

K820181 SYBRON/KERR CONTOURFeb 24, 1982
33 days to decisionK820181 · Product code: **EJJ** · DentalSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820181/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026