

K812692 FILES & REAMERSOct 23, 1981
31 days to decisionK812692 · 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

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Device record: <https://www.510kdatabase.net/k812692/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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