

Rheumatoid vasculitis: same hand, heterogeneous clinical presentation

Otávio Augusto Gomes Paz*

University Center of Pará (CESUPA), Area of Environmental, Biological and Health Sciences. Belem, Brazil

Case report

A 63-year-old non-smoker woman with a 15 years history of rheumatoid arthritis characterized by erosive polyarthrititis, rheumatoid factor elevated titers, and positive testing for anti-CCP antibodies. The treatment included methotrexate, prednisolone and hydroxicloroquine. Despite of treatment, developed ischemic ulcer at left hand dorsum and nailfold infarction and digital ischemia of the left fingers [1].

Rheumatoid vasculitis is an inflammatory process that primarily affects small to medium-sized vessels. It's highly heterogeneous clinically, with wide-spread organ involvement. The incidence has declined in the past several decades, but cutaneous remains the most common presentation. It typically occurs in patients with long-standing erosive deforming Rheumatoid Arthritis, and its manifestation is heterogeneous, depending the size of the blood vessel. The skin can present purpura, nailfold infarcts, digital gangrene and cutaneous ulcers [2].



Figure 1. A typical rheumatoid hand, with ulnar deviation, atrophy of interosseous muscles, metacarpophalangeal subluxation, and deformities (swan's neck and boutonniere).



Figure 2. There are two forms of rheumatoid vasculitis, represented by ulceration in the third metacarpophalangeal joint and periungual infarction in the figure, as well as the discrete points of ischemia in the digital pulps.

The photographs demonstrates a typical rheumatoid hand, with ulnar deviation, atrophy of interosseous muscles, metacarpophalangeal subluxation, and deformities (swan's neck and boutonniere). there are two forms of rheumatoid vasculitis, represented by ulceration in the area of the third metacarpophalangeal joint and periungual infarction in the second one, as well as the discrete points of ischemia in the digital pulps. This image reflects the heterogeneity of the clinical presentations of this single entity (Figures 1 and 2) [3].

References

1. Makol A, Matteson EL, Warrington KJ (2015) Rheumatoid vasculitis: an update. *Curr Opin Rheumatol* 27: 63-70. [[Crossref](#)]
2. Scott DG, Bacon PA, Tribe CR (1981) Systemic rheumatoid vasculitis: A clinical and laboratory study of 50 cases. *Medicine (Baltimore)* 60: 288-297.
3. Makol A, Crowson CS, Wetter DA, Sokumbi O, Matteson EL, et al. (2014) Vasculitis associated with rheumatoid arthritis: a case-control study. *Rheumatology (Oxford)* 53: 890-899. [[Crossref](#)]

Copyright: ©2018 Paz OAG. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Correspondence to: Otávio Augusto Gomes Paz, University Center of Pará (CESUPA), Area of Environmental, Biological and Health Sciences. Belem, Brazil, Tel: (91) 4005-0550; E-mail: otavionrp@gmail.com

Received: March 14, 2018; **Accepted:** March 27, 2018; **Published:** March 30, 2018