FDA 510(k) Premarket Notification

K812972 RA LATEX TEST

CLEARED

Nov 6, 1981
14 days to decision

K812972 · 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