Holy Cross
Hospital

4725 North Federal Highway
Fort Lauderdale, FL 33308
www.holy-cross.com
954-771-8000

April 13, 2017

[redacted]

RE: Item [redacted]
Lot [redacted]

Dear [redacted]

You are receiving this letter because our records indicate that you have received a medical device (femoral head) implant from Stryker on [redacted] by Dr. [redacted]

We would like to inform you that the manufacturer, Stryker, has issued a voluntary recall notification regarding this product.

The reason for the recall is that the manufacturer has received a higher than expected complaints for certain sizes of LFIT™ Anatomic CoCr V40TM Femoral Heads manufactured prior to 2011. Several potential hazards have been identified that could result in harm such as: loss of mobility, pain requiring revision, dislocation, joint instability, and periprosthetic fracture among others.

Their recommendation is that the Orthopedic Surgeon would continue to follow you at our normal scheduled appointments.

Stryker has established a dedicated call center at 1-888-644-2548 for any patient questions that you may have and/or if you are having any concerns with your implant or have any questions that we may answer for you; please do not hesitate to contact my office.

Sincerely,

From the Office of:
[redacted]