

K881274 CENTER BEND ARCH WIRESApr 12, 1988
18 days to decisionK881274 · Product code: **EJF** · DentalSource: <https://www.510kdatabase.net/k881274/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bracket, Metal, Orthodontic (EJF)
Date received	Mar 25, 1988
Decision date	Apr 12, 1988
Days to decision	18 days
Third-party review	No

APPLICANT

Company	Flexmedics
Location	Minneapolis, MN, US
Contact	SIEVERT, JR.
510(k) history	20 submissions · 20 cleared · 1986-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881274/>; 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 22, 2026