

K142248 Prostiva RF Therapy Generator, Prostiva RF Therapy Hand Piece, Prostiva RF Therapy Return Electrode, Prostiva RF Therapy Tubing System, Prostiva RF Therapy TelescopeFeb 20, 2015
190 days to decisionK142248 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k142248/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 14, 2014
Decision date	Feb 20, 2015
Days to decision	190 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Urologix, Inc.
Location	Plymouth, MN, US
Contact	Georgiann Keyport
510(k) history	1 submissions · 1 cleared · 2015-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k142248/> 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 5, 2026