

K901765 POSTEROMEDIAL GUIDEJun 25, 1990
68 days to decisionK901765 · 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

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