FDA 510(k) Premarket Notification

K830836 Q.C. TESTER

CLEARED

Apr 18, 1983
27 days to decision

K830836 · Product code: **LHI** · Microbiology
Source: https://www.510kdatabase.net/k830836/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Mar 22, 1983
Decision date	Apr 18, 1983
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Alltek Filtration, Inc.**
Location	Walker, MI, US
510(k) history	1 submissions · 1 cleared · 1983-1983