

K831348 FLUID ADMIN. SET AFS-1CMay 27, 1983
32 days to decisionK831348 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k831348/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Apr 25, 1983
Decision date	May 27, 1983
Days to decision	32 days
Third-party review	No

APPLICANT

Company	Linton Biomed Corp.
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1982-1984

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Device record: <https://www.510kdatabase.net/k831348/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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