

**K802359 DISP. FENESTRATED SURG. DRAPE SHEETS**Oct 31, 1980  
35 days to decisionK802359 · Product code: **KKX** · General Hospital  
Source: <https://www.510kdatabase.net/k802359/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Sep 26, 1980
Decision date	Oct 31, 1980
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>American Convertors Div., American Pharmaseal</b>
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1980-1984

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

Device record: <https://www.510kdatabase.net/k802359/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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