FDA 510(k) Premarket Notification

K030076 PROFESSIONAL BLOOD SAMPLE RETRIEVAL SYSTEM

CLEARED

Jul 7, 2003
180 days to decision

K030076 · Product code: **JKA** · Chemistry
Source: https://www.510kdatabase.net/k030076/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Jan 8, 2003
Decision date	Jul 7, 2003
Days to decision	180 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Biomed Personal Metabolic and Nutritional Testing**
Location	Memphis, TN, US
Contact	SUSAN B FENTRESS
510(k) history	2 submissions · 2 cleared · 2003-2003