

K820639 OBTURATION SYSTEM KITApr 9, 1982
32 days to decisionK820639 · Product code: **EKM** · Dental
Source: <https://www.510kdatabase.net/k820639/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gutta-percha (EKM)
Date received	Mar 8, 1982
Decision date	Apr 9, 1982
Days to decision	32 days
Third-party review	No

APPLICANT

Company	U.S. Shizai Corp.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1986

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Device record: <https://www.510kdatabase.net/k820639/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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