

K812972 RA LATEX TESTNov 6, 1981
14 days to decisionK812972 · Product code: **DHR** · Microbiology
Source: <https://www.510kdatabase.net/k812972/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Test, Rheumatoid Factor (DHR)
Date received	Oct 23, 1981
Decision date	Nov 6, 1981
Days to decision	14 days
Third-party review	No

APPLICANT

Company	Oxoid U.S.A., Inc.
Location	Mchenry, IL, US
510(k) history	93 submissions · 93 cleared · 1980-1989

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Device record: <https://www.510kdatabase.net/k812972/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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