

K181730 BD Vacutainer Eclipse UltraFill Blood Collection Needle

Mar 1, 2019
245 days to decisionK181730 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k181730/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Jun 29, 2018
Decision date	Mar 1, 2019
Days to decision	245 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Becton, Dickinson and Company
Location	Franklin Lakes, NJ, US
Contact	Eileen Hiller
Website	https://www.bd.com
510(k) history	134 submissions · 134 cleared · 2010-2026

Becton, Dickinson and Company is an American multinational medical technology company headquartered in Franklin Lakes, New Jersey. BD manufactures and sells medical devices, instrument systems, and reagents globally. The company maintains a strong FDA 510(k) regulatory record with FDA 510(k) clearances from total submissions spanning 2010 to 2026. BD's cleared devices span multiple categories including microbiology systems, blood collection products, and general hospital devices. The company's latest clearance in 2026 reflects continued innovation and regulatory engagement...

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Device record: <https://www.510kdatabase.net/k181730/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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