

**KSP ITALIA s.r.l**

Via dell'Artigianato, 1
06031 Bevagna (PG) - Italy
Tel. +39 0742 361947
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Capitale Sociale € 10.400,00 i.v. - REA n.160010
Iscri. Registro Imprese di Perugia n. 01793970540
Codice Fiscale / Partita Iva 01793970540
la società non è in stato di liquidazione

ksp@kspitalia.com
www.kspitalia.com

Declaration of Conformity

(in accordance with ISO/IEC 17050-1)

1) No. 040/26 - Date 02/02/2026

2) Issuer's name: **KSP ITALIA SRL**
Issuer's address: **VIA DELL'ARTIGIANATO 1, 06031 BEVAGNA (PG), ITALY Tel. 0742.36.19.47**
Fax 0742.36.19.46 www.kspitalia.com, e-mail: ksp@kspitalia.com

No. EUDAMED SRN: **IT-MF-000009316**

3) Object of the declaration: **WHEELED STAIR CLIMBER YACK, models N971.**

4) The Manufacturer KSP Italia declares under his sole responsibility that the medical device above described complies with all applicable requirements of the following legislation and fulfils all applicable provisions thereof (and any other relevant UE legislation providing for the issuance of EU declaration of conformity):

Documents No.	Title	Edition/Date of issue
5) Regulation (EU) 2017/745	Medical Devices Regulation	Emission: 5 April 2017
Directive 2011/65/EU	Restriction of the use of certain hazardous substances in electrical and electronic equipment	Emission: 8 June 2011

Additional information:

6) Medical devices designed and manufactured with quality management system compliant to ISO 13485.
CE Marked Medical device in accordance with Annex II and III, Regulation (UE) 2017/745.
Class I medical device as for rule I and rule 13, Regulation (UE) 2017/745, Annex VIII.
Registered at the Italian Ministry of Health with respective model number:
N971: 2384036.

BASIC UDI (GMN): 8055773180000YACK-N971T9

Signed for and on behalf of:

KSP Italia Srl

Bevagna, 02/02/2026

7) Maria Claudia Emanuelli,
Legal Representative

KSP ITALIA srl

Via dell'Artigianato, 1
06031 BEVAGNA (PG)

~~Tel. 0742.361947~~ Fax 0742.361946
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Declaration of Conformity (in accordance with ISO/IEC 17050-1)

1) No. 039/26 - Date 02/02/2026

2) Issuer's name: KSP ITALIA SRL
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Fax 0742.36.19.46 www.kspitalia.com, e-mail: ksp@kspitalia.com

No. EUDAMED SRN: IT-MF-000009316

3) Object of the declaration: WHEELED STAIR CLIMBER YACK, models N971H-N971H1.

4) The Manufacturer KSP Italia declares under his sole responsibility that the medical device above described complies with all applicable requirements of the following legislation and fulfils all applicable provisions thereof (and any other relevant UE legislation providing for the issuance of EU declaration of conformity):

Documents No.	Title	Edition/Date of issue
5) Regulation (EU) 2017/745	Medical Devices Regulation	Emission: 5 April 2017
Directive 2011/65/EU	Restriction of the use of certain hazardous substances in electrical and electronic equipment	Emission: 8 June 2011

Additional information:

6) Medical devices designed and manufactured with quality management system compliant to ISO 13485.
CE Marked Medical device in accordance with Annex II and III, Regulation (UE) 2017/745.
Class I medical device as for rule 1 and rule 13, Regulation (UE) 2017/745, Annex VIII.
Registered at the Italian Ministry of Health with respective model number:
N971H: 2384047; N971H1: 2384055.

BASIC UDI (GMN): 8055773180000YACK-N971T9

Signed for and on behalf of:

KSP Italia Srl

Bevagna, 02/02/2026

7) Maria Claudia Emanuelli,
Legal Representative

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Declaration of Conformity

(in accordance with ISO/IEC 17050-1)

1) No. 041/26 - Date 02/02/2026

2) Issuer's name: **KSP ITALIA SRL**
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No. EUDAMED SRN: **IT-MF-000009316**

3) Object of the declaration: **WHEELED STAIR CLIMBER YACK, models N972.**

4) The Manufacturer KSP Italia declares under his sole responsibility that the medical device above described complies with all applicable requirements of the following legislation and fulfils all applicable provisions thereof (and any other relevant UE legislation providing for the issuance of EU declaration of conformity):

Documents No.	Title	Edition/Date of issue
5) Regulation (EU) 2017/745	Medical Devices Regulation	Emission: 5 April 2017
Directive 2011/65/EU	Restriction of the use of certain hazardous substances in electrical and electronic equipment	Emission: 8 June 2011

Additional information:

6) Medical devices designed and manufactured with quality management system compliant to ISO 13485.
CE Marked Medical device in accordance with Annex II and III, Regulation (UE) 2017/745.
Class I medical device as for rule 1 and rule 13, Regulation (UE) 2017/745, Annex VIII.
Registered at the Italian Ministry of Health with respective model number:
N972: 2386614.

BASIC UDI (GMN): 805577318YACK-N972-973ZC

Signed for and on behalf of:

KSP Italia Srl

Bevagna, 02/02/2026

7) Maria Claudia Emanuelli,
Legal Representative


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Declaration of Conformity (in accordance with ISO/IEC 17050-1)

1) No. 038/26 - Date 02/02/2026

2) Issuer's name: KSP ITALIA SRL
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No. EUDAMED SRN: IT-MF-000009316

3) Object of the declaration: WHEELED STAIR CLIMBER YACK, models N972H-N972H1.

4) The Manufacturer KSP Italia declares under his sole responsibility that the medical device above described complies with all applicable requirements of the following legislation and fulfils all applicable provisions thereof (and any other relevant UE legislation providing for the issuance of EU declaration of conformity):

Documents No.	Title	Edition/Date of issue
5) Regulation (EU) 2017/745	Medical Devices Regulation	Emission: 5 April 2017
Directive 2011/65/EU	Restriction of the use of certain hazardous substances in electrical and electronic equipment	Emission: 8 June 2011

Additional information:

6) Medical devices designed and manufactured with quality management system compliant to ISO 13485.
CE Marked Medical device in accordance with Annex II and III, Regulation (UE) 2017/745.
Class I medical device as for rule I and rule 13, Regulation (UE) 2017/745, Annex VIII.
Registered at the Italian Ministry of Health with respective model number:
N972H: 2386617; N972H1: 2386618.

BASIC UDI (GMN): 805577318YACK-N972-973ZC

Signed for and on behalf of:

KSP Italia Srl

Bevagna, 02/02/2026

7) Maria Claudia Emanuelli,
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No. EUDAMED SRN: IT-MF-000009316

3) Object of the declaration: WHEELED STAIR CLIMBER YACK, models N973.

4) The Manufacturer KSP Italia declares under his sole responsibility that the medical device above described complies with all applicable requirements of the following legislation and fulfils all applicable provisions thereof (and any other relevant UE legislation providing for the issuance of EU declaration of conformity):

Documents No.	Title	Edition/Date of issue
5) Regulation (EU) 2017/745	Medical Devices Regulation	Emission: 5 April 2017
Directive 2011/65/EU	Restriction of the use of certain hazardous substances in electrical and electronic equipment	Emission: 8 June 2011

Additional information:

- 6) Medical devices designed and manufactured with quality management system compliant to ISO 13485.
CE Marked Medical device in accordance with Annex II and III, Regulation (UE) 2017/745.
Class I medical device as for rule 1 and rule 13, Regulation (UE) 2017/745, Annex VIII.
Registered at the Italian Ministry of Health with respective model number:
N973: 2386620.

BASIC UDI (GMN): 805577318YACK-N972-973ZC

Signed for and on behalf of:

KSP Italia Srl

Bevagna, 02/02/2026

7) Maria Claudia Emanuelli,
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Declaration of Conformity (in accordance with ISO/IEC 17050-1)

1) No. 037/26 - Date 02/02/2026

2) Issuer's name: KSP ITALIA SRL
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Fax 0742.36.19.46 www.kspitalia.com, e-mail: ksp@kspitalia.com

No. EUDAMED SRN: IT-MF-000009316

3) Object of the declaration: WHEELED STAIR CLIMBER YACK, models N973H-N973H1.

4) The Manufacturer KSP Italia declares under his sole responsibility that the medical device above described complies with all applicable requirements of the following legislation and fulfils all applicable provisions thereof (and any other relevant UE legislation providing for the issuance of EU declaration of conformity):

Documents No.	Title	Edition/Date of issue
5) Regulation (EU) 2017/745	Medical Devices Regulation	Emission: 5 April 2017
Directive 2011/65/EU	Restriction of the use of certain hazardous substances in electrical and electronic equipment	Emission: 8 June 2011

Additional information:

6) Medical devices designed and manufactured with quality management system compliant to ISO 13485.
CE Marked Medical device in accordance with Annex II and III, Regulation (UE) 2017/745.
Class I medical device as for rule 1 and rule 13, Regulation (UE) 2017/745, Annex VIII.
Registered at the Italian Ministry of Health with respective model number:
N973H: 2386626; N973H1: 2386627.

BASIC UDI (GMN): 805577318YACK-N972-973ZC

Signed for and on behalf of:

KSP Italia Srl

Bevagna, 02/02/2026

7) Maria Claudia Emanuelli,
Legal Representative

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