

**K022304 ACCUMAP AUTOMATIC PERIMETER**Nov 14, 2003  
486 days to decisionK022304 · Product code: **GWE** · Neurology  
Source: <https://www.510kdatabase.net/k022304/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Photic, Evoked Response (GWE)
Date received	Jul 16, 2002
Decision date	Nov 14, 2003
Days to decision	486 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Objectivision Pty , Ltd.</b>
Location	Troy, NY, US
Contact	FRED HENRY
510(k) history	1 submissions · 1 cleared · 2003-2003

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022304/> 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