

K022314 SANARUS VISICA TREATMENT SYSTEMOct 15, 2002
90 days to decisionK022314 · Product code: **GEH** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k022314/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryosurgical, Accessories (GEH)
Date received	Jul 17, 2002
Decision date	Oct 15, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sanarus Medical, Inc.
Location	Pleasanton, CA, US
Contact	SETH STABINSKY
510(k) history	14 submissions · 14 cleared · 2002-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k022314/> 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 25, 2026