

**K802340 SPECIMEN SET**Oct 10, 1980  
14 days to decisionK802340 · Product code: **FMH** · General Hospital  
Source: <https://www.510kdatabase.net/k802340/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Specimen, Sterile (FMH)
Date received	Sep 26, 1980
Decision date	Oct 10, 1980
Days to decision	14 days
Third-party review	No

**APPLICANT**

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Company	<b>American Pharmaseal Div. Ahsc</b>
Location	Mchenry, IL, US
510(k) history	64 submissions · 61 cleared · 1980-1987

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

Device record: <https://www.510kdatabase.net/k802340/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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