FDA 510(k) Premarket Notification

K880096 VAN-TEC ASPIRATION BIOPSY SYSTEM W/COLLECTION CUP

CLEARED

Mar 2, 1988
50 days to decision

K880096 · Product code: **KNW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k880096/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
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/k880096/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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