SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3050

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

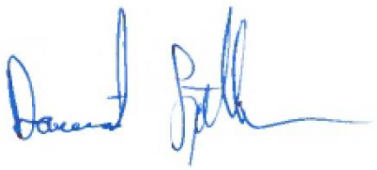
Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

This EU declaration of conformity is issued under the sole responsibility of Sjöbloms as manufacturer.

Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3110

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

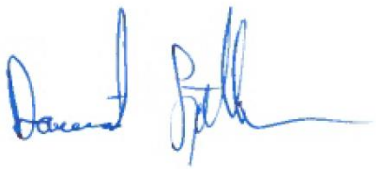
Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

This EU declaration of conformity is issued under the sole responsibility of Sjöbloms as manufacturer.

Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3500-000

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

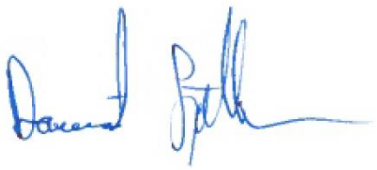
Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

This EU declaration of conformity is issued under the sole responsibility of Sjöbloms as manufacturer.

Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3500-001

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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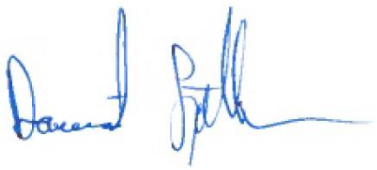
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3500-010

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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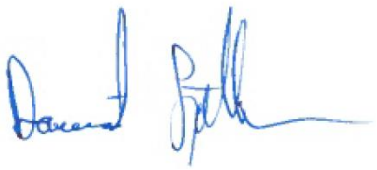
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3500-011

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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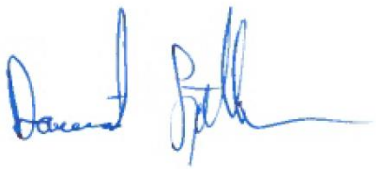
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3500-100

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

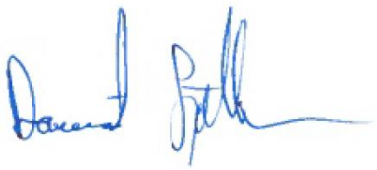
Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

This EU declaration of conformity is issued under the sole responsibility of Sjöbloms as manufacturer.

Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3500-101

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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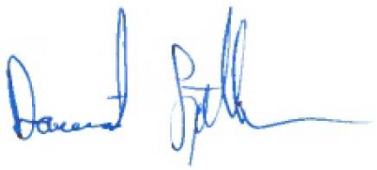
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3500-110

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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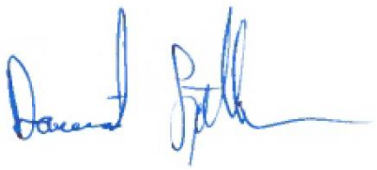
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3500-111

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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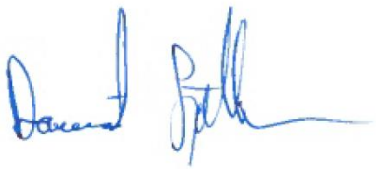
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3600

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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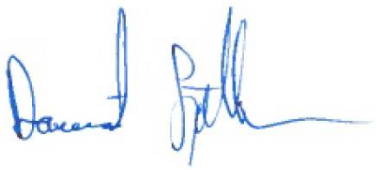
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3675

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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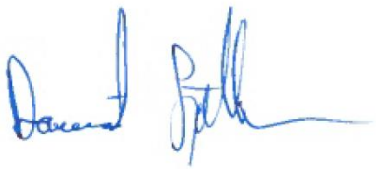
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3700

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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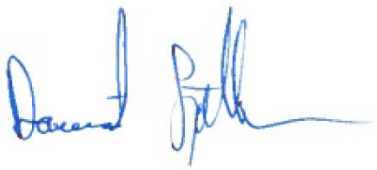
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand

Art no
080-3775

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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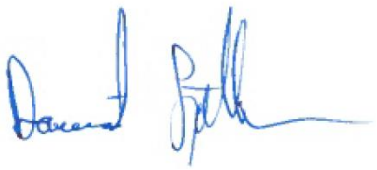
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand heavy duty

Art no
080-3900

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

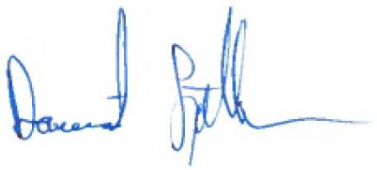
Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

This EU declaration of conformity is issued under the sole responsibility of Sjöbloms as manufacturer.

Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand heavy duty

Art no
080-3910

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

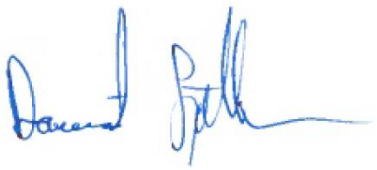
Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

This EU declaration of conformity is issued under the sole responsibility of Sjöbloms as manufacturer.

Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand heavy duty

Art no
080-3920

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

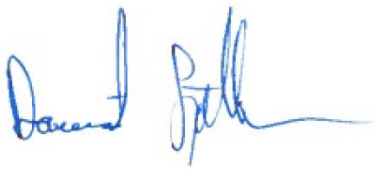
Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

This EU declaration of conformity is issued under the sole responsibility of Sjöbloms as manufacturer.

Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO

SJÖBLOMS

EU DECLARATION OF CONFORMITY

Product name
IV stand heavy duty

Art no
080-3930

Basic UDI-DI
73315087AZ

SRN
SE-MF-000005205

Intended purpose
The product is to be used in an examination room by a patient and is adjusted by the patient or member of the medical staff in a care situation for adults and children.

Manufacturer
Sjöbloms Sjukvårdsutrustning AB
Kavelvägen 4
S-894 35 SJÄLEVAD, SWEDEN

Risk class
MDR Class I

We hereby declare that the device covered by this declaration is in conformity with the EU Medical Device Regulation (2017/745, MDR) and, if applicable, with any other relevant Union legislation which contains provisions concerning the issuing of an EU declaration of conformity.

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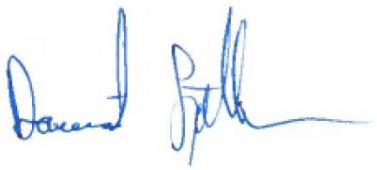
Örnsköldsvik, Sweden 2023-06-08



David Sjöblom
CEO