# **CRYOSTORE<sup>™</sup> MULTI-CHAMBER BAG**



The OriGen CryoStore™ Multi-Chamber Bag is intended for freezing blood components.

# It provides you with the traceability and durability that you need.

## TRACEABILITY

Each chamber of the bag is printed with unique numbers for easy identification. The numbers on the chambers match those on the tube set ensuring traceability.

#### DURABILITY

The OriGen CryoStore<sup>™</sup> Bag is the industry-preferred cryogenic freezing bag. The spike port and film used in every CryoStore<sup>™</sup> Multi-Chamber Bag are made from a robust EVA.

#### LABORATORY SAFETY

Tube sets have luer-actuated needle-free injection ports. These swabable ports help maintain sterility of the sample through numerous actuations and reduce the risk of needle sticks during processing.

#### **EASE OF USE**

The tube set has a syringe included for removal of air prior to filling the bag with any fluid. This will help expedite the debubbling process. Each multi-chamber bag has tubing compatible for sterile welding, needle-free access, and a male and a female luer connection.

### EFFICIENCY

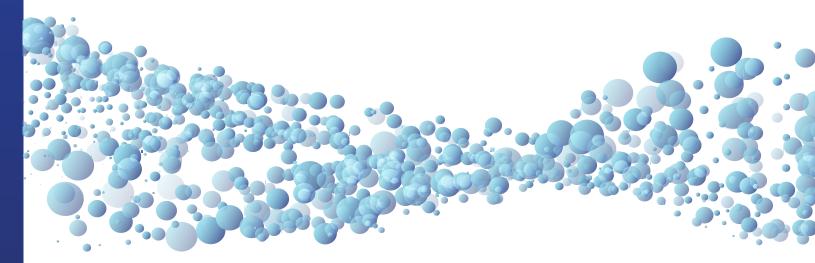
Multiple chambers are filled from a single location and the channel located at the bottom of the bag allows for easy heat seals between each chamber with a standard tube sealer. This separation of individual chambers allows multiple post thaw applications without compromising the integrity of the rest of the cellular product.

7000 Burleson Rd, Bldg D | Austin, Texas 78744 | 512.474.7278 | Toll Free 800.233.9014 | sales.us@origen.com | origen.com

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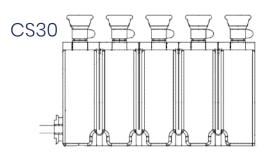
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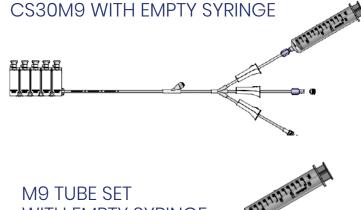


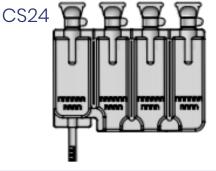
## **TECHNICAL INFORMATION**

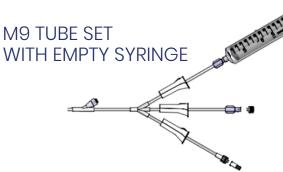
#### FOUR AND FIVE CHAMBER BAGS

PART CODE	NUMBER OF CHAMBERS	WIDTH (cm)	HEIGHT (cm)	DEPTH (cm)	FREEZE/FILL VOLUME PER CHAMBER	FREEZE/FILL VOLUME PER BAG	BAG PER TYVEK POUCH	POUCHES PER CASE
CS30M9	5	10.8	5.5	0.9	4-6ml	30ml	1	12
CS24M9	4	8.4	5.5	0.9	4-6ml	24ml	1	12









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CE Marked | FDA Cleared | ISO 13485 Certified | MDSAP Certified | Manufactured to GMP Guidelines ML21.02.R02; 2021-03

