BigLEN-GPR171 Modulates Opioid Analgesia

Chronic pain is a huge problem that affects about one-third of the U.S. population. Despite many years of research for alternatives, opioids remain the most prescribed drugs to treat many types of pain, even though they possess many side effects including addiction. We are exploring the novel neuropeptide-receptor system (BigLEN-GPR171) as a pain therapeutic. To date we find that GPR171 is highly expressed in the periaqueductal gray, a region crucial to the pain-relieving actions of opioids, and is found in cells that co-express mu opioid receptors. We found that in mice a GPR171 antagonist reduces the anaglyxic effects of morphine, while a GPR171 agonist increases those effects. This leads us to suggest that GPR171 is modulating the effects of morphine, perhaps in the periaqueductal grey, through the actions of the mu opioid receptor.

GPR171 Agonist Decreases Chronic Pain

First-line chemotherapies against solid tumors are highly efficacious in reducing the tumor burden, but have many adverse side-effects including nerve damage, leading to chronic pain. Non-addictive, efficacious pain relievers are an area of active interest, and we propose a novel target to address this pressing issue. Mice in chronic pain experience allodynia, in which hypersensitivity to small applications of pressure elicits a pain response. We found that 5 days of GPR171 agonist treatment led to an improvement in their hypersensitivity. While there is a decrease in GPR171 receptors in the PAG of mice that have chronic pain, the agonist can bind to the available receptors to produce pain relief.

GPR171 agonist can relieve touch hypersensitivity caused by neuropathic pain

BigLEN-GPR171 in Opioid Tolerance, Withdrawal and Addiction

The BigLEN-GPR171 system shows high expression in many brain regions involved in reward, such as basolateral amygdala, hippocampus, and prefrontal cortex. In order for GPR171 compounds to be used as a therapeutic for pain they must show reduced side effects compared to opioids. We find that GPR171 agonist decreases morphine tolerance and has no effect on morphine withdrawal or reward. This suggests that when combined with morphine we can enhance the positive effects (pain relief) but decreases the unwanted adverse effects after long-term administration.