FDA 510(k) Premarket Notification

K882358 PULSED FLUOROSCOPIC CONTROL MODEL:PF-02

CLEARED

Oct 19, 1988
134 days to decision

K882358 · Product code: **IZI** · Radiology
Source: https://www.510kdatabase.net/k882358/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Angiographic (IZI)
Date received	Jun 7, 1988
Decision date	Oct 19, 1988
Days to decision	134 days
Third-party review	No

APPLICANT

Company	**U.S. Imaging, Inc.**
Location	Greensburg, OH, US
Contact	MARTIN J KULIS
510(k) history	4 submissions · 4 cleared · 1988-1990