

**K854511 ELECATH PERCUTANEOUS LEFT ATRIAL  
CANNULATION SET**Feb 10, 1986  
90 days to decisionK854511 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k854511/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Nov 12, 1985
Decision date	Feb 10, 1986
Days to decision	90 days
Third-party review	No

**APPLICANT**

---

Company	<b>Electro-Catheter Corp.</b>
Location	Mchenry, IL, US
Contact	SILPE
510(k) history	35 submissions · 35 cleared · 1976-1995

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k854511/> 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 27, 2026