

K791960 ROTH-KENNEDY PASSEROct 4, 1979
3 days to decisionK791960 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k791960/>, 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 23, 2026