

**K843054 DEKA SOLUTION ADMIN. SET DK 302-A**Sep 13, 1984  
41 days to decisionK843054 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k843054/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 3, 1984
Decision date	Sep 13, 1984
Days to decision	41 days
Third-party review	No

**APPLICANT**

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Company	<b>Deka Research &amp; Development Corp.</b>
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1984-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k843054/>; 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 May 25, 2026