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Inhibitors, Agonists, Screening Libraries

# GPR55

## G protein-coupled receptor 55

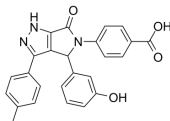
GPR55 (G protein-coupled receptor 55) is a G protein-coupled receptor that in humans is encoded by the GPR55 gene. GPR55, along with GPR119 and GPR18, have been implicated as novel cannabinoid receptors. GPR55 is activated by the plant cannabinoids 9-THC and cannabidiol, and the endocannabinoids anandamide, 2-AG, noladin ether in the low nanomolar range. Recent research suggests that lysophosphatidylinositol and its 2-arachidonoyl derivative may be the endogenous ligands for GPR55, and the receptor appears likely to be a possible target for treatment of inflammation and pain as with the other cannabinoid receptors. The physiological role of GPR55 is unclear. GPR55 has been proposed as a new potential drug target for the treatment of diabetes, Parkinson's disease, neuropathic pain, and cancer.

## GPR55 Antagonists

### CID 16020046

Cat. No.: HY-16697

CID 16020046 is a potent and selective **GPR55** antagonist and inhibits GPR55 constitutive activity with an  $IC_{50}$  of 0.15  $\mu$ M. CID 16020046 inhibits GPR55-mediated  $Ca^{2+}$  signaling and GPR55-mediated ERK1/2 phosphorylation.



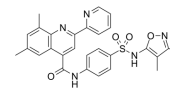
**Purity:** 99.92%  
**Clinical Data:** No Development Reported  
**Size:** 10 mM  $\times$  1 mL, 10 mg, 50 mg

### ML-193

(CID 1261822)

Cat. No.: HY-110125

ML-193 (CID 1261822) is a potent and selective antagonist of **GPR55**, with an  $IC_{50}$  of 221 nM. ML-193 shows more than 27-fold selectivity for GPR55 over GPR35, CB1 and CB2. ML-193 can improve the motor and the sensorimotor deficits of Parkinson's disease (PD) rats.



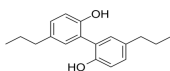
**Purity:** >98%  
**Clinical Data:** No Development Reported  
**Size:** 5 mg, 10 mg, 25 mg, 50 mg, 100 mg

### Tetrahydromagnolol

(Magnolignan)

Cat. No.: HY-116637

Tetrahydromagnolol (Magnolignan), a main metabolite of Magnolol, is a potent and selective **cannabinoid CB2 receptor** agonist with an  $EC_{50}$  of 170 nM and a  $K_i$  of 416 nM. Tetrahydromagnolol possesses 20-fold more selective for **CB2 receptor** than CB1 receptor.



**Purity:** >98%  
**Clinical Data:** No Development Reported  
**Size:** 1 mg