FDA 510(k) Premarket Notification

K911996 SURGICUTT-BLEEDING TIME DEVICE, MODIFICATION

CLEARED

Jul 17, 1991
86 days to decision

K911996 · Product code: **JCA** · Hematology
Source: https://www.510kdatabase.net/k911996/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Bleeding Time (JCA)
Date received	Apr 22, 1991
Decision date	Jul 17, 1991
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**International Technidyne Corp.**
Location	Walker, MI, US
Contact	LES HEIMANN
510(k) history	47 submissions · 47 cleared · 1983-2013