

**K801831 EVALUATOR #9500 DYNAMIC ELEC/SCANNER**Sep 9, 1980  
39 days to decisionK801831 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k801831/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 1, 1980
Decision date	Sep 9, 1980
Days to decision	39 days
Third-party review	No

**APPLICANT**

---

Company	<b>Del Mar Avionics</b>
Location	Mchenry, IL, US
510(k) history	54 submissions · 54 cleared · 1977-1999

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k801831/> 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 June 29, 2026