FDA 510(k) Premarket Notification

K954710 VAPOR-TECH ROLLER BALL ELECTRODE 24 FR.

CLEARED

Apr 24, 1996
195 days to decision

K954710 · Product code: **FAS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k954710/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrosurgical, Active, Urological (FAS)
Date received	Oct 12, 1995
Decision date	Apr 24, 1996
Days to decision	195 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Northgate Technologies, Inc.**
Location	Arlington Heights, IL, US
Contact	CASEY KUREK
510(k) history	55 submissions · 55 cleared · 1991-2021

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k954710/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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