

K802924 KONTRON PERCUTAEIOUS DOUBLE-LUMENMar 11, 1981
113 days to decisionK802924 · Product code: **DSP** · CardiovascularSource: <https://www.510kdatabase.net/k802924/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Balloon, Intra-aortic And Control (DSP)
Date received	Nov 18, 1980
Decision date	Mar 11, 1981
Days to decision	113 days
Third-party review	No

APPLICANT

Company	Avco Medical Products
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1981

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Device record: <https://www.510kdatabase.net/k802924/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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