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We performed the same analysis on S-TM cells. The results showed a response to S-IFN* α * plus PL-P, and S-IFN* α * plus PL-P and S-MT. We did not observe an additional response in S-TM cells with S-IFN* α * and PL (S-IFN* α * plus

PL). * indicates a significant difference between treatments.](IJC-146-2879-g004){#ijc 31045-fig-0004} In summary, although our

preclinical model was not representative of the *in vivo* situation, our results indicated that either pure PL was effective as single agent to stimulate a tumour-specific response or that both components interact, resulting in an enhanced response, in addition to the cytotoxicity. The IFN α antibody used in the study was raised against non-modified human IFN α and therefore our study cannot totally rule out the interference of the antibody itself. However, the antibodies used in the clinic are specific against the recombinant protein and this should be the case also for the antibodies used in our study. Our next step will be to develop a clinical study combining PL with S-IFN α to treat patients with advanced HNSCC, either by using PL and S-IFN α alone or in combination. The work presented here was funded by the La Caixa Foundation (n. LCF/BQ/E0379/2012), the Fundación Ramón Areces and the IdIS2RD programme from the

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