

Supplementary Information

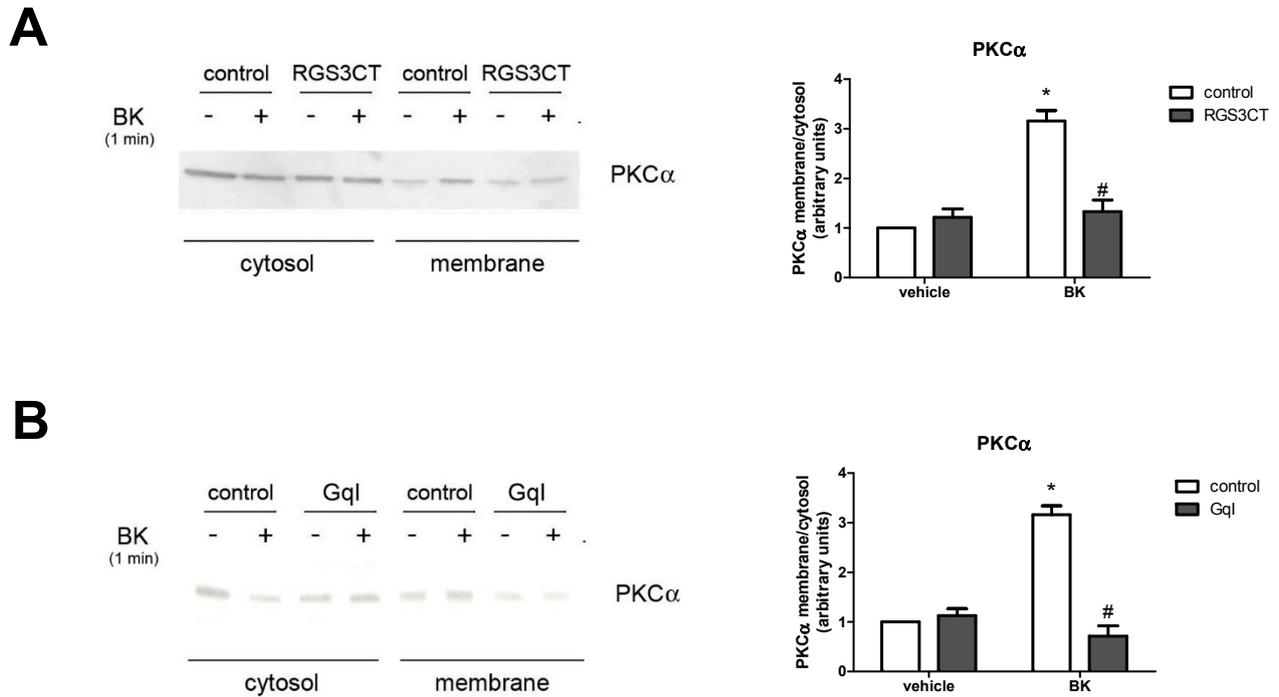


Figure S1. Bradykinin-induced PKC α translocation to membrane fraction relies on G $\alpha_{q/11}$ engagement.

C2C12 cells were transiently transfected with pcDNA3.1-RGS3CT or empty vector (control) (A) or with pRK5-GqI or empty vector (control) (B). Cells were overnight serum-starved prior to be stimulated with 1 μ M bradykinin (BK) for 1 min. Left panels: PKC α activation was evaluated as translocation of the enzyme to the membrane. Western blot analysis of PKC α were performed in membrane and cytosolic fractions. Blots representative of at least three independent experiments are shown. Right panels: the histograms represent densitometric analysis of three independent experiments. Data reported are expressed as fold increase of the membrane:cytosol ratio. The increase of PKC α membrane content induced by BK was statistically significant by Student's *t* test ($*p < 0.05$); the effect of G $\alpha_{q/11}$ inhibition on BK-induced PKC α activation was statistically significant by two-way ANOVA followed by Bonferroni's post hoc test ($\#p < 0.05$).

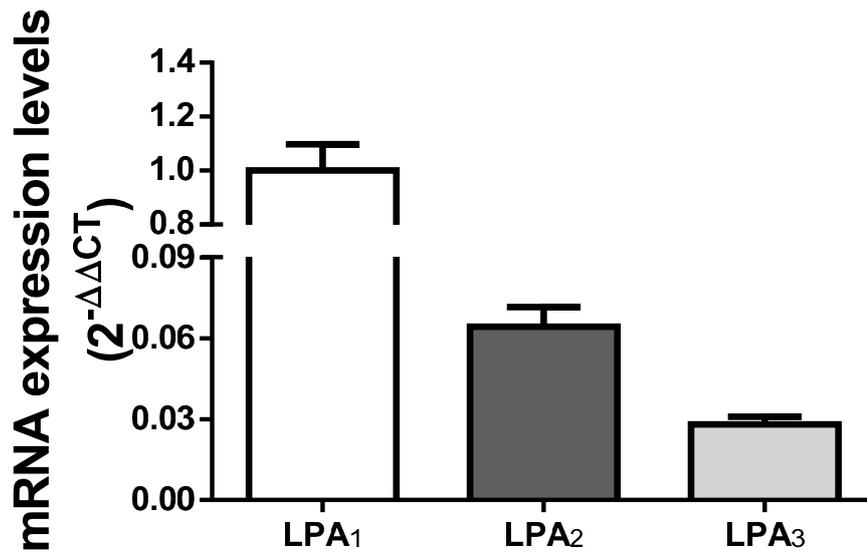


Figure S2. Expression of LPA receptors at the mRNA level.

Quantitative mRNA analysis was performed by real-time PCR by concurrent amplification of the target sequence of murine LPA₁, LPA₂ and LPA₃ genes together with that of 18S rRNA. The results are expressed as fold changes according to the $2^{-\Delta\Delta CT}$ method, using LPA₁ as calibrator. Values are means \pm SEM of three independent experiments performed in triplicate.

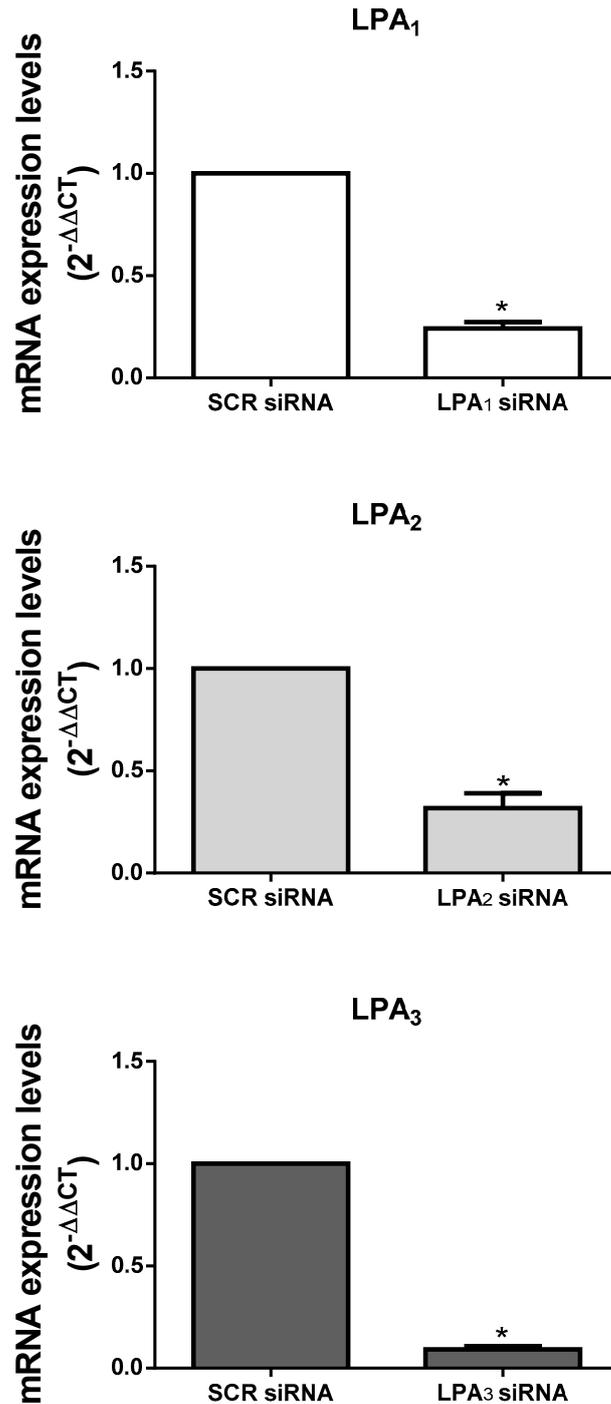


Figure S3. Down-regulation of LPAR.

C2C12 myoblasts, transfected with scrambled (SCR) or with specific siRNA for individual LPA receptors, were checked for downregulation by real-time PCR by concurrent amplification of the target sequence of murine LPA₁, LPA₂ and LPA₃ genes together with that of 18S rRNA. The results are expressed as fold changes according to the $2^{-\Delta\Delta CT}$ method. Values are means \pm SEM of three independent experiments performed in triplicate. The effect of siRNA transfection on LPAR expression is statistically significant by Student's *t* test (* p <0.05).