FDA 510(k) Premarket Notification

K132815 PUREGRAFT 50 SYSTEM

CLEARED

Jan 24, 2014
137 days to decision

K132815 · Product code: **MUU** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k132815/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Suction, Lipoplasty (MUU)
Date received	Sep 9, 2013
Decision date	Jan 24, 2014
Days to decision	137 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cytori Therapeutics, Inc.**
Location	San Diego, CA, US
Contact	KENNETH K KLEINHENZ
510(k) history	6 submissions · 6 cleared · 2006-2014