FDA 510(k) Premarket Notification

K813607 ACKRAD N-DO VAGE TUBE

CLEARED

Feb 5, 1982
38 days to decision

K813607 · 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