

BLOOD COLLECTION TUBES TECHNICAL DATA SHEET

VACUSERA® Coagulation 9NC 3.2% Sodium Citrate Blood Collection Tubes



Ref No	GMDN Code	Draw Vol (ml)	Label Type	Additive	Tube Size (mm)	Tube Type	Color	Cap Color
234501	42585	1	Paper	Sodium Citrate 3.2%	13x75	Double Tube	Blue	0
2345015	42585	1	Paper	Sodium Citrate 3.2%	13x75	Single Tube		
234502	42585	1.8	Paper	Sodium Citrate 3.2%	13x75	Double Tube		
2345025	42585	1.8	Paper	Sodium Citrate 3.2%	13x75	Single Tube		
234503	42585	2.7	Paper	Sodium Citrate 3.2%	13x75	Double Tube		
234503S	42585	2.7	Paper	Sodium Citrate 3.2%	13x75	Single Tube		
234504	42585	3.6	Paper	Sodium Citrate 3.2%	13x75	Single Tube		
234505	42585	4.5	Paper	Sodium Citrate 3.2%	13x75	Single Tube		
235505	42585	5	Paper	Sodium Citrate 3.2%	13x100	Single Tube		
434501	42585	1	Transparent	Sodium Citrate 3.2%	13x75	Double Tube		
434501S	42585	1	Transparent	Sodium Citrate 3.2%	13x75	Single Tube		
434502	42585	1.8	Transparent	Sodium Citrate 3.2%	13x75	Double Tube		
434502S	42585	1.8	Transparent	Sodium Citrate 3.2%	13x75	Single Tube		

DİSERA TIBBİ MALZEME LOJİSTİK SANAYİ VE TİCARET A.

GAZİEMİR VD.: 301 053 3601 Tic.Sic.No: Merkez 123171 K - 10769 Mersis No: 0301053360100013

 Merkez : Karabağlar Mah. 5758 Sk. No:4 H/11 - Karabağlar / İZMİR
 Tel : +90 (0232) 264 66 68 info@disera.com.tr

 Fabrika : İbni Melek Mah. Tosbi Yol 5 Sk. No: 46 Tire / İZMİR
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434503	42585	2.7	Transparent	Sodium Citrate 3.2%	13x75	Double Tube
434503S	42585	2.7	Transparent	Sodium Citrate 3.2%	13x75	Single Tube
434504	42585	3.6	Transparent	Sodium Citrate 3.2%	13x75	Single Tube
434505	42585	4.5	Transparent	Sodium Citrate 3.2%	13x75	Single Tube
435505	42585	5	Transparent	Sodium Citrate 3.2%	13x100	Single Tube

Ref No	GMDN Code	Draw Vol (ml)	Label Type	Additive	Tube Size (mm)	Tube Type	Color	Cap Color
234502DLB	42585	1.8	Paper	Sodium Citrate 3.2%	13x75	Double Tube	Light Blue	
234503DB	42585	2.7	Paper	Sodium Citrate 3.2%	13x75	Double Tube	Blue	0
234504DMB	42585	3.6	Paper	Sodium Citrate 3.2%	13x75	Double Tube	Dark Blue	0

1. Intended Use:

VACUSERA® Coagulation Tubes are filled with buffered tri-sodium citrate solution. Citrate concentrations of either 0.109 mol/l (3.2 %) or 0.129 mol/l (3.8 %) are available. The choice of the concentration depends upon the policies of the laboratories. The mixing ratio is 1 part citrate to 9 parts blood.

VACUSERA® Coagulation Tubes – Double Tube combines different capabilities of different plastics. The outer tube is made out of PET (polyethylenetherephtalate) and it guarantees protection of vacuum until the expiry date. The inner tube is made out of PP (polypropylene) which prevents gradual evaporation of liquid additives.

VACUSERA® Coagulation Tubes are used for coagulation tests.

2. Information About The Manufacturer

Manufacturer: Disera Tıbbi Malzeme Lojistik Sanayi ve Ticaret A.Ş

Country of Origin: Turkey

EN ISO 13485:2016 Cert. n 31723701 Released by Szutest

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3. Applicable Standards

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No	EXPLANATION OF STANDARD
1	EN ISO 13485:2016 Medical devices — Quality Management Systems — Requirements for Regulatory Purposes
2	EN ISO 15223-1:2021 Medical Devices — Symbols to be Used with Information to be Supplied by The Manufacturer — Part 1: General Requirements
3	EN ISO 20417:2021 Medical Devices - Information to be Supplied by the Manufacturer
4	EN ISO 14971:2019 Medical Devices — Application of Risk Management to Medical Devices
5	EN 62366-1:2015 Medical Devices - Part 1: Application of Usability Engineering to Medical Devices
6	EN ISO 11137-1:2015 Sterilization of Health Care Products - Radiation - Part 1: Requirements for Development, Validation and Routine Control of a Sterilization Process for Medical Devices
7	EN ISO 11137-2:2015 Sterilization of Health Care Products — Radiation — Part 2: Establishing the Sterilization Dose
8	EN ISO 11137-3:2017 Sterilization of health care products - Radiation - Part 3: Guidance on dosimetric aspects of development, validation and routine control
9	EN ISO 11737-1:2018 Sterilization of health care products - Microbiological methods - Part 1: Determination of a population of microorganisms on products
10	EN ISO 11737-2:2020 Sterilization of Health Care Products — Microbiological Methods — Part 2: Tests of Sterility Performed in The Definition, Validation and Maintenance of a Sterilization Process
11	ASTM F 1980-16 Standard Guide for Accelerated Aging of Sterile Barrier Systems and Medical Devices

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12	EN ISO 6710 :2017 Single-Use Containers for Human Venous Blood Specimen Collection
13	EN 556-1:2001 Sterilization of medical devices - Requirements for medical devices to be designated "Sterile" - Part 1: Requirements for terminally sterilised medical devices
14	EN ISO 14644-1:2015 Cleanrooms and associated controlled environments - Part 1: Classification of air cleanliness by particle concentration
15	EN ISO 14644-2:2015 Cleanrooms And Associated Controlled Environments — Part 2: Monitoring to Provide Evidence of Cleanroom Performance Related to Air Cleanliness by Particle Concentration
16	EN ISO 14644-3:2019 Cleanrooms And Associated Controlled Environments — Part 3: Test Methods
17	98/79/EC Directive of In vitro diagnostic medical devices
18	EN 13612: 2002 Performance evaluation of in vitro diagnostic medical devices
19	EN ISO 18113 -1 In vitro diagnostic medical devices - Information supplied by the manufacturer (labelling)

4. **Product Specifications**

Tube material:	Polyethylene teraphthalate (PET) Polypropylene (PP)
Cap material (outer shield):	Polyethylene
Rubber material (inner cap):	Bromobutyl elastomers
Additives:	Sodium Citrate 3.2% (The blood to additive ratio is 9:1)
Additive Filling Form	Liquid
Fill indicator:	Yes
Expiry (Single Tubes):	4 months
Expiry (Double Tubes):	9 months
Storage conditions:	Do not expose to direct sunlight. Store between +4 and 35 °C.

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5. Physical Properties

a) Raw Materials:

Outer tubes are made from PET (polyethylene terephthalate) material. They should be colorless and transparent.

Tube caps are made from PE (polyethylene) material. Their colors may differ according to the additive that the tube contains.

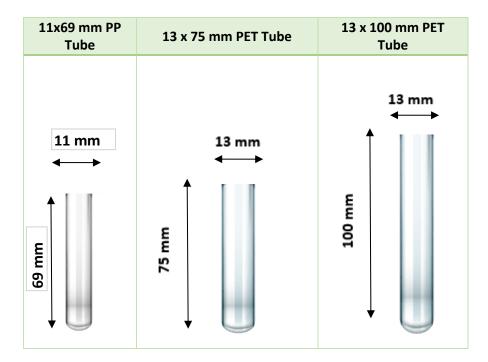
Inner tubes are made from polypropylene material.

Rubber stoppers are made from butyl elastomers.

All components are latex-free.

b) Dimensions

There are two different types of PET tubes according to their dimensions and there is one type of PP tube.



There are PE caps with two different diameters (13mm & 16mm) that could fit into these PET tubes.

Accordingly, there are stoppers with 13 mm and 16 mm diameter that could fit in these PE caps and PET tubes.

All detailed dimensions are represented in the technical drawings.

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6. Packaging

100 pieces of VACUSERA® Blood Collection Tubes are lined up in a styrofoam and wrapped with a shrink film. Each of these packages is called as rack. 12 racks are placed in a carton box and each box contains 1200 tubes.

Outer box dimensions for 13x75mm tubes: 36.0 X 53.5 X 18.5 mm Outer box dimensions for 13x100mm tubes: 36.0 X 53.5 X 24.0 mm

7. Sterilization

VACUSERA® Blood Collection Tubes are placed in the market as sterile. Products are sterilized using radiation sterilization. Sterilization process shall be validated.

Internamente sterility: SAL 10^{-6} (SAL= sterility assurance level) Standards: EN ISO 11137-2:2015; EN 556:2001/AC:2006

8. Label Information

	Tube Label	Rack Label	Carton Label
Company name and address of the manufacturer	+	+	+
Ref No (Product Code)	+	+	+
Lot No	+	+	+
Sterile (symbol) and method of sterilization	+	+	+
CE mark and single use symbol	+	+	+
Fill volume	+	+	+
Expiration Date	+	+	+
Product name and short description	+	+	+
Instructions for use (pictograms)	-	+	+
Quantity per package	-	+	+
Storage instructions	-	+	+
Primary barcode	-	+	+
Secondary barcode	-	+	+
Sterility indicator	-	-	+

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Symbols on the label	Description of the Symbols	Tube Label	Rack Label	Box Label
REF	Reference Number	~	~	~
LOT	Batch Code	~	~	~
\square	Use by	~	~	~
②	Single Use	~	~	~
[巻]	Keep away from sunlight		~	~
STERILE R	Radiation Sterilization	~	~	~
Ωi	See the Instruction for Use		~	~
IVD	In Vitro Diagnostic Device	~	~	~
*	Temperature Limits		~	~
C€	CE symbol	~	~	~

9. Instructions For Use

For more detailed information and usage instructions, please check the IFU document.

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