FDA 510(k) Premarket Notification

K963012 BIOJECT NEEDLE-FREE VIAL ADAPTER (13MM)

CLEARED

Oct 15, 1996
74 days to decision

K963012 · Product code: **LHI** · General Hospital
Source: https://www.510kdatabase.net/k963012/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, I.v. Fluid Transfer (LHI)
Date received	Aug 2, 1996
Decision date	Oct 15, 1996
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Bioject, Inc.**
Location	Portland, OR, US
Contact	JEFF SNOW
510(k) history	9 submissions · 9 cleared · 1987-2009