

Patient satisfaction was greater in the botulinum toxin arm, with about 29% and 50% reporting that they had excellent and good satisfaction, respectively, compared with 0% and 7% in the placebo arm (Table 1). There were no reports of injection-related complications in either study arm.

In conclusion, Dr Ahmad stated that, in his opinion, the data from this study indicate that treatment of plantar fasciitis with botulinum toxin resulted in greater functional scores and patient satisfaction, as well as lower pain scores, when compared with placebo. However, the study was limited by a small sample size, potential differences in physical therapy regimens, and potential use of patient-directed treatments.

c-hAM Comparable to Corticosteroids in Plantar Fasciitis

Written by Emma Hitt Nichols, PhD

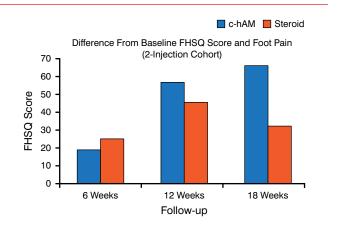
In the treatment of plantar fasciitis, micronized amniotic membrane (c-hAM) injection demonstrated similar efficacy to corticosteroid injection without adverse events. Robert D. Santrock, MD, West Virginia University, Morgantown, West Virginia, USA, presented data from a randomized, controlled, double-blinded, single-center, prospective study comparing plantar fasciitis injection of c-hAM to steroid injection [Hanselman AE et al. *Foot Ankle Int.* 2014].

Fetal tissues, including the amnion, are currently used in ophthalmology, orthopedics, and other surgical specialties. Use of fetal tissues in surgery is appealing because the tissue is able to regenerate without inflammation or scarring. The study hypothesis was that c-hAM is a safe treatment option for plantar fasciitis and is non-inferior to corticosteroids.

In this study, 23 patients diagnosed with plantar fasciitis and symptomatic for > 3 months but < 1 year were randomly assigned to receive an initial injection of either 1 mL (40 mg) of corticosteroid plus 4 mL 0.5% bupivacaine, or 1 mL c-hAM plus 4 mL 0.5% bupivacaine. At 6-week follow-up, all patients were given the option to receive a repeat injection of either study drug if needed. Patients were followed for 12 weeks after the most recent injection. Activities were not restricted, but all patients were advised to perform foot and ankle-stretching exercises 5 times a day.

Exclusion criteria included previous plantar fasciitis injection or treatment within 3 months, previous foot surgery or injury, lower extremity neuropathy, lack of ambulation, or unwillingness to receive human tissue. The primary end point was the Foot Health Status Questionnaire

Figure 1. Effect of 2 Injections of c-hAM on FHSQ Score



 $c-hAM, micronized\ amniotic\ membrane; FHSQ, Foot\ Health\ Status\ Questionnaire.$ Reproduced with permission from R Santrock, MD.

(FHSQ). Secondary end points included the visual analog scale (VAS) and patient-reported improvement.

There was no significant difference between the 2 arms in terms of foot pain, foot function, and VAS. However, in the 1-injection cohort, patients who received the corticosteroid demonstrated greater FHSQ foot pain scores at 6 and 12 weeks compared with patients who received c-hAM. Interestingly, in the 2-injection cohort, patients who received c-hAM demonstrated a trend toward a greater FHSQ foot pain score compared with the corticosteroid arm (Figure 1). A similar trend was observed in FHSQ foot function, in which 2 injections of c-hAM demonstrated increased effectiveness compared with 1 injection. After 2 injections, the c-hAM arm trended toward a greater improvement in VAS and patient-reported improvements at 18 weeks compared with the corticosteroid arm. There were no adverse events in this study.

According to Dr Santrock, the data from this study suggest that c-hAM has comparable efficacy to corticosteroids in the treatment of plantar fasciitis, and there may be a double-dose effect associated with c-hAM. He indicated that that a larger, longer-term, multicenter trial is needed to further evaluate c-hAM for plantar fasciitis.

Differences in Component Revision and Reoperation for Ankle Arthroscopy Implants

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A study of 4 implants used for total ankle arthroscopy (TAA) found that the Agility and Mobility implants required higher rates of metal component revisions