

**K897120 EQUIP COVERS, STERILE**Jan 4, 1990  
9 days to decisionK897120 · Product code: **KKX** · General Hospital  
Source: <https://www.510kdatabase.net/k897120/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Dec 26, 1989
Decision date	Jan 4, 1990
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>Amedic USA</b>
Location	Phoenix, AZ, US
Contact	THEO RUSSO-LARSSON
510(k) history	10 submissions · 10 cleared · 1987-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k897120/>, 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 May 21, 2026