

K781234 TUBES, LUKENS, TOMASAug 10, 1978
22 days to decisionK781234 · 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

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Device record: <https://www.510kdatabase.net/k781234/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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