

**K894433 BATES BLANKET**Aug 11, 1989  
25 days to decisionK894433 · Product code: **KKX** · General Hospital  
Source: <https://www.510kdatabase.net/k894433/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Jul 17, 1989
Decision date	Aug 11, 1989
Days to decision	25 days
Third-party review	No

**APPLICANT**

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Company	<b>Joe B. Bates, MD, FAAP</b>
Location	Tyler, TX, US
Contact	BATES, MD
510(k) history	1 submissions · 1 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894433/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 8, 2026