

**K870045 SKYTRON MODEL 610-2**Feb 4, 1987  
30 days to decisionK870045 · Product code: **INX** · Physical MedicineSource: <https://www.510kdatabase.net/k870045/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Air Fluidized (INX)
Date received	Jan 5, 1987
Decision date	Feb 4, 1987
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Skytron, Div. the Kmw Group, Inc.</b>
Location	Mchenry, IL, US
Contact	LARRY J PURCEY
510(k) history	19 submissions · 19 cleared · 1981-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k870045/>; 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 June 28, 2026