

**K770112 SPECIMEN COLLECTOR**Mar 14, 1977  
54 days to decisionK770112 · Product code: **FMH** · General Hospital  
Source: <https://www.510kdatabase.net/k770112/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Specimen, Sterile (FMH)
Date received	Jan 19, 1977
Decision date	Mar 14, 1977
Days to decision	54 days
Third-party review	No

**APPLICANT**

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Company	<b>Portex, Inc.</b>
Location	Walker, MI, US
510(k) history	20 submissions · 20 cleared · 1977-2004

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

Device record: <https://www.510kdatabase.net/k770112/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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