

March 15, 2007

Federal Communications commission  
Attn: Ms. Linda Elliott  
7435 Oakland Mills Rd.  
Columbia, MD 21036

Re: Response to FCC for Grant Certification under FCC ID RIASJMRF

Dear Ms. Elliott:

In response to the following request from March 15, 2007, I offer the following:

FCC Question:

“The FCC ID is not shown on the ID label. Submit a sketch or photograph showing the FCC ID on the ID label, as appropriate”.

SJM Response:

The FCC ID is included on the ID label submitted on March 6<sup>th</sup> as shown on the preceding page. The ID has been highlighted for easy reference.

If there are any questions, please feel free to contact me at 818-493-2629.

Sincerely,



Deanna Hughes  
Regulatory Affairs  
St. Jude Medical

RF Implantable Medical Device - Label

DRAFT

<p>                    SN SAMPLE                  6000XXXX-XXXX                       SN SAMPLE                  6000XXXX-XXXX             </p>	<p> <b>CURRENT™ DR RF 2207-30</b>                  Tiered-therapy cardioverter/defibrillator             </p>	<p>   <b>ST. JUDE MEDICAL</b>                  California   New York   London   Paris   Mexico City   Seoul   Singapore   Sydney   Taipei   Toronto             </p>	<p>     <b>VVED DDR</b> </p>
<p>                    SN SAMPLE                  6000XXXX-XXXX             </p>	<p> <b>CURRENT™ DR RF 2207-30</b>                  Tiered-therapy cardioverter/defibrillator             </p>	<p> <b>2207-30</b>      Use Before: <b>20xx/xx/xx</b> </p> <p> <b>SHIPPED SETTINGS</b>                  The pulse generator is shipped with all functions off.             </p> <p> <b>CONTENTS</b>                  One pulse generator, accessories and product literature.             </p> <p> <b>STERILE EO</b>                  Contents of this package have been ethylene oxide sterilized. Sterility cannot be guaranteed if sterile package is damaged or opened. Refer to the accompanying manual.             </p> <p> <b>STORAGE CONDITIONS</b>                  Store at 10°C (50°F) to 45°C (113°F).             </p> <p> <b>CAUTION</b>                  Rx only             </p> <p>                 Because of the numerous possible configurations, lead/pulse generator and/or lead/defibrillator, compatibility should be confirmed with the pulse generator/defibrillator and/or lead manufacturer prior to implantation of the pacing/defibrillation system.             </p> <p> <b>Proposition 65:</b> A State of California water initiative, requires the following notice:  <b>WARNING:</b> This product and its packaging have been sterilized with ethylene oxide. This packaging may expose you to ethylene oxide, a chemical known to the state of California to cause cancer or birth defects or other reproductive harm.             </p>	<p>                 Serial number:  <b>SN SAMPLE</b>                  Manufacturing date:  <b>20xx/xx/xx</b> </p> <p> <b>6000XXXX-XXXX</b> </p> <p> <b>FCC ID: RIAS/JMRF</b> </p> <p>  Single Use Only             </p>
<p>                 6000yyyy-yyy             </p>	<p> <b>CURRENT™ DR RF 2207-30</b>                  Tiered-therapy cardioverter/defibrillator             </p>	<p>   <b>ST. JUDE MEDICAL</b> </p> <p>                 Use Before: <b>20xx/xx/xx</b> </p> <p>                 Serial number:  <b>SN SAMPLE</b>                  6000XXXX-XXXX             </p>	<p>                   XXXXXXXXXXXXXXXXXXXXXXXX                    XXXXXXXXXXXXXXXXXXXXXXXX             </p>

FCC ID