

K830573 PRE VUEMar 29, 1983
34 days to decisionK830573 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k830573/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Feb 23, 1983
Decision date	Mar 29, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Taut, Inc.
Location	Walker, MI, US
510(k) history	16 submissions · 16 cleared · 1983-2002

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Device record: <https://www.510kdatabase.net/k830573/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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