FDA 510(k) Premarket Notification

K820738 LUSTRALLOY

CLEARED

Apr 9, 1982
22 days to decision

K820738 · Product code: **EJJ** · Dental
Source: 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