



Product PN GSS015A03S GSS015B03S	Mod. 984e-ext Rev. 07
Description HI-FLO Extension Set Adult 0.2 µm Non-Vented	

HI-FLO Extension Set Adult with 0.2 µm Non-vented Filter

PRODUCT DESCRIPTION	<p>Infusion extension set for infusion. The device is sterile and can be used in combination with an infusion set or an infusion device (pump or syringe). The connections are a female luer lock with vented cap inlet and a rotating male luer lock with cap outlet, compliant with ISO 80369-7. The set is provided with a non-vented Speedflow® Adult filter, with an hydrophilic PES membrane with 0.2 µm pore size for particles retention. The 'A' model set is provided with a male luer lock between the tube and the filter, while the 'B' model is provided with a male luer lock with ring between the tube and the filter. The approximate total length of the set is 282 mm. The set includes the following components:</p> <ul style="list-style-type: none"> ▪ Female luer lock with vented cap inlet; ▪ Male luer Lock / Male luer lock with ring; ▪ Non-vented Speedflow Adult 0.2 µm IV Filter with male luer lock; ▪ Protective cap.
CONFIGURATIONS	<p>GSS015A03S 0.2 µm with male luer lock GSS015B03S 0.2 µm with male luer lock with ring</p>
MANUFACTURER NAME	<p>GVS SpA Via Roma, 50 40069 Zola Predosa (BO) – ITALY Phone: +39.051.6176311 – Fax: +39.051.6176200 e-mail: gvs@gvs.it – website: www.gvs.com</p>
INTENDED USE / APPLICATION	<p>Infusion of Drugs (including chemotherapy), Anesthetics, Nutritional Solutions and Saline Solutions. The set can be connected via Luer Locks to all commercially available infusion administration sets compliant with ISO 80369-7. The set can be used either for gravity or in combination with an infusion set or an infusion device (pump or syringe). The set is intended specifically to eliminate air bubbles and to retain particles and bacteria. The set is a single-use device that can be used applications that last up to 96 hour. The set has to be disposed after each therapy. Sets with 0.2 µm filter cannot be used for infusion of hyper alimentation liquids, lipids, blood and/or blood derivatives. The device should only be supervised and used by qualified healthcare personnel.</p>
CLASS OF THE PRODUCT	<p>Class IIa – sterile – single use Rule 2 and 3 Annex VIII 2017/472/UE</p>
EMDN	<p>A03020101 LOW PRESSURE EXTENSION LINES</p>
MATERIALS	<p>Filter media: PES Hydrophilic membrane Filter housing: MBS Tubes: PVC (DEHP Free) Luer Lock: PVC / MABS Caps: HDPE</p> <p>Regulatory Compliance:</p> <ul style="list-style-type: none"> ▪ Biocompatibility according to ISO 10993-1 ▪ Rohs directive 2011/65/UE ▪ DEHP plasticizer Free ▪ Latex free ▪ Reach 1907/2006/CE (hazardous substances regulation) ▪ Reg. 1272/2008/CE (classification, labelling and packaging of substances and mixtures)



PRODUCT SPECIFICATION



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PRODUCT CHARACTERISTICS	Physical/Mechanical: Approximate total length: 282 mm Total internal volume of the set: 6 ml Weight: 10.8 g Input/output connectors: Female luer lock with vented cap inlet compliant with ISO 80369-7 Rotating male luer lock with cap outlet compliant with ISO 80369-7 Operating temperature Range: From 5 °C to 40 °C Storage temperature Range: From 0 °C to 40 °C Maximum applicable pressure: 8 bar Biological: Biocompatibility: Compliant with ISO 10993-1 Features: Type of administration: gravity / pressure Duration of the application: up to 96 hours Filter: Speedflow Adult 0.2 µm non-vented Filter pore size: 0.2 µm Filter internal volume: 1.2 ml Filter media: hydrophilic PES membrane Flow Regulator: No Drip Chamber: No Roller: No Y-injection site: No Clamp: No Tubing: PVC tube – inner Ø 3.0 mm / outer Ø 4.1 mm
	PRODUCT SHELF LIFE 5 years
STERILIZATION	Sterile: Yes – EtO Suitable for Sterilization/Re-sterilization: No
APPLICABLE STANDARDS AND REGULATIONS	Product Certification: CE mark Applicable Standards and Technical Regulations: EN 556-1 Sterilization of medical devices — Requirements for medical devices to be designated "STERILE" — Part 1: Requirements for terminally sterilized medical devices EN ISO 8536-4 Infusion equipment for medical use — Part 4: Infusion sets for single use, gravity feed EN ISO 8536-9 Infusion equipment for medical use — Part 9: Fluid lines for single use with pressure infusion equipment EN ISO 8536-11 Infusion equipment for medical use — Part 11: Infusion filters for single use with pressure infusion equipment EN ISO 10993-1 Biological evaluation of medical devices — Part 1: Evaluation and testing within a risk management process EN ISO 10993-4 Biological evaluation of medical devices — Part 4: Selection of tests for interactions with blood EN ISO 10993-5 Biological evaluation of medical devices — Part 5: Tests for in vitro cytotoxicity EN ISO 10993-7 Biological evaluation of medical devices — Part 7: Ethylene oxide sterilization residuals EN ISO 10993-10 Biological evaluation of medical devices — Part 10: Tests for irritation and skin sensitization EN ISO 10993-11 Biological evaluation of medical devices — Part 11: Tests for systemic toxicity EN ISO 10993-18 Biological evaluation of medical devices — Part 18: Chemical characterization of materials EN ISO 10993-23 Biological evaluation of medical devices — Part 23: Tests for irritation EN ISO 11135 Sterilization of health-care products — Ethylene oxide — Requirements for the development, validation and routine control of a sterilization process for medical devices EN ISO 11607-1 Packaging for terminally sterilized medical devices — Part 1: Requirements for materials, sterile barrier systems and packaging systems



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	<p>EN ISO 11607-2 Packaging for terminally sterilized medical devices — Part 2: Validation requirements for forming, sealing and assembly processes</p> <p>EN ISO 13485 Medical devices — Quality management systems — Requirements for regulatory purposes</p> <p>EN ISO 14971 Medical devices — Application of risk management to medical devices</p> <p>EN ISO 15223-1 Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied - Part 1: General requirements</p> <p>EN ISO 20417 Medical devices — Information to be supplied by the manufacturer</p> <p>IEC 62366-1 Medical devices — Part 1: Application of usability engineering to medical devices</p> <p>EN ISO 80369-1 Small-bore connectors for liquids and gases in healthcare applications — Part 1: General requirements</p> <p>EN ISO 80369-7 Small-bore connectors for liquids and gases in healthcare applications — Part 7: Connectors for intravascular or hypodermic applications</p> <p>EN ISO 80369-20 Small-bore connectors for liquids and gases in healthcare applications — Part 20: Common test methods</p>
INSTRUCTIONS FOR USE	Available languages: English / Italian / German / French / Spanish
PACKAGING	<p>Primary Packaging: Devices are individually packed and label in medical paper bags. Bag Size: 130 X 200 mm</p> <p>Secondary Packaging: Bags are placed inside a microperforated sack.</p> <p>Tertiary Packaging: Carton Box. Box Size: 60 x 40 x 20 cm Box Weight: 1.8 Kg Devices per box: 100</p>
CERTIFICATIONS	ISO 9001:2015 ISO 13485:2016 CE Certificate (2017/745/UE)

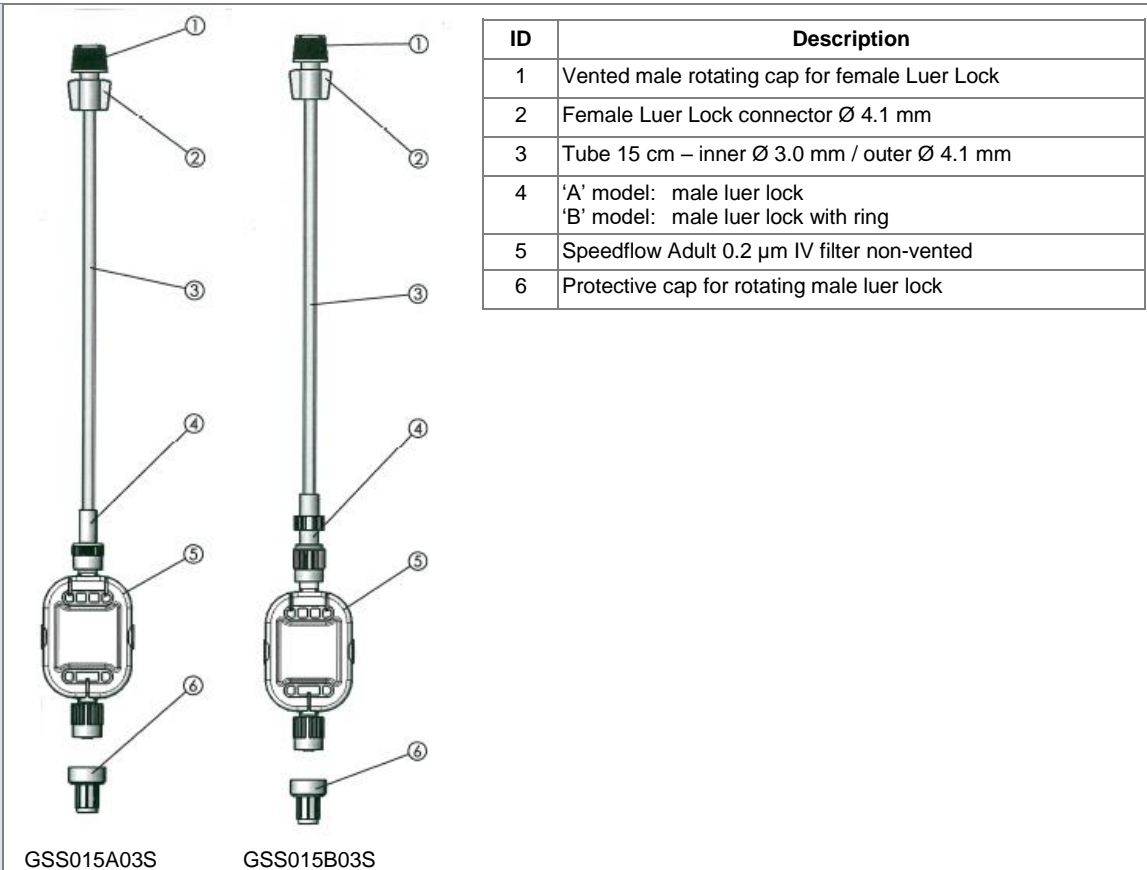
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Rev. 07

DRAWING



REVISIONS AND APPROVALS:

DATE	REV.	REASON FOR CHANGE	ISSUED BY: (name/function/signature)	VERIFIED BY: (name/function/signature)	APPROVED BY: (name/function/signature)
26/02/2023	00	First issue	Barbara Palmieri RA Specialist 	Barbara Finessi QA Manage 	Luca Zanini VP Healthcare and Lifesience