

# EC Declaration of Conformity

according to Directive 98/79/EC, on in vitro diagnostic medical devices

Ref. No.:20220513-A07

**Manufacturer**  
(Name, Address)

**Getein Biotech, Inc.**  
No. 9 Bofu Road, Luhe District, Nanjing, 211505, China

**Authorized Representative**  
(Name, Address)

**CMC Medical Devices & Drugs S.L.**  
Add: C/ Horacio Lengo N° 18, CP 29006, Málaga, Spain

**Medical device**

| No. | Product Name  |
|-----|---|
| 1   | Getein 1600 Immunofluorescence Quantitative Analyzer        |
| 2   | Cardiac Troponin I Fast Test Kit (Immunofluorescence Assay) |
| 3   | NT-proBNP Fast Test Kit (Immunofluorescence Assay)          |
| 4   | hs-CRP+CRP Fast Test Kit (Immunofluorescence Assay)         |
| 5   | NT-proBNP/cTnI Fast Test Kit (Immunofluorescence Assay)     |
| 6   | CK-MB/cTnI/Myo Fast Test Kit (Immunofluorescence Assay)     |
| 7   | D-Dimer Fast Test Kit (Immunofluorescence Assay)            |
| 8   | PCT Fast Test Kit (Immunofluorescence Assay)                |
| 9   | CysC Fast Test Kit (Immunofluorescence Assay)               |
| 10  | mAlb Fast Test Kit (Immunofluorescence Assay)               |
| 11  | NGAL Fast Test Kit (Immunofluorescence Assay)               |
| 12  | $\beta$ 2-MG Fast Test Kit (Immunofluorescence Assay)       |
| 13  | CK-MB/cTnI Fast Test Kit (Immunofluorescence Assay)         |
| 14  | HCG+ $\beta$ Fast Test Kit (Immunofluorescence Assay)       |
| 15  | H-FABP Fast Test Kit (Immunofluorescence Assay)             |
| 16  | PCT/CRP Fast Test Kit (Immunofluorescence Assay)            |
| 17  | CK-MB/cTnI/H-FABP Fast Test Kit (Immunofluorescence Assay)  |
| 18  | HbA1c Fast Test Kit (Immunofluorescence Assay)              |
| 19  | NT-proBNP/NGAL Fast Test Kit (Immunofluorescence Assay)     |
| 20  | CK-MB Fast Test Kit (Immunofluorescence Assay)              |
| 21  | hs-cTnI Fast Test Kit (Immunofluorescence Assay)            |
| 22  | T3 Fast Test Kit (Immunofluorescence Assay)                 |
| 23  | T4 Fast Test Kit (Immunofluorescence Assay)                 |
| 24  | TSH Fast Test Kit (Immunofluorescence Assay)                |
| 25  | Scr Fast Test Kit (Immunofluorescence Assay)                |
| 26  | PLGF Fast Test Kit (Immunofluorescence Assay)               |

- 27 HCY Fast Test Kit (Immunofluorescence Assay)
- 28 Anti-CCP Fast Test Kit (Immunofluorescence Assay)
- 29 25-OH-VD Fast Test Kit (Immunofluorescence Assay)
- 30 Lp-PLA2 Fast Test Kit (Immunofluorescence Assay)
- 31 FOB Fast Test Kit (Immunofluorescence Assay)
- 32 SAA Fast Test Kit (Immunofluorescence Assay)
- 33 H. pylori Fast Test Kit (Immunofluorescence Assay)
- 34 PRL Fast Test Kit (Immunofluorescence Assay)
- 35 Transferrin Fast Test Kit (Immunofluorescence Assay)
- 36 Insulin Fast Test Kit (Immunofluorescence Assay)
- 37 PG I /PG II Fast Test Kit (Immunofluorescence Assay)
- 38 LH Fast Test Kit (Immunofluorescence Assay)
- 39 FSH Fast Test Kit (Immunofluorescence Assay)
- 40 Anti-TP Fast Test Kit (Immunofluorescence Assay)
- 41 AFP/CEA Fast Test Kit (Immunofluorescence Assay)
- 42 AMH Fast Test Kit (Immunofluorescence Assay)
- 43 fT3 Fast Test Kit (Immunofluorescence Assay)
- 44 fT4 Fast Test Kit (Immunofluorescence Assay)
- 45 Total IgE Fast Test Kit (Immunofluorescence Assay)
- 46 Vit-B12 Fast Test Kit (Immunofluorescence Assay)
- 47 Prog Fast Test Kit (Immunofluorescence Assay)
- 48 Testosterone Fast Test Kit (Immunofluorescence Assay)
- 49 E2 Fast Test Kit (Immunofluorescence Assay)
- 50 RF Fast Test Kit (Immunofluorescence Assay)
- 51 ASO Fast Test Kit (Immunofluorescence Assay)
- 52 Ferritin Fast Test Kit (Immunofluorescence Assay)
- 53 ST2 Fast Test Kit (Immunofluorescence Assay)
- 54 CA125 Fast Test Kit (Immunofluorescence Assay)
- 55 CA19-9 Fast Test Kit (Immunofluorescence Assay)
- 56 CA15-3 Fast Test Kit (Immunofluorescence Assay)
- 57 RSV/Influenza A/B Fast Test Kit (Immunofluorescence Assay)
- 58 Influenza A/B Fast Test Kit (Immunofluorescence Assay)
- 59 RSV Fast Test Kit (Immunofluorescence Assay)
- 60 IL-6 Fast Test Kit (Immunofluorescence Assay)
- 61 BNP Fast Test Kit (Immunofluorescence Assay)
- 62 SAA/CRP Fast Test Kit (Immunofluorescence Assay)
- 63 Folate acid Fast Test Kit (Immunofluorescence Assay)
- 64 hs-CRP Fast Test Kit (Immunofluorescence Assay)
- 65 TnT Fast Test Kit (Immunofluorescence Assay)
- 66 PCT/IL-6 Fast Test Kit (Immunofluorescence Assay)



- 67 HBP Fast Test Kit (Immunofluorescence Assay)
- 68 S100-β Fast Test Kit (Immunofluorescence Assay)
- 69 CK-MB/hs-cTnI/Myo Fast Test Kit (Immunofluorescence Assay)
- 70 Cortisol Fast Test Kit (Immunofluorescence Assay)
- 71 CEA Fast Test Kit (Immunofluorescence Assay)
- 72 AFP/CEA Fast Test Kit (Immunofluorescence Assay)

**Classification** Other device (according to Annex II of the directive 98/79/EC)

**Conformity assessment route** Annex III of the 98/79/EC

|                     |                     |                     |                     |
|---------------------|---------------------|---------------------|---------------------|
| <b>Applicable</b>   | EN 13612:2002       | EN ISO 14971:2019   | EN ISO15223-1:2016  |
| <b>coordination</b> | EN ISO 18113-1:2011 | EN ISO 18113-2:2011 | EN ISO 18113-3:2011 |
| <b>standards</b>    | EN ISO 23640:2015   | EN ISO 13485:2016   | ISO 780:2015        |
|                     | EN 61326-2-6:2006   | IEC 61326-1:2013    |                     |
|                     | EN 61010-2-101:2002 | IEC 61010-1:2010    |                     |

Signatory representative declares herein the above-mentioned device meets the basic requirements of the European Parliament and the Council's in vitro diagnostic medical devices directive: 98/79/EC Annex I.

This declaration of conformity is based on European Parliament and the Council's 98/79/EC directive Annex III. The compiled technical file and quality system document according to 98/79/EC directive Annex III are testified and the quality system certificate has issued by BSI Group The Netherlands B. V. The manufacturer is exclusively responsible for the declaration of conformity.

**General Manager** Enben Su

Nanjing

13<sup>th</sup>, May, 2022

(place and date of issue)

(name and signature or equivalent marking of authorized person)

