

K811412 NEISSERIA-KWIKJun 16, 1981
28 days to decisionK811412 · Product code: **JSX** · Chemistry
Source: <https://www.510kdatabase.net/k811412/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Identification, Neisseria Gonorrhoeae (JSX)
Date received	May 19, 1981
Decision date	Jun 16, 1981
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Windsor Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1981-1987

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k811412/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026