FDA 510(k) Premarket Notification

K912007 BAXTER SINGLE PATIENT USE BLOOD PRESSURE CUFF

CLEARED

Jul 22, 1991
77 days to decision

K912007 · Product code: **DXQ** · Cardiovascular
Source: https://www.510kdatabase.net/k912007/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	May 6, 1991
Decision date	Jul 22, 1991
Days to decision	77 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Tecnadyne Scientific, Inc.**
Location	Vero Beach, FL, US
Contact	JOHN GARRETT
510(k) history	5 submissions · 5 cleared · 1990-1994