

K880002 MULTI-ALLERGEN IGE FAST SCREEN (MOLDS)Feb 17, 1988
44 days to decisionK880002 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880002/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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