

## **Inflammation leads through PGE/EP<sub>3</sub> signaling to HDAC5/MEF2-dependent transcription in cardiac myocytes**

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**Appendix Table S1** – Concentrations, descriptions and supporting references of the used compounds.

<b>Compound (used concentration)</b>	<b>Description with reference</b>
<b><u>L798106</u> (200 nM)</b>	Potent and highly selective EP <sub>3</sub> receptor antagonist (K <sub>i</sub> values are 0.3, 916, > 5000 and > 5000 nM at EP <sub>3</sub> , EP <sub>4</sub> , EP <sub>1</sub> and EP <sub>2</sub> receptors, respectively) (Clarke et al, 2004; Juteau et al, 2001).
<b><u>L161,982</u> (2 μM)</b>	Potent and selective EP <sub>4</sub> receptor antagonist (K <sub>i</sub> values are 32, 7100, 5800, 6100 nM for rat EP <sub>4</sub> , EP <sub>3</sub> , EP <sub>2</sub> and EP <sub>1</sub> receptors, respectively) (Balzary et al, 2006; Machwate et al, 2001).
<b><u>AH6809</u> (10 μM)</b>	EP <sub>1</sub> (pA <sub>2</sub> = 6.8) and EP <sub>2</sub> (K <sub>i</sub> = 350 nM) receptor antagonist (Kiriya et al, 1997).
<b><u>Bisindolylmaleimide I</u> (3 μM)</b>	Selective inhibitor of protein kinase C. IC <sub>50</sub> values are 0.0084, 0.0180, 0.210, 0.132, and 5.8 μM for α, β1, δ, ε and ζ PKC isoforms, respectively (Jacobson et al, 1995; Martiny-Baron et al, 1993).
<b><u>PTX</u> (100 ng/ml)</b>	Bacterial toxin that inhibits G <sub>i/o</sub> and G <sub>t</sub> via catalyzation of ADP-ribosylation (Vettel et al, 2012).
<b><u>NSC 23766</u> (100 μM)</b>	Selective inhibitor of Rac1 activation by preventing Rac1-GEF (TrioN and Tiam1, IC <sub>50</sub> ~ 50 μM) interaction (Gao et al, 2004; Levay et al, 2013; Vettel et al, 2012).
<b><u>SQ 22536</u> (100 μM)</b>	Cell-permeable adenylate cyclase inhibitor (Harris et al, 1979; Mitra et al, 2006).
<b><u>AIP</u> (1 μM)</b>	Selective and potent CaMKII inhibitor (IC <sub>50</sub> : 40 nM) (Ishida et al, 1995).
<b><u>BPKDi</u> (3 μM)</b>	Selective inhibitor of the three PKD isoforms with low nanomolar potency but it failed to inhibit members of the CaMKII and MARK families or SIK1 and GRK5 (Haworth et al, 2012; Meredith et al, 2010; Monovich et al, 2010).
<b><u>IPA-3</u> (30 μM)</b>	A selective small-molecule inhibitor that targets the autoregulatory mechanism of group 1 PAKs (PAK1,2,3) and promotes their inactive conformation. Group 1 PAKs were shown to be inhibited at 10 μM (Deacon et al, 2008; Vettel et al, 2012).

**Appendix Table S2** – The used primer sequences for quantitative real-time PCR

<b>Gene</b>		<b>Primer sequence</b>
<i>IL6</i>	sense	5'-gatggatgctaccaaactggat-3'
	antisense	5'-ccaggtagctatggactccaga-3'
<i>TNF<math>\alpha</math></i>	sense	5'-ctgtagcccacgtcgtagc-3'
	antisense	5'-ttgagatccatgccgttg-3'
<i>Nur77</i>	sense	5'-tgttgatgttctgcctttg-3'
	antisense	5'-ggaggccatgtcgatcag-3'
<i>Myomaxin</i>	sense	5'-tctcctagaggcgctgtcc-3'
	antisense	5'-tgcacaagtctgattcttccat-3'
<i>AdamTS1</i>	sense	5'-aaaggcattggctacttctttg-3'
	antisense	5'-ggactacagggagtgccatc-3'
<i>ANP</i>	sense	5'-cacagatctgatggattcaaga-3'
	antisense	5'-cctcatcttctaccggcatc-3'
<i>BNP</i>	sense	5'-gtcagtcgcttgggctft-3'
	antisense	5'-ccagagctggggaagaag-3'
<i>18S</i>	sense	5'-gcaattattcccatgaacg-3'
	antisense	5'-gggacttaatcaacgcaagc-3'

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