





## **EU Declaration of Conformity**

## **KLEEYA® TRIGGER SOLUTIONS PACK**

solutions for the chemiluminescence signal generation in the fully automated chemiluminescence immunoassay analyzer KleeYa

Product Code/ Article Number:	226491-TRP
Basic UDI-DI:	5999860094KTSTJ
Classification:	☑ A □ B □ C □ D   Rule 5, Annex VIII, IVD Regulation
Conformity Route	<ul> <li>□ Annex IX Technical documentation Examination</li> <li>□ ANNEX IX Full Quality System</li> <li>□ ANNEX XI Production Quality System</li> <li>⋈ ANNEX I &amp; II + III</li> </ul>
Legal manufacturer:	Diatron MI Plc. H-1097 Táblás u. 39, Budapest, Hungary
SRN:	HU-MF-000020416

We, as the manufacturer of the device specified above, declare under our sole responsibility that the product meets all applicable requirements of the following regulation and directive(s):

Legal Requirement	Title
IVD Regulation (IVDR) (EU) 2017/746	REGULATION (EU) 2017/746 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on in vitro diagnostic medical devices and repealing Directive 98/79/EC and Commission Decision 2010/227/EU

Conformity assessment was successfully conducted according to the stipulations of the respective regulation / directive(s).

The application of the Trigger solutions for in vitro diagnostic purposes requires a separate conformity assessment according to Regulation (EU) 2017/746 for the full system into which it will be incorporated and / or is used in combination with (e.g. assay).



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This Declaration of Conformity is valid for the device's configuration and the regulatory requirements effectual at the date the Declaration was issued. Changes affecting the device, and / or the applicable regulations trigger a review of the conformity assessment the Declaration is based on, and the issuance of a new version of the document.

Diatron MI Zrt.

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2024-01-17 Date:

**Budapest** Place:

Signature:

Péter Zsóka

Director of finance