FDA 510(k) Premarket Notification

K000639 VISIONARY 2000

CLEARED

May 25, 2000
90 days to decision

K000639 · Product code: **FSY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k000639/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Surgical, Ceiling Mounted (FSY)
Date received	Feb 25, 2000
Decision date	May 25, 2000
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Burton Medical Products Corp.**
Location	Van Nuys, CA, US
Contact	DORIAN SWARTZ
510(k) history	15 submissions · 15 cleared · 1985-2010