

K820738 LUSTRALLOYApr 9, 1982
22 days to decisionK820738 · Product code: **EJJ** · DentalSource: <https://www.510kdatabase.net/k820738/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alloy, Amalgam (EJJ)
Date received	Mar 18, 1982
Decision date	Apr 9, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Hoyt Laboratories
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1977-1982

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Device record: <https://www.510kdatabase.net/k820738/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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