

**K870654 EFER ARTHROSCOPE**May 14, 1987  
90 days to decisionK870654 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k870654/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Feb 13, 1987
Decision date	May 14, 1987
Days to decision	90 days
Third-party review	No

**APPLICANT**

---

Company	<b>Roseburg SA</b>
Location	Washington, DC, US
Contact	DARIENNE MOYER
510(k) history	24 submissions · 23 cleared · 1985-1994

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k870654/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 22, 2026