FDA 510(k) Premarket Notification

K812692 FILES & REAMERS

CLEARED

Oct 23, 1981
31 days to decision

K812692 · Product code: **EKS** · Dental
Source: https://www.510kdatabase.net/k812692/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	File, Pulp Canal, Endodontic (EKS)
Date received	Sep 22, 1981
Decision date	Oct 23, 1981
Days to decision	31 days
Third-party review	No

APPLICANT

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