VIASYS HEALTHCARE
MEDSYSTEMS

CORFLO Enteral Access Devices
PULMANEX Airway Management Devices
STACKHOUSE Surgical Systems

August 4, 2008

FCC Authorization & Evaluation Division
7435 Oakland Mills Road
Columbia, Maryland 21046

Subject: Revision Declaration

To Whom It May Concern,

We hereby declare under our sole responsibility that no modifications have been made to the product under evaluation, Viasys Navigator BioNavigation Device, Model Number Navigator (1.5), such that the transmit frequency and/or output power have been affected. The model number for the product has changed from **Navigator (1.5)** to **60-3020**. Model number **Navigator (1.5)** was used an initial pre-production prototype designator. Model number **60-3020** was assigned as the final reorder code / model for the product. These model numbers are references to the same product, which was tested at DLS Electronic Systems, Inc. in August of 2007.

Submission of these documents and test results were delayed approximately eleven months following the reporting of all data as a result of the following business driven considerations:

(1) It was a project management decision that all performance and environmental validation testing shall be completed and approved internally prior to certification submissions.
(2) Prioritization of engineering resources and budgetary dollars were shifted throughout Fiscal Year 2008.

Please contact Shawn Purnell with any questions you may have.

Dated this 6TH day of AUGUST, 2008.

By: _____ SHAWN PURNELL
(Signature) (Print name)

Title: SENIOR ENGINEER - PRODUCT DEVELOPMENT

On behalf of: VIASYS MEDSYSTEMS
(Company Name)

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