FDA 510(k) Premarket Notification

K896384 FIBEROPTIC LIGHT SOURCE, FOI-1A

CLEARED

Jan 23, 1990
78 days to decision

K896384 · Product code: **EQH** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k896384/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Source, Carrier, Fiberoptic Light (EQH)
Date received	Nov 6, 1989
Decision date	Jan 23, 1990
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Jedmed Instrument Co.**
Location	Mchenry, IL, US
Contact	CRAIG RAPP
510(k) history	92 submissions · 91 cleared · 1979-2014