

K830783 VERI-STAPHApr 8, 1983
28 days to decisionK830783 · 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

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Device record: <https://www.510kdatabase.net/k830783/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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