

**K812331 VTS-1000 VISUAL TESTING SYSTEM**Nov 16, 1981  
90 days to decisionK812331 · Product code: **HLX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k812331/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photostimulator, Ac-powered (HLX)
Date received	Aug 18, 1981
Decision date	Nov 16, 1981
Days to decision	90 days
Third-party review	No

**APPLICANT**

---

Company	<b>Nicolet Biomedical Instruments</b>
Location	Mchenry, IL, US
510(k) history	26 submissions · 26 cleared · 1977-1988

---

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

Device record: <https://www.510kdatabase.net/k812331/> 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