Modular Neck Stems

Information About the Voluntary Product Recall

Stryker Orthopaedics voluntarily recalled its Rejuvenate Modular and ABG II modular-neck hip stems.

This voluntary recall was initiated due to the potential risks associated with modular-neck stems. These risks include the potential for fretting and/or corrosion at or about the modular-neck junction, which may result in adverse local tissue reactions manifesting with pain and/or swelling.

Please see the FAQs below for more information.

Your Particular Implant

Your surgeon can tell you if you received a Rejuvenate Modular or ABG II modular-neck hip stem, or you may review your medical records for implant identification information. If you have further questions, please contact the call center at 1-888-317-0200.

If you have symptoms of pain and/or swelling in or around your replaced hip, you should schedule an office visit with your surgeon and discuss your symptoms. If you have no symptoms, you should continue to follow the post-operative plan that your surgeon has outlined for you.