

K822568 EAR PROBESSep 14, 1982
21 days to decisionK822568 · Product code: **EKL** · DentalSource: <https://www.510kdatabase.net/k822568/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Point, Silver, Endodontic (EKL)
Date received	Aug 24, 1982
Decision date	Sep 14, 1982
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Kelleher Corp.
Location	Mchenry, IL, US
510(k) history	94 submissions · 94 cleared · 1982-1983

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Device record: <https://www.510kdatabase.net/k822568/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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