FDA 510(k) Premarket Notification

K982133 OLYMPUS ENDOSCOPIC SYSTEM LUMBAR HERNIA DISCECTOMY AND ITS ANCILLARY EQUIPMENT

CLEARED

Oct 22, 1998
127 days to decision

K982133 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k982133/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Jun 17, 1998
Decision date	Oct 22, 1998
Days to decision	127 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**The Olympus Optical Co.**
Location	Melville, NY, US
Contact	RUTH C FORSTADT
510(k) history	22 submissions · 22 cleared · 1998-2003