FDA 510(k) Premarket Notification

K810115 DOPPLEX-PLB & PXB

CLEARED

Jul 16, 1981
181 days to decision

K810115 · Product code: **KNG** · Radiology
Source: https://www.510kdatabase.net/k810115/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Ultrasonic, Fetal (KNG)
Date received	Jan 16, 1981
Decision date	Jul 16, 1981
Days to decision	181 days
Third-party review	No

APPLICANT

Company	**Huntleigh Technology, Inc.**
Location	Walker, MI, US
510(k) history	23 submissions · 23 cleared · 1981-1999