

**K200932 Blood Collection Needles**Dec 14, 2021  
616 days to decisionK200932 · Product code: **FMI** · General HospitalSource: <https://www.510kdatabase.net/k200932/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Apr 7, 2020
Decision date	Dec 14, 2021
Days to decision	616 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vacutest Kima Srl</b>
Location	Arzegrande, IT
Contact	Monica Vianello
510(k) history	1 submissions · 1 cleared · 2021-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k200932/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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