

MethoCult™ H4230

Methylcellulose-Based Medium Without Cytokines for Human Cells

Catalog #04230

80 mL



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FOR RESEARCH USE ONLY. NOT INTENDED FOR HUMAN OR ANIMAL DIAGNOSTIC OR THERAPEUTIC USES.

Product Description

Incomplete Methylcellulose-Based Medium for Colony-Forming Unit (CFU) Assays for Human Cells

MethoCult™ H4230 is recommended as a base medium for the culture of human cells to detect and quantify hematopoietic progenitors in human bone marrow (BM), mobilized peripheral blood (MPB), peripheral blood (PB,) and cord blood (CB) samples using CFU assays. This formulation allows for the addition of an exogenous source of erythropoietin (EPO) and other cytokines to meet the specific requirements of investigators.

MethoCult™ H4230 is also useful for other applications, including cloning of cell lines.

Properties

Storage: Store at -20°C.

Shelf Life: Stable until expiry date (EXP) on label.

Contains:

- Methylcellulose in Iscove's MDM
- Fetal bovine serum
- Bovine serum albumin
- 2-Mercaptoethanol
- Supplements

Directions for Use

NOTE: If product is received partially thawed, place immediately at -20°C or thaw and aliquot as described below.

PREPARATION OF COMPLETE METHOCULT™ H4230 MEDIUM

MethoCult™ H4230 does not contain EPO or other cytokines. These can be added directly to the bottle or to each tube after aliquoting. Refer to Table 1 for volumes required to prepare complete MethoCult™ H4230 medium per bottle or per tube. The ratio (v:v) of MethoCult™ to other components in liquid medium (e.g. cytokines) is important for viscosity, which ensures optimal CFU growth and morphology.

Use sterile techniques to prepare complete MethoCult™ H4230 medium (MethoCult™ H4230 base medium + desired components).

NOTE: Do not use pipettes to dispense methylcellulose as the volume dispensed will not be accurate. Syringes and large bore blunt-end needles should be used for accurate dispensing of viscous methylcellulose medium and to prevent needle-stick injuries.

A. TO PREPARE 100 ML BOTTLE

1. Thaw 80 mL bottle at room temperature (15 - 25°C) or overnight at 2 - 8°C. Do not thaw MethoCult™ at 37°C.
2. Prepare desired growth factors and supplements in Iscove's MDM (Catalog #36150) in a volume of 20 mL and add to MethoCult™ (for a total volume of 100 mL). Shake vigorously for 1 - 2 minutes and then let stand for at least 5 minutes to allow bubbles to rise to the top before aliquoting.
3. Using a 3 or 6 mL luer lock syringe attached to a 16 gauge Blunt-End Needle (Catalog #28110), aliquot 3 mL per tube for 1.1 mL duplicate cultures or 4 mL per tube for 1.1 mL triplicate cultures. Complete MethoCult™ medium is now ready for use.

B. TO PREPARE INDIVIDUAL TUBES

1. Thaw 80 mL bottle at room temperature (15 - 25°C) or overnight at 2 - 8°C. Do not thaw MethoCult™ at 37°C.
2. Shake vigorously for 1 - 2 minutes and then let stand for at least 5 minutes to allow bubbles to rise to the top before aliquoting.
3. Using a 3 or 6 mL luer lock syringe attached to a 16 gauge Blunt-End Needle (Catalog #28110), aliquot 2.4 mL per tube for 1.1 mL duplicate cultures or 3.2 mL per tube for 1.1 mL triplicate cultures.
NOTE: Before adding components, tubes of incomplete MethoCult™ medium may be stored at -20°C until expiry date as indicated on label. After thawing aliquoted tubes, add desired components (see step 4) and mix well.
4. Add desired growth factors and supplements in Iscove's MDM (Catalog #36150) to tubes of MethoCult™ H4230 (see Table 1 for required volumes).
5. Vortex tubes to mix well. Complete MethoCult™ medium is now ready for use.
6. Aliquot any remaining MethoCult™ H4230 base medium for duplicate or triplicate cultures (see Table 1 for required volumes), store at -20°C, then add desired components after thawing. Mix well before use.

Table 1: Volumes Required for Preparation of Complete MethoCult™ H4230 Medium

COMPONENT	PER BOTTLE	PER TUBE (duplicate 1.1 ml cultures)	PER TUBE (triplicate 1.1 ml cultures)
MethoCult™ H4230	80 mL	2.4 mL	3.2 mL
IMDM with cytokines*	20 mL	0.6 mL	0.8 mL
TOTAL VOLUME	100 mL	3.0 mL	4.0 mL

*For a complete list of available cytokines, refer to our website at www.stemcell.com.

For recommended cell plating concentrations, set-up of human CFU assays, and counting and classification of colonies, refer to the Technical Manual: Human Colony-Forming Unit Assays Using MethoCult™ (Document #28404), available on our website at www.stemcell.com or contact us to request a copy.

References

- Eaves CJ, Eaves AC: Anatomy and physiology of hematopoiesis. In: Childhood Leukemias (2nd Edition) (CH Pui, ed.), Cambridge University Press, Cambridge, pp 69-105, 2006
- Eaves C, Lambie K: Atlas of Human Hematopoietic Colonies, STEMCELL Technologies Inc., Vancouver, 1995 (Catalog #28700)
- Wognum B, Yuan N, Lai B, & Miller CL: Colony Forming Cell Assays for Human Hematopoietic Progenitor Cells. In: Basic Cell Culture Protocols (CD Helgason, CL Miller, eds.), Humana Press Inc., Clifton, New Jersey, p267-83, 2013.
- Nissen-Druey C, Tichelli A, Meyer-Monard S: Human Hematopoietic Colonies in Health and Disease, S. Karger Medical and Scientific Publishers, Basel, 2005. Reprint of Acta Haematol 113: 5-96, 2005 (Catalog #28760)

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