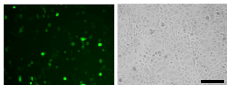
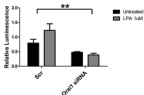


GFP

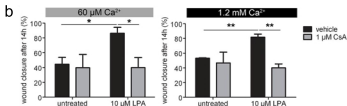
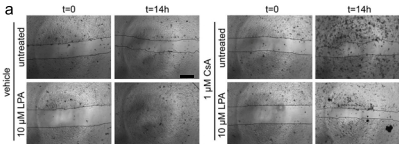
Brightfield



Supplementary figure 1. Transfection efficiency of siRNA following nucleofection. Primary keratinocytes were transfected with GFP using nucleofection, 24hours post transfection cells were imaged using a fluorescence microscope to determine transfection efficiency. Scale bar=100 μ m.



Supplementary figure 2. Dual luciferase reporter assay shows significantly less LPA-triggered activation of NFAT2 after transfection with Orai1-directed siRNA compared to scrambled siRNA. Cells were treated with 10uM LPA for 24h where indicated (** $p < 0.01$, 2-way ANOVA)



Supplementary fig.3 : LPA-induced keratinocyte migration into scratch wounds requires calcineurin activity. (A) Keratinocyte cultures were subjected to 2D scratch wounding assays, which were performed in medium containing 60 μ M Ca²⁺ or physiological 1.2 mM Ca²⁺. The average percentages of wound closure (B) are represented as bar graphs \pm SEM. Pre-incubation of cells with 1 μ M cyclosporin A significantly impaired LPA-induced migration in both assays ($n \geq 3$, ** $p < 0.01$, * $p < 0.05$, t test and 2-way ANOVA). Scale bar=100 μ m.