

K142304 Viberect Penile Vibratory Stimulation DeviceOct 3, 2014
46 days to decisionK142304 · Product code: **KXQ** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k142304/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Vibrator For Therapeutic Use, Genital (KXQ)
Date received	Aug 18, 2014
Decision date	Oct 3, 2014
Days to decision	46 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Reflexonic, LLC
Location	Austin, TX, US
Contact	Kambiz Tajkarimi
510(k) history	2 submissions · 2 cleared · 2011-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k142304/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 6, 2026