

**K242126 I.V. Administration Set**Aug 22, 2024  
34 days to decisionK242126 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k242126/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 19, 2024
Decision date	Aug 22, 2024
Days to decision	34 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

**APPLICANT**

---

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**

---

Consulting firm	<b>Shanghai SUNGO Management Consulting Co., Ltd.</b>
Contact	Eva Li

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

---

510k Database - [www.510kdatabase.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k242126/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated May 13, 2026