

**K830868 COMBISET OR INFUSION SET**Apr 14, 1983  
27 days to decisionK830868 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k830868/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 18, 1983
Decision date	Apr 14, 1983
Days to decision	27 days
Third-party review	No

**APPLICANT**

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Company	<b>Pan-Tek Intl., Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k830868/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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