

K812639 UNICUFFOct 2, 1981
16 days to decisionK812639 · Product code: **DXP** · CardiovascularSource: <https://www.510kdatabase.net/k812639/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Vessel Occlusion (DXP)
Date received	Sep 16, 1981
Decision date	Oct 2, 1981
Days to decision	16 days
Third-party review	No

APPLICANT

Company	Frontier Products
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1983

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Device record: <https://www.510kdatabase.net/k812639/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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