

MEETING ABSTRACT

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# Signalling and function of the human G protein-coupled receptor 55

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From 16th Scientific Symposium of the Austrian Pharmacological Society (APHAR)  
Vienna, Austria. 25-27 November 2010

## Background

We have recently shown that the G protein-coupled receptor 55 (GPR55) responds to some of the cannabinoid and non-cannabinoid ligands in addition to the classical cannabinoid 1 (CB<sub>1</sub>) and 2 (CB<sub>2</sub>) receptors. Here we show multiple signaling pathways triggered by GPR55 in response to its agonists. In addition the cytoskeleton rearrangement mediated by GPR55 is investigated.

## Materials and methods

HEK-293 cells stably expressing the human GPR55 receptor were characterized in terms of signaling properties. To this end, reporter gene, dynamic mass redistribution (DMR), mitogen-activated protein kinases (MAPK) activation and phalloidin actin staining assays have been performed.

## Results

Here we show that GPR55 is activated by lysophosphatidylinositol (LPI), AM251, SR141716A (rimonabant) and AM281. GPR55 activation induces NF- $\kappa$ B, NFAT and CREB activation. Stimulation of GPR55 induces F-actin formation under the control of G $\alpha$ 13, RhoA and ROCK. We also show the suitability of Corning<sup>®</sup> Epic<sup>®</sup> DMR assay for GPR55 ligand screening. Furthermore, GPR55 activation leads to phosphorylation of extracellular signal-regulated kinase 1/2 (ERK 1/2).

## Conclusions

GPR55 as the novel cannabinoid receptor triggers distinct signaling pathways in response to LPI and some classical CB<sub>1</sub> receptor inverse agonists/antagonists.

Stress fiber formation mediated by GPR55 might indicate the probable function of this receptor *in vivo*.

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Published: 16 November 2010

doi:10.1186/1471-2210-10-S1-A8

**Cite this article as:** Balenga et al.: Signalling and function of the human G protein-coupled receptor 55. *BMC Pharmacology* 2010 **10**(Suppl 1):A8.

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