

**K093702 REPROCESSED ULTRASONIC COAGULATING SHEARS**

Mar 4, 2010  
93 days to decision

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

**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k093702/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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