

K832378 VIBRATORY ENDODONTIC SYSTEMOct 4, 1983
77 days to decisionK832378 · 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

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Device record: <https://www.510kdatabase.net/k832378/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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