

Electronic Supplementary Information

**Synthesis of Poly(lactide-co-glycerol) as a Biodegradable and Biocompatible
Polymer with High Loading Capacity for Dermal Drug Delivery**

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Synthesis of PLG11 and PLG12

PLG11, PLG12 was synthesized according to previously reported method. Briefly, 1/1 and 1/2 molar ratio of glycidol/L-lactide were mixed and stirred at 100 °C for 24 h. Afterwards, the product was dissolved in DMF and dialyzed against DMF and water for 6 and 72 h, respectively. Finally, the solvent was evaporated by vacuum and the product was lyophilized. The product was analyzed using $^1\text{H NMR}$ (S1) and GPC (S2).

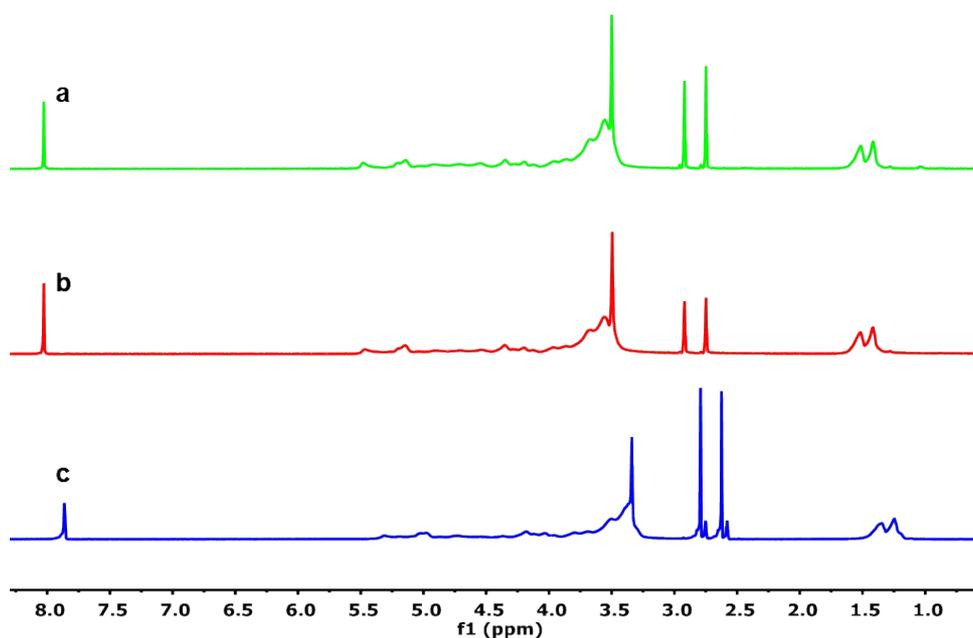


Figure S1. $^1\text{H NMR}$ spectrum of PLG11 (a), PLG21 (b) and PLG (c) recorded in DMF-d_7 solvent.

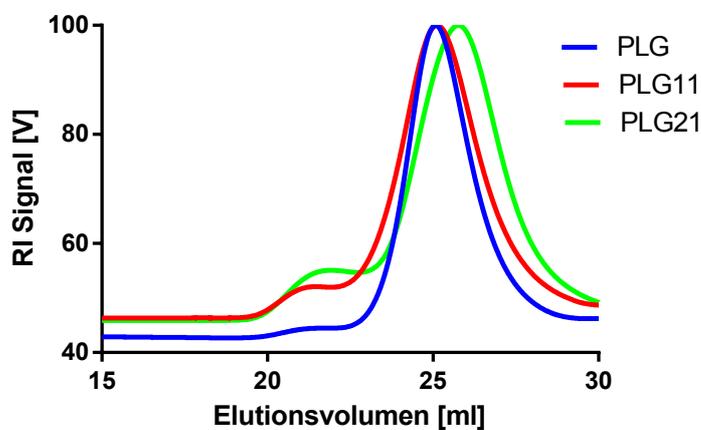


Figure S2. GPC spectrum of PLG11 (red), PLG21 (green) and PLG (blue)

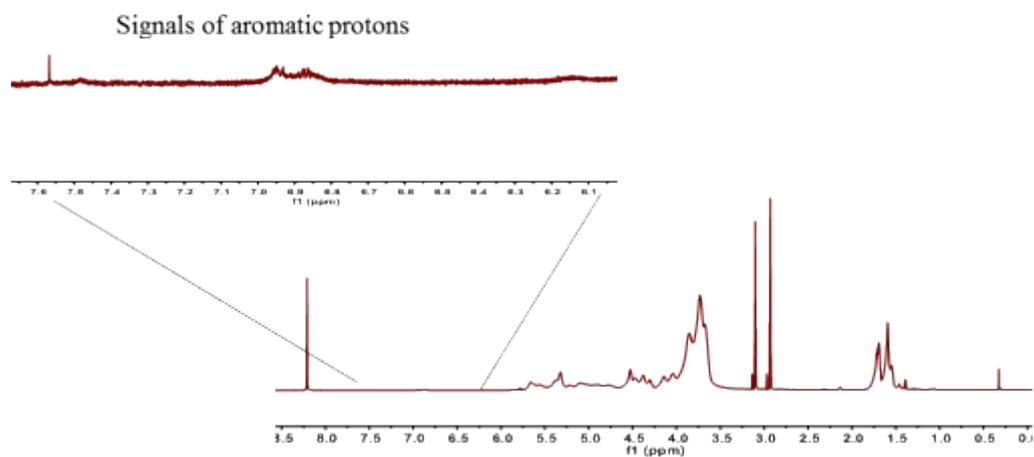


Figure S3. ¹H NMR spectrum of FITC conjugated PLG.

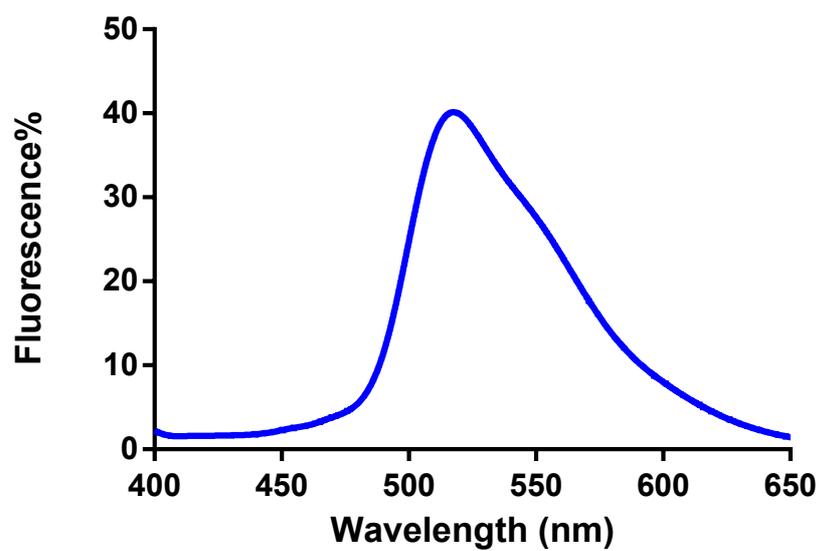


Figure S4. Fluorescence spectrum of FITC conjugated PLG.

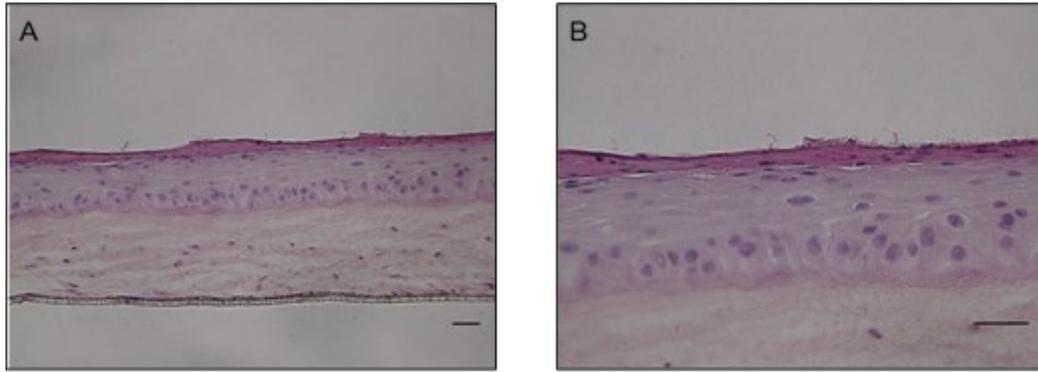


Figure S5. Representative pictures of hematoxylin & eosin staining of normal skin models. **A** magnification 10x. **B** Magnification 20x. Scale bar = 50 μ m.

Gene	Primer sense 5'-3'	Primer antisense 5'-3'
<i>FLG</i>	AAGGAACTTCTGGAAAAGGAATTT C	TTGTGGTCTATATCCAAGTGATCC AT
<i>GAPDH</i>	CTCTCTGCTCCTCCTGTTTCGAC	TGAGCGATGTGGCTCGGCT

Table S1. Primer sequences for qRT-PCR

Antibody	Isotype	Clone	WB	Company
TSLP	rabbit IgG	polyclonal	1:1000	Abcam, Cambridge, United Kingdom
β -actin	mouse IgG1	monoclonal	1:10000	Sigma-Aldrich, Munich, Germany

Table S2. Antibody dilutions for Western Blot (WB)

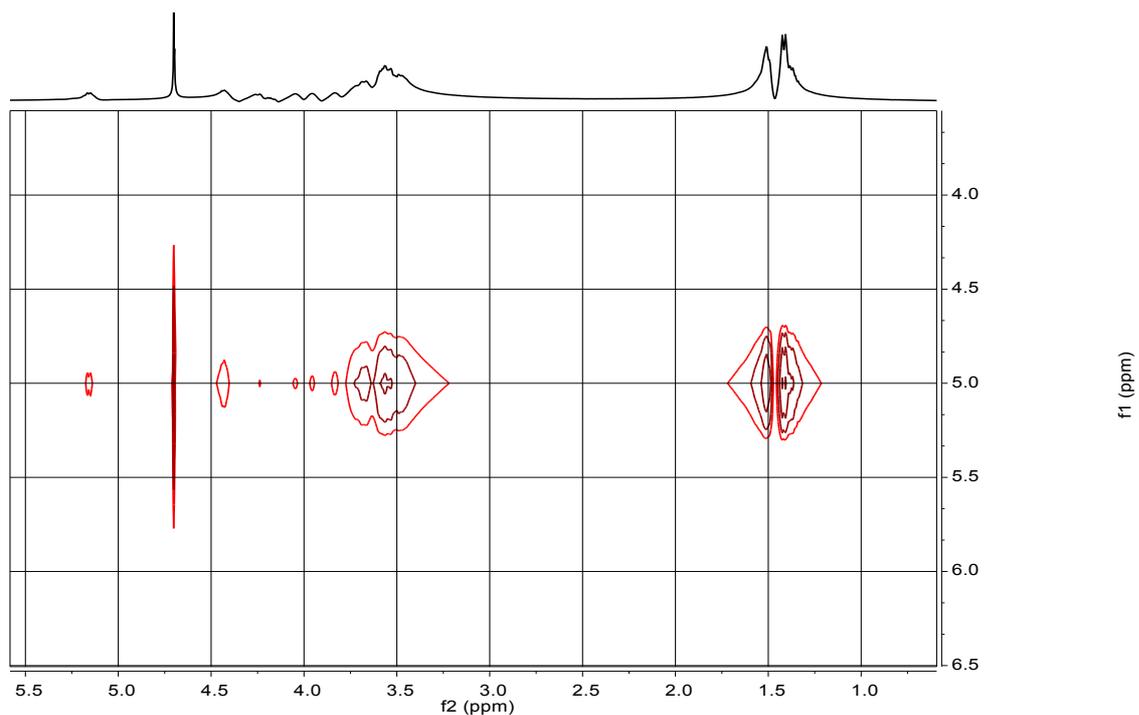


Figure S6. DOSY NMR of PLG measured in D₂O

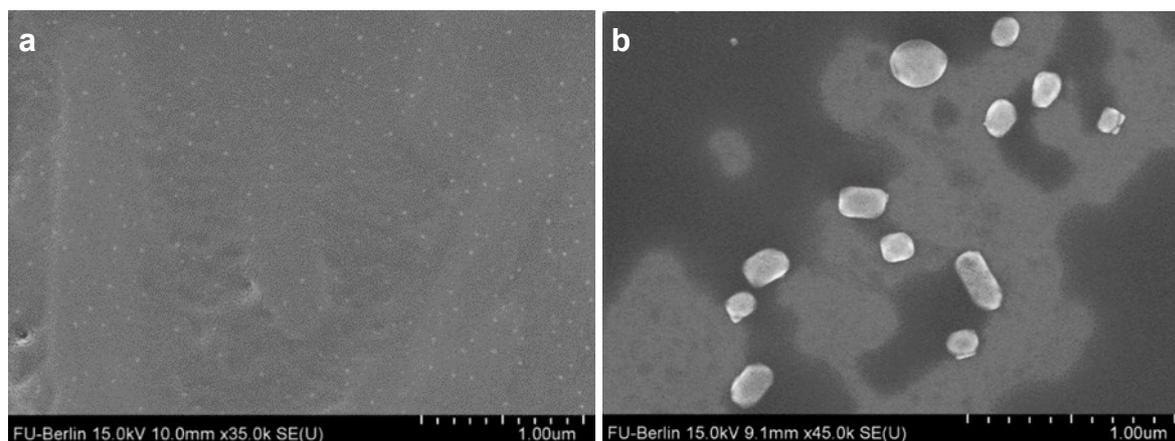


Figure S7. SEM images of PLG before loading TAC (a) after loading TAC with higher resolution.

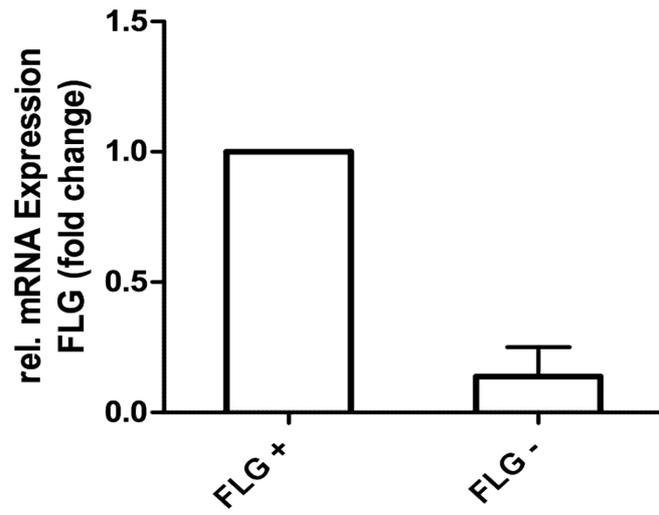


Figure S8. Relative mRNA expression of filaggrin in normal (FLG+) and filaggrin-deficient (FLG-) skin models (n=2, mean \pm SEM). GAPDH served as housekeeping gene.