

**K812061 MODEL S10VS VISUAL STIMULATOR**Aug 20, 1981  
30 days to decisionK812061 · Product code: **GWE** · Neurology  
Source: <https://www.510kdatabase.net/k812061/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Photic, Evoked Response (GWE)
Date received	Jul 21, 1981
Decision date	Aug 20, 1981
Days to decision	30 days
Third-party review	No

**APPLICANT**

---

Company	<b>Grass Instrument Co.</b>
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1977-2002

---

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

Device record: <https://www.510kdatabase.net/k812061/> 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 30, 2026