

K843389 CLINTRON AIR FLUIDIZED SUPPORT J-101UDSep 5, 1984
7 days to decisionK843389 · Product code: **INX** · Physical MedicineSource: <https://www.510kdatabase.net/k843389/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Air Fluidized (INX)
Date received	Aug 29, 1984
Decision date	Sep 5, 1984
Days to decision	7 days
Third-party review	No

APPLICANT

Company	Uhi Corp.
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1984

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k843389/> Data sourced from FDA 510(k) public records (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. - Generated June 30, 2026