FDA 510(k) Premarket Notification

K030394 FLUID SAMPLE RETRIEVAL SYSTEM

CLEARED

Oct 27, 2003
264 days to decision

K030394 · Product code: **JKA** · Chemistry
Source: https://www.510kdatabase.net/k030394/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Vials, Systems, Serum Separators, Blood Collection (JKA)
Date received	Feb 5, 2003
Decision date	Oct 27, 2003
Days to decision	264 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