

**K933848 PRESPO 7, SELF-SUFFICIENT DRIP DEVICE**Mar 3, 1995  
574 days to decisionK933848 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k933848/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 6, 1993
Decision date	Mar 3, 1995
Days to decision	574 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Bonding Corp.</b>
Location	Coraopolis, PA, US
Contact	JEAN-PIERRE LA RUDULIER
510(k) history	1 submissions · 1 cleared · 1995-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933848/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 6, 2026