FDA 510(k) Premarket Notification

K811679 SILASTIC TENDON PASSER H.P.

CLEARED

Jul 10, 1981
25 days to decision

K811679 · Product code: **HWQ** · Orthopedic
Source: https://www.510kdatabase.net/k811679/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Passer (HWQ)
Date received	Jun 15, 1981
Decision date	Jul 10, 1981
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**Dow Corning Corp. Healthcare Industries Materials**
Location	Mchenry, IL, US
510(k) history	31 submissions · 31 cleared · 1976-1985