FDA 510(k) Premarket Notification

CLEARED

K880095 BERGER PROSTATE CULTURE CYTOLOGY BRUSH

Mar 2, 1988
50 days to decision

K880095 · Product code: **FDX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k880095/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Cytology Brush (FDX)
Date received	Jan 12, 1988
Decision date	Mar 2, 1988
Days to decision	50 days
Third-party review	No

APPLICANT

Company	**Van-Tec, Inc.**
Location	Walker, MI, US
Contact	VANCE, JR.
510(k) history	52 submissions · 52 cleared · 1982-1988

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k880095/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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