

K842759 MODEL 503FBJan 18, 1985
186 days to decisionK842759 · Product code: **LOS** · Cardiovascular
Source: <https://www.510kdatabase.net/k842759/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jul 16, 1984
Decision date	Jan 18, 1985
Days to decision	186 days
Third-party review	No

APPLICANT

Company	Atrix
Location	Portland, OR, US
Contact	KIM F DUNCAN
510(k) history	2 submissions · 2 cleared · 1985-1986

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Device record: <https://www.510kdatabase.net/k842759/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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