

**K912843 FLEXMEDICS RETRACT ORTH PROBE, PETTINE BONE PROBE**

Sep 24, 1991  
90 days to decision

K912843 · Product code: **HXB** · Orthopedic  
Source: <https://www.510kdatabase.net/k912843/>

**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe (HXB)
Date received	Jun 26, 1991
Decision date	Sep 24, 1991
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Flexmedics</b>
Location	Minneapolis, MN, US
Contact	TIMOTHY LEY
510(k) history	20 submissions · 20 cleared · 1986-1999

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k912843/> 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