FDA 510(k) Premarket Notification

K881274 CENTER BEND ARCH WIRES

CLEARED

Apr 12, 1988
18 days to decision

K881274 · Product code: **EJF** · Dental
Source: 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