FDA 510(k) Premarket Notification

K781234 TUBES, LUKENS, TOMAS

CLEARED

Aug 10, 1978
22 days to decision

K781234 · Product code: **KDQ** · General Hospital
Source: https://www.510kdatabase.net/k781234/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bottle, Collection, Vacuum (KDQ)
Date received	Jul 19, 1978
Decision date	Aug 10, 1978
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Ludwig Medical, Inc.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1978-1978