Dulbecco's Modified Eagle Medium (DMEM)

With 4.5gms Glucose per litre, and Sodium pyruvate
Without L-Glutamine, Sodium bicarbonate and Phenol red

Product Code: AT190A

Product Description:
Dulbecco's Modified Eagle Medium (DMEM) is one of the most widely used modification of Eagle's medium. DMEM is a modification of Basal Medium Eagle (BME) that contains four fold concentration of amino acids and vitamins. Additionally, the formulation also includes glycine, serine and ferric nitrate. The original formulation contains 1000mgs/L of glucose and was originally used to culture embryonic mouse cells.

DMEM high glucose is a further modification of original DMEM and contains 4500mgs glucose per litre. The additional glucose has proved to be useful in cultivating various other cell lines including primary cultures of mouse and chicken cells as well as various normal and transformed cell lines.

Composition:

Ingredients          mg/L

INORGANIC SALTS
Calcium chloride dihydrate     265.000
Ferric nitrate nonahydrate     0.100
Magnesium sulphate anhydrous   97.720
Potassium chloride             400.000
Sodium chloride               6400.000
Sodium dihydrogen phosphate anhydrous     109.000

AMINO ACIDS
Glycine                  30.000
L-Arginine hydrochloride  84.000
L-Cystine dihydrochloride 62.570
L-Histidine hydrochloride monohydrate 42.000
L-Isoleucine            105.000
L-Leucine               105.000
L-Lysine hydrochloride    146.000
L-Methionine            30.000
L-Phenylalanine          66.000
L-Serine                42.000
L-Threonine             95.000
L-Tryptophan            16.000
L-Tyro sine disodium salt 103.790

VITAMINS
Choline chloride        4.000
D-Ca-Pantothenate       4.000
Folic acid              4.000
Nicotinamide            4.000
Pyridoxal hydrochloride 4.000
Riboflavin              0.400
Thiamine hydrochloride  4.000
i-Inositol              7.200

OTHERS
D-Glucose              4500.000
Sodium pyruvate        110.000

Directions:
1. Suspend 13.0gms in 900ml tissue culture grade water with constant and gentle stirring until the powder is completely dissolved. Do not heat the water.
2. Add 3.7gms of sodium bicarbonate powder (TC230) or 49.3ml of 7.5% sodium bicarbonate solution (TCL013) and 0.584gms of L-Glutamine powder (TC243) or 20ml of 200mM L-Glutamine solution (TCL012) for 1 litre of medium and stir until dissolved.
3. Adjust the pH to 0.2-0.3 pH units below the desired pH using 1N HCl or 1N NaOH since the pH tends to rise during filtration.
4. Make up the final volume to 1000ml with tissue culture grade water.
5. Sterilize the medium immediately by filtering through a sterile membrane filter with a porosity of 0.22 micron or less, using positive pressure rather than vacuum to minimize the loss of carbon dioxide.
6. Aseptically add sterile supplements as required and dispense the desired amount of sterile medium into sterile containers.
7. Store liquid medium at 2-8°C and in dark till use.

Please refer disclaimer overleaf
Materials required but not provided:
- Tissue culture grade water (TCL010)
- Sodium bicarbonate (TC230)
- Sodium bicarbonate solution 7.5% (TCL013)
- 1N Hydrochloric acid (TCL003)
- 1N Sodium hydroxide (TCL002)
- L-Glutamine powder (TC243)
- L-Glutamine solution 200mM (TCL012)
- Foetal bovine serum (RM1112/RM10432)

Quality Control:

Appearance
White to light pink, homogenous powder

Solubility
Clear solution at 13.0gms/L.

pH without Sodium Bicarbonate
5.90 - 6.50

pH with Sodium Bicarbonate
7.40 - 8.00

Osmolality without Sodium Bicarbonate
250.00 - 290.00

Osmolality with Sodium Bicarbonate
330.00 - 370.00

Cultural Response
The growth promotion capacity of the medium is assessed qualitatively by analyzing the cells for the morphology and quantitatively by estimating the cell counts and comparing it with a control medium.

Endotoxin Content
NMT 1EU/ml

Storage and Shelf Life:

1. All the powdered media and prepared liquid culture media should be stored at 2-8°C. Use before the expiry date. In spite of above recommended signs of deterioration /degradation in certain instances. This can be indicated by change in colour, change in appearance and presence of particulate matter and haziness after dissolution.

2. Preparation of concentrated medium is not recommended since free base amino acids and salt complexes having low solubility may precipitate in concentrated medium.

3. pH and sodium bicarbonate concentration of the prepared medium are critical factors affecting cell growth. This is also influenced by surface-to-volume ratio. For example, in large bottles, such as Roux bottles pH tends to rise perceptibly as significant volume of carbon dioxide is released. Therefore, optimal conditions of pH, sodium bicarbonate concentration, surface-to-volume ratio must be determined for each cell type. We recommend stringent monitoring of pH. If needed, pH can be adjusted by using sterile 1N HCl or 1N NaOH or by bubbling in carbon dioxide.

4. If required, supplements can be added to the medium prior to or after filter sterilization observing sterility precautions. Shelf life of the medium will depend on the nature of supplement added to the medium.

Revision: 2/2016

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