FDA 510(k) Premarket Notification

K781392 CYTOMEGILISH TEST KIT

CLEARED

Oct 24, 1978
75 days to decision

K781392 · Product code: **GOK** · Microbiology
Source: https://www.510kdatabase.net/k781392/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Hai (including Hai Control), Rubella (GOK)
Date received	Aug 10, 1978
Decision date	Oct 24, 1978
Days to decision	75 days
Third-party review	No

APPLICANT

Company	**Microbiological Assoc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1977-1980