FDA 510(k) Premarket Notification

K761152 S-UREA-N, IN VITRO DIAG FOR BUN

CLEARED

Dec 9, 1976
10 days to decision

K761152 · Product code: **CDL** · Chemistry
Source: https://www.510kdatabase.net/k761152/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Berthelot Indophenol, Urea Nitrogen (CDL)
Date received	Nov 29, 1976
Decision date	Dec 9, 1976
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Advanced Biomedical Methods, Inc.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1976-1994