

Appendix A. Supplementary material

Development of resiquimod-loaded modified PLA-based nanoparticles for cancer immunotherapy: a kinetic study

Cédric Thauvin^a, Jérôme Widmer^b, Inès Mottas^{a,b}, Sandra Hocevar^a, Eric Allémann^a, Carole Bourquin^{a,b,c,*}, Florence Delie^{a,*}

^a*School of Pharmaceutical Sciences, University of Geneva, University of Lausanne, Rue Michel-Servet 1, 1211 Geneva, Switzerland*

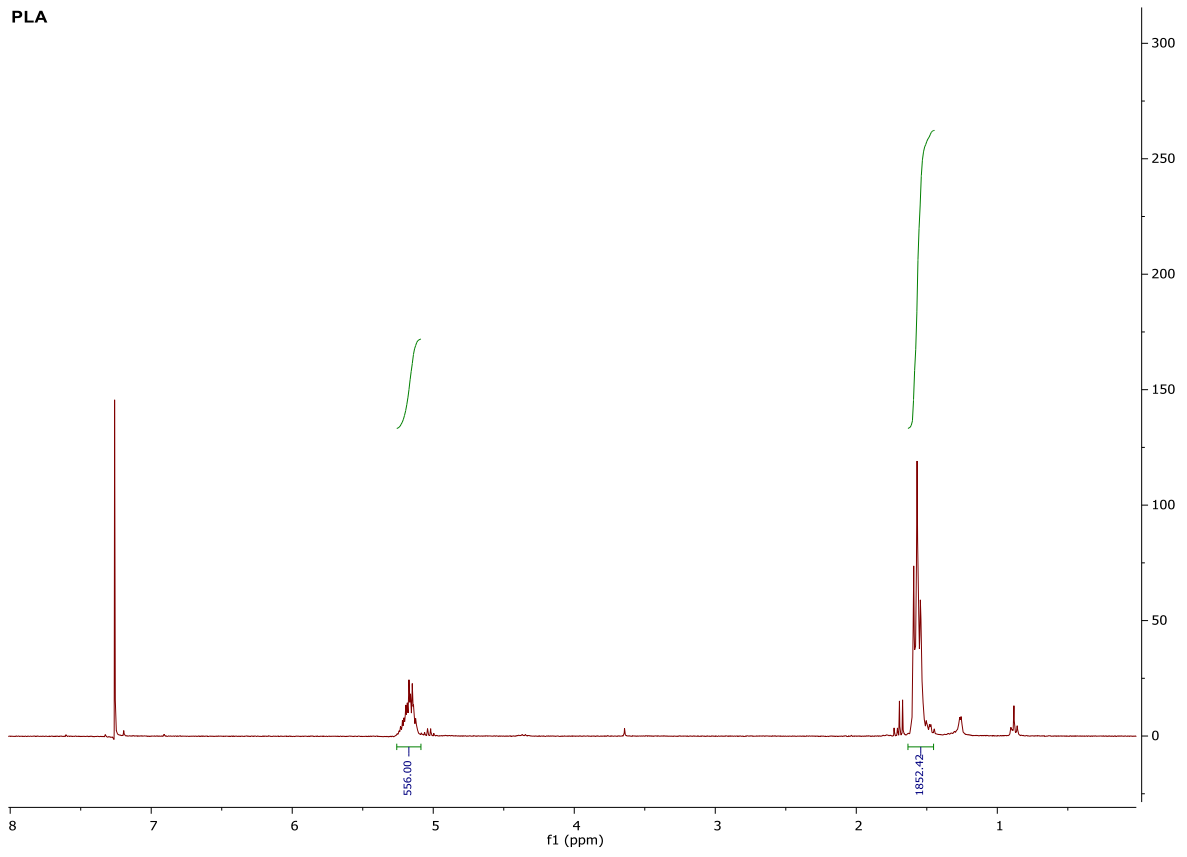
^b*Chair of Pharmacology, Faculty of Science and Medicine, University of Fribourg, Chemin du Musée 5, 1700 Fribourg, Switzerland*

^c*Faculty of Medicine, University of Geneva, Rue Michel-Servet 1, 1211 Geneva, Switzerland*

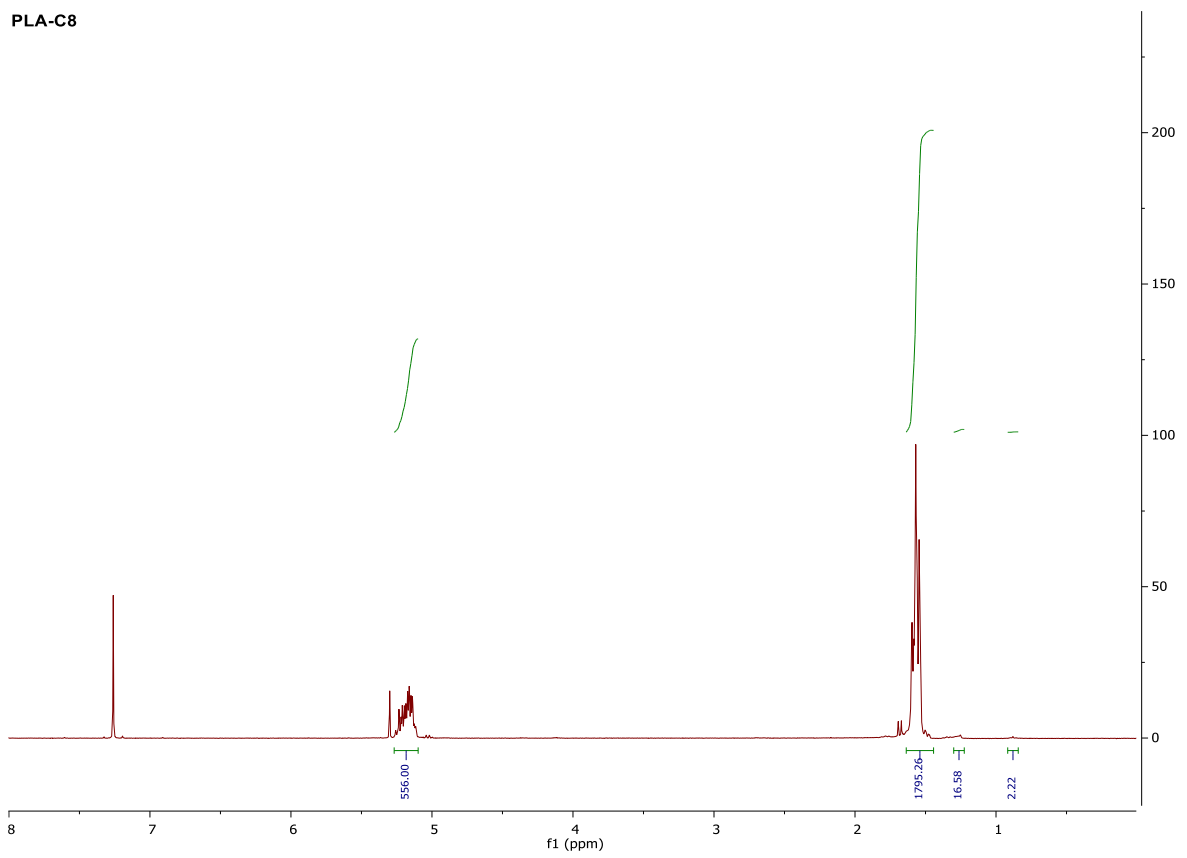
* Corresponding authors, the authors contributed equally to this work.

E-mail address: florence.delie@unige.ch (F. Delie) and carole.bourquin@unige.ch (C. Bourquin)

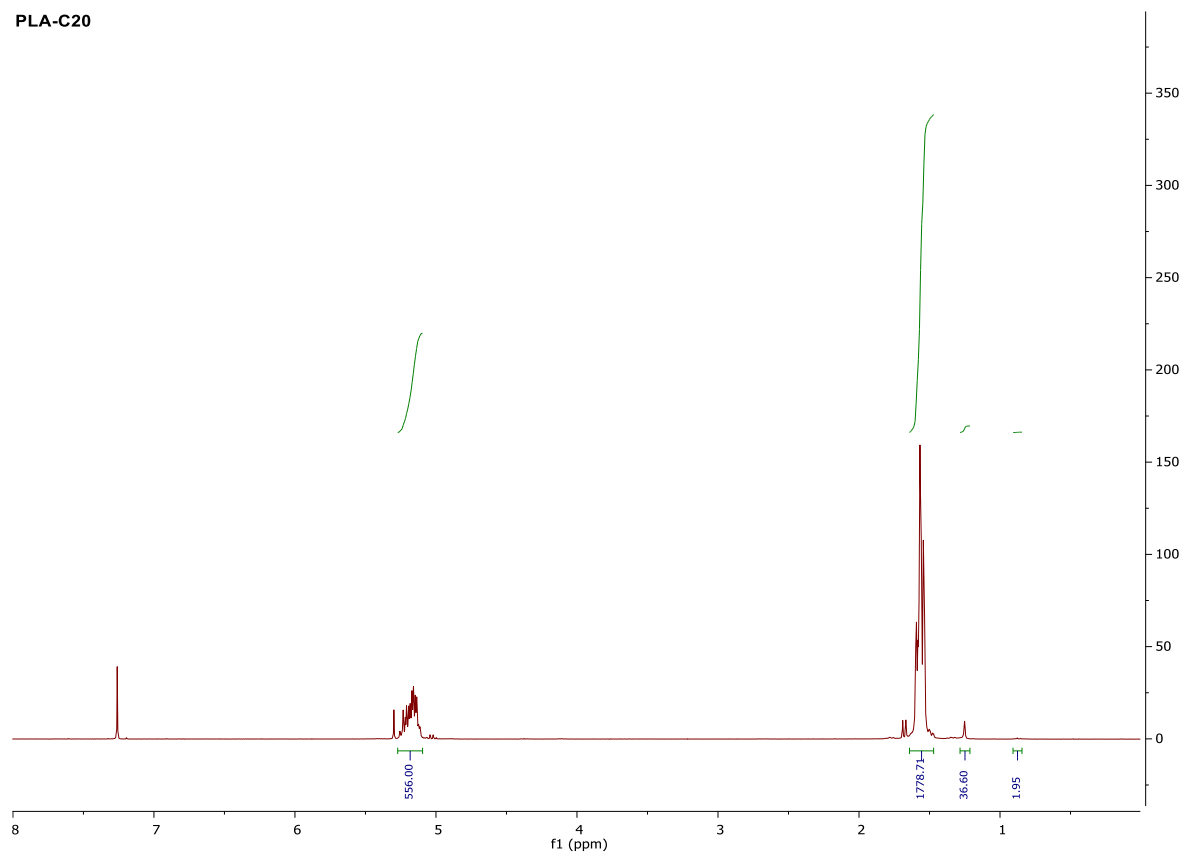
PLA



PLA-C8



PLA-C20



PLA-mPEG2000

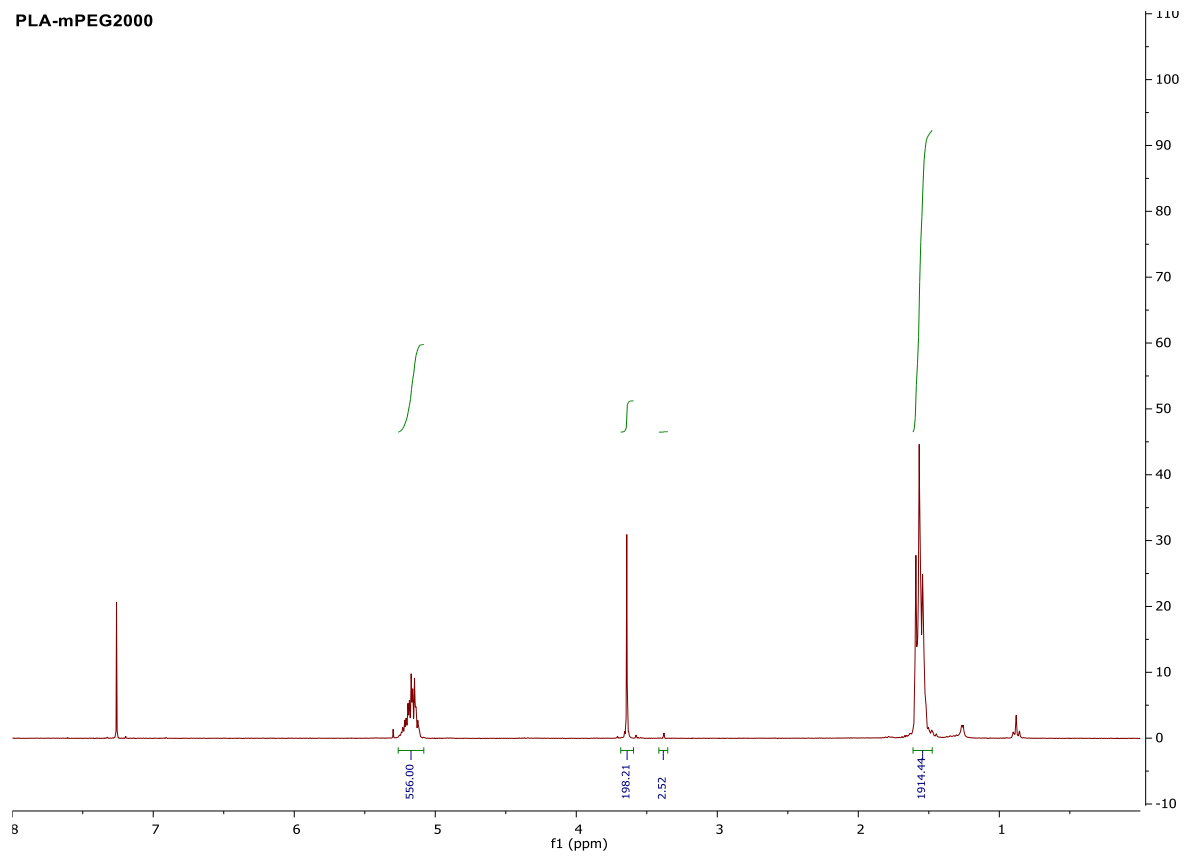


Fig. S1. ^1H NMR spectra of PLA-based polymers.

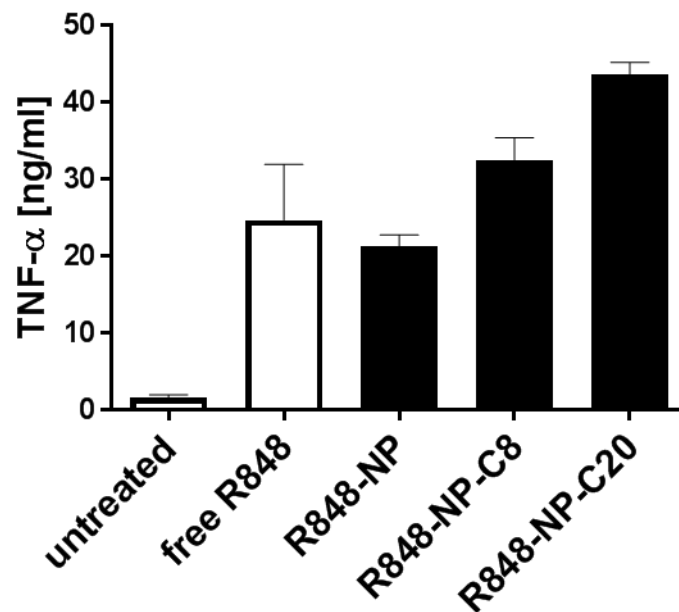


Fig. S2. R848-loaded PLA-NP activation of TNF- α release in macrophages. The release of TNF- α from J774 cells was assessed after incubation for 24 h with R848-loaded NP (R848 concentration: 0.1 $\mu\text{g}/\text{ml}$ for free R848 and all R848-loaded NP conditions). Each bar represents mean \pm SD, $n=3$. Data show one representative experiment out of three.