IDE Final Report

IDE Gxxxxx

IDE Title (if title being used)

Gxxxx/R00X

Name of Sponsor-Investigator, MD

X Professor, Department

Institution

Date of Submission

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# General Information

*Please state your:*

1. *IDE number*
2. *Device name and indication(s) for use*
3. *Sponsor’s name address, phone numbers, and fax*
4. *Sponsor’s email address*
5. *Contact person*

# Study Progress

*(Data from the beginning of the study should be reported, unless otherwise indicated)*

## Brief Summary of study progress in relation to the investigational plan

## Number of Investigators/Investigational Sites

*Include a list of investigators.*

## Number of Subjects Enrolled

## Number of Devices Shipped

## Disposition of All Devices Shipped

## Brief Summary of the Results

## Summary of Anticipated and Unanticipated Adverse Effects

## Deviations from the Investigational Plan

*Please, describe all the deviations from the investigational plan since the last progress report.*

# Risk Analysis

*A thorough risk analysis and risk mitigation strategies are critical for the FDA’s decision to allow a study to continue. Update the risk analysis from your initial application with any relevant changes. Include a summary of any new adverse information (since the last progress report) that may affect the risk analysis. This includes preclinical data, animal studies, foreign data, clinical studies, etc. For more details on what to include in the risk analysis, please see the DTMI original IDE template.*

*Also, please attach the reprints of any articles published from data collection from this study.*

## Summary of any new adverse information

*Summary of any new adverse information (since last progress report) that may affect the risk analysis; this includes preclinical data, animal studies, foreign data, clinical studies, etc.*

## Reprints of any Articles

*Also, please attach the reprints of any articles published from data collection from this study.*

# Other Changes

## Summary of any changes in Manufacturing

*Summary of any changes in the manufacturing process and quality control, including changes that have not been submitted as a supplemental application.*

## Summary of all changes not required to be submitted in a prior supplement

*Summary of all changes in the investigational plan that were not required to be submitted in a supplemental application.*

# marketing application or Future Plans

## Progress towards approval/clearance

*Progress towards product approval/clearance, with date (or projected date) of PMA or 510(k) submission; or indication that marketing of device is not planned*

## Plans to submit another IDE

*Any plans to submit another IDE application for this device or a modification of this device.*

*References*

*IDE Reports*

* *Suggested Format for IDE Progress Report*
* *https://www.fda.gov/medical-devices/investigational-device-exemption-ide/ide-reports*
* [*https://www.fda.gov/medical-devices/investigational-device-exemption-ide/ide-reports#sugforforidepro*](https://www.fda.gov/medical-devices/investigational-device-exemption-ide/ide-reports#sugforforidepro)