

K220083 LDV((Low Dead Volume)SyringeMay 25, 2022
135 days to decisionK220083 · Product code: **QNG** · General Hospital
Source: <https://www.510kdatabase.net/k220083/>**SUBMISSION DETAILS**

| | |
|-----------------------|-------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Low Dead Space Piston Syringe (QNG) |
| Date received | Jan 10, 2022 |
| Decision date | May 25, 2022 |
| Days to decision | 135 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Anhui Hongyu Wuzhou Medical Manufacturer Co., Ltd. |
| Location | Taihu, Anqing City, CN |
| Contact | Bingyi Xiang |
| 510(k) history | 12 submissions · 12 cleared · 2019-2024 |

REGULATORY CONSULTANT

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|-----------------|--|
| Consulting firm | Shanghai Mind-Link Consulting Co., Ltd. |
| Contact | Evan Hu |

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.netDevice record: <https://www.510kdatabase.net/k220083/> Data sourced from FDA 510(k) public records (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. - Generated May 26, 2026