

K020985 MODIFIED TFX MEDICAL SAFETY NEEDLE WITH INTRODUCERJun 25, 2002
90 days to decisionK020985 · Product code: **KNT** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k020985/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Mar 27, 2002
Decision date	Jun 25, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Tfx Medical, Inc.
Location	Jeffrey, NH, US
Contact	RICK LYKINS
510(k) history	4 submissions · 4 cleared · 2000-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020985/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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