FDA 510(k) Premarket Notification

K901765 POSTEROMEDIAL GUIDE

CLEARED

Jun 25, 1990
68 days to decision

K901765 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k901765/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Apr 18, 1990
Decision date	Jun 25, 1990
Days to decision	68 days
Third-party review	No

APPLICANT

Company	**Smith & Nephew Dyonics, Inc.**
Location	Andover, MA, US
Contact	ERIC BANNON
510(k) history	22 submissions · 19 cleared · 1990-1996