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### 510(k) Premarket Notification Database

<b>Device Classification Name</b>	<a href="#">Stimulator, Nerve, Transcutaneous, For Pain Relief</a>
<b>510(K) Number</b>	K062641
<b>Regulation Number</b>	<a href="#">882.5890</a>
<b>Device Name</b>	BODY-STIM, BIOMODULATOR, BEST-RSI, BEST PRO, MODE
<b>Applicant</b>	AVAZZIA, INC. 13154 Coit Rd., Ste. 200 Dallas, TX 75240
<b>Contact</b>	Catherine Tone
<b>Classification Product Code</b>	<a href="#">GZJ</a>
<b>Date Received</b>	09/06/2006
<b>Decision Date</b>	04/30/2007
<b>Decision</b>	Substantially Equivalent (SE)
<b>Classification Advisory Committee</b>	Neurology
<b>Review Advisory Committee</b>	Physical Medicine
<b>Statement/Summary/Purged Status</b>	Summary Only
<b>Summary</b>	<a href="#">Summary</a>
<b>Type</b>	Traditional
<b>Reviewed By Third Party</b>	No
<b>Expedited Review</b>	No

Database Updated 5/08/2007

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