

K792695 ANTI-GLOBULIN CONTROLJan 24, 1980
28 days to decisionK792695 · Product code: **DHR** · Immunology
Source: <https://www.510kdatabase.net/k792695/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Test, Rheumatoid Factor (DHR)
Date received	Dec 27, 1979
Decision date	Jan 24, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Meridian Diagnostics, Inc.
Location	Walker, MI, US
510(k) history	92 submissions · 92 cleared · 1980-1999

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Device record: <https://www.510kdatabase.net/k792695/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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