

K963012 BIOJECT NEEDLE-FREE VIAL ADAPTER (13MM)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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k963012/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 23, 2026