VIAMED

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Letter of Authorization for Authorized Representatives

Medical Device Control Division
Thai Food and Drug Administration
Ministry of Public Health

6th May 2025

Dear Sir/Madam,

Subject: Letter of Authorization for Intega (Thailand) Co., Ltd.

We, Viamed Ltd., as the Product Owner, hereby authorize Intega (Thailand) Co., Ltd., as the Registrant to prepare and submit applications for the evaluation and registration of medical devices to the Thai Food and Drug Administration on our behalf.

This authorization shall apply to the following medical devices:

Product's Code	Name	Physical Manufacturer
8090121313V	Gas Sampling Line H	bluepoint medical GmbH&Co.KG

We also authorize Intega (Thailand) Co., Ltd. to make declarations and to submit documents on our behalf, regarding the above medical devices, in support of this application. These declarations and submissions are made pursuant to the requirements of the Medical Device Act B.E. 2551 (2008), the amended Medical Device Act B.E. 2562 (2nd edition) and any other applicable laws that may also be in force.

This authorization shall remain in effect until our notification to the Thai Food and Drug Administration in writing that the authorization is revoked.

We undertake to provide post-market support and assistance to the Registrant as may be required in relation to any matter involving the above medical devices.

We acknowledge that any non-compliance with any registration condition issued by the Thai Food and Drug Administration in relation to medical devices registered with the Medical Device Control Division may result in the suspension or cancellation of the medical device registration.

We agree to assist the Thai Food and Drug Administration with any request for information on the above medical devices.

Yours Sincerely,

Derek Lamb
Managing Director

VIAMED Company registered in England Company Registration Number 01291765