

K811582 THE PLACERJul 28, 1981
53 days to decisionK811582 · Product code: **EID** · DentalSource: <https://www.510kdatabase.net/k811582/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Restorative And Impression Material (EID)
Date received	Jun 5, 1981
Decision date	Jul 28, 1981
Days to decision	53 days
Third-party review	No

APPLICANT

Company	Cavitron Corp.
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1976-1981

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Device record: <https://www.510kdatabase.net/k811582/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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