FDA 510(k) Premarket Notification

K792695 ANTI-GLOBULIN CONTROL

CLEARED

Jan 24, 1980
28 days to decision

K792695 · 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