FDA 510(k) Premarket Notification

K924981 OPERATIVE CHOLANGIOGRAM CATHETER

CLEARED

Apr 6, 1993
187 days to decision

K924981 · Product code: **GBZ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k924981/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cholangiography (GBZ)
Date received	Oct 1, 1992
Decision date	Apr 6, 1993
Days to decision	187 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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