

K232759 Viewer (5.4)May 21, 2024
256 days to decisionK232759 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k232759/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Sep 8, 2023
Decision date	May 21, 2024
Days to decision	256 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Elements Viewer; Mixed Reality Viewer; Smart Layout; Elements Viewer Smart Layout

APPLICANT

Company	Brainlab AG
Location	Heimstetten, DE
Contact	Sadwini Suresh
Website	http://brainlab.com
510(k) history	135 submissions · 135 cleared · 1996-2025

Brainlab AG is a digital medical technology company specializing in image-guided surgery and neurology devices. The company operates with a manufacturing facility in Heimstetten, Germany, and develops innovative surgical navigation and alignment systems for cranial, spine, and neurosurgical procedures. Brainlab has received FDA 510(k) clearances from total submissions since its first clearance in 1996. The company maintains a strong regulatory presence in neurology devices, with recent clearances spanning surgical navigation systems, robotic alignment platforms, trajector...