

# Puroindoline-a and $\beta$ 1-purothionin form ion channels in giant liposomes but exert different toxic actions on murine cells

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Puroindoline-a (PIN-a) and  $\beta$ 1-purothionin ( $\beta$ 1-PTH), isolated from wheat endosperm of *Triticum aestivum* sp., have been suggested to play a role in plant defence mechanisms against phytopathogenic organisms. We investigated their ability to form pores when incorporated into giant liposomes using the patch-clamp technique. PIN-a formed cationic channels ( $\approx$  15 pS) with the following selectivity  $K^+ > Na^+ \approx Cl^-$ . Also,  $\beta$ 1-PTH formed channels of  $\approx$  46 pS and 125 pS at +100 mV, the selectivity of which was  $Ca^{2+} > Na^+ \approx K^+ \approx Cl^-$  and  $Cl^- \approx Na^+$ , respectively. In isolated mouse neuromuscular preparations,  $\beta$ 1-PTH induced muscle membrane depolarization, leading to blockade of synaptic transmission and directly elicited muscle twitches. Also,  $\beta$ 1-PTH caused swelling of differentiated neuroblastoma NG108-15 cells, membrane bleb formation, and disorganization of F-actin. In contrast, similar concentrations of PIN-a had no detectable effects. The cytotoxic actions of  $\beta$ 1-PTH on mammalian cells may be explained