FDA 510(k) Premarket Notification

K830970 DOUBLE LUMEN TRANSFER DEVICE

CLEARED

Apr 12, 1983
15 days to decision

K830970 · Product code: **LHI** · General Hospital
Source: https://www.510kdatabase.net/k830970/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Mar 28, 1983
Decision date	Apr 12, 1983
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**Codan Trading Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983