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stryker®
Medical

Attention: Application Examiner
Re: Request for Class II Permissive Change
Applicant: Stryker medical
FCC ID: Z7A-6506 and Z7A 6516

To whom it may concern,

This request is submitted by Stryker Medical for the purpose of seeking approval for a Class II Permissive Change for the Power PRO XT (6506) and the Power PRO IT (6516). Both products are currently authorized under FCC ID Z7A-6506 and Z7A-6516 respectively. This is pursuant to 47 CFR 2.1043 for changes to an RF device. The permissive change rules in Section 2.1043 describe the modifications that may be made to an RF device without filing for a new equipment authorization.

The wireless communication device used on both products has not changed form or function. The original PCB configuration, as well as the original schematic still applies for the wireless communication device. The modification only pertains to increase in distance for wireless transmission per our test specifications and filtering the cot main board to improve EMC emissions.

We have recently tested our 6506 product and determined that a Class II Permissive Change is required due to an increase in radiated spurious emissions within the 840MHz to 850MHz frequency range. The test results have been provided for consideration. The increase in radiated emissions is still within the allowable limits for compliance and the maximum output power rating has not changed from the original authorizationd. Therefore, we believe it is acceptable to request a Class II Permissive Change for both products.

Sincerely,

Carmen Davis

Carmen Davis
Sr. Approvals Engineer
Stryker Medical
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Portage, Mi. 49002