

K081102 VSU - VISIONSENSE STEREOSCOPIC VISION SYSTEMMay 29, 2008
42 days to decisionK081102 · Product code: **GWG** · Neurology
Source: <https://www.510kdatabase.net/k081102/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscope, Neurological (GWG)
Date received	Apr 17, 2008
Decision date	May 29, 2008
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Visionsense, Ltd.
Location	Washington, DC, US
Contact	GERARD J PRUD'OMME
510(k) history	13 submissions · 13 cleared · 2008-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081102/> Data sourced from FDA 510(k) public records (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. - Generated May 26, 2026