

**K891180 TRANSESOPHAGEAL ULTRASOUND ACCESSORIES
KIT**Jun 15, 1989
101 days to decisionK891180 · Product code: **DXP** · Cardiovascular
Source: <https://www.510kdatabase.net/k891180/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Vessel Occlusion (DXP)
Date received	Mar 6, 1989
Decision date	Jun 15, 1989
Days to decision	101 days
Third-party review	No

APPLICANT

Company	Ingress Technologies, Inc.
Location	Phoenix, AZ, US
Contact	SHANE H ABOWITT
510(k) history	1 submissions · 1 cleared · 1989-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k891180/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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