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**stryker**

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**Orthopaedics**

Dear Patient,

You are receiving this letter because your surgeon has identified that you are a recipient of a Rejuvenate or ABG II modular-neck hip stem. Stryker Orthopaedics, the manufacturer, initiated a voluntary recall of these 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 ALTR (adverse local tissue reactions) manifesting with pain and/or swelling.

If you have no symptoms, you should continue to follow the post-operative plan that your surgeon has outlined for you. However, 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.

We understand that this information may raise questions. To help address your questions, Stryker has the following resources available:

- **Patient Call Center - 1-888-317-0200**
- **Web Resource - [www.AboutStryker.com/ModularNeckStems](http://www.AboutStryker.com/ModularNeckStems)**