FDA 510(k) Premarket Notification

K984466 DILON 2000, MODEL 2000-6X8 AND DILON 2000, MODEL 2000-5X5

CLEARED

Mar 16, 1999
90 days to decision

K984466 · Product code: **IYX** · Radiology
Source: https://www.510kdatabase.net/k984466/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Scintillation (gamma) (IYX)
Date received	Dec 16, 1998
Decision date	Mar 16, 1999
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dilon Technologies, Inc.**
Location	Stillwater, MN, US
Contact	LEE H FAIRCHILD
510(k) history	2 submissions · 2 cleared · 1999-2011