FDA 510(k) Premarket Notification

K022314 SANARUS VISICA TREATMENT SYSTEM

CLEARED

Oct 15, 2002
90 days to decision

K022314 · Product code: **GEH** · General & Plastic Surgery
Source: 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