



Cardiac Rhythm Management

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December 19, 2007

Federal Communications Commission
Office of Engineering and Technology
445 12TH ST SW
Washington DC 20554

VIA FEDERAL EXPRESS AND FACSIMILE

Re: Waiver Order DA 07-3160 and notice of submission for FDA approval

Federal Communications Commission:

Pursuant to the waiver Order (DA 07-3160) released on July 11, 2007, Boston Scientific is required to notify the FCC of the submission of the Cognis and Teligen products for Food and Drug Administration (FDA) approval within 10 business days of that event.

The Cognis and Teligen FDA submission was sent on December 6, 2007 and was received by the FDA on December 7, 2007.¹

Please contact me with any questions.

Sincerely,

Kurt Wheeler
Manager²
Engineering Compliance
Boston Scientific CRM
(651) 582 5051

cc: Julius Knapp, Chief, Office of Engineering and Technology
FCC Office of Engineering and Technology Laboratory Division

¹ Under the ordering clause in paragraph 14 of the waiver Order, the three year term of the waiver shall commence on FDA approval because the FDA submission was made prior to March 31, 2008. Pursuant to paragraph 15 of the waiver Order, Boston Scientific will notify the FCC of the date of FDA approval as part of its submission for equipment certification of the Cognis and Teligen devices.

² I am now the Manager with oversight of FCC regulatory compliance for CRM devices.