FDA 510(k) Premarket Notification

K830783 VERI-STAPH

CLEARED

Apr 8, 1983
28 days to decision

K830783 · Product code: **JWX** · Microbiology
Source: https://www.510kdatabase.net/k830783/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Screening, Staphylococcus Aureus (JWX)
Date received	Mar 11, 1983
Decision date	Apr 8, 1983
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Zeus Technologies**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1983-1984