FDA 510(k) Premarket Notification

K791960 ROTH-KENNEDY PASSER

CLEARED

Oct 4, 1979
3 days to decision

K791960 · Product code: **HWQ** · Orthopedic
Source: https://www.510kdatabase.net/k791960/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Passer (HWQ)
Date received	Oct 1, 1979
Decision date	Oct 4, 1979
Days to decision	3 days
Third-party review	No

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025