FDA 510(k) Premarket Notification

K213223 Multi-Band Ligator

CLEARED

Jun 6, 2022
250 days to decision

K213223 · Product code: **FHN** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k213223/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ligator, Hemorrhoidal (FHN)
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

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

REGULATORY CONSULTANT

Consulting firm	**Shanghai Truthful Information Technology Co., Ltd.**
Contact	Boyle Wang

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