FDA 510(k) Premarket Notification

K060573 BURBANK TENACULUM

CLEARED

Apr 21, 2006
46 days to decision

K060573 · Product code: **HDC** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k060573/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tenaculum, Uterine (HDC)
Date received	Mar 6, 2006
Decision date	Apr 21, 2006
Days to decision	46 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Vascular Control Systems, Inc.**
Location	San Juan Capistrano, CA, US
Contact	KATHLEEN ROBERTS
510(k) history	9 submissions · 9 cleared · 2002-2006