

K212407 Gastric Lightguide (GLG)Jul 26, 2022
357 days to decisionK212407 · Product code: **KNT** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k212407/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Aug 3, 2021
Decision date	Jul 26, 2022
Days to decision	357 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Naser Dib Gabinet Lekarki Nasmed
Location	Warszawa, PL
Contact	Naser Dib
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	Aztech Regulatory & Quality, LLC
Contact	Joseph Azary

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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