FDA 510(k) Premarket Notification

K093702 REPROCESSED ULTRASONIC COAGULATING SHEARS

CLEARED

Mar 4, 2010
93 days to decision

K093702 · Product code: **NLQ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k093702/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single-use Reprocessed Ultrasonic Surgical Instruments (NLQ)
Date received	Dec 1, 2009
Decision date	Mar 4, 2010
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ascent Healthcare Solutions**
Location	Phoenix, AZ, US
Contact	MOIRA BARTON VARTY
510(k) history	21 submissions · 21 cleared · 2006-2011