FDA 510(k) Premarket Notification

K823536 INTRALIG

CLEARED

Dec 28, 1982
28 days to decision

K823536 · Product code: **EJI** · Dental
Source: https://www.510kdatabase.net/k823536/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Cartridge (EJI)
Date received	Nov 30, 1982
Decision date	Dec 28, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Miltex Instrument Co.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1992