FDA 510(k) Premarket Notification

K070871 STARION UNIVERSAL POWER SUPPLY

CLEARED

Apr 13, 2007
15 days to decision

K070871 · Product code: **HQO** · Ophthalmic
Source: https://www.510kdatabase.net/k070871/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Unit, Cautery, Thermal, Ac-powered (HQO)
Date received	Mar 29, 2007
Decision date	Apr 13, 2007
Days to decision	15 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Starion Instruments**
Location	Saratoga, CA, US
Contact	BRIAN GRIGSBY
510(k) history	13 submissions · 13 cleared · 1999-2007

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k070871/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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