

Philips Medical Systems

June 12, 2007

Federal Communications Commission Authorization and Evaluation Division

FCC ID: PQC-WMTS-MODULE:

Confidentiality Request regarding application for Class II Permissive Change certification

Pursuant to Sections 0.457 and 0.459 of the Commission's Rules, we hereby request confidential treatment of information accompanying this application as outlined below:

Exhibit TypeFile NameBlock DiagramWMTS w D & Triband BlkDia.pdfSchematicsWMTS w D & Triband Schem.pdfOperational DescriptionWMTS w D & Triband OpDes.pdfParts ListWMTS w D & Triband PartsLst.pdfTuning ProcedureWMTS w D & Triband TunPro

The above materials contain trade secrets and proprietary information not customarily released to the public. The public disclosure of these materials may be harmful to the applicant and provide unjustified benefits to its competitors.

The applicant understands that pursuant to Section 0.457 of the Rules, disclosure of this application and all accompanying documentation will not be made before the date of the Grant for this application.

Pursuant to DA04-1705 June 15, 2004 of the Commission's public notice, we also request temporary confidential treatment of information accompanying this application as outlined below:

Exhibit TypeFile NameExternal PhotosWMTS w D & Triband ExtPho.pdfInternal PhotosWMTS w Dband IntPho.pdfInternal PhotosWMTS w Triband IntPho.pdf

Temporary confidentiality from public disclosure is important for Philips Medical Systems from a commercial perspective. It enables the company to complete its development and regulatory efforts prior to introducing the product to the marketplace. Releasing information on the product via the FCC website prior to formal market introduction can be confusing to our customers and to the medical community, and places the company at a competitive disadvantage.

Sincerely.

Barry Wyshogrool
Regulatory Engineer

