FDA 510(k) Premarket Notification

K955150 MAXI-FLOW

CLEARED

Jan 23, 1996
71 days to decision

K955150 · Product code: **BYY** · General Hospital
Source: https://www.510kdatabase.net/k955150/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Aspirating, Flexible, Connecting (BYY)
Date received	Nov 13, 1995
Decision date	Jan 23, 1996
Days to decision	71 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Bio-Medical Devices, Inc.**
Location	Irvine, CA, US
Contact	NICK HERBERT
510(k) history	7 submissions · 7 cleared · 1994-2004