FDA 510(k) Premarket Notification

K911913 PRECISION GLIDE - ENHANCED SURGICAL BLADE

CLEARED

May 13, 1991
13 days to decision

K911913 · Product code: **GES** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k911913/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blade, Scalpel (GES)
Date received	Apr 30, 1991
Decision date	May 13, 1991
Days to decision	13 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Bd Becton Dickinson Vacutainer Systems Preanalytic**
Location	Washington, DC, US
Contact	J ARNSBERGER
510(k) history	632 submissions · 625 cleared · 1976-2001