

K012976 TWIN PLUSFeb 12, 2002
160 days to decisionK012976 · Product code: **OLV** · Neurology
Source: <https://www.510kdatabase.net/k012976/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Standard Polysomnograph With Electroencephalograph (OLV)
Date received	Sep 5, 2001
Decision date	Feb 12, 2002
Days to decision	160 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Grass-Telefactor Divison
Location	West-Warwick, RI, US
Contact	JOHN B CHATTEN
510(k) history	1 submissions · 1 cleared · 2002-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012976/> 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 June 30, 2026