

**K132011 APOTECA DUG COMPOUNDING DOSING  
DEVICE, APOTECA I.V. TRANSFER SET**Mar 21, 2014  
263 days to decisionK132011 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k132011/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Syringe, Piston (FMF)
Date received	Jul 1, 2013
Decision date	Mar 21, 2014
Days to decision	263 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aea Srl</b>
Location	Angeli Di Rosora, IT
Contact	MICHELE MENGONI
510(k) history	3 submissions · 3 cleared · 2014-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132011/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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