FDA 510(k) Premarket Notification

K072410 KSEA CLEARVISION II, MODEL 40334120

CLEARED

May 9, 2008
256 days to decision

K072410 · Product code: **GWG** · Neurology
Source: https://www.510kdatabase.net/k072410/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscope, Neurological (GWG)
Date received	Aug 27, 2007
Decision date	May 9, 2008
Days to decision	256 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**KARL STORZ Endoscopy-America, Inc.**
Location	Mchenry, IL, US
Contact	CRYSTAL DIZOL
510(k) history	361 submissions · 361 cleared · 1980-2024