

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

Document No.: GP-GMSQ-2023121301

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 (Address: Republic of Moldova, Chisinau, MD-2059, Petricani street, 88/1, office 10) as our official and non-exclusive distributor for registration, promoting, selling, distributing and providing after-sale services of under-mentioned product in the territory of Moldova only:

FIA8000 Quantitative Immunoassay Analyzer and Reagents

Getein1100 Immunofluorescence Quantitative Analyzer and Reagents

Getein1160 Immunofluorescence Quantitative Analyzer and Reagents

Getein 1600 Immunofluorescence Quantitative Analyzer and Reagents

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.

Sanmedico SRL will carry out marketing efforts to fulfill service and maintenance for above mentioned products and will provide with users benefits of having a local stock of above mentioned products and on-time delivery with every order, supported by a local service in local language.

This authorization starts from 1st Jan, 2024 and will be valid to 31th, December, 2024.

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

基蛋生物科技股份有限公司 Getein Biotech, Inc. GET EN BIOTECH, INC. Seat & Signature

(大. www. Than

Authority Person Name: Steven Zhou

Authority Person Position: Regional Manager

Date: 2023.12.13





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:

Please see scope page.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2020-05-29

Latest Revision Date: 2023-04-26

bsi.



Effective Date: 2023-07-26 Expiry Date: 2026-07-25

Page: 1 of 3

...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.

Certificate No: MD 728432

Registered Scope:

Design & Development, Manufacture and Distribution of Chemiluminescence Immunoassay, Biochemistry Assay, Point of Care Assay (including Colloidal Gold Assay, Immunofluorescence Assay, Dry Chemistry Assay), PCR Assay and Colloidal Gold self-testing Assay to detect infectious disease. 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), PCR Assay to detect infectious disease, Immunofluorescence self-testing Assay to detect dyslipidemia disease, Blood Coagulation Assay to detect thrombotic disease.

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,传染病相关PCR分子诊断试剂和胶体金自测试剂。 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,传染病相关PCR分子诊断试剂,血脂异常疾病相关免疫荧光自测试剂,血栓疾病相关血凝试剂配套使用的分析仪。



Original Registration Date: 2020-05-29 Effective Date: 2023-07-26 Latest Revision Date: 2023-04-26 Expiry Date: 2026-07-25

Page: 2 of 3

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.

Certificate No: MD 728432

Location

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

Getein Biotech, Inc. No. 6 KeFeng Road Jiangbei New District Nanjing Jiangsu

Jiangsu 211505 China

基蛋生物科技股份有限公司

中国 江苏省 南京 江北新区 科丰路6号 邮编: 211505

Registered Activities

Design & Development, Manufacture and Distribution of Chemiluminescence Immunoassay, Biochemistry Assay, Point of Care Assay (including Colloidal Gold Assay, Immunofluorescence Assay, Dry Chemistry Assay), PCR Assay and Colloidal Gold self-testing Assay to detect infectious disease. 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), PCR Assay to detect infectious disease, Immunofluorescence self-testing Assay to detect dyslipidemia disease, Blood Coagulation Assay to detect thrombotic disease.

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,传染病相关PCR分子诊断试剂和胶体金自测试剂。 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,传染病相关PCR分子诊断试剂,血脂异常疾病相关免疫荧光自测试剂,血栓疾病相关血凝试剂配套使用的分析仪。

Manufacture of Chemiluminescence Immunoassay, Biochemistry Assay, Point of Care Assay (including Colloidal Gold Assay, Immunofluorescence Assay, Dry Chemistry Assay), Colloidal Gold self-testing Assay to detect infectious disease. Manufacture of Analyzers in use of Chemiluminescence Immunoassay, Biochemistry Assay, Point of Care Assay (including Colloidal Gold Assay, Immunofluorescence Assay, Dry Chemistry Assay), PCR Assay to detect infectious disease, Immunofluorescence self-testing Assay to detect dyslipidemia disease, Blood Coagulation Assay to detect thrombotic disease.

生产化学发光法试剂,生化试剂,即时诊断(包括胶体金法,免疫荧光法,干式化学法)试剂和传染病相关胶体金自测试剂。生产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,传染病相关PCR分子诊断试剂,血脂异常疾病相关免疫荧光自测试剂,血栓疾病相关血凝试剂配套使用的分析仪。

Original Registration Date: 2020-05-29 Effective Date: 2023-07-26 Latest Revision Date: 2023-04-26 Expiry Date: 2026-07-25

Page: 3 of 3

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.

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)
11/11/11/11	3 4 5 6 7 8 9 10 11	NT-proBNP Fast Test Kit (Immunofluorescence Assay)
		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)



Commence of the second of the				***
	67	HBP Fast Tes	t Kit (Immunofluorescence Assa	ay)
	68	S100-β Fast T	est Kit (Immunofluorescence A	ssay)
	69	CK-MB/hs-c7	InI/Myo Fast Test Kit (Immuno	fluorescence Assay)
	70	Cortisol Fast	Test Kit (Immunofluorescence A	Assay)
	/71	CEA Fast Tes	t Kit (Immunofluorescence Ass	ay)
	72	AFP/CEA Fa	st Test Kit (Immunofluorescer	nce Assay)
Classification	Other device ((according to	o Annex II of the directive 98	8/79/EC)
Conformity assessment route	Annex III of th	e 98/79/EC		
Applicable	EN 13612:20	02	EN ISO 14971:2019	EN ISO15223-1:2016
coordination	EN ISO 1811	3-1:2011	EN ISO 18113-2:2011	EN ISO 18113-3:2011
standards	EN ISO 2364		EN ISO 13485:2016	ISO 780:2015
	EN 61326-2-6		IEC 61326-1:2013	
	EN 61010-2-	101:2002	IEC 61010-1:2010	
Annex III. The comp	oiled technical file ed and the quality er is exclusively re	ed on Europe and quality system cert	vitro diagnostic medical deve ean Parliament and the Cou system document accordin tificate has issued by BSI Gor the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B.
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager	oiled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B.
This declaration of of Annex III. The companies III are testified V. The manufacture	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. nity.
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. nity.
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)
This declaration of of Annex III. The companies Annex III are testified V. The manufacture General Manager Naming 13th, May,	biled technical file ed and the quality er is exclusively re Enben Su	ed on Europe and quality system cert	ean Parliament and the Cou system document accordin tificate has issued by BSI G or the declaration of conform	uncil's 98/79/EC directive g to 98/79/EC directive roup The Netherlands B. hity. e or equivalent zed person)

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 Transfer



Test Card Insert



Click "Start" Icon



Result Show and Print

Outside Mode (Mass samples rapid test mode)



Sample Transfer



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 optional lithium battery



USER-FRIENDLY INTERFACE

Android system

7-inch touch screen



- 3 Test Card Slot
- 2 SD Card Recognition Zone 4 SD Card Slot



- **5** USB Slot
- **6** Built-in Thermal Printer

- **1** 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

10, 000 data

Language

English/Chinese/Spanish/Portuguese

Screen

7-inch touch screen

Power Supply

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

Working Environment

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

Dimensions

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	QUALIFI	ICATIC
Cardi	ac Markers								
IF1001	cTnl	Myocardial infarction	0.10 ng/mL	S/P/WB	0.10-50.00 ng/mL	100 μL	10 min	NMPA	CE
IF1098	TnT	Myocardial infarction	14.0 pg/mL	S/P/WB	10.0-10000.0 pg/mL	100 μL	15 min	NMPA	CE
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
IF1004	NT-proBNP/cTnI	Heart failure/acute myocardial infarction/acute coronary syndrome	NT-proBNP: 185 pg/mL cTnl: 0.10 ng/mL	S/P/WB	100-15000 pg/mL 0.10-50.00 ng/mL	100 μL	10 min	NMPA	CE
IF1005	CK-MB/cTnI/Myo	Myocardial damage /infarction	CK-MB: 5.00 ng/mL cTnl: 0.10 ng/mL Myo: 70.0 ng/mL	S/P/WB	2.50-80.00 ng/mL 0.10-50.00 ng/mL 30.0-600.0 ng/mL	100 μL	10 min	NMPA	CE
IF1012	CK-MB/cTnI	Myocardial damage /infarction	CK-MB: 5.00 ng/mL cTnl: 0.10 ng/mL	S/P/WB	2.50-80.00 ng/mL 0.10-50.00 ng/mL	100 μL	10 min	NMPA	CE
IF1014	H-FABP	Myocardial damage	6.36 ng/mL	S/P/WB	1.00-120.00 ng/mL	100 μL	3 min	NMPA	CE
IF1016	CK-MB/cTnI/H-FABP	Myocardial damage /infarction	CK-MB: 5.00 ng/mL cTnl: 0.10 ng/mL H-FABP: 6.36 ng/mL	S/P/WB	2.50-80.00 ng/mL 0.10-50.00 ng/mL 2.00-100.00 ng/mL	100 μL	10 min	NMPA	CE
IF1018	CK-MB	Myocardial injury	5.00 ng/mL	S/P/WB	2.50-80.00 ng/mL	100 μL	10 min	NMPA	CE
IF1087	ST2	AMI and chronic heart failure	35.0 ng/mL	S/P/WB	3.0-200.0 ng/mL	100 μL	15 min		CE
Coagu	ulation Marker								
IF1006	D-Dimer	Venous thromboembolism	0.50 mg/L	P/WB	0.10-10.00 mg/L	100 μL	10 min	NMPA	C
	nmation			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		,			
IF1003	hs-CRP+CRP	Cardiovascular inflammation /normal inflammation	3.0 mg/L 10.0 mg/L	S/P/WB/ Fingertip blood	0.5-200.0 mg/L	10 μL	3 min	NMPA	C
IF1007	PCT	Sepsis, bacterial infection	0.10 ng/mL	S/P/WB	0.05-50.00 ng/mL	100 μL	15 min	NMPA	C
IF1015	PCT/CRP	Sepsis, bacterial infection	PCT: 0.10 ng/mL CRP: 3.0 mg/L	S/P/WB	0.10-50.00 ng/mL 0.5-200.0 mg/L	20 μL	15 min	NMPA	
IF1044	SAA	Bacterial/Virus infection	10.0 mg/L	S/P/WB/ Fingertip blood	5.0-200.0 mg/L	10 μL	5 min	NMPA	C
IF1090	SAA/CRP	Neonatal sepsis, Bacterial/virus infection	SAA: 10.0 mg/L CRP: 10.0 mg/L	S/P/WB/ Peripheral blood	5.0-200.0 mg/L 0.5-200.0 mg/L	10 μL	5 min	NMPA	C
IF1088	IL-6	Acute inflammation	7.0 pg/mL	S/P/WB/ Peripheral blood	1.5-4000.0 pg/mL	40 μL	15 min	NMPA	C
IF1096	hs-CRP	Cardiovascular disease, routine inflammation	Refer to user manual	S/P	0.5-200.0 mg/L	100 μL	3 min	NMPA	C
IF1139	Calprotectin	Inflammatory bowel disease	<50.0 μg/g	Fecal specimen	10.0-600.0 μg/g	100 μL	15 min		C
Renal	Function								
IF1008	CysC	Acute and chronic renal diseases	0.51-1.09 mg/L	S/P/WB	0.50-10.00 mg/L	10 μL	3 min	NMPA	C
IF1009	mAlb	Diabetic nephropathy, hypertensive nephropathy	20.0 mg/L	Urine	10.0-200.0 mg/L	100 μL	3 min	NMPA	C
IF1010	NGAL	Acute kidney injury	Serum: 200.0 ng/mL Urine: 100.0 ng/mL	S/Urine	50.0-5000.0 ng/mL	10 μL	10 min	NMPA	C
IF1011	β_2 -MG	Acute and chronic kidney diseases/tumours	0.80-3.00 mg/L	S/P/WB	0.50-20.00 mg/L	10 μL	3 min	NMPA	C
Diabe	etes Mellitus								
IF1017	HbA1c	Diabetes mellitus	3.80%-5.80%	WB	2.00%-14.00%	10 μL	5 min	NGSP IFCC	NM C
Metab	oolic Marker								
IF1031	25-OH-VD	Osteomalacia, osteoporosis	30.00-50.00 ng/mL	S/P/WB/ Capillary blood	8.00-70.00 ng/mL	20 μL	20 min	NMPA	C
IF1112	Osteocalcin	Osteoporosis	Male: 14-70 ng/mL Female:11-48 ng/mL	S/P/WB	1.5-300.0 ng/mL	100 μL	15 min		C
Thyro	id Function								
IF1024	TSH	Thyroid malfunction	0.27-4.20 μlU/mL	S/P	0.10-50.00 μIU/mL	100 μL	15 min	NMPA	(
IF1022	Т3	Hyperthyroidism, hypothyroidism	1.30-3.10 nmol/L	S/P	0.30-10.00 nmol/L	100 μL	15 min	NMPA	
IF1023	T4	Hyperthyroidism, hypothyroidism	59.00-154.00 nmol/L	S/P	5.40-320.00 nmol/L	100 μL	15 min	NMPA	(
		11 d 12			2.2				
IF1067	fT3	Hyperthyroidism, hypothyroidism	3.10-6.80 pmol/L	S/P/WB	0.60-50.00 pmol/L	100 μL	15 min	NMPA	C

Cat. # TEST ITEMS				SAMPLE TYPES			REACTION TIME	QUALIFICATION	
Reproduction/Fertility									
IF1013	HCG+β	Fertility	5.1 mIU/mL	S/P	5.0-100000.0 mIU/mL	100 μL	10 min	NMPA	CE
IF1055	LH	PCOS, infertility evaluation	Refer to User Manual	S/P	0.20-150.00 mIU/mL	100 μL	15 min	NMPA	CE
IF1056	FSH	PCOS, infertility evaluation and pituitary disorders	Refer to User Manual	S/P	0.20-150.00 mIU/mL	100 μL	15 min	NMPA	CE
IF1066	АМН	Fertility, PCOS, gonadal function, precocious/late puberty	Refer to User Manual	S/P	0.10-20.00 ng/mL	100 μL	15 min	NMPA	CE
IF1048	PRL	Infertility, gonadal disorders	Refer to User Manual	S/P	0.50-200.00 ng/mL	100 μL	15 min	NMPA	C€
IF1071	Prog	Infertility, evaluation of ovulation	Refer to User Manual	S/P	0.10-40.00 ng/mL	100 μL	15 min		CE
w IF1073	Testosterone	Female polycystic ovary syndrome, male testosterone insufficiency	Male: 1.75-7.81 ng/mL Female: 0.10-0.75 ng/mL	S/P	0.10-16.00 ng/mL	100 μL	15 min		C€
WIF1138	Estradiol	Ovarian function	Refer to User Manual	S/P	40.0-4800.0 pg/mL	100 μL	15 min		C€
Tumor	Markers								
IF1053	tPSA	Prostate cancer	4.00 ng/mL	S/P	0.40-100.00 ng/mL	100 μL	15 min	NMPA	
IF1072	fPSA	Prostate cancer	1.00 ng/mL	S/P	0.03-30.00 ng/mL	100 μL	10 min	NMPA	
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	NMPA	C€
IF1051	CEA	Cancer marker: colon cancer etc.	4.7 ng/mL	S/P	2.0-500.0 ng/mL	100 μL	15 min	NMPA	C€
WIF1079	CA125	Ovarian cancer	35.0 U/mL	S/P/WB	2-500.0 U/mL	100 μL	15 min		CE
WIF1080	CA19-9	Pancreatic cancer	27.0 U/mL	S/P/WB	2-1000.0 U/mL	100 μL	15 min		C€
WIF1081	CA15-3	Breast cancer	26.2 U/mL	S/P/WB	1.5-300.0 U/mL	100 μL	15 min		CE
Infection	ous Disease								
IF1057	Anti-HCV	Hepatitis C	1.00 S/CO	S/P	1.00-20.00 S/CO	100 μL	15 min		
IF1058	Anti-TP	Syphilis	1.00 S/CO	S/P	1.00-50.00 S/CO	100 μL	15 min		CE
IF1059	Anti-HIV	AIDS	1.00 S/CO	S/P	1.00-1000.00 S/CO	100 μL	15 min		
IF1064	HBsAg	Hepatitis B	1.00 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/mL	100 μL	15 min		
IF1091	SARS-CoV-2 Antigen	COVID-19	1.00 COI	Nasal swab		100 μL	15 min		C€
IF1047	H. pylori	H. pylori infection	5.0 ng/mL	Stool	1.0-200.0 ng/mL	10-50 mg	10 min		CE
IF1086	Influenza A/B	Respiratory viral infection	1.00 COI	Nasal swab		100 μL	15 min		C€
IF1136	Dengue NS1 Ag	Dengue virus infection	1.00 S/CO	S/P/WB	1.00-50.00 S/CO	100 μL	15 min		C€
Specifi	ic Protein and Rh	neumatism							
IF1075	RF	Rheumatoid arthritis	15.9 IU/mL	S/P/WB	10.0-640.0 IU/mL	10 μL	10 min	NMPA	C€
IF1076	ASO	Rheumatic fever, acute glomerulonephritis, group A streptococcal infection	408.0 IU/mL	S/P/WB	60.0-1370.0 IU/mL	10 μL	10 min	NMPA	C€
IF1029	Anti-CCP	Rheumatoid arthritis	25.0 U/mL	S/P/WB	10.0-400.0 U/mL	10 μL	15 min		C€
Others									
IF1077	Ferritin	Anemia/tumors	Male: 30.00-400.00 ng/mL Female: 13.00-150.00 ng/mL	S/P	0.50-1000.00 ng/mL	10 μL	15 min	NMPA	C€
IF1069	Total IgE	Allergic disorders	Refer to User Manual	S/P/WB	1.00-2000.00 IU/mL	100 μL	15 min		C€
_	PG I/PG II	Atrophic gastritis, stomach cancer	PG I < 70.0 ng/mL PG I/PG II < 3.0 ng/mL	S/P	PG I: 1.0-200.0 ng/mL PG II: 1.0-100.0 ng/mL	100 μL	15 min		C€
W IF1052		Storrider earreer	,		O/				

Coming soon: VB12, Folate...



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

Fax: +86-25-68568500

 $\hbox{E-mail: overseas@getein.com.cn; marketing@getein.com.cn}\\$

Web.: www.getein.com





















