FDA 510(k) Premarket Notification

K832312 I.V. TRANSFER NEEDLE

CLEARED

Sep 12, 1983
60 days to decision

K832312 · Product code: **LHI** · General Hospital
Source: https://www.510kdatabase.net/k832312/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Jul 14, 1983
Decision date	Sep 12, 1983
Days to decision	60 days
Third-party review	No

APPLICANT

Company	**Baxa Corp., Sub. of Cook Group, Inc.**
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1978-1991