FDA 510(k) Premarket Notification

K093556 VCARE DX

CLEARED

Mar 12, 2010
115 days to decision

K093556 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k093556/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Nov 17, 2009
Decision date	Mar 12, 2010
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Conmed Corporation**
Location	Utica, NY, US
Contact	SARAH RIZK
Website	https://www.conmed.com
510(k) history	83 submissions · 83 cleared · 2004-2026

Conmed Corporation is a global medical device manufacturer specializing in surgical equipment and operating room solutions. The company operates with a manufacturing facility in Utica, US, and serves multiple surgical specialties including general surgery, orthopedics, and patient monitoring. Conmed has received FDA 510(k) clearances from total submissions since its first clearance in 2004. The company maintains active regulatory engagement, with its most recent clearance in 2026. Its cleared devices focus primarily on General & Plastic Surgery applications, including ele...