Rheumatoid Cachexia: What Is It and Why Is It Important?

To the Editor:

We were delighted that rheumatoid cachexia (RC), the condition of reduced muscle mass and increased fat mass associated with rheumatoid arthritis (RA), received editorial attention in the March issue of The Journal of Rheumatology. As pointed out by Rajbhandary, et al, while RC is evident in the majority of patients with RA, and is a key contributor to the patient’s disability, increased morbidity, and premature mortality, this important feature of RA receives disproportionately little clinical consideration. Thus, all efforts to highlight the clinical significance of RC are to be applauded. However, we consider it necessary to comment on several conclusions in this editorial, which, left unchallenged, will mislead readers unfamiliar with the subject.

Rajbhandary, et al state that “anti-TNFα [tumor necrosis factor-α] therapy is an intervention that seems to improve RC,” and substantiate this claim by reference to results from one of our studies. However, this conclusion is not supported by either our findings or those of subsequent investigations, as none of these studies observed improvements in lean body mass in patients with RA following anti-TNF therapy. Further, the investigations of Metsios, et al6,7,8, among others, have shown substantial increases in lean body mass and total fat mass in patients treated with anti-TNF agents. Thus, current findings do not support the use of anti-TNF therapy specifically to counter RC.

The editorial also claims that “studies have shown that increased protein intake would be helpful for overcoming the catabolic process in RA, particularly when combined with an exercise intervention,” and refers to another of our studies as the supporting reference. While we agree that the combination of protein supplementation and exercise is likely to be successful in reversing the adverse body composition changes of RC, this effect has not been shown, as our intervention did not include exercise, and to our knowledge this combined intervention has yet to be trialled in patients with RA. The authors and title of our report were also incorrectly transcribed in the editorial — readers are directed to the reference list below to find the correct biographical details.

Additionally, although correctly stating that high-intensity progressive resistance training (HI PRT; i.e., weight training) is efficacious in reversing the adverse effects of RC on body composition, Rajbhandary, et al refer to any of the studies (e.g., Marcara, et al9, Hakkinen, et al10, Lemmey, et al11), including a randomized controlled trial performed by our group9, that have shown this to be the case. [Each of the studies referenced6,7,8, among others, has shown substantial increases in lean body mass and reductions in fat mass following HI PRT.] Instead, the study they do refer to is that of Rall, et al11, which, alone among studies investigating the effects of HI PRT on RC, reports no changes in body composition following training. We do, however, agree with their conclusion that HI PRT “is the best way to improve muscle strength and physical functioning in patients with RA” and “should be routinely prescribed and maintained.”

Speculating on the etiology of RC, the editorial refers to one of our reports12 that described reduced serum insulin-like growth factor-I (IGF-I) levels, and echoes our conclusion at the time that these diminished levels may contribute to RC. However, these earlier postulations have been superseded by our finding that attenuated muscle levels of IGF (mIGF-I, mIGF binding protein-3) are much more likely to be involved in the mechanism underlying RC8.

Finally, we were surprised and disappointed that, in an area of research where there has been a recent upsurge in interest, the editorial relied almost exclusively on old references (i.e., 18 out of 19 references were from 2006 or before; 13 of 19 were from 2002 or before), and thus overlooked important recent developments in the field.

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