PRODUCT SPECIFICATION

CE 0051

Product PN	**GSS005A03S GSS005B03S**	**Mod. 984e-ext**
Description	**HI-FLO Extension Set Adult 0.2 µm Non-Vented Y-Infusion connector**	**Rev. 07**

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

PRODUCT DESCRIPTION	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 a 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 set is provided with a y-injection site, for administration of additional drugs by means of a syringe using the same infusion line. The approximate total length of the set is 480 mm. The set includes the following components: ▪ Female luer lock with vented cap inlet; ▪ Y-Injection point; ▪ Male luer Lock / Male luer lock with ring; ▪ Non-vented Speedflow Adult 0.2 µm IV Filter with male luer lock; ▪ Protective cap.
CONFIGURATIONS	**GSS005A03S** 0.2 µm with male luer lock **GSS005B03S** 0.2 µm with male luer lock with ring
MANUFACTURER NAME	**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
INTENDED USE / APPLICATION	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.
CLASS OF THE PRODUCT	Class IIa – sterile – single use Rule 2 and 3 Annex VIII 2017/472/UE
EMDN	A03020101 LOW PRESSURE EXTENSION LINES
MATERIALS	**Filter media:** PES Hydrophilic membrane **Filter housing:** MBS **Tubes:** PVC (DEHP Free) **Y-injection:** SEBS / MABS **Luer Lock:** PVC / MABS **Caps:** HDPE **Regulatory Compliance:** ▪ Biocompatibility according to ISO 10993-1 ▪ Rohs directive 2011/65/EU ▪ DEHP plasticizer Free

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	▪ Latex free ▪ Reach 1907/2006/CE (hazardous substances regulation) ▪ Reg. 1272/2008/CE (classification, labelling and packaging of substances and mixtures)
PRODUCT CHARACTERISTICS	**Physical/Mechanical:** **Approximate total length:** 480 mm **Total internal volume of the set:** 10 ml **Weight (approx.):** 12 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 0.2/1.2 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:** Yes **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

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	EN ISO 11607-1	Packaging for terminally sterilized medical devices — Part 1: Requirements for materials, sterile barrier systems and packaging systems
	EN ISO 11607-2	Packaging for terminally sterilized medical devices — Part 2: Validation requirements for forming, sealing and assembly processes
	EN ISO 13485	Medical devices — Quality management systems — Requirements for regulatory purposes
	EN ISO 14971	Medical devices — Application of risk management to medical devices
	EN ISO 15223-1	Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied - Part 1: General requirements
	EN ISO 20417	Medical devices — Information to be supplied by the manufacturer
	IEC 62366-1	Medical devices — Part 1: Application of usability engineering to medical devices
	EN ISO 80369-1	Small-bore connectors for liquids and gases in healthcare applications — Part 1: General requirements
	EN ISO 80369-7	Small-bore connectors for liquids and gases in healthcare applications — Part 7: Connectors for intravascular or hypodermic applications
	EN ISO 80369-20	Small-bore connectors for liquids and gases in healthcare applications — Part 20: Common test methods
INSTRUCTIONS FOR USE	Available languages: English / Italian / German / French / Spanish	
PACKAGING	**Primary Packaging:** Devices are individually packed and label in medical paper bags. Bag Size: 130 X 200 mm **Secondary Packaging:** Bags are placed inside a microperforated sack. **Tertiary Packaging:** Carton Box. Box Size: 60 x 40 x 20 cm Box Weight: 1.9 Kg Devices per box: 100	
CERTIFICATIONS	**ISO 9001:2015** **ISO 13485:2016** **CE Certificate (2017/745/UE)**	

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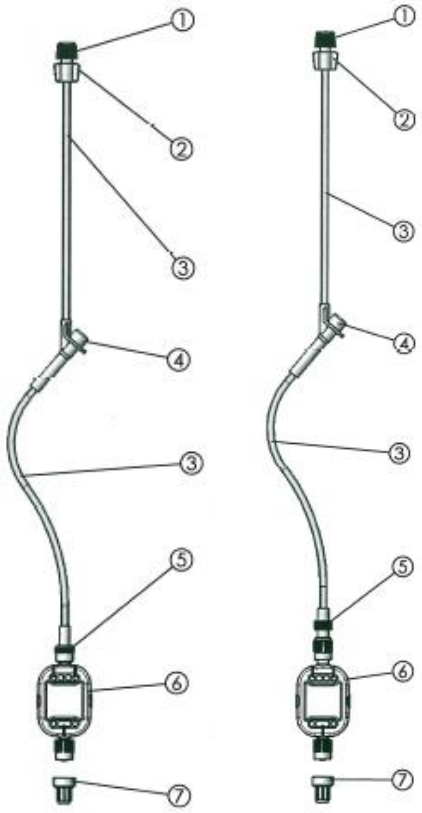
DRAWING



GSS005A03S GSS005B03S

ID	Description
1	Vented male rotating cap for female Luer Lock
2	Female Luer Lock connector Ø 4.1 mm
3	Tube 15 cm – inner Ø 3.0 mm / outer Ø 4.1 mm
4	Y-Injection site
5	'A' model: male luer lock 'B' model: male luer lock with ring
6	Speedflow Adult 0.2 µm IV filter non-vented
6	Protective cap for rotating male luer lock

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