

K181871 EliA Celikey IgG ImmunoassayMar 1, 2019
232 days to decisionK181871 · Product code: **MVM** · Immunology
Source: <https://www.510kdatabase.net/k181871/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Autoantibodies, Endomysial(tissue Transglutaminase) (MVM)
Date received	Jul 12, 2018
Decision date	Mar 1, 2019
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary
Other names	EliA GliadinDP IgA Immunoassay; EliA GliadinDP IgG Immunoassay

APPLICANT

Company	Phadia AB
Location	Uppsala, SE
Contact	Carina Magnusson
Website	http://www.phadia.com
510(k) history	32 submissions · 32 cleared · 2007-2022

Phadia AB is a medical products company headquartered in Uppsala, Sweden. The company develops, manufactures, and markets blood test systems for clinical diagnosis and monitoring of allergy, asthma, and autoimmune diseases. Phadia AB received FDA 510(k) clearances from total submissions, with no denied submissions on record. The company's regulatory focus is entirely on Immunology devices. Clearances span from 2007 to 2022, establishing a consistent track record in immunoassay and allergen testing technologies. The company's cleared devices include immunoassay systems for...