EasySep™ Human CD56 Positive Selection Kit II

For processing 5 x 10⁷ cells from muscle cultures

Catalog #17855

Positive Selection

Document #1000000694 | Version 01



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Description

Isolate highly purified CD56+ cells from fresh or previously frozen human muscle cultures (myoblasts and fibroblasts) in as little as 24 minutes by immunomagnetic positive selection.

- · Fast and easy-to-use
- Up to 95% purity
- No columns required

This kit targets CD56+ cells for positive selection with antibodies recognizing the CD56 surface marker. Desired cells are labeled with antibodies and magnetic particles, and separated without columns using an EasySep[™] magnet. Unwanted cells are simply poured off, while desired cells remain in the tube. Isolated cells are immediately available for downstream applications such as flow cytometry, culture, or DNA/RNA extraction.

• This is the Product Information Sheet (PIS) for isolating CD56+ cells from human muscle cultures. If isolating CD56+ cells from human peripheral blood mononuclear cells (PBMCs), refer to the applicable PIS (Document #1000005559).

Component Descriptions

COMPONENT NAME	COMPONENT #	QUANTITY	STORAGE	SHELF LIFE	FORMAT
EasySep™ Human CD56 Positive Selection Cocktail II	17855C	1 x 1 mL	Store at 2 - 8°C. Do not freeze.	Stable until expiry date (EXP) on label.	A combination of monoclonal antibodies in PBS. Includes an Fc receptor blocking antibody.
EasySep™ Dextran RapidSpheres™ 50100	50100	1 x 1 mL	Store at 2 - 8°C. Do not freeze.	Stable until expiry date (EXP) on label.	A suspension of magnetic particles in water.

PBS - phosphate-buffered saline

Components may be shipped at room temperature (15 - 25°C) but should be stored as indicated above.

Sample Preparation

MUSCLE CULTURES

Muscle cultures can be established from human skeletal muscle tissue using MyoCult™-SF Expansion Supplement Kit (Human; Catalog #05980). For complete instructions, refer to the applicable PIS, available at www.stemcell.com or contact us to request a copy.

Alternatively, human skeletal muscle cultures can be generated using the methods published by Agley CC et al. and Soriano-Arroquia A et al.¹⁻²

After preparation, resuspend cells at 5 x 10^6 cells/mL in recommended medium.

PERIPHERAL BLOOD OR LEUKAPHERESIS

If processing peripheral blood or leukapheresis samples, refer to the applicable PIS (Document #10000005559).

Recommended Medium

EasySep™ Buffer (Catalog #20144), or PBS containing 2% FBS and 1 mM EDTA. Medium should be free of Ca++ and Mg++.



Directions for Use – Manual EasySep™ Protocols

See page 1 for Sample Preparation and Recommended Medium. Refer to Table 1 for detailed instructions regarding the manual EasySep™ procedure for each magnet.

Table 1. EasySep™ Human CD56 Positive Selection Kit II Protocol

STEP	INSTRUCTIONS	EasySep™ (Catalog #18000)		
Prepare sample at the indicated cell concentration within the volume range.		5 x 10^6 cells/mL 0.2 - 1 mL		
•	Add sample to required tube.	5 mL (12 x 75 mm) polystyrene round-bottom tube (e.g. Catalog #38007)		
2	Add Selection Cocktail to sample.	100 µL/mL of sample		
2	Mix and incubate.	RT for 10 minutes		
3	Vortex RapidSpheres™. NOTE: Particles should appear evenly dispersed.	30 seconds		
4 Add Ra Mix and	Add RapidSpheres™ to sample.	50 µL/mL of sample		
	Mix and incubate.	RT for 5 minutes		
5	Add recommended medium to top up the sample to the indicated volume. Mix by gently pipetting up and down 2 - 3 times.	Top up to 2.5 mL		
	Place the tube (without lid) into the magnet and incubate.	RT for 3 minutes		
6	Pick up the magnet, and in one continuous motion invert the magnet and tube,** pouring off the supernatant. Remove the tube from the magnet; this tube contains the isolated cells.	Discard supernatant		
7	Repeat steps as indicated.	Steps 5 and 6, two more times (total of 3 x 3-minute separations)		
8	Resuspend cells in desired medium. Be sure to collect cells from the sides of the tube.	Isolated cells are ready for use		

RT - room temperature (15 - 25°C) * Leave the magnet and tube inverted for 2 - 3 seconds, then return upright. Do not shake or blot off any drops that may remain hanging from the mouth of the tube.

EasySep™ Human CD56 Positive Selection Kit II



Notes and Tips

ASSESSING PURITY

For purity assessment of CD56+ cells by flow cytometry, use one of the following fluorochrome-conjugated antibody clones:

- · Anti-Human CD56 Antibody, Clone HDC56 (Catalog #60021; partial blocking), or
- Anti-human CD56 antibody, clone CMSSB (partial blocking), or
- Anti-human CD56 antibody, clone NCAM16.2 (partial blocking)

The following method can also be used:

• Use a fluorochrome-conjugated secondary antibody, such as Goat Anti-Mouse Ig (H+L) Antibody, Polyclonal (Catalog #60138).

Data



Starting with muscle cultures at 5 - 10 days after derivation, the CD56+ cell content of the isolated fraction is typically 95 ± 4.3% (mean ± SD using the purple EasySep[™] Magnet). In the above example, the purities of the start and final isolated fractions are 57.38% and 95.18%, respectively.

References

- 1. Agley CC et al. (2015) Isolation and quantitative immunocytochemical characterization of primary myogenic cells and fibroblasts from human skeletal muscle. J Vis Exp (95): 52049.
- 2. Soriano-Arroquia A et al. (2017) Preparation and culture of myogenic precursor cells/primary myoblasts from skeletal muscle of adult and aged humans. J Vis Exp (120): e55047.

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