

Add: No.9 Bofu Road, Luhe District, Nanjing, 211505, China Tel: 86-25-68568508 Email: overseas@geteincom.cn Web: www.bio-GP.com.cn

Document No.: GP-GMSQ-2022-110

Letter of Authorization

To whom it may concern,

We, Getein Biotech, Inc. (No.9 BoFu Road, Luhe District, Nanjing, 211505, China), hereby authorize Sanmedico SRL. as our official distributor for registering, promoting, selling, distributing, taking part in tenders, maintaining & after sale technical services of under-mentioned product in the territory of Moldova:

Sanmedico SRL will comply with the laws and regulations of the countries and regions where they are located in and where they are selling mentioned product to, otherwise, the risks and losses arising therefrom shall be undertaken by Sanmedico SRL

This authorization starts from Jan 1, 2022 and will be valid to December 31 2023

Getein Biotech, Inc. has the right to terminate the authorization before validity and will inform Sanmedico SRL with 10 days in advance.

Getein Biotech, Inc.

Name: Steven Zhou

Position: Overseas Sales Director

基蛋生物科技股份有限公司 GETEIN BIOTECH, INC.

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CE acco	Declaration of Conformity rding to Directive 98/79/EC, on in vitro diagnostic medical devices
Maker (Name, Address)	Getein Biotech, Inc. No. 9 Bofu Road, Luhe District, Nanjing, 211505, China
Authorized Representative (Name, Address)	Lotus Global Co., Ltd 1 Four Seasons Terrace West Draylon, Middlesex London, UB7 9GG United Kingdom
Medical device	Getein 100 Immunofluorescence Quantitative Analyzer Cardiac Troponin I Fast Test Kit (Immunofluorescence Assay) NT-prosNP Fast Test Kit (Immunofluorescence Assay) NS-CRP+CRP Fast Test Kit (Immunofluorescence Assay) NT-prosNP Fast Test Kit (Immunofluorescence Assay) NT-prosNP Fast Test Kit (Immunofluorescence Assay) D-Dimer Fast Test Kit (Immunofluorescence Assay) D-Dimer Fast Test Kit (Immunofluorescence Assay) PCT Fast Test Kit (Immunofluorescence Assay) MAIb Fast Test Kit (Immunofluorescence Assay) CysC Fast Test Kit (Immunofluorescence Assay) CysC Fast Test Kit (Immunofluorescence Assay) HCG+β Fast Test Kit (Immunofluorescence Assay) TSH Fast Test Kit (Immunofluorescence Assay) TSH Fast Test Kit (Immunofluorescence Assay) PCT/CRP Fast Test Kit (Immunofluorescence Assay) PCT/CRP Fast Test Kit (Immunofluorescence Assay) H-FABP Fast Test Kit (Immunofluorescence Assay)
	Classification of products according to directive : Others
Applicable coordination standards:	Batch/serial No. Type, production term (if applicable) : EN ISO 14971:2012 EN ISO 23640:2015 EN ISO 13485:2016 EN 980:2008 EN 13612:2002 EN ISO15223-1:2012 EN 1041:2008 EN ISO 18113-1:2011 EN ISO 18113-2:2011 EN ISO 18113-3:2011 EN-IEC 61326-1:2013 EN-IEC 61010-1:2010 IEC 61010-2-101:2015 EN-IEC 61326-2-2:2013
European Parliam This declaration of Annex III. The com III are testified and	ntative declares herein the above mentioned device meets the basic requirements of the ent and the Council's in vitro diagnostic medical devices directive: 98/79/EC Annex III. conformity is based on European Parliament and the Council's 98/79/EC directive apiled technical file and quality system document according to 98/79/EC directive Annex I the quality system certificate has issued by TÜV Rheinland (Shanghai) Co., Ltd.
Manager: MomjM, 3 (place and date	rd, Jan, 2017



Declaration of Conformity

C

Maker (Name, Address) Getein Biotech, Inc. No. 9 Bofu Road, Luhe District, Nanjing, 211505, China
Authorized Representative (Name, Address) Lotus NL B.V. Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague, Netherlands. FIA8000 Quantitative Immunoassay Analyzer FIA8600 Quantitative Immunoassay Analyzer Cardiac Troponin I Fast Test Kit One Step Test for cTnl (Colloidal Gold) cTnl Rapid Test (Colloidal Gold Gold Assay) One Step Test for NT-proBNP (Colloidal Gold) One Step Test for NT-proBNP/cTnl (Colloidal Gold) One Step Test for NT-proBNP/cTnl (Colloidal Gold) One Step Test for D-Dimer (Colloidal Gold) One Step Test for D-Dimer (Colloidal Gold) One Step Test for PCT (Colloidal Gold) One Step Test for NGAL (Colloidal Gold) One Step Test for NGAL (Colloidal Gold) One Step Test for HbA1c (Colloidal Gold) One Step Test for PCT/CRP (Colloidal Gold) One Step Test for CK-MB/cTnl/H-FABP (Colloidal Gold) One Step Test for CK-MB/cTnl/Colloidal Gold) One Step Test for CK-MB/cTnl (Colloidal Gold) One Step Test for CK-MB/cTnl (Colloidal Gold) One Step Test for CK-MB/cTnl (Colloidal Gold)
Representative (Name, Address) Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague, Netherlands.
FIA8600 Quantitative Immunoassay Analyzer Cardiac Troponin I Fast Test Kit One Step Test for cTnl (Colloidal Gold) cTnl Rapid Test (Colloidal Gold Assay) One Step Test for NT-proBNP (Colloidal Gold) One Step Test for NT-proBNP/cTnl (Colloidal Gold) One Step Test for CK-MB/cTnl/Myo (Colloidal Gold) One Step Test for b-Dimer (Colloidal Gold) One Step Test for D-Dimer (Colloidal Gold) One Step Test for PCT (Colloidal Gold) One Step Test for β2-MG (Colloidal Gold) One Step Test for mAlb (Colloidal Gold) One Step Test for NGAL (Colloidal Gold) One Step Test for HCG+β (Colloidal Gold) One Step Test for HCG+β (Colloidal Gold) One Step Test for HCG+β (Colloidal Gold) One Step Test for PCT/CRP (Colloidal Gold) One Step Test for PCT/CRP (Colloidal Gold) One Step Test for CK-MB/cTnl/H-FABP (Colloidal Gold) One Step Test for CK-MB/cTnl/H-FABP (Colloidal Gold) One Step Test for CK-MB/cTnl (Colloidal Gold)
One Step Test for T4/T3 (Colloidal Gold) One Step Test for T3 (Colloidal Gold) One Step Test for T4 (Colloidal Gold) One Step Test for 25-OH-VD (Colloidal Gold) One Step Test for FOB (Colloidal Gold) One Step Test for FOB (Colloidal Gold) One Step Test for H. pylori (Colloidal Gold) One Step Test for SAA (Colloidal Gold) Getein1100 Immunofluorescence Quantitative Analyzer Getein1600 Immunofluorescence Quantitative Analyzer Getein1180 Immunofluorescence Quantitative Analyzer

D-Dimer Fast Test Kit (Immunofluorescence Assay)

PCT Fast Test Kit (Immunofluorescence Assay) β2-MG Fast Test Kit (Immunofluorescence Assay) mAlb Fast Test Kit (Immunofluorescence Assay) NGAL Fast Test Kit (Immunofluorescence Assay) CysC Fast Test Kit (Immunofluorescence Assay) CK-MB Fast Test Kit (Immunofluorescence Assay) CK-MB/cTnl Fast Test Kit (Immunofluorescence Assay) HCG+β Fast Test Kit (Immunofluorescence Assay) HbA1c Fast Test Kit (Immunofluorescence Assay) PCT/CRP Fast Test Kit (Immunofluorescence Assay) CK-MB/cTnl/H-FABP Fast Test Kit (Immunofluorescence Assay) H-FABP Fast Test Kit (Immunofluorescence Assay) 25-OH-VD Fast Test Kit (Immunofluorescence Assay) TSH Fast Test Kit (Immunofluorescence Assay) T3 Fast Test Kit (Immunofluorescence Assay) T4 Fast Test Kit (Immunofluorescence Assay 25-OH-VD Fast Test Kit (Immunofluorescence Assay) FOB Fast Test Kit (Immunofluorescence Assay) H. pylori Fast Test Kit (Immunofluorescence Assay) SAA Fast Test Kit (Immunofluorescence Assay) LH Fast Test Kit (Immunofluorescence Assay) FSH Fast Test Kit (Immunofluorescence Assay) AMH Fast Test Kit (Immunofluorescence Assay) PRL Fast Test Kit (Immunofluorescence Assay) **CK-MB Control** cTnl Control Myo Control NT-proBNP Control **D-Dimer Control CRP Control PCT Control** β2-MG Control mAlb Control NGAL Control CysC Control H-FABP Control HbA1c Control HCG+B Control CK-MB/cTnl/Myo Control CK-MB/cTnl Control NT-proBNP/cTnl Control **TSH Control** T4/T3 Control T3 Control T4 Control Others Classification of products according to directive Batch/serial No. Type, production term (if applicable)

Applicable	EN ISO 14971:2012 EN 13612:2002	EN ISO 23640:2015 EN ISO15223-1:2012	EN ISO 13485:2016 EN ISO 18113-2:2011
coordination	EN 1041:2008	EN ISO 18113-1:2011	EN ISO 18113-3:2011
standards:	IEC 61010-1:2010 IEC 61326-1:2013	IEC 61010-2-081:2015 IEC 61326-2-2:2013	IEC 61010-2-101:2015

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 III. 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 TÜV Rheinland (Shanghai) Co., Ltd.

General Manager: Enben Su

(place and date of issue)

(name and signature or equivalent

marking of authorized person)



EC Declaration of Conformity

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

Ref. No.:20220513-A05

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 Nº 18, CP 29006, Málaga, Spain

	No.	Product Name			
	1	Getein 1100 Immunofluorescence Quantitative Analyzer			
	// 2	Cardiac Troponin I Fast Test Kit (Immunofluorescence Assay)			
4 1175 3	//3	NT-proBNP Fast Test Kit (Immunofluorescence Assay)			
H 1/// 78%	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)			
Medical device	12	β2-MG Fast Test Kit (Immunofluorescence Assay)			
wiedical device	13	CK-MB/cTnI Fast Test Kit (Immunofluorescence Assay)			
	14	HCG+β 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)			



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)	
	%
69 CK-MB/hs-cTnI/Myo Fast Test Kit (Immunofluorescence Assay)	
	The second of th
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	
Applicable EN 13612:2002 EN ISO 14971:2019 EN ISO15223-	1:2016
coordination EN ISO 18113-1:2011 EN ISO 18113-2:2011 EN ISO 18113-	-3:2011
standards 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	
Nowsing 13th, May, 2022 (place and date of issue) (pame and signature or equivalent	
(place and date of issue) (name and signature or equivalent	2
marking of authorized person)	
	1 1
marking of authorized person)	





Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that: Getein Biotech, Inc.

No.9 Bofu Road Luhe District Nanjing Jiangsu 211505 China 基蛋生物科技股份有限公司

中国 江苏省 南京市 六合区

沿江工业开发区 博富路9号 邮编: 211505

Holds Certificate No: MD 728432

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 for the following scope:

Design & Development, Manufacture and Distribution of Chemiluminescence Immunoassay, Biochemistry Assay, Point of Care Assay (including Colloidal Gold Assay, Immunofluorescence Assay, Dry Chemistry Assay). Design & Development, Manufacture and Distribution of Analyzers in use of Chemiluminescence Immunoassay, Biochemistry Assay, Point of Care Assay (including Colloidal Gold Assay, Immunofluorescence Assay, Dry Chemistry Assay). 研发,生产和销售化学发光法试剂,生化试剂,即时诊断(包括胶体金法,免疫荧光法,干式化

学法)试剂。 研发,生产和销售用于化学发光法试剂,生化试剂,即时诊断(包括胶体金法,免疫荧光法, 干式化学法)试剂配套使用的分析仪。

For and on behalf of BSI:

Gary E Slack, Senior Vice President - Medical Devices

jany C Stade

Original Registration Date: 2020-05-29 Effective Date: 2020-07-26 Latest Revision Date: 2020-07-22 Expiry Date: 2023-07-25

Page: 1 of 1

bsi.



...making excellence a habit."

This certificate was issued electronically and remains the property of BSI and is bound by the conditions of contract. An electronic certificate can be authenticated <u>online</u>.

Printed copies can be validated at www.bsi-global.com/ClientDirectory or telephone +86 10 8507 3000.

Reference Code: GP-DT-018-07-19 Issued by 07/26/2019

CERTIFICATE

Getein Biotech

hereby certifies

Mr. Vitalie Goreacii

from Sanmedico SRL.

Completion of Getein Products Technical and Operational Training & Qualification of After-sales Service

基蛋生物科技股份有限公司 GETEIN BIOTECH, INC.









HIGHLY EFFICIENT & ACCURATE

Advanced fluorescence immunoassay

Multiple quality control



REAL-TIME AND RAPID TEST

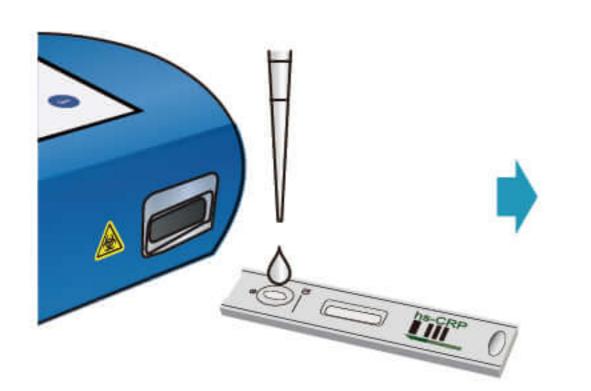
One-step test

3-15 min/test

5 sec/test for multiple tests

OPERATION MODES

Inside Mode (single sample rapid test mode)



Sample Dispense



Test Card Insert

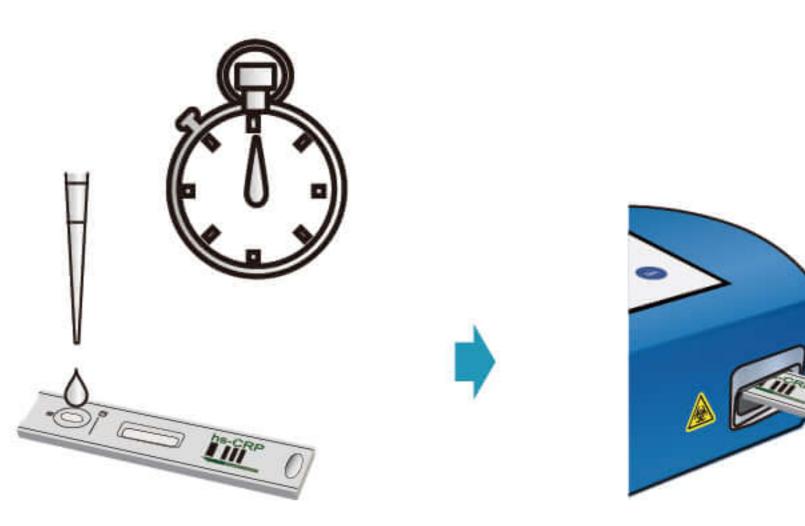


Click "Start" Icon



Result Show and Print

Quick mode (mass samples rapid test mode)



Sample Dispense



Timing the Reaction Manually



Click "Start" Icon



Result Show and Print





CONVENIENT OPERATION

RFID card calibration

Keyboard and mouse connectivity through USB port

Handwriting input available

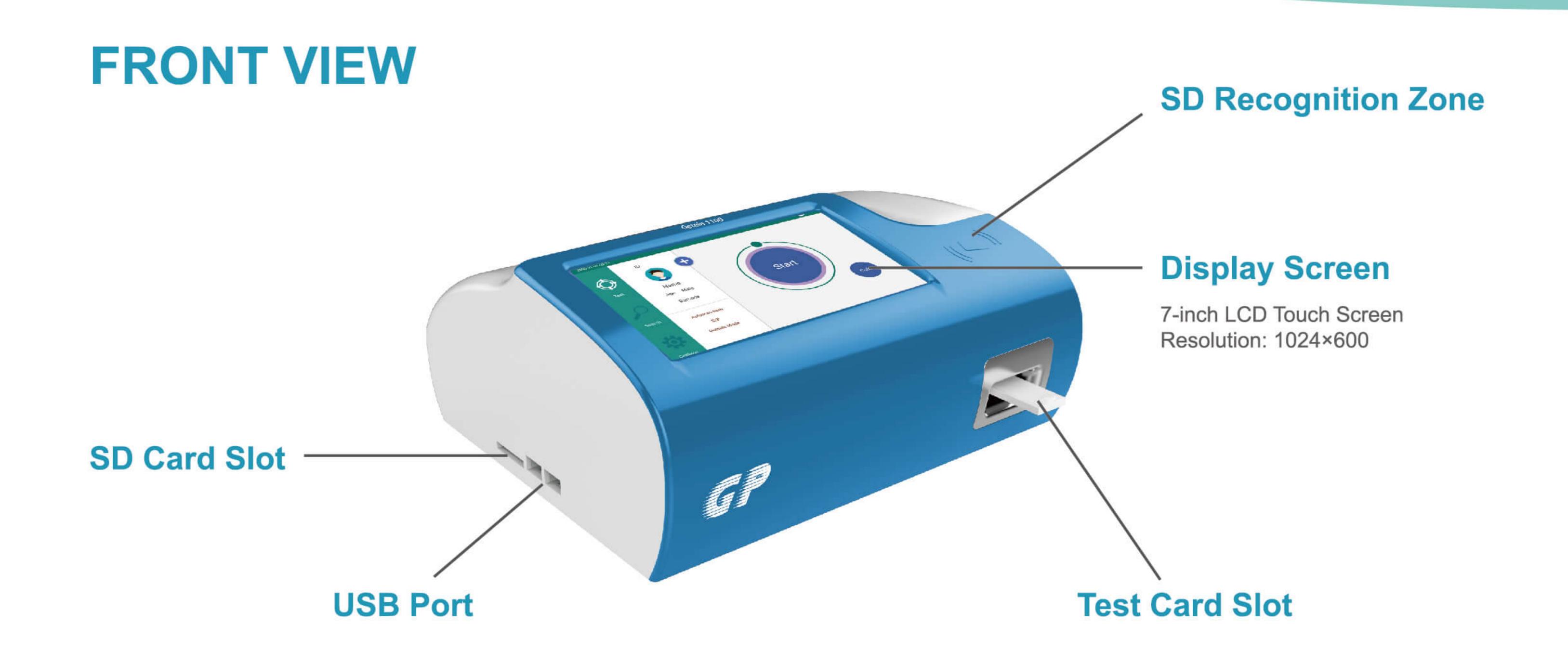
Continuous test for 3 hours with lithium battery



USER-FRIENDLY INTERFACE

Android system

7 inch touch screen







PORTABLE DESIGN

Small in size: 261 ×241 ×115 mm

Light in weight: 2.0 kg



LARGE MEMORY

Up to 10,000 results storage capacity

TECHNICAL PARAMETERS

Methodology

Immunofluorescence

Result

Quantitative

Sample Type

WB, plasma, serum, urine, Stool, Nasal swab, Saliva, Capillary blood

Storage Capacity

10000 data

Language

English/Chinese/Spanish/Portuguese

Screen

7 inch touch screen

Power Supply

100-240 V~50 Hz/60 Hz, 60 VA

Working Environment

Tempreture: 10-35 °C
Relative humidity ≤70%
Air pressure 70.0~106.0 kpa

Dimension

261 mm×241 mm×115 mm (D×W×H)

Weight

2.0 kg

TEST ITEMS

Cat.#	TEST ITEMS	DISEASES	CUT-OFF VALUE	SAMPLE TYPES	MEASURING RANGE	SAMPLE VOLUME	REACTION TIME	QUALIFIC	CATIOI
Cardia	ac Markers								
IF1001	cTnI	Myocardial infarction	0.1 ng/mL	S/P/WB	0.1-50.0 ng/mL	100 µL	10 min	NMPA	C€
IF1089	BNP	Heart failure	100.0 pg/mL	P/WB	5.0-5000.0 pg/mL	100 µL	10 min	NMPA	CE
IF1002	NT-proBNP	Heart failure	300 pg/mL	S/P/WB	100-35000 pg/mL	100 µL	10 min	NMPA	CE
IF1005	CK-MB/cTnl/Myo	Myocardial damage /infarction	CK-MB: 5.0 ng/mL cTnl: 0.1 ng/mL Myo: 70 ng/mL	S/P/WB	2.5-80.0 ng/mL 0.1-50.0 ng/mL 30.0-600.0 ng/mL	100 µL	10 min	NMPA	CE
IF1012	CK-MB/cTnI	Myocardial damage /infarction	CK-MB: 5.0 ng/mL cTnl: 0.1 ng/mL	S/P/WB	2.5-80.0 ng/mL 0.1-50.0 ng/mL	100 µL	10 min	C	ξ
IF1014	H-FABP	Myocardial damage	6.36 ng/mL	S/P/WB	1.0-120.0 ng/mL	100 µL	3 min	NMPA	C€
F1016	CK-MB/cTnI/H-FABP	Myocardial damage /infarction	CK-MB: 5.0 ng/mL cTnl: 0.1 ng/mL H-FABP: 6.36 ng/m	S/P/WB L	2.5-80.0 ng/mL 0.1-50.0 ng/mL 2.0-100.0 ng/mL	100 µL	10 min	NMPA	CE
F1018	CK-MB	Myocardial injury	5.0 ng/mL	S/P/WB	2.5-80.0 ng/mL	100 µL	10 min	C	ξ
F1078	ST2	Heart failure	35.0 ng/mL	S/P/WB	3.1-200.0 ng/mL	100 µL	10 min	C	ξ
Coag	ulation Markers								
F1006	D-Dimer	Venous thromboembolism	0.5 mg/L	P/WB	0.1-10.0 mg/L	100 µL	10 min	NMPA	CE
Inflam	nmation								
F1003	hs-CRP+CRP	Cardiovascular inflammation /normal inflammation	3 mg/L 10 mg/L	S/P/WB/ Fingertip blood	0.5-200 mg/L	10 µL	3 min	NMPA	CΕ
IF1007	PCT	Sepsis, bacterial infection	0.1 ng/mL	S/P/WB	0.1-50.0 ng/mL	100 µL	15 min	NMPA	CE
F1015	PCT/CRP	Sepsis, bacterial infection	PCT: 0.1 ng/mL CRP: 3.0 mg/L	S/P/WB	0.1-50.0 ng/mL 0.5-200.0 mg/L	100 µL	15 min	NMPA	CΕ
F1044	SAA	Bacterial/Virus infection	10.0 mg/L	S/P/WB/ Fingertip blood	5.0-200.0 mg/L	10 µL	5 min	NMPA	CE
F1090	SAA/CRP	Neonatal sepsis, Bacterial/virus infection	SAA: 10.0 mg/L CRP: 10.0 mg/L	S/P/WB/ Capillary blood	5.0-200.0 mg/L 0.5-200.0 mg/L	10 µL	5 min	NMPA	C€
F1088	IL-6	Acute inflammation	7.0 pg/mL	S/P/WB/ Peripheral blood	1.5-4000.0 pg/mL	100 µL	15 min	(ξ
Renal	Function								
F1008	CysC	Acute and chronic renal diseases	0.51-1.09 mg/L	S/P/WB	0.5-10.0 mg/L	10 µL	3 min	NMPA	CE
F1009	mAlb	Diabetic nephropathy, hypertensive nephropathy	20.0 mg/L	Urine	10.0-200.0 mg/L	100 µL	3 min	NMPA	C€
F1010	NGAL	Acute kidney injury	Serum: 200 ng/mL Urine: 100 ng/mL	S/Urine	50-5000 ng/mL	10 µL	10 min	NMPA	CE
IF1011	β ₂ -MG	Acute and chronic kidney diseases/tumours	0.8-3.0 mg/L	S/P/WB	0.5-20.0 mg/L	10 µL	3 min	NMPA	CE
Diabe	tes Mellitus								
F1017	HbA1c	Diabetes mellitus	3.8%-5.8%	WB	2%-14%	10 µL	5 min	NGSP	NMP/
Metab	oolic Marker								
F1031	25-OH-VD	Osteomalacia, osteoporosis	30.0-50.0 ng/mL	S/P	8.0-70.0 ng/mL	40 µL	15 min	NMPA	CE
Thyro	id Function								
F1024	TSH	Thyroid malfunction	0.27-4.20 µIU/mL	S/P	0.10-50.00 μIU/mL	100 µL	15 min	NMPA	CE
F1022	Т3	Hyperthyroidism, hypothyroidism	1.30-3.10 nmol/L	S/P	0.30-10.00 nmol/L	40 µL	15 min	C	ξ
F1023	T4	Hyperthyroidism, hypothyroidism	59.0-154.0 nmol/L	S/P	5.4-320.0 nmol/L	40 µL	15 min	(ξ
F1067	fT3	Hyperthyroidism, hypothyroidism	3.1-6.8 pmol/L	S/P	0.4-50.0 pmol/L	100 µL	15 min	C	ξ
	fT4	Hyperthyroidism,	12.0-22.0 pmol/L	S/P	0.3-100.0 pmol/L	100	15 min		

Cat.#	TEST ITEMS	DISEASES	CUT-OFF VALUE	SAMPLE TYPES		SAMPLE VOLUME	REACTION TIME	QUALIFICATION
Repro	duction/Fertility							
IF1013	HCG+β	Fertility	5.1 mIU/mL	S/P	5-100000 mIU/mL	100 µL	10 min	NMPA CE
IF1055	LH	Homeostasis fertility regualtion	Refer to User Manual	S/P	0.2-150.0 mIU/mL	100 µL	15 min	CE
IF1056	FSH	PCOS, infertility evaluation and pituitary disorders	Refer to User Manual	S/P	0.2-150.0 mIU/mL	100 µL	15 min	CE
IF1066	AMH	Fertility, PCOS, gonadal function, precocious/late puberty	Refer to User Manual	S/P	0.10-20.00 ng/mL	200 µL	15 min	CE
IF1048	PRL	Infertility, gonadal disorders	Refer to User Manual	S/P	0.50-200.0 ng/mL	100 µL	15 min	CE
IF1071	Prog	Infertility, evaluation of ovulation	Refer to User Manual	S/P	0.10-40.00 ng/mL	100 µL	15 min	CE
Tumor	Markers							
IF1053	tPSA	Prostate cancer	4.0 ng/mL	S/P	0.50-100.00 ng/mL	100 µL	15 min	
IF1072	fPSA	Prostate cancer	1.0 ng/mL	S/P	0.10-30.00 ng/ml	100 µL	10 min	
IF1050	AFP	Liver cancer, cancer of ovaries or testicles, etc.	7.0 ng/mL	S/P	2.0-500.0 ng/mL	100 µL	15 min	C€
IF1051	CEA	Cancer marker: colon cancer etc.	4.7 ng/mL	S/P/WB	2.0-500.0 ng/mL	100 µL	15 min	CE
Infecti	ous Disease							
IF1057	Anti-HCV	Hepatitis C	1 S/CO	S/P	1.00-20.00 S/CO	100 µL	15 min	
IF1058	Anti-TP	Syphilis	1 S/CO	S/P	1.00-50.00 S/CO	100 µL	15 min	C€
IF1059	Anti-HIV	AIDS	1 S/CO	S/P	1.00-1000.00 S/CO	100 µL	15 min	
IF1064	HBsAg	Hepatitis B	1 IU/mL	S/P	1.00-100.00 IU/mL	100 µL	15 min	
IF1063	Anti-HBs	Hepatitis B	10.00 mIU/mL	S/P/WB	10.00-1000.00 mIU/m	L100 µL	15 min	
IF1084	2019-nCoV lgM/lgG	COVID-19	1 COI	S/P/WB		100 µL	10 min	CE
IF1091	SARS-CoV-2 Antigen	COVID-19	1 COI Nasa	al swab/Sali	va	100 µL	15 min	CE
IF1092	SARS-CoV-2 Total Antibody/ Neutralizing Antibody	COVID-19	Refer to User Manual	S/P/WB		100 µL	10 min	CE
IF1095	SARS-CoV-2 Neutralizing Antibody	COVID-19	Refer to User Manual	S/P/WB		100 µL	10 min	
IF1047	H. pylori	H. pylori infection	5 ng/mL	Stool	1-200 ng/mL	150 mg	10 min	C€
Others	S							
IF1077	Ferritin	Anemia/tumors	Male: 30-400 ng/mL Female: 13-150 ng/mL	S/P	0.50-2000.00 ng/mL	100 µL	15 min	CE
IF1069	Total IgE	Allergic disorders	Refer to User Manual	S/P/WB	1.00-2000.00 IU/mL	100 µL	15 min	CE

Coming Soon: FOB, ASO, RF, anti-CCP, Folate...



Add: No.9 Bofu Road, Luhe District, Nanjing, 211505, China Tel: +86-25-68568508/68568594

Fax: +86-25-68568500

E-mail: sales@getein.com.cn; overseas@getein.com.cn

Web: en.bio-gp.com.cn







