

**K982133 OLYMPUS ENDOSCOPIC SYSTEM LUMBAR HERNIA
DISCECTOMY AND ITS ANCILLARY EQUIPMENT**Oct 22, 1998
127 days to decisionK982133 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982133/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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