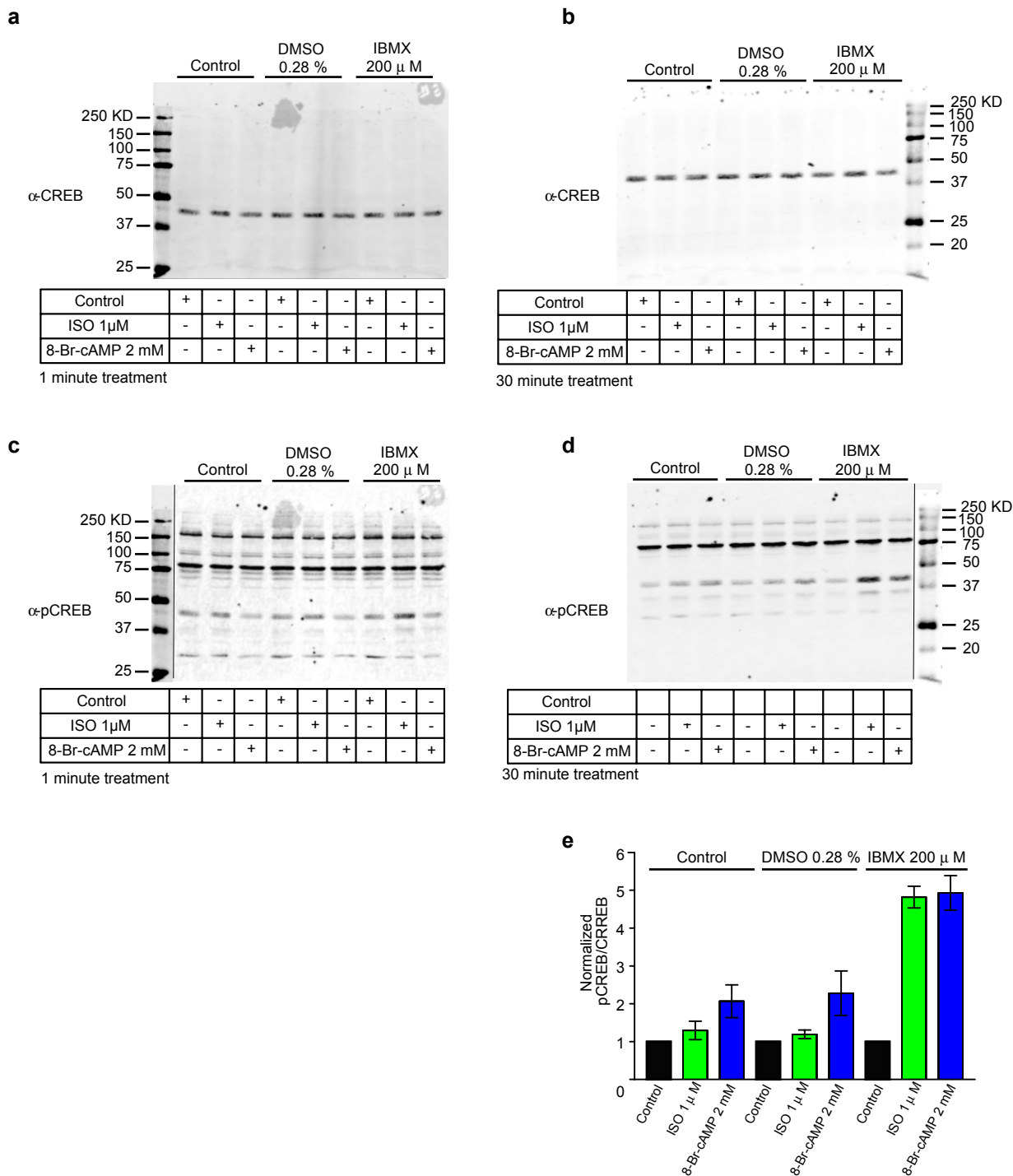


**Table S1 | Summary of drug targets and observed effect.**

<b>Drug</b>	<b>Target</b>	<b>Molecular Function</b>	<b>Observed Effect on <math>\beta_2</math>AR <math>Ca^{2+}</math> signal</b>
<b><i>Norepinephrine</i></b>	All ARs	Agonist	Calcium mobilization
<b><i>Epinephrine</i></b>	All ARs	Agonist	Calcium mobilization
<b><i>Isoproterenol</i></b>	All ARs	Agonist	Calcium mobilization
<b><i>Terbutaline</i></b>	All ARs	Agonist	Calcium mobilization
<b><i>Prazosin</i></b>	$\alpha_1$ ARs	Antagonist	No inhibition
<b><i>Yohimbine</i></b>	$\alpha_{1/2}$ ARs	Antagonist	No inhibition
<b><i>Propranolol</i></b>	$\beta_1$ and $\beta_2$ ARs	Antagonist	Suppression
<b><i>ICI 118,551</i></b>	$\beta_2$ AR	Antagonist	Suppression
<b><i>ATP</i></b>	P <sub>2</sub> YR	Agonist	N/A
<b><i>8-bromo-cAMP</i></b>	PKA	Activator	No $Ca^{2+}$ mobilization
<b><i>H-89</i></b>	PKA	Inhibitor	No suppression
<b><i>KT 5720</i></b>	PKA	Inhibitor	No suppression
<b><i>IBMX</i></b>	PDE	Inhibitor	No effect
<b><i>ddAd</i></b>	AC	Inhibitor	No effect
<b><i>SQ 22,536</i></b>	AC	Inhibitor	No effect
<b><i>DAMGO</i></b>	$\mu$ OR ( $G_{ai/o}$ coupled)	Agonist	No effect
<b><i>CTX</i></b>	$G_{as}$	Activator	No $Ca^{2+}$ mobilization
<b><i>PTX</i></b>	$G_{ai/o}$	Inhibitor	No inhibition
<b><i>EGTA</i></b>	$Ca^{2+}$	Chelator	No suppression of peak response
<b><i>TG</i></b>	SERCA-ATPase pump on ER membrane	Deplete ER of $Ca^{2+}$	Suppression
<b><i>U73122</i></b>	PLC	Inhibitor	Suppression
<b><i>2-APB</i></b>	InsP <sub>3</sub> R	Inhibitor	Suppression
<b><i>Carbachol</i></b>	M <sub>3</sub> R ( $G_{oq}$ coupled)	Agonist	N/A

**Figure S1**



**Supplemental Figure S1 | Full western blot scans from figure 3.** Infrared fluorescence scans of immunoblots, each representative of three independent experiments. a,b) staining for total CREB (43 kDa) after 1 and 30 minutes treatment with ISO and 8-Br-cAMP. c,d) staining for pCREB (43 kDa) after 1 and 30 minutes treatment with ISO and 8-Br-cAMP. e) pCREB/CREB ratios normalized to control averaged from 3 independent experiments following treatments with ISO and 8-Br-cAMP for 30 minutes. Bar graphs are averages of three or more independent experiments, error bars indicate SEM. Panels a and c are from the same gel, as are panels c and d, and in each case an image of the  $M_r$  marker lane imaged using the optics for the dye used to detect total CREB in panel a or b is pasted adjacent to the images of the lanes containing the samples for each gel.