FDA 510(k) Premarket Notification

K832378 VIBRATORY ENDODONTIC SYSTEM

CLEARED

Oct 4, 1983
77 days to decision

K832378 · Product code: **EKQ** · Dental
Source: https://www.510kdatabase.net/k832378/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Preparer, Root Canal Endodontic (EKQ)
Date received	Jul 19, 1983
Decision date	Oct 4, 1983
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Syntex Dental Products, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1983