TÜV SÜD Product Service GmbH· Ridlerstr. 65 · 80339 Munich · Germany

Add value.
Inspire trust.

DiaMed GmbH
Pra Rond 23
1785 Cressier FR
Switzerland

Digitally signed by Levința Alexandru
Date: 2025.07.21 12:13:59 EEST
Reason: MoldSign Signature
Location: Moldova
MOLDOVA EUROPEANĂ

Your reference/letter of	Our reference/name	Tel. extension/Email	Fax extension	Date	Page
CBW 040330	713330580-03b	--- medical_devices@tuvsud.com	---	2024-10-09	1 of 11

TÜV SÜD Product Service GmbH
Confirmation Letter
CLI 040330 0196 Rev. 00

Reference: 713267047 | 713281388 | 713283488 | 713309238 | 713309242 | 713313459 | 713313460 | 713329914 | 713332971 | 713332972 | 713333137 | 713334057 | 713339857 | 713339864 | 713344735 | 713344759 | 713344779 | 713345479 | 713346822 | 713330580-03b

To whom it may concern,

Confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation EU 2024/1860 amending Regulations (EU) 2017/746 (in the following referenced as IVDR) as regards the transitional provisions for certain in vitro diagnostic medical devices.

With this letter TÜV SÜD Product Service GmbH, designated under IVDR and identified by the number 0123 on NANDO, confirms that we have received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of IVDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of IVDR with the above stated manufacturer with the following Single Registration Number (SRN)

Single Registration Number: CH-MF-000020826

The devices covered by the formal application and the written agreement mentioned above are identified in the Tables below.

- Table 1 identifies the devices for which an IVDR application has been received, written agreement concluded and for which TÜV SÜD Product Service GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive.

- Table 2 identifies the devices for which an IVDR application has been received and a written agreement concluded, but TÜV SÜD Product Service GmbH has not yet taken the responsibility for appropriate

surveillance of the corresponding devices under the applicable Directive or these devices did not require a Notified Body certificate under Directives.

If devices covered by certificates issued under Directive Directive 98/79/EC (IVDD) that expired after 26. May 2022 and before 09. July 2024, without having been withdrawn, this letter also confirms that

- the manufacturer signed the written agreement under IVDR by the date of IVDD certificate expiry; or
- provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 54(1) of IVDR or Article 92(1) of the IVDR respectively.

The transition timelines in accordance Article 110 (3) of IVDR that apply to the devices covered by this letter, subject to the manufacturer's continued compliance to the other conditions specified in Article 110 (3c) of IVDR, are shown below:

- 31. December 2027, for devices certified under IVDD
- 31. December 2027, for class D devices;
- 31. December 2028, for class C devices;
- 31. December 2029, for class B devices and for class A devices placed on the market in sterile condition

We reserve the right to invoice any issuance, copies, amendments and / or changes of the confirmation letter according to effort.

For confirmation letter validity see www.tuvsud.com/ps-cert?q=cert:CLI 040330 0196

In case of inquiries please contact medical_devices@tuvsud.com.

On behalf of the Notified Body TÜV SÜD Product Service GmbH,
2024-10-09

TÜV SÜD Product Service GmbH
Medical and Health Services

Juergen Puels
Juergen Puels (9. Oktober 2024 14:17 GMT+2)

Dr. Jürgen Püls
Conformity Assessment Responsible (CARE)

TÜV SÜD Product Service GmbH
Medical and Health Services

Christian Ullmann
Christian Ullmann (9. Oktober 2024 15:16 GMT+2)

Dr. Christian Ullmann
Application Reviewer

Table 1: Devices covered by this letter and for which TÜV SÜD Product Service GmbH is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive:

Device name or Basic UDI-DI (under IVDR application)	IVDR Device classification (as proposed by the manufacturer and verified during application review)	If the IVDR device is a substitute device, identification of the corresponding IVDD device	IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification
DiaClon Anti-K	Class D incl. ST/NPT	N/A	V7 040330 0153 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon Rh + K Pheno II	Class D incl. ST/NPT	N/A	V7 040330 0153 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon Rh-Subgroups + Cw + K	Class D incl. ST/NPT	N/A	V7 040330 0153 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon Rh-Subgroups + K	Class D incl. ST/NPT	N/A	V7 040330 0153 Rev. 01 V1 040330 0182 Rev. 01 NB #012§
ID-DiaClon Anti-c	Class D incl. ST/NPT	N/A	V7 040330 0153 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell ABO A1, A2, B, O	Class D incl. ST/NPT	N/A	V7 040330 0157 Rev. 02 V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell ABO A1, B	Class D incl. ST/NPT	N/A	V7 040330 0157 Rev. 02 V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell ABO A1, B, O	Class D incl. ST/NPT	N/A	V7 040330 0157 Rev. 02 V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell ABO A2	Class D incl. ST/NPT	N/A	V7 040330 0157 Rev. 02 V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell ABO O	Class D incl. ST/NPT	N/A	V7 040330 0157 Rev. 02 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/D (DVI-, DVI-) + Reverse Grouping	Class D incl. ST/NPT	N/A	V7 040330 0158 Rev. 00 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/D (DVI+, DVI-) + Reverse Grouping	Class D incl. ST/NPT	N/A	V7 040330 0158 Rev. 00 V1 040330 0182 Rev. 01 NB #0123

Device name or Basic UDI-DI (under IVDR application)	IVDR Device classification (as proposed by the manufacturer and verified during application review)	If the IVDR device is a substitute device, identification of the corresponding IVDD device	IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification
DiaClon ABO/Rh + Epreuve Sérique et DiaClon Rh-Sousgroupes + K	Class D incl. ST/NPT	N/A	V7 040330 0161 Rev.01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/DVI+/DVI- + DAT	Class D incl. ST/NPT	N/A	V7 040330 0165 Rev.00 V1 040330 0182 Rev. 01 NB #0123
DiaClon RhD + Phenotype	Class D incl. ST/NPT	N/A	V7 040330 0167 Rev. 02 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABD (DVI-) Confirmation for Patients	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABD-Confirmation for Donors	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABD-Confirmation for Patients	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/D	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/D + DAT	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/D + Reverse Grouping	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/Rh for Donors	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/Rh for Newborns DVI+	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/Rh for Patients	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon Complete Crossmatch	Class D incl. ST/NPT	N/A	V7 040330 0171 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/D + Reverse Grouping	Class D incl. ST/NPT	N/A	V7 040330 0174 Rev. 01 V1 040330 0182 Rev. 01 NB #0123

Device name or Basic UDI-DI (under IVDR application)	IVDR Device classification (as proposed by the manufacturer and verified during application review)	If the IVDR device is a substitute device, identification of the corresponding IVDD device	IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification
DiaClon ABO/D + Reverse Grouping for Patients	Class D incl. ST/NPT	N/A	V7 040330 0174 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon BAS-Test IgG	Class D incl. ST/NPT	N/A	V7 040330 0177 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/Rh for Newborns DVI-	Class D incl. ST/NPT	N/A	V7 040330 0178 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/D + Reverse Grouping for Donors	Class D incl. ST/NPT	N/A	V7 040330 0179 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
IH-QC1	Class D incl. ST/NPT	N/A	V7 040330 0183 rev.01 V1 040330 0182 Rev. 01 NB #0123
IH-QC2	Class D incl. ST/NPT	N/A	V7 040330 0183 rev.01 V1 040330 0182 Rev. 01 NB #0123
IH-QC3	Class D incl. ST/NPT	N/A	V7 040330 0183 rev.01 V1 040330 0182 Rev. 01 NB #0123
IH-QC4	Class D incl. ST/NPT	N/A	V7 040330 0183 rev.01 V1 040330 0182 Rev. 01 NB #0123
IH-QC5	Class D incl. ST/NPT	N/A	V7 040330 0183 rev.01 V1 040330 0182 Rev. 01 NB #0123
IH-QC6	Class D incl. ST/NPT	N/A	V7 040330 0183 rev.01 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/DVI-	Class D incl. ST/NPT	N/A	V7 040330 0185 Rev. 00 V1 040330 0182 Rev. 01 NB #0123
DiaClon ABO/DVI- for Patients	Class D incl. ST/NPT	N/A	V7 040330 0185 Rev. 00 V1 040330 0182 Rev. 01 NB #0123
Control Card A	Class D incl. ST/NPT	N/A	V7 040330 0159 Rev. 01 V1 040330 0182 Rev. 01 NB #0123
ID-Internal Quality Control	Class D incl. ST/NPT	N/A	V7 040330 0166 Rev. 01 V1 040330 0182 Rev. 01 NB #0123

Device name or Basic UDI-DI (under IVDR application)	IVDR Device classification (as proposed by the manufacturer and verified during application review)	If the IVDR device is a substitute device, identification of the corresponding IVDD device	IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification
Coombs Anti-IgG	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
DiaScreen	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-Antigen Profile II	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-Antigen Profile III	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-Antigen Profile III (ID-Testsera Anti-M,-N,-S,s,-Fya,-Fyb (6x5,0ml))	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-Card Fya	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-Card Fya/Fyb	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-Card Fyb	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-Dia (Diego) Positive	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell I-II	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell I-II-III	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell I-II-III Asia	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell IP-IIP-IIIP	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell Pool	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaCell SF	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaPanel	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaPanel Plus 6	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaPanel-P	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaScreen I-II-III-IV	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaScreen I-II-III-IV-VP-VIP	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
ID-DiaScreen Prophylax	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123

Device name or Basic UDI-DI (under IVDR application)	IVDR Device classification (as proposed by the manufacturer and verified during application review)	If the IVDR device is a substitute device, identification of the corresponding IVDD device	IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification
LISS/Coombs	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
Test Serum ID-Anti-Fya	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123
Test Serum ID-Anti-Fyb	Class D incl. ST/NPT	N/A	V1 040330 0182 Rev. 01 NB #0123

Legend: ST – self-testing; NPT – near-patient testing; CDx – companion diagnostics

Table 2: Devices covered by this letter and for which TÜV SÜD Product Service GmbH is NOT responsible for appropriate surveillance of the corresponding devices under the applicable Directive:

Device name or Basic UDI-DI (under IVDR application)	IVDR Device classification (as proposed by the manufacturer and verified during application review)	If the IVDR device is a substitute device, identification of the corresponding IVDD device	IVDD Certificate Reference(s) of the devices under IVDR application, and the NB Identification
Anti-D Reference Reagent	Class D incl. ST/NPT	N/A	N/A - Device did not require a Notified Body certificate under Directives
ID-DiaClon Anti-D	Class D incl. ST/NPT	N/A	N/A - Device did not require a Notified Body certificate under Directives
NaCl, Enzyme Test and Cold Agglutinins	Class D incl. ST/NPT	N/A	N/A - Device did not require a Notified Body certificate under Directives

Confirmation Letter Version History

Date	TÜV SÜD Product Service GmbH internal reference traceable to each version of the letter	Action
2024-10-09	713330580-03b	Initial issue

Attachment

Additional Information for the devices listed in the table(s) above:

Device name or Basic UDI-DI (under IVDR application)	REF / Article Number (under IVDD)	Article name (under IVDD)
361052A0037087	002121	DiaClon Anti-K
361052A003758H	002224	DiaClon Rh + K Pheno II
361052A003758H	002226	DiaClon Rh + K Pheno II
361052A003758H	002227	DiaClon Rh + K Pheno II
361052A003758H	002225	DiaClon Rh + K Pheno II
361052A003738D	002134	DiaClon Rh-Subgroups + Cw + K
361052A003738D	002136	DiaClon Rh-Subgroups + Cw + K
361052A003738D	002137	DiaClon Rh-Subgroups + Cw + K
361052A003738D	002135	DiaClon Rh-Subgroups + Cw + K
361052A0037189	002124	DiaClon Rh-Subgroups + K
361052A0037189	002126	DiaClon Rh-Subgroups + K
361052A0037189	002127	DiaClon Rh-Subgroups + K
361052A0037189	002125	DiaClon Rh-Subgroups + K
361052A003748F	002154	ID-DiaClon Anti-c
361052A003748F	002151	ID-DiaClon Anti-c
361052A0033687	003619	ID-DiaCell ABO A1, A2, B, O
361052A003327X	003624	ID-DiaCell ABO A1, B
361052A0033585	003615	ID-DiaCell ABO A1, B, O
361052A0033483	003621	ID-DiaCell ABO A2
361052A003337Z	003623	ID-DiaCell ABO O
361052A003668G	001275	DiaClon ABO/D (DVI-, DVI-) + Reverse Grouping
361052A003668G	001276	DiaClon ABO/D (DVI-, DVI-) + Reverse Grouping
361052A003688L	001386	DiaClon ABO/D (DVI+, DVI-) + Reverse Grouping
361052A003938K	001397	DiaClon ABO/Rh + Epreuve Sérique et DiaClon Rh-Sousgroupes + K
361052A003938K	001398	DiaClon ABO/Rh + Epreuve Sérique et DiaClon Rh-Sousgroupes + K
361052A003638A	001374	DiaClon ABO/DVI+/DVI- + DAT
361052A003638A	001376	DiaClon ABO/DVI+/DVI- + DAT
361052A003768K	002234	DiaClon RhD + Phenotype
361052A003768K	002237	DiaClon RhD + Phenotype
361052A0035387	001284	DiaClon ABD (DVI-) Confirmation for Patients
361052A0035387	001286	DiaClon ABD (DVI-) Confirmation for Patients
361052A0035387	001287	DiaClon ABD (DVI-) Confirmation for Patients
361052A0035387	001285	DiaClon ABD (DVI-) Confirmation for Patients
361052A003568D	001134	DiaClon ABD-Confirmation for Donors
361052A003568D	001136	DiaClon ABD-Confirmation for Donors
361052A003568D	001133	DiaClon ABD-Confirmation for Donors
361052A003568D	001135	DiaClon ABD-Confirmation for Donors
361052A0035489	001254	DiaClon ABD-Confirmation for Patients
361052A0035489	001256	DiaClon ABD-Confirmation for Patients

361052A0035489	001257	DiaClon ABD-Confirmation for Patients
361052A0035489	001255	DiaClon ABD-Confirmation for Patients
361052A003558B	001324	DiaClon ABO/D
361052A003558B	001326	DiaClon ABO/D
361052A003558B	001323	DiaClon ABO/D
361052A003558B	001325	DiaClon ABO/D
361052A0036186	001346	DiaClon ABO/D + DAT
361052A0036186	001344	DiaClon ABO/D + DAT
361052A0036186	001347	DiaClon ABO/D + DAT
361052A0036186	001345	DiaClon ABO/D + DAT
361052A003658E	001234	DiaClon ABO/D + Reverse Grouping
361052A003658E	001236	DiaClon ABO/D + Reverse Grouping
361052A003658E	001237	DiaClon ABO/D + Reverse Grouping
361052A003658E	001235	DiaClon ABO/D + Reverse Grouping
361052A003578F	001037	DiaClon ABO/Rh for Donors
361052A003578F	001039	DiaClon ABO/Rh for Donors
361052A003578F	001033	DiaClon ABO/Rh for Donors
361052A003578F	001038	DiaClon ABO/Rh for Donors
361052A0036288	001047	DiaClon ABO/Rh for Newborns DVI+
361052A0036288	001049	DiaClon ABO/Rh for Newborns DVI+
361052A0036288	001048	DiaClon ABO/Rh for Newborns DVI+
361052A0036288	001050	DiaClon ABO/Rh for Newborns DVI+
361052A0035285	001044	DiaClon ABO/Rh for Patients
361052A0035285	001046	DiaClon ABO/Rh for Patients
361052A0035285	001043	DiaClon ABO/Rh for Patients
361052A0035285	001045	DiaClon ABO/Rh for Patients
361052A003808A	004614	DiaClon Complete Crossmatch
361052A003808A	004616	DiaClon Complete Crossmatch
361052A003808A	004617	DiaClon Complete Crossmatch
361052A003808A	004615	DiaClon Complete Crossmatch
361052A003678J	001248	DiaClon ABO/D + Reverse Grouping
361052A003678J	001249	DiaClon ABO/D + Reverse Grouping
361052A003648C	001264	DiaClon ABO/D + Reverse Grouping for Patients
361052A003648C	001266	DiaClon ABO/D + Reverse Grouping for Patients
361052A003648C	001267	DiaClon ABO/D + Reverse Grouping for Patients
361052A003648C	001265	DiaClon ABO/D + Reverse Grouping for Patients
361052A003818C	001965	DiaClon BAS-Test IgG
361052A003818C	001966	DiaClon BAS-Test IgG
361052A0036084	001027	DiaClon ABO/Rh for Newborns DVI-
361052A0036084	001028	DiaClon ABO/Rh for Newborns DVI-
361052A0036084	001030	DiaClon ABO/Rh for Newborns DVI-
361052A0036084	001029	DiaClon ABO/Rh for Newborns DVI-
361052A003698N	001365	DiaClon ABO/D + Reverse Grouping for Donors
361052A003698N	12010791	DiaClon ABO/D + Reverse Grouping for Donors
361052A0034282	009321	IH-QC1
361052A003417Y	009322	IH-QC2
361052A003407W	009323	IH-QC3
361052A003468A	009324	IH-QC4
361052A0034588	009325	IH-QC5

361052A0034486	009326	IH-QC6
361052A003588H	001424	DiaClon ABO/DVI-
361052A003588H	001425	DiaClon ABO/DVI-
361052A003588H	001427	DiaClon ABO/DVI-
361052A003598K	001294	DiaClon ABO/DVI- for Patients
361052A003598K	001296	DiaClon ABO/DVI- for Patients
361052A003598K	001297	DiaClon ABO/DVI- for Patients
361052A003728B	001711	Control Card A
361052A003728B	001714	Control Card A
361052A0034384	009925	ID-Internal Quality Control
361052A003798R	004024	Coombs Anti-IgG
361052A003798R	004026	Coombs Anti-IgG
361052A003798R	004027	Coombs Anti-IgG
361052A003798R	004025	Coombs Anti-IgG
361052A003778M	004704	DiaScreen
361052A003778M	004705	DiaScreen
361052A003778M	004707	DiaScreen
361052A003778M	004706	DiaScreen
361052A003498G	008610	ID-Antigen Profile II
361052A003478C	008701	ID-Antigen Profile III
361052A003488E	008712	ID-Antigen Profile III (ID-Testsera Anti-M,-N,-S,s,-Fya,-Fyb (6x5,0ml))
361052A003127R	007270	ID-Card Fya
361052A003157X	006110	ID-Card Fya/Fyb
361052A003147V	007280	ID-Card Fyb
361052A0031885	004134	ID-Dia (Diego) Positive
361052A003237W	003613	ID-DiaCell I-II
361052A003227U	004310	ID-DiaCell I-II-III
361052A0032786	003614	ID-DiaCell I-II-III Asia
361052A003217S	005310	ID-DiaCell IP-IIP-IIIP
361052A003207Q	003631	ID-DiaCell Pool
361052A003207Q	003630	ID-DiaCell Pool
361052A0031987	003640	ID-DiaCell SF
361052A0032582	004114	ID-DiaPanel
361052A003307T	004414	ID-DiaPanel Plus 6
361052A0032684	004214	ID-DiaPanel-P
361052A003298A	004311	ID-DiaScreen I-II-III-IV
361052A003247Y	004316	ID-DiaScreen I-II-III-IV-VP-VIP
361052A003317V	004330	ID-DiaScreen Prophylax
361052A003788P	004014	LISS/Coombs
361052A003788P	004016	LISS/Coombs
361052A003788P	004017	LISS/Coombs
361052A003788P	004015	LISS/Coombs
361052A003167Z	007272	Test Serum ID-Anti-Fya
361052A0031783	007282	Test Serum ID-Anti-Fyb
361052A0008789	105500	Anti-D Reference Reagent
361052A0009282	007531	ID-DiaClon Anti-D
361052A0026589	005014	NaCl, Enzyme Test and Cold Agglutinins
361052A0026589	005016	NaCl, Enzyme Test and Cold Agglutinins

361052A0026589	005017	NaCl, Enzyme Test and Cold Agglutinins
361052A0026589	005015	NaCl, Enzyme Test and Cold Agglutinins

2024-10-09

TÜV SÜD Product Service GmbH
Medical and Health Services

Juergen Puels
Juergen Puels (9. Oktober 2024 14:17 GMT+2)

Dr. Jürgen Püls
Conformity Assessment Responsible (CARE)