## Dr. Lubrano, et al reply

To the Editor:

We have read the letter of Dr. Scarpa<sup>1</sup>. Our editorial was written as a proposal for a different vision of psoriatic arthritis (PsA), a multifaceted syndrome, in the light of a better treatment stratification, and more in general, personalized medicine<sup>2</sup>.

In our editorial, we pointed out that PsA, because of its phenotypic complexity, could be better identified and managed as a syndrome, stimulating the reader to think of this condition as a combination of symptoms and signs that together represent a disease process<sup>2</sup>.

Indeed, we are aware that the pathogenesis, at present, is unique for all manifestations of PsA<sup>3</sup> and we did not mention that the term *syndrome* means different pathogenesis of the various manifestations of PsA.

However, PsA belongs to the group of spondyloarthritis (SpA), a wide spectrum or a constellation of diseases with some common genetic background<sup>4.5</sup> and it is potentially a separate disease of this group or a major characteristic of the same disease<sup>6</sup>.

As a similar condition, Behçet disease has been identified as a syndrome because of its potential complexity, and using this definition can help with disease management<sup>7</sup>.

Finally, we hope that our proposal to identify PsA as a syndrome could be of some interest, to provide better treatment management toward a target-to-treat strategy<sup>8</sup>. This is not a step back but a step forward to implement our knowledge and management of PsA.

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