

**Nailfold capillaroscopy examination in patients
with systemic lupus erythematosus (SLE) and
correlation with its various clinical and
laboratory data**

Thesis

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ABSTRACT

This work aimed to study the prevalence of nailfold capillaroscopy changes in patients with systemic lupus erythematosus, find out the patterns of these changes and to correlate these findings with different clinical and laboratory parameters.

Fourty patients with SLE, all fulfilling the 1997 revised criteria for the classification of SLE. All patients included in this study were subjected to full history taking, clinical examination, laboratory investigations as well as nailfold capillaroscopy examination.

Nail fold capillaroscopic abnormalities were significantly found in SLE patients in relation to controls. Some of these abnormalities showed statistical significant correlations with different clinical and laboratory parameters.

Keywords: - Systemic lupus erythematosus - Nailfold capillaroscopy

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LIST OF ABBREVIATIONS

ACA	Anticentromere antibodies
aCL	anti-cardiolipin antibodies
ACLE	Acute cutaneous lupus erythematosus
ACR	American College of Rheumatology
AECA	Antiendothelial cell antibody
ANA	Antinuclear antibody
Anti-ds DNA	Anti- double stranded deoxyribonucleic acid
aPL	antiphospholipid antibodies
CAD	Coronary artery disease
CBC	Complete blood count
CCLE	Chronic cutaneous lupus erythematosus
CECs	circulating endothelial cells
CTD	Connective tissue diseases
CVA	Cardiovascular accident
DLCO	Diffusing lung capacity for carbon monoxide
DLE	discoid lupus erythematosus
DM	dermatomyositis
EC	endothelial cell
ECG	Electrocardiogram
ECs	Endothelial cells
EMC	Essential mixed cryoglobulinemia
ESR	Erythrocyte sedimentation rate

EULAR	European League Against Rheumatism
FM	Fibromyalgia
gm	gram
HB	Hemoglobin
HSCT	Hemopoietic stem cell transplantation
ICAM 1	Intercellular adhesion molecule 1
IFN-γ	Interferon-gamma
IL-1	Interleukin-1
JIA	Juvenile idiopathic arthritis
LE	Lupus erythematosus
Max	Maximum
MCTD	Mixed connective tissue disease
mm	Millimeter
mRNA	Messenger ribonucleic acid
NFC	Nailfold capillaroscopy
NO	Nitrous oxide
OA	Osteoarthritis
PAPS	Primary antiphospholipid syndrome
PMNs	Polymorphonuclear leukocytes
PRP	Primary Raynaud's phenomenon
PSS	Primary Sjögren's syndrome
RA	Rheumatoid arthritis
RP	Raynaud's phenomenon
S.D.	Standard deviation

SCLE	Subacute cutaneous lupus erythematosus
SD-pattern	The scleroderma dermatomyositis pattern
SLE	Systemic lupus erythematosus
SLEDAI	Systemic Lupus Erythematosus Disease Activity Index
SPVP	Subpapillary venous plexus
SRP	Secondary Raynaud's phenomenon
SS	Sjögren's syndrome
SSc	Systemic sclerosis
SSD	Scleroderma spectrum disorder
SSS	Secondary Sjögren's syndrome
TF	Tissue factor
TLC	Total leucocytic count
TNF-α	Tumor necrosis factor-α
TTP	Thrombotic thrombocytopenia purpura
UCTD	Undifferentiated connective tissue disease
VCAM 1	Vascular cell adhesion molecule-1
vWF	Von Willebrand factor
WBC	White blood cells
μm	Micrometer

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INTRODUCTION

Systemic lupus erythematosus (SLE) is a complex disease with variable presentations, course and prognosis (**Bertsias et al., 2008**).

SLE is the most diverse of the systemic autoimmune diseases because it may affect any organ of the body and displays a broad spectrum of clinical manifestations (**Cervera et al., 1999**).

A pathologic hallmark of SLE is the appearance of diverse vascular lesions (**Font et al., 2001**).

Over the past 25 years, nailfold capillaroscopy (NFC) has gained a diagnostic value in the field of rheumatology based on descriptive data from patients with distinct CTD (**Anders et al., 2000**).

Few diagnostic techniques can combine all the positive features typical of capillaroscopy (low cost, uninvassiveness, repeatability, high sensitivity, good specificity and easy interpretation of results) (**Grassi and De Angelis, 2007**).

NFC may be a useful method to evaluate the microvascular changes in patients with SLE (**Ingegnoli et al., 2005**) and it was proved to be an easy-to-perform noninvasive technique, able to achieve useful data to better evaluate such a pleomorphic disease especially concerning its outcome and prognosis (**Ricciari et al., 2005**).

Capillaroscopy can give a clue for early diagnosis of patients, since the presence of typical capillaroscopic abnormalities seems to be related to the development of lupus erythematosus (**Facina et al., 2006**).

Abnormalities in nailfold capillaroscopy may reflect the extent of microvascular involvement in SLE. Nailfold capillaroscopy can be useful in the diagnostic procedure of the disease and lead to more effective strategies in the treatment of systemic organ dysfunction in SLE (**Kuryliszyn-Moskal et al., 2007**).

Kuryliszyn-Moskal and colleagues (2007) reported that all SLE patients with internal organ manifestations showed severe or moderate pathological changes under NFC. Mild changes in capillaroscopy in patients with systemic involvement were not observed.

Also, **Ingegnoli and colleagues (2005)** reported that the presence of major capillary abnormalities in NFC examination in SLE patients seems to herald a more severe clinical course of the illness.

AIM OF THE WORK

- 1- To study the prevalence of nailfold capillaroscopic changes in patients with SLE.
- 2- To find out the patterns of these changes
- 3- To correlate these findings with different clinical and laboratory parameters.