

K911974 PGK STEREOTACTIC DEVICENov 5, 1991
186 days to decisionK911974 · Product code: **HRX** · Orthopedic
Source: <https://www.510kdatabase.net/k911974/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	May 3, 1991
Decision date	Nov 5, 1991
Days to decision	186 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Northern Virginia Radiology and Nuclear Medicine
Location	Falls Church, VA, US
Contact	PANOS KOUTROUVELIS
510(k) history	1 submissions · 1 cleared · 1991-1991

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k911974/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026