

For research use only  
Not for use in diagnostic procedures



# iMatrix-511

Product No. 892 011 350 µg  
Product No. 892 012 1,050 µg

Version 010  
Store at 2-15°C

**Product description:** iMatrix-511 is a recombinant human laminin-511 E8 fragment protein expressed in Chinese Hamster Ovary (CHO)-S cells. iMatrix-511 contains the integrin-binding site of the laminin-511 molecule. iMatrix-511 is a useful cell culture substrate for feeder-free culture and single-cell passage of ES cells and iPS cells, facilitating stable culture expansion. iMatrix-511 is also useful for the culture of other cells adhering to laminin-511.

**Content:** Recombinant human laminin-511 E8 fragment protein in PBS(-)

**Concentration:** 0.5 mg/mL

**Amount:** 175 µg / 0.35 mL / tube  
Product No. 892 011 350 µg / 2 tubes  
Product No. 892 012 1,050 µg / 6 tubes

**Storage:** Store at 2°C to 15°C, protect from light.

**Expiration date:** The shelf life is 2 years from the date of manufacture. The expiration date is printed on the outer carton.

**Activity:** The dissociation constant (Kd) for the binding with integrin  $\alpha 6\beta 1$  is 10 nM or less.

**Methods of use:** By either of the following methods, iMatrix-511 can be coated onto a culture vessel. **The optimum coating density may differ by cell-type, cell-line, medium selected, or purpose.** Insufficient coating density may result in the detachment of cells and varied cell conditions while the excessive coating density may lead to difficulty in detaching cells for passage.

## A. Pre-coating method

Determine the optimal coating density. 0.5 µg/cm<sup>2</sup> is a standard but test between 0.1 and 1.5 µg/cm<sup>2</sup>.

- 1) Dilute iMatrix-511 with PBS(-). Use the diluted iMatrix-511 immediately. To coat with 0.5 µg/cm<sup>2</sup> onto a 6-well plate with 9.6 cm<sup>2</sup>/well, dilute 9.6 µL of iMatrix-511 with 2 mL of PBS(-) per well.
- 2) Place the diluted iMatrix-511 into a culture vessel and incubate either at 37°C for 1 h, or at room temperature for 3 h, or at 4°C overnight.
- 3) Aspirate the coating solution. Then, immediately seed your cells. **Do not allow the coated surface to dry.**

## B. Pre-mixing method

Determine the optimal coating density for cell culture. The standard density is 0.25 µg/cm<sup>2</sup> but test between 0.1 and 1.5 µg/cm<sup>2</sup>. The optimal coating density may be affected by the medium and cell density of the cell suspension.

- 1) Add iMatrix-511 to the cell suspension. To coat with 0.25 µg/cm<sup>2</sup> onto a 6-well plate with 9.6 cm<sup>2</sup>/well, add 4.8 µL of iMatrix-511 to 2 mL of the cell suspension per well.
- 2) Place the cell suspension containing iMatrix-511 into a culture vessel.

\*If you face difficulties in detaching cells for passage, re-adjust the conditions (e.g., reduce the coating density).

## References:

Taniguchi Y. *et al.* (2009), *J. Biol. Chem.* **284** (12): 7820-31  
Miyazaki T. *et al.* (2012), *Nat. Commun.* **3**: 1236  
Nakagawa M. *et al.* (2014), *Sci. Rep.* **4**: 3594  
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Goparaju S.K. *et al.* (2017), *Sci. Rep.* **7**: 42367  
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**Caution:** For research use only. Not intended for human use. In the event of accidental ingestion or contact with the eyes, immediately wash the affected area and seek medical attention.

**Product information:** Current information including references and Q&A are available on the website of Matrixome, Inc. Please use the URL or QR code below.

**Designed by:** Matrixome Inc.

3-2 Yamadaoka, Suita, Osaka 565-0871, Japan  
Institute for Protein Research, Osaka University  
Tel: +81-6-6877-0222 Fax: +81-6-6877-0002  
E-mail: [info@matrixome.co.jp](mailto:info@matrixome.co.jp)  
URL: <http://www.matrixome.co.jp/en/>



**Manufactured by:** Nippi, Incorporated  
1-1-1 Senju Midori-cho, Adachi, Tokyo 120-8601, Japan