

K892665 PRO-JECT(TM) SAFETY NEEDLE HOLDEROct 4, 1989
174 days to decisionK892665 · Product code: **JKA** · Chemistry
Source: <https://www.510kdatabase.net/k892665/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Apr 13, 1989
Decision date	Oct 4, 1989
Days to decision	174 days
Third-party review	No

APPLICANT

Company	Pro-Tec Containers, Inc.
Location	Irvine, CA, US
Contact	RALPH T HAGINS
510(k) history	6 submissions · 6 cleared · 1989-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k892665/>; 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 30, 2026