Periungual Capillaroscopy Findings in a Series of Patients with Granulomatosis with Polyangiitis (Wegener): An Observational Study

Uceda Montañes Julia, Martínez Pérez Rosalía, Velloso Feijoo María Luisa, Marenco de la Fuente José Luis

Rheumatology Unit, Valme University Hospital, Seville, Spain.
Email: juliauced@gmail.com

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ABSTRACT

Introduction: Periungual capillaroscopy studies skin microcirculation. Its main indication is in the differential diagnosis of Raynaud’s phenomenon as well as diagnosis and prognosis of systemic sclerosis (SS). Many rheumatic diseases present characteristic capillaroscopic alterations, but little is known at present about the alterations in skin microcirculation of systemic vasculitis. Objective: Our main objective was to study the alterations in skin microcirculation by capillaroscopy in patients with granulomatosis with polyangiitis (GPA). The secondary objective was to evaluate the association of capillaroscopic alterations with systemic alteration. Material and Methods: We studied patients in our unit with GPA, and we collected demographic and clinical variables. Periungual capillaroscopy was performed on all patients, and always by the same investigator. Results: We reviewed 10 patients with a mean age of 55.7. They were predominantly women. We found capillaroscopic alterations in 8 of the 10 patients. Morphological alteration was the most common finding (62.5%), followed by bleeding (50%). We found no mega-capillaries or neovessels. These findings were not related to any specific organ system involvement. Discussion: The tortuous pattern with bleeding was predominant in our series, with lower capillary density, findings which were unrelated to any specific organ system involvement. A single study describes capillaroscopic alterations in GPA, detecting a high percentage (92%) of avascular areas, in contrast with the predominance of a twisting pattern and bleeding found in our study. Further studies are warranted to confirm these results.

Keywords: Periungual Capillaroscopy; Vasculitis; Granulomatosis with Polyangiitis

1. Introduction

Periungual capillaroscopy dates from 1911 [1], it is a simple and non-invasive technique for studying cutaneous microcirculation. It is performed on the periungual bed as this area is where the capillaries are arranged parallel to the skin surface, thus enabling the study of their entire length.

It is useful in both the diagnosis and prognosis of certain rheumatic diseases that present with microcirculation damage. Nowadays the main indication of this technique is in the differential diagnosis of Raynaud’s phenomenon [2,3] and prognosis of scleroderma.

Currently we only have one specific pattern of involvement of cutaneous microcirculation: sclerodermiform pattern, typical of scleroderma. This pattern was first defined by Maricq in 1973 [4], and then redefined in 2000 by Cutolo [5].

In other rheumatic diseases such as dermatomyositis/polymyositis [6], mixed connective tissue disease, systemic lupus erythematosus—antiphospholipid syndrome [7,8], Sjögren’s syndrome [9], rheumatoid arthritis [7] and psoriatic arthritis [10], we find capillaroscopic alterations suggestive of each of these rheumatic diseases, but specific capillaroscopic patterns have not yet been described for each of them. Further research studies are warranted in order to define specific patterns.

Currently there are limited publications on possible capillaroscopic alterations in patients with vasculitis, and to date there is no known specific capillaroscopic pattern as is the case in other rheumatic diseases [11-13].

As with other types of vasculitis, in granulomatosis with polyangiitis in (GPA) (Wegener) there have been...
Table 1. Baseline characteristics of the patient.

<table>
<thead>
<tr>
<th>Patient</th>
<th>Sex</th>
<th>Fever</th>
<th>Arthritis</th>
<th>Oral and upper airway involvement</th>
<th>Oral and upper respiratory involvement</th>
<th>Cutaneous involvement</th>
<th>Pulmonary involvement</th>
<th>Pulmonary involvement present</th>
<th>Renal involvement</th>
<th>Renal involvement present</th>
<th>Neurological involvement</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Female</td>
<td>Yes</td>
<td>Yes</td>
<td>Nasal ulcers and vocal cord granulomas</td>
<td>Yes</td>
<td>Purpura</td>
<td>Cavitary nodules in LII</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Axonal mononeuropathy</td>
<td>Rituximab, cyclophosphamide, prednisone, azathioprine</td>
</tr>
<tr>
<td>2</td>
<td>Male</td>
<td>Yes</td>
<td>Yes</td>
<td>Chronic sinusitis</td>
<td>Yes</td>
<td>Purpura</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Cyclophosphamide and prednisone</td>
</tr>
<tr>
<td>3</td>
<td>Male</td>
<td>No</td>
<td>No</td>
<td>Maxillary sinusitis</td>
<td>Yes</td>
<td>Purpura</td>
<td>Pulmonary haemorrhage</td>
<td>Yes</td>
<td>Yes mild renal impairment, developing</td>
<td>Yes</td>
<td>Multiple mononeuropathy</td>
<td>Prednisone and cyclophosphamide</td>
</tr>
<tr>
<td>4</td>
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<td>No</td>
<td>No</td>
<td>Maxillary sinusitis</td>
<td>Yes</td>
<td>Purpura</td>
<td>No</td>
<td>No</td>
<td>Mild CKF</td>
<td>Yes</td>
<td>Paraesthesias?</td>
<td>Prednisone</td>
</tr>
<tr>
<td>5</td>
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<td>No</td>
<td>Maxillary sinusitis</td>
<td>Yes</td>
<td>No</td>
<td>Alveolar haemorrhage</td>
<td>Yes</td>
<td>Yes nephrotic</td>
<td>Yes</td>
<td>No</td>
<td>Prednisone and cyclophosphamide</td>
</tr>
<tr>
<td>6</td>
<td>Female</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Purpura</td>
<td>Cavitary nodules in LII</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Prednisone</td>
</tr>
<tr>
<td>7</td>
<td>Male</td>
<td>Yes</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>Ischemic lesions in lower limbs</td>
<td>Cavitary bilateral nodules</td>
<td>Yes</td>
<td>Proliferative focal and (illegible) GN</td>
<td>Yes</td>
<td>Cyclophosphamide and prednisone</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>Female</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Necrotic ulcer in lower limbs</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Cyclophosphamide and prednisone</td>
</tr>
<tr>
<td>9</td>
<td>Male</td>
<td>Yes</td>
<td>Yes</td>
<td>Maxillary sinusitis</td>
<td>Yes</td>
<td>No</td>
<td>Cavitary bilateral nodules</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>10</td>
<td>Female</td>
<td>Yes</td>
<td>Yes</td>
<td>Maxillary sinusitis</td>
<td>Yes</td>
<td>No</td>
<td>Cavitary bilateral nodules</td>
<td>Yes</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>Axonal mononeuropathy</td>
</tr>
</tbody>
</table>
few studies on the possible capillaroscopic alterations that these patients may present. Anders et al., studied 12 patients with GPA and found capillaroscopic alterations in 92%, with avascular areas being the characteristic finding in these patients. Capillary dilatations were not observed [14].

With further studies on the findings of periungual capillaroscopy and comparison with what is described in the literature to date, we might obtain capillaroscopic alterations specific to this type of vasculitis as we have in other rheumatic diseases, and therefore use this technique as another tool for a given diagnosis and prognosis of this pathology.

The aim of the study is to describe the findings in periungual capillaroscopy in our series of patients with GPA and compare them with those in the literature today.

2. Methods
A descriptive cross-sectional study on alterations detected in periungual capillaroscopy in patients with GPA was carried out.

2.1. Subjects/Patients
The study was carried out at the Rheumatology Unit of the Virgen de Valme Hospital, Seville, Spain.

We reviewed all patients diagnosed with GPA under the ACR criteria [15], which are currently seen in our unit. Demographic and clinical variables were recorded.

2.2. Measurements and Data Collection
The capillaroscope employed was a stereomicroscope (stereoscope) with trinocular head, zoom range from 1x to 4x magnification, with cold light illuminator and high resolution camcorder eyepiece.

Periungual capillaroscopy was performed on the 3rd, 4th and 5th fingers of the right and left hands in each patient and always by the same rheumatologist. In each capillaroscopy the following were assessed:
1) The morphology of the loops: predominant type and presence of neovessels;
2) Dilatation of the loops: size, frequency of dilated loops and location of the dilatation;
3) Capillary density and presence of avascular areas;
4) Haemorrhages: number and form;
5) Visibility of venular plexus;
6) General appearance.

2.3. Analysis
The sample is described in terms of the distribution of variables using central tendency measures. The capillaroscopic findings in each patient are described and correlated with the organ system involvement of the patients.

3. Results
We reviewed 10 patients with GPA. The current average age was $55.7 \pm 16.5$ years. They were predominantly women (60%). The average age at diagnosis was $49.4 \pm 15.86$ years. Seventy percent of patients presented involvement of the upper respiratory tract (sinusitis, nasal ulcers); the same percentage presented pulmonary involvement (cavitary nodules or alveolar haemorrhage); skin symptoms such as purpura or necrotic ulcers were present in 70%. Forty percent had renal involvement (kidney failure, proliferative glomerulonephritis), and 40% had peripheral neurological involvement (Table 1).

All patients were able to undergo the capillaroscopic study. Capillaroscopic alterations were found in 8 of the 10 patients. Among patients with pathological capillaroscopy 62.5% had alterations in capillary morphology (tortuous capillaries), 50% haemorrhage (single or multiple), 37.5% avascular areas and 75% lower capillary density. There was no capillary dilatation, megacapillaries, and formation of new vessels (Table 2).
Table 3. Capillaroscopic alterations according to organ system involvement.

<table>
<thead>
<tr>
<th>Organ system involvement</th>
<th>Capillaroscopic pathology</th>
<th>Abnormal morphology</th>
<th>Bleeding</th>
<th>Avascular areas</th>
<th>Lower capillary density</th>
<th>Dilatation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Respiratory (7)</td>
<td>5</td>
<td>3</td>
<td>3</td>
<td>2</td>
<td>5</td>
<td>0</td>
</tr>
<tr>
<td>Renal (4)</td>
<td>3</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td>Neurological (4)</td>
<td>3</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td>Skin (7)</td>
<td>6</td>
<td>3</td>
<td>3</td>
<td>2</td>
<td>4</td>
<td>0</td>
</tr>
</tbody>
</table>

When the data were analysed in terms of organ system involvement, the following were found (Table 3).

4. Discussion

Periungual capillaroscopy is most useful for the study of Raynaud’s phenomenon, especially in SS, because of its prognostic value and correlation with disease severity, as there are clearly defined patterns.

In our series, we observed a higher frequency of twisting capillaries than in the healthy population, and an increase in the presence of bleeding in the periungual bed. In the patients studied, we found less capillary density and presence of avascular areas. None of these findings in periungual capillaroscopy could be related to specific organ system involvement. Our results contrast with Anders et al., published in 2000. They studied 12 patients with GPA and found capillaroscopic alterations in 92%, with avascular areas being the characteristic finding in these patients. No capillary dilatations were observed [14]. Both studies are on a small number of patients.

In summary, the data obtained in our study and in other previous series, leads us to believe that patients with GPA present alterations in skin microcirculation. These are preliminary results and could not be used to draw conclusions. Further studies are warranted with larger numbers of patients to compare the alterations detected.

REFERENCES


