

PRODUCT CATALOGUE

SepaSperm®

- For sperm preparation
 by the density gradient centrifugation.
- O Free from animal derivatives.
- O SepaSperm® Wash Solution is HEPES buffered.



	REF	Code	Contents
SepaSperm® Solution 100%	92143	SE100G-100	100mL
SepaSperm [®] Solution 80%	92073	SE80G-100	100mL
	92079	SE80G-50	50mL
SepaSperm [®] Solution 60%	92074	SE60G-100	100mL
	92080	SE60G-50	50mL
SepaSperm® Solution 40%	92075	SE40G-100	100ml
	92081	SE40G-50	50ml
SepaSperm® Wash Solution	92147	SEWSG-100	100ml

COMPONENTS

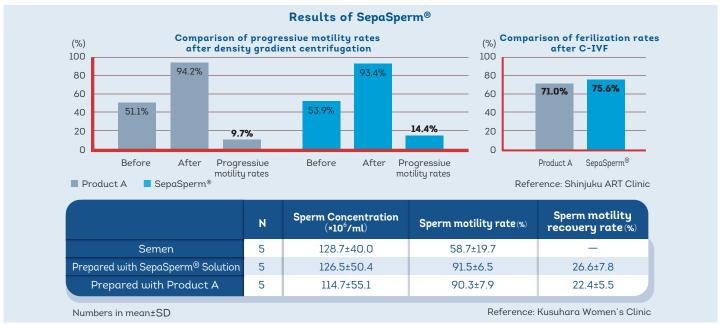
Calcium chloride / Gentamicin sulfate / Glucose / HEPES /
Magnesium sulfate / Potassium chloride / Potassium phosphate /
Recombinant human albumin / Silica particles / Sodium bicarbonate /
Sodium chloride / Sodium lactate / Sodium pyruvate

QUALITY CONTROL

pH 7.2-7.6 / Osmolarity 270-300 mOsm/L / Endotoxin <0.25EU/mL / Mouse Embryo Assay \ge 80% / Sperm Survival (24h) \ge 80% / Sperm Penetration \ge 3 / Sterility Test

Storage: 2-8℃ Shelf life: 18 months

RESULT



Specification may change without pre-notice for purpose of product improvement.

Kitazato Corporation

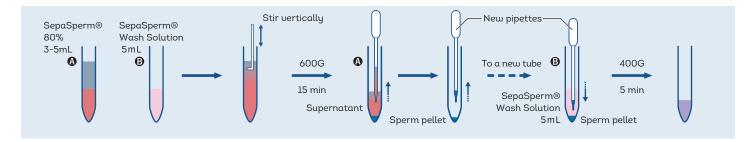
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An example of single-layer gradient centrifugation

- Bring all solutions to room temperature (20-25°C) before use. Prepare two 15mL centrifuge tubes.

 One of them with 3-5mL of SepaSperm® 80%. The other with 5mL of SepaSperm® Wash Solution.
- 2 Check the semen volume with a 5mL pipette or similar and evaluate the semen sample with a sperm count chamber.
- 3 Pipette semen on top of the prepared SepaSperm® 80%. Stir 2cm vertically around the layers between the semen and SepaSperm® 80%
- 4 Centrifuge for 15 minutes at approximately 600 x g. (Separation of sperm and seminal fluid)
 Depending on the state of collected sperm, adjust xg or centrifugation time accordingly.
- 5 After centrifugation, remove the supernatant with a new pipette.
- 6 Add 5mL of SepaSperm® Wash Solution (SEWS) to the sperm pellet and resuspend. Centrifuge at 400 x g for 5 minutes for washing. Depending on the state of collected sperm, adjust xg or centrifugation time accordingly.
- 7 Remove the supernatant leaving the sperm pellet only. Dilute the sperm pellet with SepaSperm® Wash Solution according to the following treatment.



An example of two-layer density gradient centrifugation

- Bring all the solutions to room temperature $(20-25^{\circ}C)$ before use. Prepare two 15mL centrifuge tubes. One of them with layers in the next order from bottom to up;
 - SepaSperm® 80% 1.5-2.0mL ②SepaSperm® 40% 1.5-2.0mL ③Liquified semen The other with 5mL of SepaSperm® Wash Solution.
- 2 Check the semen volume with a 5mL pipette or similar and evaluate the semen sample with a sperm count chamber.
- 3 Centrifuge the tube at 600 x g for 15 minutes. (Separation of sperm and seminal fluid)

 Depending on the state of collected sperm, adjust xg or centrifugation time accordingly.
- 4 Aspirate the sperm pellet.
- 5 Transfer the sperm pellet to a new centrifuge tube prepared with 5mL of SepaSperm® Wash Solution, and mix well.
- 6 Centrifuge at 400 x g for 5 minutes for washing.

 Depending on the state of collected sperm, adjust xg or centrifugation time accordingly.
- 7 Remove the supernatant leaving the sperm pellet only. Dilute the sperm pellet with SepaSperm® Wash Solution according to the following treatment.

