FDA 510(k) Premarket Notification

K790122 RUBANON

CLEARED

Feb 15, 1979
23 days to decision

K790122 · Product code: **GOK** · Microbiology
Source: https://www.510kdatabase.net/k790122/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Hai (including Hai Control), Rubella (GOK)
Date received	Jan 23, 1979
Decision date	Feb 15, 1979
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Organon, Inc.**
Location	Mchenry, IL, US
510(k) history	41 submissions · 41 cleared · 1977-1995