

K963682 GENIE'S; SINGLE CEILING AND DOUBLE CEILINGNov 13, 1996
71 days to decisionK963682 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k963682/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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