

**K800319 ALPHA DETECTOR**Mar 3, 1980  
20 days to decisionK800319 · Product code: **HCC** · Neurology  
Source: <https://www.510kdatabase.net/k800319/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Feb 12, 1980
Decision date	Mar 3, 1980
Days to decision	20 days
Third-party review	No

**APPLICANT**

---

Company	<b>M.O.E Electronics Research &amp; Prod.</b>
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1980-1985

---

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

Device record: <https://www.510kdatabase.net/k800319/> 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 July 1, 2026