



cell signaling solutions

Certificate of Analysis

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E-C-L Cell Attachment Matrix

(entactin-collagen IV-laminin)

Catalog # 08-110

Lot # 24051

Source: Engelbreth-Holm-Swarm (EHS) mouse tumor.

Use and Handling: Thaw frozen product at 4°C and keep on ice thereafter. Dilute with sterile serum-free medium to approximately 20µg/ml. Added directly to culture vessels to achieve 5-10µg/cm². Allow the matrix proteins to adsorb to the vessel for one hour at 37°C or overnight at 4°C prior to adding a cell suspension.

Sterility: This lot of E-C-L was tested and found negative for the presence of bacteria, fungi and mycoplasma.

Storage and Stability: Stable for 1 year at -20°C from date of shipment. Aliquot to avoid repeated freezing and thawing.

Formulation: 5mg in 5ml of 0.05M Tris-HCl, pH 7.4, 0.15M NaCl. Protein determined by the Bradford dye binding assay using gamma globulin as the standard. Frozen Solution.

**FOR RESEARCH USE ONLY
NOT FOR USE IN HUMANS**

Quality Control Testing

Neurite Outgrowth Assay: This lot of E-C-L was tested at 5µg/cm² and found to promote neurite outgrowth using PC12 rat pheochromocytoma cells.

IMPORTANT NOTE

Color variations from yellow to dark red may occur in frozen or thawed vials of E-C-L. These color variations are caused by the interaction of CO₂ with the buffer and phenol red present in the product. Such color variations are normal and do not affect the efficacy of the product.

References:

1. Kleinman, H.K., *et al.*, *Biochemistry* **25**: 312, 1986.
2. Kleinman, H.K., *et al.*, *Biochemistry* **21**: 6188, 1982.
3. Hadley, M.A., *et al.*, *J. Cell. Biol.* **101**: 1511, 1985.

Neurite Outgrowth Assay Protocol

1. Coat 6 well dishes with **E-C-L**, at a concentration of 2.5-10µg/cm². Incubate 1 hour at 37°C or overnight at 4°C. Aspirate and rinse 2 times with PBS before adding cell suspension.
2. Prepare a cell suspension at 2x10⁴cells/ml in RPMI-1640/10% horse serum/5% FBS. Add 2ml of cell suspension to each well.
3. Prepare NGF (Catalog # 01-170) as described on certificate of analysis to a final concentration of 1µg/ml. Add 20ng/ml NGF.
4. Incubate plates at 37°C for 3-5 days.
5. Record neurite outgrowth at each E-C-L concentration.