

**K895431 JOHNSON & JOHNSON TUBE SECUREMENT DEVICE**Oct 13, 1989  
37 days to decisionK895431 · Product code: **KMK** · General Hospital  
Source: <https://www.510kdatabase.net/k895431/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Sep 6, 1989
Decision date	Oct 13, 1989
Days to decision	37 days
Third-party review	No

**APPLICANT**

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Company	<b>Johnson &amp; Johnson Patient Care, Inc.</b>
Location	New Brunswick, NJ, US
Contact	MC CANN
510(k) history	4 submissions · 2 cleared · 1988-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895431/>; 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 7, 2026