

**K841361 SYRINGE ADDITIVE IV ADMIN. SETS**May 21, 1984  
49 days to decisionK841361 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k841361/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 2, 1984
Decision date	May 21, 1984
Days to decision	49 days
Third-party review	No

**APPLICANT**

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Company	<b>Quest Medical, Inc.</b>
Location	Walker, MI, US
510(k) history	39 submissions · 39 cleared · 1980-2024

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

Device record: <https://www.510kdatabase.net/k841361/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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