

K813607 ACKRAD N-DO VAGE TUBEFeb 5, 1982
38 days to decisionK813607 · Product code: **FQH** · General Hospital
Source: <https://www.510kdatabase.net/k813607/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lavage, Jet (FQH)
Date received	Dec 29, 1981
Decision date	Feb 5, 1982
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Ackrad Laboratories
Location	Mchenry, IL, US
510(k) history	42 submissions · 41 cleared · 1979-2002

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Device record: <https://www.510kdatabase.net/k813607/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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