FDA 510(k) Premarket Notification

K952084 HUMID-VENT FILTER PEDI

CLEARED

Jun 12, 1995
40 days to decision

K952084 · Product code: **BYD** · Anesthesiology
Source: https://www.510kdatabase.net/k952084/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condenser, Heat And Moisture (artificial Nose) (BYD)
Date received	May 3, 1995
Decision date	Jun 12, 1995
Days to decision	40 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Gibeck, Inc.**
Location	Indianapolis, IN, US
Contact	BRIAN GRIGSBY
510(k) history	15 submissions · 15 cleared · 1990-1997