UNITED STATES DISTRICT COURT
DISTRICT OF MINNESOTA

IN RE: STRYKER REJUVENATE AND ABG II HIP IMPLANT PRODUCTS LIABILITY LITIGATION	MDL No. 13-2441 (DWF/DJF)
This Document Relates to: Robert T. Griffin, Plaintiff, v. Civil No. 15-2412 (DWF/DJF) Howmedica Osteonics, *d/b/a* Stryker Orthopaedics, Stryker Corp., Stryker Sales Corporation, and Stryker Ireland Limited, Defendants.	**ORDER**

Pending before the Court is the above captioned *Robert T. Griffin* matter (Civil No. 15-2412 (DWF/DJF); MDL No. 13-2441 (DWF/DJF)). By way of prior order, counsel for Plaintiff Robert T. Griffin was permitted to withdraw. As part of that same order (MDL No. 13-2441, (Doc. No. 1752); Civil No. 15-2412, (Doc. No. 22)), product identification was required to be served on lead MDL counsel for Plaintiffs and Defendants to establish that Plaintiff has one of the subject MDL Affected Products. The Court has now been informed that the product identification documentation confirms that Plaintiff does *not* have one of the Affected Products at issue in this MDL litigation.

Based upon the foregoing and the record in this case, **IT IS HEREBY ORDERED** that:

1. The action is **DISMISSED WITH PREJUDICE** and without fees or costs to any party.

Dated: April 16, 2025

s/Donovan W. Frank
DONOVAN W. FRANK
United States District Judge