

CERTIFICATE

DIRECTIVE 98/79/EC
FULL QUALITY ASSURANCE SYSTEM

CeCert Sp. z o.o. hereby confirms that manufactured by

Tianjin Galenus
Medical Co., Ltd.

Room 403, 4th Floor, Building C02,
Pioneering Headquarters Base,
Wuqing District, Tianjin,
P.R.China, 301700

in vitro diagnostic medical device for

HBsAg/HCV/HIV/Syphilis combo Rapid Test
(Whole Blood/Serum/Plasma)

catalogue number: GL094

in terms of full quality assurance system, comply with requirements of Annex IV
to Directive 98/79/EC (as amended), as evidenced by the
assessment conducted by CeCert Sp. z o.o.

CE **2934**

Validity date: 06.05.2022 – 26.05.2025

Issue date: 06.05.2022

Check it



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