

**K812852 SET FOR ASEPTIC DECANTING OF A FLEX I.V**Nov 2, 1981  
20 days to decisionK812852 · Product code: **KPE** · General HospitalSource: <https://www.510kdatabase.net/k812852/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Oct 13, 1981
Decision date	Nov 2, 1981
Days to decision	20 days
Third-party review	No

**APPLICANT**

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Company	<b>Dip, Inc.</b>
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1981-1981

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k812852/> 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 9, 2026