FDA 510(k) Premarket Notification

K901108 FIBERLUME

CLEARED

Mar 28, 1990
19 days to decision

K901108 · Product code: **HBI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k901108/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Illuminator, Fiberoptic, Surgical Field (HBI)
Date received	Mar 9, 1990
Decision date	Mar 28, 1990
Days to decision	19 days
Third-party review	No

APPLICANT

Company	**Burton**
Location	Van Nuys, CA, US
Contact	GRETEL ANDERSON
Website	http://www.burtonalbionfc.co.uk/
510(k) history	1 submissions · 1 cleared · 1990-1990