SOP: M027.2

Updated: 7/14/22 KE

Preparation of Complete DMEM/F12 Media for Hybridoma Growth

Materials and Reagents:

- 1. DMEM/F12 + Glutamax without HEPES (ThermoFisher/Fisher Scientific, Gibco Invitrogen #10565018)
- 2. L-glutamine, 200mM (100X), 100 mL bottle (Fisher Scientific Cat # 25030081)
- 3. Antibiotic-Antimycotic 100X liquid, 100 mL bottle (Fisher Scientific Cat # 15240-062)
- 4. β-Mercaptoethanol 1000X liquid, 50 mL bottle (Life Technologies Corporation Cat # 21985-023)
- 5. Fetal Bovine Serum, Not heat inactivated (Sigma Cat# F2442-500mL, also available through PMF/ARCBIO)
- 6. NCTC-109 media supplement, 500mL bottle (Fisher Scientific Cat# 21340-039)
- 7. Sodium Pyruvate, 100 mM, 100mL bottle (Hyclone, GE Healthcare Life Sciences Cat# SH30239.01)
- 8. Oxaloacetic Acid, 1 mM, 5G bottle (Sigma Cat# O9504)
- 9. Insulin, Bovine Pancreas, 5 mL bottle (Sigma Cat# I0516-5mL)
- 10. HAT media supplement, 50X, 100 mL bottle (Gibco, Fisher Scientific Cat # 21060017)
- 11. HT media supplement, 100X, 50 mL bottle (Gibco, ThermoFisher Cat # 11067-030)
- 12. 0.22 µm sterile filter unit, PES membrane (500 mL or 1 L)
- 13. Serological pipettes (assorted volumes)
- 14. P1000 pipet with *filter* tips
- 15. Automated pipetman
- 16. 2.5 % Vesphene (or other disinfectant of choice)
- 17. 70% Ethanol
- 18. Biosafety cabinet

Protocol:

- 1. Prepare biosafety cabinet (BSC) for work under aseptic conditions (wipe down cabinet surface with 2.5% Vesphene (or disinfectant of choice) and then 70% ethanol. 70% isopropanol may be used if fungal contamination is of concern.
- 2.____ Prepare 500mL or 1L of media as follows (Notes 1-3 & 8) and mix well:

Component	Qty./1 L DMEM	Qty/500 mL DMEM	Final Conc
DMEM/F12	Q.S. to 1L	Q.S. to 500 mL	N/A
*Antibiotic-antimycotic (100X)	10 mL	5 mL	1X
NCTC-109 media	100 mL	50 mL	10 %
FBS	100 mL/200 mL	50 mL/100 mL	10%/20%
*2-Mercaptoethanol (1000X)	1 mL	0.5 mL	1X
*Sodium Pyruvate (100 mM)	10 mL	5 mL	1 mM
*Oxaloacetic acid (1 mM)	2 mL	1 mL	2 μΜ
Insulin Bovine Pancreas	0.5 mL	0.25 mL	0.05%
L-glutamine (200 mM, 100X)	10 mL	5 mL	2 mM
HT (100X)	10 mL	5 mL	1X

*Indicates hazardous substance (complete media must go in hazardous waste). See document: Hybridoma Media Haz Waste_List of Components in "MAB_HYB_SOPs" folder to generate hazardous waste labels for complete media.

See Note 8 for information on HAT and HT supplements

3	In the BSC, transfer media to a 500 mL or 1 L $0.22~\mu m$ sterile filter unit (PES membrane) and connect to the vacuum. Filter the complete media (Note 4).
4	Aliquot media into smaller volumes (Note 5).
5	Store media at 4°C for up to 1 month (Notes 6 & 7).

Notes:

- 1. After FBS has been added, the medium is considered "complete".
- 2. Add up volume of supplements (reagents to be added to media) and remove the calculated volume from the 500 mL base media. Keep the "incomplete" base media in a sterile storage container for future use, if needed. This incomplete media can be dumped down the drain, if NOT hazardous waste.
- 3. The DMEM/F12 media can be used for culturing SP2/0 cells. Grow SP2/0 cells in 10% FBS up to fusion. Make sure to check which media SP2/0 cells were grown in (as some were grown in IMDM media).
- 4. Media can be filtered using a benchtop vacuum. However, make sure that the filter unit is sealed and not opened outside of BSC.
- 5. Aliquoting media allows for ease of warming up and saves media from contamination or becoming too alkaline (red color) due to repeated/constant use of big storage bottle.
- 6. Antibiotics have very short half-lives, meaning that they need to be replenished frequently (at least once every week).

Table 1 – Half-lives of Antibiotics and Antimycotics (Harlow and Lane).

Antibiotic/Antimycotic	Target	Half-life at 37°C	Mechanism of action
	Gram positive		
Penicillin G sodium	bacteria	2 days	Cell wall inhibitor
	Gram negative		Inhibits translation in 30S
Streptomycin sulfate	bacteria	4 days	subunit
			Alters membrane
Amphotericin B	Fungi, yeast	4 days	permeability

- 7. L-glutamine supplement should be replenished in the media after one month at 4°C and one week at 37°C.
- 8. For establishing hybridoma cell lines (fusion, parentals), add HAT supplement to a 1X concentration to the complete IMDM media with 20% FBS above. When parental hybridoma lines have been established, cells should be weaned off the HAT supplement. Parentals can be frozen in HAT media, but ideally weaned off and frozen in HT media (See SOP: AB104.6). Parentals should be thawed and grown in complete IMDM media, 20% FBS, with 1X HT (and all other additional supplements provided in this IMDM media SOP).

Subcloned hybridomas (*established hybridoma cell lines*) should be grown with the 1X HT supplement, unless otherwise noted for a particular cell line. 10% or 20% FBS can be used for *established hybridoma cell line* growth (generally thaw/grow lines in 10% FBS – more cost effective and they grow slower).

Component	Qty/1 L IMDM	Qty/500 mL IMDM	Final Concentration
HAT (50X)	20 mL	10 mL	1X

Tips:

To mix media, inverting the bottle is sufficient. Too vigorous of mixing creates bubbles (FBS).

References:

- 1. *Antibodies: A Laboratory Manual*. Ed Harlow and David Lane. Cold Spring Harbor Laboratory, New York. 1988. pp. 247-281.
- 2. Kohler, G., and Milstein, C. *Nature* 256, 495-497. August 7, 1975. Continuous Cultures of Fused Cells Secreting Antibody of Predefined Specificity.
- 3. Wayne M. Yokoyama, Michelle Christensen, Gary Dos Santos, and Diane Miller. Current Protocols in Immunology. UNIT 2.5 Production of Monoclonal Antibodies. Published Online: September 1, 2006.