FDA 510(k) Premarket Notification

K891180 TRANSESOPHAGEAL ULTRASOUND ACCESSORIES KIT

CLEARED

Jun 15, 1989
101 days to decision

K891180 · 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