

**K243032 EHL Probe (SCDG-AS)**Mar 21, 2025  
175 days to decisionK243032 · Product code: **FFK** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k243032/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotripter, Electro-hydraulic (FFK)
Date received	Sep 27, 2024
Decision date	Mar 21, 2025
Days to decision	175 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	EHL Probe (SCDG-BS); EHL Probe (SCDG-CS)

**APPLICANT**

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Company	<b>Guangzhou Potent Medical Equipment Joint-Stock Co., Ltd.</b>
Location	Guangzhou, CN
Contact	Zhengzhou Li
510(k) history	7 submissions · 7 cleared · 2025-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k243032/> 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 14, 2026