

**K210381 I.V. Administration Set**Aug 14, 2021  
186 days to decisionK210381 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k210381/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 9, 2021
Decision date	Aug 14, 2021
Days to decision	186 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bq Plus Medical Co., Ltd.</b>
Location	Shanghai, CN
Contact	Jin Zhang
510(k) history	3 submissions · 3 cleared · 2021-2024

**REGULATORY CONSULTANT**

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Consulting firm	<b>Mid-Link Consulting Co, Ltd.</b>
Contact	Diana Hong

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k210381/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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