

**K803214 ACMI BIPOLAR HEMOSTATIC ELECTRODE**Jan 15, 1981  
28 days to decisionK803214 · Product code: **FEH** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k803214/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Flexible Suction Coagulator (FEH)
Date received	Dec 18, 1980
Decision date	Jan 15, 1981
Days to decision	28 days
Third-party review	No

**APPLICANT**

---

Company	<b>American Cystoscope Makers, Inc.</b>
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1976-1983

---

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

Device record: <https://www.510kdatabase.net/k803214/> 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 28, 2026