



# Severe subacromial-subdeltoid inflammation with rice bodies associated with implantation of a bio-inductive collagen scaffold after rotator cuff repair



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## Case report

A 48-year-old man underwent rotator cuff repair augmented with a Regeneten bio-inductive implant (Smith & Nephew, Andover, MA, USA). At 4 months after the procedure, an acute, massive swelling developed in the subacromial space associated with intense pain. The patient had no chills or fever, and the complete blood count showed a normal white blood cell (WBC) count. The sedimentation rate was normal, at 19 mm/h, but the C-reactive protein level was abnormal, at 24.9 mg/L (normal level < 8 mg/L). The finding of the rheumatoid factor test was negative. Aspiration of the bursa revealed bloody, turbid fluid that was submitted for cytologic analysis and Gram staining. The cell count included a markedly elevated WBC count (14,449 WBCs/ $\mu$ L, 54% neutrophils). Magnetic resonance imaging (MRI) showed a healed rotator cuff tendon with significant swelling with particulate debris in the subacromial-subdeltoid bursa (Fig. 1 and Fig. 2). The particulate debris resembled rice bodies.

Because infection was suggested, the patient underwent surgery to perform washout, débridement, and placement of tubes for drainage. The material retrieved during the

surgical procedure looked purulent, containing multiple dense debris (Fig. 3). The patient received intravenous antibiotics while final cultures were awaited. Both the initial aspirate obtained in the office and the second aspirate obtained at the time of surgery were held for 14 days. Both failed to grow any organisms. With negative culture findings, the antibiotics were stopped and the patient began physical therapy. At 4 weeks, the swelling was completely gone. The C-reactive protein level at 6 weeks' follow-up had normalized. At 3 months' follow-up (7 months after rotator cuff repair), the pain had nearly completely disappeared and the patient had returned to work.

## Discussion

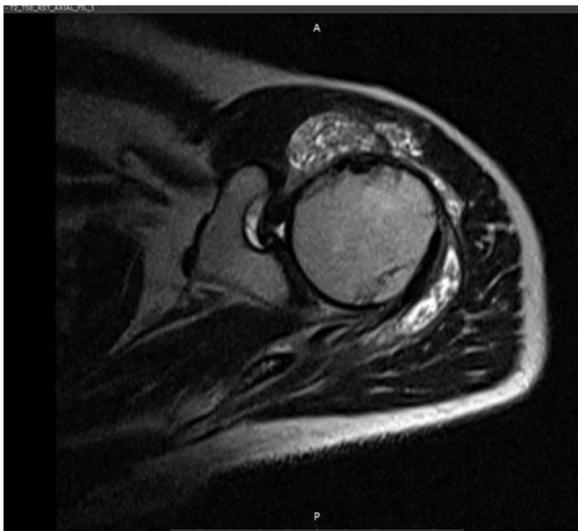
The Regeneten implant has 3 essential components. The implant itself is highly purified collagen made of bovine tendon. It is made of collagen fibers processed to create a highly oriented and highly porous collagen scaffold cleansed of animal DNA.<sup>1,15</sup> By incorporating onto the host's rotator cuff tendon, thickness is added and strain is reduced.<sup>3</sup> Healing of the tendon is therefore thought to be facilitated. The construct has 2 other parts. Staples made of polyether ether ketone are used laterally on the patch to hold it in place along the greater tuberosity. Polyether ether ketone does not dissolve. However, allergic responses to

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**Figure 1** Coronal image in patient with rice body formation with bio-inductive collagen.



**Figure 2** Axial image in patient with rice body formation with bio-inductive collagen.

implants made of this material have been reported in the literature.<sup>9,12</sup>

The medial side of the implant is held in place with several staples made of polylactic acid, a combination of both the L- and D-isomers of lactic acid. At 4 to 6 months, the staples begin to dissolve by a process of hydrolysis. The D-isomer delays the reaction as a material made only of the L form of lactic acid would dissolve more quickly. The process is usually complete by 1 year (C. Van Kampen, personal communication). Regardless of the time to degradation, the process of degradation is associated with a foreign body response that can evoke an inflammatory reaction.<sup>5</sup> Reactions to implants and suture material made of similar substances have been researched and well reported in the literature.<sup>7,10,12,13,19</sup> In a case report on a rotator cuff repair by Sivaloganathan et al,<sup>16</sup> a suture anchor made of poly-L-lactic acid appeared to cause



**Figure 3** Fluid from subacromial bursa in patient with rice body formation with bio-inductive collagen.

severe inflammatory bursitis with rice body formation. They termed the MRI appearance the “risotto sign.” Rice body formation may result from de novo formation within the synovial fluid.<sup>11</sup> Other authors have suggested that formation occurs as a result of microinfarction of the synovium and shedding, followed by fibrin encasement.<sup>4</sup> Formation of multiple rice bodies with massive subacromial-subdeltoid bursitis is an associated MRI finding in rheumatoid disease, chondrocalcinosis, seronegative arthritis, and septic conditions.<sup>2,8,17,18</sup> Rice body formation simply occurring as a result of chronic bursitis absent all the aforementioned conditions has been reported.<sup>6,14</sup> All these conditions seem to have inflammation as a common theme. In preliminary research regarding use of the Regeneten bio-inductive implant, acute bursal swelling did occur in 1 of 13 patients.<sup>1</sup> This patient was treated with an arthroscopic washout. No rice bodies were found, and the rotator cuff had healed. In a multicenter trial involving use of this biological scaffold to treat partial cuff tears, “significant bursal swelling” did develop in 1 of 33 patients at 3 months after implantation.<sup>15</sup> No rice bodies were reported in the MRI findings.

In this case report, multiple rice body-like particulate debris was noted on MRI as well as within the fluid retrieved at the time of washout. A T2 image showed iso-intense nodules consistent with rice body formation. The differential diagnosis of chondromatosis would have had a different MRI appearance.<sup>6</sup> Eosinophils were absent in the first aspirated fluid, which probably ruled out some type of allergic phenomena. Infection was ruled out by negative culture findings.

## Conclusion

Augmentation of rotator cuff injuries with the Regeneten bio-inductive implant has been shown to facilitate healing<sup>1,15</sup> and can be very useful in the surgeon’s treatment of rotator cuff pathology. However, the

potential for an acute and very painful inflammatory phase exists during the course of recovery. Although the exact cause of this severe inflammation remains unknown, the timing might suggest some type of exaggerated host response during the degradation of the polylactic acid staples. The potential for an acute, significant painful phase in a patient's recovery should be part of the informed-consent process when using the Regeneten implant.

## Disclaimer

The author, his immediate family, and any research foundations with which he is affiliated have not received any financial payments or other benefits from any commercial entity related to the subject of this article.

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