distress in patients with psoriasis: A cluster analysis. J Invest Dermatol, 122: 602–607.
- 12. Bardazzi F, Balestri R, Baldi E, et al. (2010). Correlation between BMI and PASI in patients affected by moderate to severe psoriasis undergoing biological therapy. Dermatol Ther, 1: 14-9
- 13. Miller IM, Skaaby T, Ellervik C, et al. (2013). Quantifying cardiovascular disease risk factors in patients with psoriasis: a meta-analysis. Br J Dermatol, 169:1180–7.
- 14. Owczarczyk-Saczonek A, Nowicki R. (2015). Prevalence of cardiovascular disease risk factors, and metabolic syndrome and its components in patients with psoriasis aged 30 to 49 years. Postep Dermatol Alergol, 32:290-5.
- 15. Szponar-Bojda A, Krasowska D, Pietrzak A, Chodorowska G. (2012). Metabolic syndrome in psoriasis. Post Dermatol Alergol, 29:356-62.
- 16. Huskic J, Alendar F. (2007). Tissue angiotensinconverting enzyme in patients with various clinical forms of psoriasis. Bosn J Basic Med Sci, 7:103-6.

ISSN: 1827-8620

PhOL Farhood, et al. 5 (pag 1-6)

Table (1): Comparison of parameters between patients and controls by unpaired t test

Parameter	Patients N=60 Mean+SD	Controls N=60 Mean+SD	P value
Age (yr)	43.58+10.13	41.52+8.47	0.228
BMI (kg/m²)	28.29+3.78	26.4+4.83	0.019
HDL (mg/dl)	54.93+5.99	59.17+1.07	<0.001
TG (mg/dl)	146.68+13.91	123.07+6.02	<0.001
LDL (mg/dl)	114.6+27.61	71.35+12.83	<0.001
Glucose (mg/dl)	141.58+43.94	109.17+33.27	<0.001
Duration (yr)	8.78+6.58		
PSAI	14.36+1.63		

Table (2): Comparison of parameters according to gender by unpaired t test

Parameter	Females N=27 Mean+SD	Males N=33 Mean+SD	P value
Age (yr)	41.78+7.39	45.06+11.83	0.195
BMI (kg/m²)	28.76+3.38	27.91+4.09	0.385
HDL (mg/dl)	56.08+5.6	53.98+6.22	0.180
TG (mg/dl)	142.44+14.23	150.15+12.84	0.031
LDL (mg/dl)	111.34+24.05	117.26+30.32	0.413
Glucose (mg/dl)	142.59+34.93	140.76+50.67	0.874
Duration (yr)	6.93+3.77	10.29+7.94	0.036
PSI	13.61+0.98	14.97+1.81	0.001

Table (3): Correlation of PASI with other parameters within patients group

Parameters	PASI		
	r	Р	
Age (yr)	0.561	<0.001	
Duration (yr)	0.619	<0.001	
BMI (kg/m²)	0.180	0.169	
HDL	-0.283	0.029	
TG	0.376	0.003	
LDL	0.468	<0.001	
Glucose	-0.156	0.235	

PhOL Farhood, et al. 6 (pag 1-6)

Table (4): Correlation of duration of psoriasis with other parameters within patients group

Parameters	Duration		
	r	Р	
Age (yr)	0.875	<0.001	
PASI	0.619	<0.001	
BMI (kg/m²)	0.162	0.218	
HDL	-0.478	<0.001	
TG	0.679	<0.001	
LDL	0.679	<0.001	
Glucose	-0.097	0.460	

Table (5): Comparison between patients and controls by Fisher exact test

Parameter		Patients N=60 No. (%)	Controls N=60 No. (%)	P value
Sex	Females	27 (45.0)	32 (53.3)	0.465
	Males	33 (55.0)	28 (46.7)	
Hypertension	No	40 (66.7)	60 (100)	<0.001
	Yes	20 (33.3)	0 (0.0)	

Table (6): Comparison of parameters according to hypertension by unpaired t test

Parameter	Negative N=40 Mean+SD	Positive N=20 Mean+SD	P value
Age (yr)	37.65+6.18	55.45+4.24	<0.001
BMI (kg/m²)	28.04+3.84	28.8+3.7	0.468
HDL (mg/dl)	56.51+5.57	51.76+5.66	0.003
TG (mg/dl)	139.98+10.7	160.09+9.09	<0.001
LDL (mg/dl)	99.96+20.66	143.87+11.75	<0.001
Glucose (mg/dl)	145.38+46.35	134+38.68	0.349
Duration (yr)	5.21+3.13	15.9+5.85	<0.001
PSI	13.78+1.49	15.52+1.25	<0.001