

**K213222 Disposable Polypectomy Snare**Jun 6, 2022  
250 days to decisionK213222 · Product code: **FDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k213222/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Snare, Flexible (FDI)
Date received	Sep 29, 2021
Decision date	Jun 6, 2022
Days to decision	250 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Beijing Zksk Technology Co., Ltd.</b>
Location	Beijing, CN
Contact	Ma Li
510(k) history	6 submissions · 6 cleared · 2022-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Shanghai Truthful Information Technology Co., Ltd.</b>
Contact	Boyle Wang

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

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