FDA 510(k) Premarket Notification

K820181 SYBRON/KERR CONTOUR

CLEARED

Feb 24, 1982
33 days to decision

K820181 · Product code: **EJJ** · Dental
Source: https://www.510kdatabase.net/k820181/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Alloy, Amalgam (EJJ)
Date received	Jan 22, 1982
Decision date	Feb 24, 1982
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Sybron Corp.**
Location	Mchenry, IL, US
510(k) history	37 submissions · 37 cleared · 1977-1986