FDA 510(k) Premarket Notification

K963682 GENIE' SINGLE CEILING AND DOUBLE CEILING

CLEARED

Nov 13, 1996
71 days to decision

K963682 · Product code: **FSY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k963682/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Surgical, Ceiling Mounted (FSY)
Date received	Sep 3, 1996
Decision date	Nov 13, 1996
Days to decision	71 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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