

**K110566 VIBERECT PENILE VIBRATORY STIMULATION  
DEVICE**Jun 24, 2011  
116 days to decisionK110566 · Product code: **KXQ** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k110566/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Vibrator For Therapeutic Use, Genital (KXQ)
Date received	Feb 28, 2011
Decision date	Jun 24, 2011
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Reflexonic, LLC</b>
Location	Austin, TX, US
Contact	CAROLINE TONTINI
510(k) history	2 submissions · 2 cleared · 2011-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110566/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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