FDA 510(k) Premarket Notification

K982409 VES 1501-M

CLEARED

Aug 21, 1998
42 days to decision

K982409 · Product code: **FWF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k982409/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Television, Endoscopic, Without Audio (FWF)
Date received	Jul 10, 1998
Decision date	Aug 21, 1998
Days to decision	42 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Angiolaz, Inc.**
Location	Rockingham, VT, US
Contact	JOHN D PLUMADORE
510(k) history	10 submissions · 10 cleared · 1990-2000