FDA 510(k) Premarket Notification

K800312 150 DRIVER/REAMER

CLEARED

Feb 22, 1980
10 days to decision

K800312 · Product code: **KIJ** · Orthopedic
Source: https://www.510kdatabase.net/k800312/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Orthopedic, Dc-powered Motor And Accessory/attachment (KIJ)
Date received	Feb 12, 1980
Decision date	Feb 22, 1980
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Black & Decker(Tm)**
Location	Mchenry, IL, US
510(k) history	11 submissions · 11 cleared · 1980-1982