

AUTHORIZATION LETTER

Wr. Neudorf, 09.02.2021

TO WHOM IT MAY CONCERN

We, **DIALAB Produktion und Vertrieb von chemisch-technischen Produkten und Laborinstrumenten Gesellschaft m.b.H.**, headquarter in Austria, IZ-NÖ Süd, Hondastrasse Obj. M55, A-2351 Wr. Neudorf hereby declares that the below mentioned company is our official **representative** and is authorized to register, sell and distributor our products in the territory of Moldavia:

ECHIPAMED PLUS SRL
Valea Trandafirilor str., 24B, of.80,
MD-2001 Chisinau, Moldova

This certificate remains in force from 01.01.2021 until 31.01.2022 or is terminated during that period on the expiry of not less than 30 days' notice in writing given by either party to the other.

Signed for and on behalf of
DIALAB GmbH



Produktion und Vertrieb von chemisch-technischen Produkten und Laborinstrumenten Gesellschaft m.b.H.
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Christina ERNST
Export Manager

DIALAB Produktion und Vertrieb
von chemisch-technischen
Produkten und Laborinstrumenten
Gesellschaft m.b.H.

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Managing Director / Geschäftsführer
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UID/VAT: ATU 150 136 06 | DVR: 0130985

Raiffeisen Regionalbank Moedling
BIC / SWIFT: RLNWAT3333
IBAN AT: AT97 3725 0000 0000 6739
IBAN USD: AT52 3225 0000 0000 6739



EC Certificate

mdc medical device certification GmbH

Notified Body 0483
herewith certifies that

**Dialab Produktion und Vertrieb von
chemisch-technischen Produkten und
Laborinstrumenten Gesellschaft m.b.H.
IZ-NOE Sued, Hondastrasse, Objekt M55
2351 Wr. Neudorf
Austria**

for the scope

Reagents and reagent products for the determination
of the blood groups ABO system and Rhesus (C, D, E, e);
reagents for the determination of irregular Anti-Erythrocytic Antibodies (AHG);
reagents and reagent products for detection and
quantification of the following infectious diseases:

Rubella, Toxoplasmosis, Cytomegalovirus and Chlamydia; reagents and
reagent products for detection and quantification of tumoral marker PSA;
devices for self-diagnosis for the measurement of blood sugar;
devices for self-diagnosis for pregnancy and ovulation (hormones)
(see attachment)

has introduced and applies a

Quality System

for the design, manufacture and final inspection.

The mdc audit has proven that this quality system
meets all requirements according to

**Annex IV – excluding Section 4 and 6
of the Council Directive 98/79/EC**

of the European Parliament and of the Council of
27 October 1998 on in vitro diagnostic medical devices.

The surveillance will be held as specified in Annex IV, Section 5.

Valid from 2017-09-29
Valid until 2021-04-22
Registration no: D4000100005
Report no: P17-00988-105929
Stuttgart 2017-09-29

ALL

Head of Certification Body



mdc medical device certification GmbH
Königsplatz 3
D-70391 Stuttgart, Germany
Phone: +49 (0) 71 1-253597-0
Fax: +49 (0) 71 1-253597-10
Internet: <http://www.mdc-on.de>



Attachment of the certificate
No. D4000100005 Date 2017-09-29 Page 1 of 1

Product category	Product	Class
Reagents and reagent products for the determination of the blood groups ABO system	Anti-A monoclonal, Clone 9113D10 Anti-B monoclonal, Clone 9621A8 Anti-AB monoclonal, Clone 152D12+9113D10	List A, Annex II
Reagents and reagent products for the determination of the blood groups Rhesus system (C, c, D, E, e)	Anti-D monoclonal, Clone RUM-1+MS-26 Anti-C monoclonal, Clone MS-24 Anti-e monoclonal, Clone MS-33 Anti-E monoclonal, Clone MS-258 Anti-e monoclonal, Clone MS-16+MS-63 Anti-D negative control	List A, Annex II
Reagents for the determination of irregular Anti-Erythrocytic Antibodies	Anti-HG polyspecific/monoclonal	List B, Annex II
Reagents and reagent products for detection and quantification of the following infectious diseases: Rubella, Toxoplasmosis, Cytomegalovirus and Chlamydia (ELISA, immunochromatographic test)	Toxoplasma IgG, Toxoplasma IgM Rubella IgG, Rubella IgM CMV IgG, CMV IgM DIAQUICK Toxoplasma IgM Cassette DIAQUICK Rubella IgM Cassette DIAQUICK CMV IgM Cassette DIAQUICK Chlamydia Cassette	List B, Annex II
Reagents and reagent products for detection and quantification of tumoral marker PSA (ELISA, immunochromatographic rapid test)	PSA DIAQUICK PSA Cassette	List B, Annex II
Devices for self-diagnosis for the measurement of blood sugar	DiaCheck Pro Blood Glucose Monitoring System DiaCheck Pro Blood Glucose Test Strips DiaCheck Pro Glucose Control Solution	List B, Annex II
Devices for self-diagnosis for pregnancy and ovulation (hormones)	DIALAB Pregnancy Test (hCG) DiaFem Schwangerschaftstest (hCG)	self-testing not Annex II

ALL

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EC DECLARATION OF CONFORMITY

We



Dialab Produktion und Vertrieb von chemisch – technischen Produkten und Laborinstrumenten Gesellschaft m. b. H.
IZ NOE-Sued, Hondastrasse, Objekt M55
A-2351 Wiener Neudorf, Austria

declare, on our own responsibility, that our products specified in the enclosed addendum "Devices of List A, Annex II (TUV Süd), Rev.01 (1 page)",
classified as follows according to the directive on in vitro diagnostic medical devices
98/79/EC:

Devices of List A, Annex II

meet the applicable provisions of the European Directive 98 / 79 / EC for in-vitro-diagnostic medical devices and the Austrian Medical Product Law.

This Declaration is based on Conformity Assessment Procedures according to Annex IV, sections 3 and 4 of the aforesaid Directive in cooperation with the notified body

TÜV SÜD Product Service GmbH, code no. 0123
Ridlerstraße 65, 80339 München, Germany

This Declaration is valid until 2021-04-22.

Wiener Neudorf, 2019-04-08

DI Marlene Ramsey
Managing Director

DIALAB Produktion und Vertrieb
von chemisch-technischen
Produkten und Laborinstrumenten
Gesellschaft m.b.H.

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Managing Director | Geschäftsführer:
Marlene Ramsey
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BANK LEZ: AT52 2225 0000 0079 6739

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Addendum to the Declaration of Conformity of Devices for List A, Annex II (TUV Süd)

ABO and Rhesus Typing
Anti-A, monoclonal
REF B05405
CONT 1x 10 mL
LOT 600134-A4, 600135-A3, 600135-A2, 600135-A1, 600135-F4, 600140-J3, 600145-B4
CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0008 Rev. 00
www.dialab.at

Anti-B, monoclonal
B05406
1x 10 mL
601056-B3, 610159-A4, 610159-C3, 610160-A2, 610161-G2, 610164-H3, 610165-G3, 610170-B3

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0008 Rev. 00

Anti-AB, monoclonal
B05407
1x 10 mL
620119-C2, 620121-C2, 620124-B3, 620125-F3, 620128-B2

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0008 Rev. 00

Anti-D (IgM/IgG),
monoclonal
B05408
1x 10 mL
740156-B2, 740158-B3, 740162-A2, 740162-B4, 740162-D2, 740163-F3, 740163-H2, 740167-D3, 740168-B3, 740168-D2

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0007 Rev. 00

Anti-D Negative Control
B09936
1x 10 mL
650293-B7, 650314-A4, 650314-A5, 650314-A6, 650314-A7, 650325-A4

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0007 Rev. 00

Anti-C, monoclonal
B06411
1x 5 mL
Currently no lots released

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0007 Rev. 00

Anti-c, monoclonal
B06413
1x 5 mL
Currently no lots released

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0007 Rev. 00

Anti-E, monoclonal
B06412
1x 5 mL
Currently no lots released

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0007 Rev. 00

Anti-e, monoclonal
B06414
1x 5 mL
Currently no lots released

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0007 Rev. 00

HIV & HCV Testing

DIAQUICK HIV Plus
REF H18100
CONT -25 tests (25x H18100B)
-1x 4 mL buffer
LOT 1902064

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0006 Rev. 00

DIAQUICK HIV Plus WB
H18101
-25 tests (25x H18101B)
-1x 4 mL buffer
1902065

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0006 Rev. 00

DIAQUICK HCV Plus
H18200
25 tests (25x H18200B)
-1x 4 mL buffer
1902066

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0005 Rev. 00

DIAQUICK HCV Plus WB
H18201
25 tests (25x H18201B)
-1x 4 mL buffer
1902067

CE-Certificate No.: V1 026709 0004 Rev. 01 and V7 026709 0005 Rev. 00

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2351 Wiener Neudorf, Austria

declare, on our own responsibility, that our products specified in the enclosed
addendum "Other Products (Blood Grouping), Rev.09 (1 page)",
classified as follows according to the directive on in vitro diagnostic medical
devices 98/79/EC:

Devices other than those covered by Annex II (Blood Grouping)

meet the applicable provisions of the European Directive 98 / 79 / EC for in-vitro-
diagnostic medical devices and the Austrian Medical Product Law

CE marking has been carried out following the procedure referred to in Annex
III of the aforesaid Directive.

This declaration is valid until 2022-03-28.


DI Marlene Ramsey
Managing Director

Wiener Neudorf, 2019-04-08

DI Marlene Ramsey
Managing Director

Addendum to the Declaration of Conformity for Other Products (Blood Grouping)

BLOOD GROUPING AND RHESUS TYPING

ABO and Rhesus Typing	REF	CONT
Anti-Cv, monoclonal	B13001	1x 2 mL
Rare Blood Grouping Reagents		
Anti-H Leclin	B09934	1x 2 mL
Anti-k, monoclonal	B18908	1x 2 mL
Auxiliary Reagents		
BSA 22%	B05176	1x 10 mL
BSA 30%	B05175	1x 10 mL
LISS solution	B05178	1x 10 mL
Papain	B08000	1x 10 mL
Papain	B08001B	1x 1 L



Other Products (Blood Grouping), Rev.08

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