

K112334 VEIN-GOGH INSTRUMENTMay 8, 2012
270 days to decisionK112334 · Product code: **ONQ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k112334/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical Coagulation For Aesthetic (ONQ)
Date received	Aug 12, 2011
Decision date	May 8, 2012
Days to decision	270 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Refine USA, LLC
Location	Jacksonville Beach, FL, US
Contact	BRIAN SMITH
510(k) history	2 submissions · 2 cleared · 2012-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k112334/> 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 29, 2026