

K011190 SURX RADIOFREQUENCY ELECTROSURGICAL GENERATOR SYSTEM AND ACCESSORIES (SURX LP SYSTEM)

Jan 8, 2002
264 days to decision

K011190 · Product code: **MUK** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k011190/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical Radiofrequency System, Stress Urinary Incontinence, Female, Transvaginal Or Laparoscopic, Pelvic Tissue (MUK)
Date received	Apr 19, 2001
Decision date	Jan 8, 2002
Days to decision	264 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Surx, Inc.
Location	Pleasanton, CA, US
Contact	ALAN CURTIS
510(k) history	3 submissions · 3 cleared · 2002-2002

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k011190/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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