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### Evaluation of Therapeutic Success of Hyperhidrosis Therapy

We read with interest the article by Karamfilov et al<sup>1</sup> suggesting lower relapse rates of hyperhidrosis after high-dose botulinum toxin type A injections (BOTOX; Allergan Inc, Irvine, Calif, hereinafter, generically, botulinum toxin A).<sup>1</sup> Various protocols for treating hyperhidrosis with botulinum toxin A have been issued empirically without controlled comparison of doses, dilutions, number of injections, or pharmaceutical products. Thus, any attempt to provide evidence-based information on how to optimize botulinum toxin A treatment should be welcomed. For this purpose, however, stringent study designs, accurate measurements of sweating, and uniform follow-up schedules are indispensable. Unfortunately, Karamfilov et al<sup>1</sup> did not implement a control group receiving low-dose botulinum toxin A, which could have been easily provided by a left-vs-right comparison, with each patient being his own control.

Also, the iodine-starch test and planometry, which are helpful to visualize the active hyperhidrotic area, are not pertinent for exact quantification of sweating. In fact, positive findings on the iodine-starch test easily occur in any healthy individual. It is, however, the rate of sweating (amount per minute) that makes a person hyperhidrotic, and this can accurately be determined by gravimetry using blotting paper, a high precision scale, and a stopwatch.<sup>2</sup> In 156 patients recently screened for severe axillary hyperhidrosis, the mean ± SD active area as visualized by the iodine-starch test was 48.5 ± 4.4 cm<sup>2</sup>, ranging from 14.2 to 66.5 cm<sup>2</sup>, which showed no correlation

to actual sweat rates measured by gravimetry (52-858 mg/min). The fact that gravimetric values may vary considerably does not discredit this method but rather demonstrates the dynamics of eccrine glands in hyperhidrotics. After botulinum toxin A treatment, gravimetric sweat rates have been shown to be consistently low.<sup>2,3</sup>

Follow-up as reported by Karamfilov et al<sup>1</sup> ranged from 5 to 15 months, but it remained unclear at what intervals patients were observed—if there was any regular follow-up schedule at all. Waiting for the patient to ask for subsequent treatment is definitely too volatile a parameter for a clinical study, especially when trying to establish measurable benefits compared with already existing protocols. Finally, but not of least importance, the safety of high-dose botulinum toxin A as proclaimed by Karamfilov et al<sup>1</sup> is questionable; the authors failed to mention that the risk of antibody induction rises not only with treatment frequency but also particularly with higher doses.<sup>4</sup>

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### The Wheal: To Be or Not to Be

Sad to say, until a dictionary of dermatology equivalent to the Oxford English Dictionary comes into being, dermatology will forever be a twiglike imposter, rather than an authentic branch of knowledge.

A. Bernard Ackerman, MD

It is astonishing that the current significantly different meanings of the term *cutaneous elementary lesions* have received so little attention in dermatology journals. But it is precisely for this reason that the renewed controversy regarding basic dermatological lesions reflected in the ARCHIVES holds great interest.<sup>1-4</sup> The suggested elimination of *wheal* “from the list of basic terms”<sup>4</sup> is worthy of comment. In our opinion, there are also reasons favoring its preservation on such a list. Dermatologists have traditionally used a specific term to describe lesions of urticaria except in French dermatology. The analysis of such a tradition may contribute to a better understanding of this controversy.<sup>5</sup>

English Tradition. *Wheal* is an Anglo-Saxon word. Robert Willan and Thomas Bateman turned this into a spe-