

**K963570 SIMS ADULT LUMBAR PUNCTURE KIT  
(4800-20,4801-20,4806-22,4807-22)**

Oct 18, 1996  
43 days to decision

K963570 · Product code: **FMJ** · General Hospital  
Source: <https://www.510kdatabase.net/k963570/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manometer, Spinal-fluid (FMJ)
Date received	Sep 5, 1996
Decision date	Oct 18, 1996
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Smiths Industries Medical Systems, Inc.</b>
Location	Keene, NH, US
Contact	TIMOTHY J TALCOTT
510(k) history	22 submissions · 19 cleared · 1993-1998

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

Device record: <https://www.510kdatabase.net/k963570/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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