

**K052690 SURETEK REPROCESSED LAPAROSCOPIC INSTRUMENTS**May 12, 2006  
226 days to decisionK052690 · Product code: **NUJ** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k052690/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation Accessories, Laparoscopic & Endoscopic, Reprocessed (NUJ)
Date received	Sep 28, 2005
Decision date	May 12, 2006
Days to decision	226 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Suretek Medical</b>
Location	Greenville, SC, US
Contact	MIKE SAMMON
510(k) history	4 submissions · 4 cleared · 2006-2006

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052690/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026