



Manufacturer's Declaration of Certificate Validity:

with respect to the validity of certificates issued under Council Directive 93/42/EEC on medical devices that have been extended as a result of the REGULATION (EU) 2023/607 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 15 March 2023 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices.

Manufacturer name	Caesarea Medical Electronics Ltd.
Manufacturer address	16 Shacham Street, Industrial Park Caesarea North, Caesarea 3088900, Israel.
EUDAMED SRN (if available)	IL-MF-000030321

Authorised Representative name (if applicable)	Becton Dickinson Ireland Limited
Authorised Representative address	Donore Road, Drogheda, Co. Louth, A92 YW26, Ireland.
EUDAMED SRN (if available)	IE-AR-000007610

This declaration confirms that the devices listed in the attached schedule meet following conditions for extension of their certificates issued under Council Directive 93/42/EEC (MDD) on medical devices as stated in the regulation 2023/607:

- The certificate(s) covering the listed devices was valid on the 26 May 2021.
- The devices continue to comply with Directive 93/42/EEC (MDD)
- There are no significant changes in the design and intended purpose since 26 May 2021.
- The device(s) do not present unacceptable risks to health or safety of patients, users or other persons, or to other aspects of the protection of public health.
- A quality management system in accordance with Article 10(9), Regulation (EU) 2017/745 (MDR) has been put in place by the manufacturer.
- A formal application to the notified body in accordance with Section 4.3, first subparagraph of Annex VII, Regulation (EU) 2017/745 (MDR) for conformity assessment has been made for the device(s) listed and a signed written agreement is in place in accordance with Section 4.3, second subparagraph, of Annex VII, Regulation (EU) 2017/745 (MDR) before the expiry date of the listed certificate.
- Post-market surveillance, market surveillance, vigilance, registration of economic operators and of devices in accordance with Regulation (EU) 2017/745 (MDR) is in place for the device(s) listed.

Signed for and on behalf of the manufacturer:

Signature:



DocuSigned by:
Tom Healy
Signer Name: Tom Healy
Signing Reason: I approve this document
Signing Time: 25-May-2023 | 3:35:22 PM BST

Print name:

Tom Healy

6AB495850B1C4F8EA72F8BC377AE2C9F

Date: 25-May-2023

Title/position: Senior RA Manager

Manufacturer name: Caesarea Medical Electronics Ltd.

Schedule of devices

Model Number/Catalogue number	Device Name	Certificate number ⁱ	Expiry date on the certificate ⁱⁱ	Extended expiry date ⁱⁱⁱ
BD BodyGuard™ Infusion Pumps				
999-603BDEN	BD BodyGuard™ Infusion Pump English	2122031CE01	01-Jun-23	31-Dec-28
999-603BDES	BD BodyGuard™ Infusion Pump Spanish	2122031CE01	01-Jun-23	31-Dec-28
999-603IT	BD BodyGuard™ Infusion Pump Italian	2122031CE01	01-Jun-23	31-Dec-28
999-603DE	BD BodyGuard™ Infusion Pump German	2122031CE01	01-Jun-23	31-Dec-28
999-603FR	BD BodyGuard™ Infusion Pump French	2122031CE01	01-Jun-23	31-Dec-28
999-603NL	BD BodyGuard™ Infusion Pump Dutch	2122031CE01	01-Jun-23	31-Dec-28
999-603DK	BD BodyGuard™ Infusion Pump Danish	2122031CE01	01-Jun-23	31-Dec-28
999-603SE	BD BodyGuard™ Infusion Pump Swedish	2122031CE01	01-Jun-23	31-Dec-28
999-603FI	BD BodyGuard™ Infusion Pump Finnish	2122031CE01	01-Jun-23	31-Dec-28
999-603NO	BD BodyGuard™ Infusion Pump Norwegian	2122031CE01	01-Jun-23	31-Dec-28
999-603PT	BD BodyGuard™ Infusion Pump Portuguese	2122031CE01	01-Jun-23	31-Dec-28
999-603GR	BD BodyGuard™ Infusion Pump Greek	2122031CE01	01-Jun-23	31-Dec-28
999-603PL	BD BodyGuard™ Infusion Pump Polish	2122031CE01	01-Jun-23	31-Dec-28
999-603TU	BD BodyGuard™ Infusion Pump Turkish	2122031CE01	01-Jun-23	31-Dec-28
999-603RU	BD BodyGuard™ Infusion Pump Russian	2122031CE01	01-Jun-23	31-Dec-28
BD BodyGuard™ Epidural Infusion Pumps				
999-683BDEN	BD BodyGuard™ Epidural Infusion Pump English	2122031CE01	01-Jun-23	31-Dec-28
999-683DE	BD BodyGuard™ Epidural Infusion Pump German	2122031CE01	01-Jun-23	31-Dec-28
999-683FR	BD BodyGuard™ Epidural Infusion Pump French	2122031CE01	01-Jun-23	31-Dec-28
999-683IT	BD BodyGuard™ Epidural Infusion Pump Italian	2122031CE01	01-Jun-23	31-Dec-28
999-683BDES	BD BodyGuard™ Epidural Infusion Pump Spanish	2122031CE01	01-Jun-23	31-Dec-28
999-683NL	BD BodyGuard™ Epidural Infusion Pump Dutch	2122031CE01	01-Jun-23	31-Dec-28

Model Number/Catalogue number	Device Name	Certificate numberⁱ	Expiry date on the certificateⁱⁱ	Extended expiry dateⁱⁱⁱ
999-683DK	BD BodyGuard™ Epidural Infusion Pump Danish	2122031CE01	01-Jun-23	31-Dec-28
999-683SE	BD BodyGuard™ Epidural Infusion Pump Swedish	2122031CE01	01-Jun-23	31-Dec-28
999-683FI	BD BodyGuard™ Epidural Infusion Pump Finnish	2122031CE01	01-Jun-23	31-Dec-28
999-683NO	BD BodyGuard™ Epidural Infusion Pump Norwegian	2122031CE01	01-Jun-23	31-Dec-28
999-683BDRU	BD BodyGuard™ Epidural Infusion Pump Russian	2122031CE01	01-Jun-23	31-Dec-28
BD BodyGuard™ Pain Manager Infusion Pumps				
999-803BDEN	BD BodyGuard™ Pain Manager English	2122031CE01	01-Jun-23	31-Dec-28
999-803DE	BD BodyGuard™ Pain Manager German	2122031CE01	01-Jun-23	31-Dec-28
999-803FR	BD BodyGuard™ Pain Manager French	2122031CE01	01-Jun-23	31-Dec-28
999-803IT	BD BodyGuard™ Pain Manager Italian	2122031CE01	01-Jun-23	31-Dec-28
999-803BDES	BD BodyGuard™ Pain Manager Spanish	2122031CE01	01-Jun-23	31-Dec-28
999-803NL	BD BodyGuard™ Pain Manager Dutch	2122031CE01	01-Jun-23	31-Dec-28
999-803TU	BD BodyGuard™ Pain Manager Turkish	2122031CE01	01-Jun-23	31-Dec-28
999-803GR	BD BodyGuard™ Pain Manager Greek	2122031CE01	01-Jun-23	31-Dec-28
999-803DK	BD BodyGuard™ Pain Manager Danish	2122031CE01	01-Jun-23	31-Dec-28
999-803SE	BD BodyGuard™ Pain Manager Swedish	2122031CE01	01-Jun-23	31-Dec-28
999-803FI	BD BodyGuard™ Pain Manager Finnish	2122031CE01	01-Jun-23	31-Dec-28
999-803NO	BD BodyGuard™ Pain Manager Norwegian	2122031CE01	01-Jun-23	31-Dec-28
999-803BDRU	BD BodyGuard™ Pain Manager Russian	2122031CE01	01-Jun-23	31-Dec-28
BD BodyGuard™ Duo Infusion Pumps				
999-903EN	BD BodyGuard™ Duo Infusion Pump English	2122031CE01	01-Jun-23	31-Dec-28
999-903ES	BD BodyGuard™ Duo Infusion Pump Spanish	2122031CE01	01-Jun-23	31-Dec-28
999-903IT	BD BodyGuard™ Duo Infusion Pump Italian	2122031CE01	01-Jun-23	31-Dec-28
999-903DE	BD BodyGuard™ Duo Infusion Pump German	2122031CE01	01-Jun-23	31-Dec-28
999-903FR	BD BodyGuard™ Duo Infusion Pump French	2122031CE01	01-Jun-23	31-Dec-28

Model Number/Catalogue number	Device Name	Certificate number ⁱ	Expiry date on the certificate ⁱⁱ	Extended expiry date ⁱⁱⁱ
BD BodyGuard™ Syringe Infusion Pumps				
999-103BDEN	BD BodyGuard™ T Syringe Pump English	2122031CE01	01-Jun-23	31-Dec-28
999-103BDSE	BD BodyGuard™ T Syringe Pump Swedish	2122031CE01	01-Jun-23	31-Dec-28
999-103BDRU	BD BodyGuard™ T Syringe Pump Russian	2122031CE01	01-Jun-23	31-Dec-28
BodyGuard™ 323 Color Vision Infusion Pumps				
999-603EN	BodyGuard™ 323 Color Vision Infusion Pump English	2122031CE01	01-Jun-23	26-May-24
999-603ES	BodyGuard™ 323 Color Vision Infusion Pump Spanish	2122031CE01	01-Jun-23	26-May-24
999-603PFM	BD BodyGuard™ Infusion Pump German (PFM)	2122031CE01	01-Jun-23	26-May-24
BodyGuard™ 545 Color Vision Infusion Pumps				
999-683EN	BodyGuard™ 545 Color Vision Infusion Pump English	2122031CE01	01-Jun-23	26-May-24
999-683ES	BodyGuard™ 545 Color Vision Infusion Pump Spanish	2122031CE01	01-Jun-23	26-May-24
BodyGuard™ 595 Color Vision Infusion Pumps				
999-803EN	BodyGuard™ 595 Color Vision Infusion Pump English	2122031CE01	01-Jun-23	26-May-24
999-803ES	BodyGuard™ 595 Color Vision Infusion Pump Spanish	2122031CE01	01-Jun-23	26-May-24
T34 Multi-Purpose Syringe Infusion				
999-103EN	T34 Syringe Pump English	2122031CE01	01-Jun-23	26-May-24
999-103SE	T34 Syringe Pump Swedish	2122031CE01	01-Jun-23	26-May-24
BD BodyGuard™ MicroSets Infusion Sets				
120-000DBLSK	BD BodyGuard™ MicroSet Blood; Blood Set, Microbore Tubing, Drip Chamber, Y Port, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
100-163XN2YNKS	BD BodyGuard™ MicroSet Microbore Tubing, Female Luer Connector, Needle Free Port, Y Ports, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-124XSK	BD BodyGuard™ MicroSet Microbore Tubing, Spike, Needle-free Port, Back Check Valve, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28

Model Number/Catalogue number	Device Name	Certificate numberⁱ	Expiry date on the certificateⁱⁱ	Extended expiry dateⁱⁱⁱ
100-163XNKS	BD BodyGuard™ MicroSet Syringe; Tubing, Syringe Adapter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-112XSFK	BD BodyGuard™ MicroSet Microbore Tubing, Spike, Inline 1.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-160XYSK	BD BodyGuard™ MicroSet Microbore Tubing, Drip Chamber, Y Port, Anti Siphon Valve, Male MD Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
GEN00001	BD BodyGuard™ MicroSet Microbore Tubing, Drip Chamber, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
GEN00002	BD BodyGuard™ MicroSet Microbore Tubing, Drip Chamber, Needle Free Port, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
GEN00004	BD BodyGuard™ MicroSet Microbore Tubing, Spike, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
GEN00014	MICROSET, N-DEHP, STRAIGHT	2122031CE02	01-Jun-23	31-Dec-28
100-160XSMG90EK	BD BodyGuard™ MicroSet Grey Microbore Tubing, 90° Spike, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
100-184XSK	BD BodyGuard™ MicroSet Blue Microbore Tubing, Syringe Adapter, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
100-184XSYK	BD BodyGuard™ MicroSet Blue Microbore Tubing, Syringe Adapter, Y Port, Female Luer Connector, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-160X90SK	BD BodyGuard™ MicroSet Blue Striped Microbore Tubing, 90° Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28

Model Number/Catalogue number	Device Name	Certificate numberⁱ	Expiry date on the certificateⁱⁱ	Extended expiry dateⁱⁱⁱ
120-160XCSEK	BD BodyGuard™ MicroSet Blue Striped Microbore Tubing, Spike, Y Port, Female Luer Connector, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-160XCSK	BD BodyGuard™ MicroSet Blue Striped Microbore Tubing, Spike, T Port, Female Luer Connector, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-160XSFMK	BD BodyGuard™ MicroSet Microbore Tubing, Spike, Inline 1.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
121-160XCSEK	BD BodyGuard™ MicroSet Blue Striped Microbore Tubing, Spike, Y Port, Female Luer Connector, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
PCA00001	BD BodyGuard™ MicroSet Blue Striped Microbore Tubing, Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
PCA00002	BD BodyGuard™ MicroSet Low Sorbing Microbore Tubing, Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
PCA00003	BD BodyGuard™ MicroSet Low Sorbing Microbore Tubing, Spike, Inline 0.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
PCA00004	BD BodyGuard™ MicroSet Blue Striped Microbore Tubing, Spike, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
PCA00005	BD BodyGuard™ MicroSet Blue Striped Microbore Tubing, 150ml EVA Bag, Inline 0.2 µm Filter, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
100-160XE	BD BodyGuard™ MicroSet Yellow Striped Microbore Tubing, Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28

Model Number/Catalogue number	Device Name	Certificate numberⁱ	Expiry date on the certificateⁱⁱ	Extended expiry dateⁱⁱⁱ
100-163XE90SK	BD BodyGuard™ MicroSet Yellow Striped Microbore Tubing, 90° Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
100-163XESK	BD BodyGuard™ MicroSet Yellow Striped Microbore Tubing, Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
100-163XESVK	BD BodyGuard™ MicroSet Yellow Striped Microbore Tubing, Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
101-163XE90SK	BD BodyGuard™ MicroSet Microbore Tubing, 90° Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
EPI00001	BD BodyGuard™ MicroSet Yellow Striped Microbore Tubing, Spike, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
EPI00003	BD BodyGuard™ MicroSet Yellow Stripped Microbore Tubing, Spike, Inline 0.2 µm filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
130-163XE90SK	BD BodyGuard™ MicroSet Green Microbore Tubing, 90° Spike, Anti Siphon, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-112XPEFKY	BD BodyGuard™ MicroSet Low Sorbing Microbore Tubing, Spike, Inline 1.2 µm Filter, Y Port, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-112XSFK	BD BodyGuard™ MicroSet Microbore Tubing, Spike, Manual Priming Valve, Inline 1.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-124XSFK	BD BodyGuard™ MicroSet Microbore Tubing, Spike, Y Port, Inline 1.2 µm Filter, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
TPN00001	BD BodyGuard™ MicroSet Microbore Tubing, Spike, Inline 1.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28

Model Number/Catalogue number	Device Name	Certificate numberⁱ	Expiry date on the certificateⁱⁱ	Extended expiry dateⁱⁱⁱ
TPN00003	BD BodyGuard™ MicroSet Microbore Tubing, Spike, Inline 1.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
TPN00004	BD BodyGuard™ MicroSet Light Resistant Microbore Tubing, Inline 1.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
TPN00005	BD BodyGuard™ MicroSet Microbore Tubing, Vented Spike, Inline 1.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
ONC00002	BD BodyGuard™ MicroSet Microbore Tubing, 2 Drip Chambers, Y Port, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
ONC00004	BD BodyGuard™ MicroSet Low Sorbing Microbore Tubing, Vented Spike, Y Port, Back Check Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
ONC00004UV	BD BodyGuard™ MicroSet Sorbing Light Resistant Microbore Tubing, Vented Spike, Y Port, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
ONC00005	BD BodyGuard™ MicroSet Low Sorbing Microbore Tubing, Drip chamber, Y Port, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
ONC00006	BD BodyGuard™ MicroSet Low Sorbing Microbore Tubing, Drip Chamber, Inline 0.2 µm Filter, Y Port, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
ONC00008	BD BodyGuard™ MicroSet Low Sorbing Microbore Tubing, Anti Siphon Valve, 5 SmartSite Ports, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
ONC00010	BD BodyGuard™ MicroSet Low Sorbing Light Resistant Microbore Tubing, Anti Siphon Valve 5 SmartSite Ports, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28

Model Number/Catalogue number	Device Name	Certificate numberⁱ	Expiry date on the certificateⁱⁱ	Extended expiry dateⁱⁱⁱ
ONC00011	BD BodyGuard™ MicroSet Low Sorbing Microbore Tubing, Anti Siphon Valve, 3 SmartSite Ports, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
ONC00012	BD BodyGuard™ MicroSet Low Sorbing Light Resistant Microbore Tubing, Anti Siphon Valve 3 SmartSite Ports, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
120-000TF	BD BodyGuard™ MicroSet Microbore Tubing, 2 Spikes, Inline 1.2 µm Filter, Anti-Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	31-Dec-28
EPI00011-NRF	BD BodyGuard™ MicroSet NR Fit Yellow Striped Microbore Tubing, 90° Spike, Anti Siphon Valve, Male NRFit Connector	2122031CE02	01-Jun-23	31-Dec-28
EPI00012-NRF	BD BodyGuard™ MicroSet NR Fit Yellow Microbore Tubing, 90° Spike, Anti Siphon Valve, Male NRFit Connector	2122031CE02	01-Jun-23	31-Dec-28
7290012271045	MicroSet Blood; Blood Set, Microbore Tubing, Drip Chamber, 2 Y Ports, Anti Siphon Valve, Female Luer Connector, Male Luer Connector	2122031CE02	01-Jun-23	26-May-24
M100-172SL	BD BodyGuard™ MicroSet Syringe Microbore Tubing, Female Luer Connector, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	26-May-24
M100-172SB	BD BodyGuard™ MicroSet Syringe Microbore Tubing, Female Luer Connector, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	26-May-24
7290012271014	BD BodyGuard™ MicroSet Microbore Tubing, Drip Chamber, 2 Y Ports, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	26-May-24
7290012271069	BD BodyGuard™ MicroSet Blue Stripe Microbore Tubing, Inline 0.2 µm Filter, Anti Siphon Valve, Male Luer Connector	2122031CE02	01-Jun-23	26-May-24

ⁱ Number of certificate issued under Directive 93/42/EEC (MDD)

ⁱⁱ The expiry date on the certificate issued under Directive 93/42/EEC (MDD)

ⁱⁱⁱ Extended expiry date according to Article 120 3b and 3c, Regulation (EU) 2017/745 (MDR)

Certificate Of Completion

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You may contact us to let us know of your changes as to how we may contact you electronically, to request paper copies of certain information from us, and to withdraw your prior consent to receive notices and disclosures electronically as follows:

To contact us by email send messages to: dewey.phan@bd.com

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To let us know of a change in your email address where we should send notices and disclosures electronically to you, you must send an email message to us at dewey.phan@bd.com and in the body of such request you must state: your previous email address, your new email address. We do not require any other information from you to change your email address.

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ii. send us an email to dewey.phan@bd.com and in the body of such request you must state your email, full name, mailing address, and telephone number. We do not need any other information from you to withdraw consent.. The consequences of your withdrawing consent for online documents will be that transactions may take a longer time to process..

Required hardware and software

The minimum system requirements for using the DocuSign system may change over time. The current system requirements are found here: <https://support.docusign.com/guides/signer-guide-signing-system-requirements>.

Acknowledging your access and consent to receive and sign documents electronically

To confirm to us that you can access this information electronically, which will be similar to other electronic notices and disclosures that we will provide to you, please confirm that you have read this ERSD, and (i) that you are able to print on paper or electronically save this ERSD for your future reference and access; or (ii) that you are able to email this ERSD to an email address where you will be able to print on paper or save it for your future reference and access. Further, if you consent to receiving notices and disclosures exclusively in electronic format as described herein, then select the check-box next to 'I agree to use electronic records and signatures' before clicking 'CONTINUE' within the DocuSign system.

By selecting the check-box next to 'I agree to use electronic records and signatures', you confirm that:

- You can access and read this Electronic Record and Signature Disclosure; and
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