FDA 510(k) Premarket Notification

K880002 MULTI-ALLERGEN IGE FAST SCREEN (MOLDS)

CLEARED

Feb 17, 1988
44 days to decision

K880002 · Product code: **DHB** · Immunology
Source: https://www.510kdatabase.net/k880002/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Test, Radioallergosorbent (rast) Immunological (DHB)
Date received	Jan 4, 1988
Decision date	Feb 17, 1988
Days to decision	44 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Contact	MONTE
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025