

**K901849 LUMISCOPE EMS 8000**Aug 24, 1990  
122 days to decisionK901849 · Product code: **IPF** · Physical MedicineSource: <https://www.510kdatabase.net/k901849/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Muscle, Powered (IPF)
Date received	Apr 24, 1990
Decision date	Aug 24, 1990
Days to decision	122 days
Third-party review	No

**APPLICANT**

---

Company	<b>Lumiscop Co., Inc.</b>
Location	Edison, NJ, US
Contact	MORGENIER, III
510(k) history	13 submissions · 13 cleared · 1988-2003

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901849/>; 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 26, 2026