



Surface Immobilization Chemistry of a Laminin-Derived Peptide Affects Keratinocyte Activity

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(A)

Figure 1. (**A**) HPLC analysis and (**B**) Electrospray ionization (ESI) mass spectroscopy spectrum of S-LamLG3 (*M*w = 2037.49 Da).



Figure 2. (A) HPLC analysis and (B) ESI mass spectroscopy spectrum of D-LamLG3 (*M*w = 1935.35 Da).



Figure 3. Ratio of the N1s X-ray photoelectron spectroscopy to the C 1*s* counts (N 1*s*/C1*s*) for D-LamLG3 vs. S-LamLG3 following up to 73 days in artificial saliva (37 °C). Differences in mean N 1*s*/C 1*s* Counts between D-LamLG3 and S-LamLG3 were assessed with an unpaired *t*-test; there were no statistically significant differences at any time point (p > 0.05).



Figure 4. Representative micrographs of oral keratinocyte Col17 immunofluorescence after one day of culture. The scale bar is 100 μ m.



Figure 5. Representative micrographs of oral keratinocyte integrin β 4 immunofluorescence after one day of culture. The scale bar is 100 μ m.