



## **Product Specification**

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## **Technical Information**

#### **RPMI-1640**

With L-Glutamine, 2gms per liter Glucose, 0.165 moles per liter MOPS buffer Without Sodium bicarbonate 1X **Liquid cell Culture Medium** 

#### Product Code: AL1200A

Application:-Roswell Park Memorial Institute (RPMI) media are a series of media developed by Moore et al for the culture of human normal and neoplastic cells in vitro. RPMI 1640 is the most commonly used medium in the series. A modification of McCoy's 5A medium, the medium was specifically designed to support the growth of human lymphoblastoid cells in suspension culture. Presently the medium is extensively used for a wide range of anchorage dependant cell lines. The medium needs to be supplemented with 5-20% fetal bovine serum. The medium is also known to support growth of cells in the absence of serum.

AL1200A is RPMI 1640 with L-glutamine, 2gms per Litre glucose and 0.165M per Litre MOPS buffer. It does not contain sodium bicarbonate. \*"MOPS, a zwitterionic buffer does not antagonize antifungal agents at final concentration of 0.165mol/L for pH 7.0. Therefore, this medium is used as a diluent for antifungal agents that are water-soluble as well as water-insoluble. For waterinsoluble antifungal agents, that cannot be prepared as stock solutions in water, such as amphotericin B, anidulafungin, itraconazole, ketoconazole, posaconazole and voriconazole, a dilution series of the agent should be prepared first at 100 times final strength in an appropriate solvent. Each of these non-aqueous solutions should then be diluted tenfold in RPMI-1640 broth". Users are advised to review the literature for recommendations regarding medium supplementation and physiological growth requirements specific for different cell lines.

Composition**	
Ingredients	mg/Litre
INORGANIC SALTS	
Calcium nitrate tetrahydrate	100.000
Magnesium sulphate anhydrous	48.840
Potassium chloride	400.000
Sodium chloride	6000.000
Sodium phosphate dibasic anhydrous	800.000
AMINO ACIDS	
Glycine	10.000
L-Arginine hydrochloride	241.000
L-Asparagine	50.000
L-Aspartic acid	20.000
L-Cystinedihydrochloride	65.200
L-Glutamic acid	20.000
L-Glutamine	300.000
L-Histidine hydrochloride monohydrate	20.960
L-Hydroxyproline	20.000
L-Isoleucine	50.000
L-Leucine	50.000
L-Lysine hydrochloride	40.000
L-Methionine	15.000
L-Phenylalanine	15.000
L-Proline	20.000
L-Serine	30.000
L-Threonine	20.000
L-Tryptophan	5.000
L-Tyrosine disodium salt	28.830
L-Valine	20.000





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#### VITAMINS

Choline chloride 3.000 D-Biotin 0.200 D-Ca-Pantothenate 0.250 Folic acid 1.000 Niacinamide 1.000 Pyridoxine hydrochloride 1.000 Riboflavin 0.200 Thiamine hydrochloride 1.000 Vitamin B12 0.005 i-Inositol 35.000 p-Amino benzoic acid (PABA) 1.000

**OTHERS** 

D-Glucose 2000.000
Glutathione reduced 1.000
MOPS buffer, free acid 34500.000
Phenol red sodium salt 5.300

### Methodology

1. Add 26.67ml of 7.5% sodium bicarbonate solution (TCL1013) for 1 litre of medium.

### Material required but not provided

Sodium bicarbonate solution 7.5% (TCL1013)

## **Quality control**

#### Appearance

Yellow colored, clear solution.

#### рΗ

5.30 -5.90

#### Osmolality in mOsm/Kg H<sub>2</sub>O

390.00 -430.00

#### Sterility

No bacterial or fungal growth is observed after 14 days of incubation, as per USP specification.

#### 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 through minimum three subcultures.

#### **Endotoxin Content**

NMT 5EU/ml

### Storage and Shelf Life

Store at 2-8°C away from bright light. Shelf life is 12 months. Use before expiry date given on the product label.





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## Disclaimer:

- User must ensure suitability of the product(s) in their application prior to use.
- The product conforms solely to the technical information provided in this booklet and to the best of knowledge research and development work carried at **CDH** is true and accurate.
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