**The Role of Allopregnanolone in Neuroinflammation**

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**Introduction**

Neurosteroids are part of a new class of steroid hormones that regulate neuronal functions by acting as ligands for neurotransmitter receptors like GABA₆ and NMDA receptors (1). The mechanisms through which these compounds operate are still largely unknown. Previous research has shown that allopregnanolone (ALLO), a metabolic derivative of the more common steroid hormone progesterone, plays several neuroprotective roles, one of which is the reduction of neuroinflammation in certain neurodegenerative disorders (2).

The goal of our project was to determine the molecular mechanism—that is, the receptor—through which ALLO is working to exert its neuroprotective effects. The two candidate receptors we studied are the ligand-gated ion channel GABA₆ receptor, for which ALLO is already a known modulator, and the bile acid G-protein coupled receptor TGR5. Both receptors have been implicated in cell-mediated immunity (3).

**Synthesis of Allopregnanolone**

- CHOLESTEROL
- P450scc
- PREGNENOLONE
- 3α-HSD
- PROGESTERONE
- 5α-DIHYDROXYPROGESTERONE
- 3α-HSD
- 3α,5α-TETRAHYDROPROGESTERONE (ALLOPREGNANOLONE)

**Data & Results**

**Part 1: GABA₆ Receptor and Cell Proliferation Assays**

![Graph showing effect of allopregnanolone and GABA on RAW 264.7 mouse macrophage BrdU incorporation. Methods: 5x10⁵ cells were plated. ALLO, GABA, or ALLO & GABA was added for 72 hours. BrdU was added in the last 24 hours. Cells were fixed and incubated with a BrdU antibody and a colorimetric substrate. Absorbance, a measure of cell proliferation, was read at 370 nm (y-axis). N = average of 12 wells/treatment ± SEM. Results: ALLO and GABA reduce RAW cell proliferation.](image)

**Part 2: ALLO Stimulation of TGR5 Bile Acid Receptor**

![Graph showing effect of ALLO on 293 kidney cells that were transiently transfected with hTGR5 receptor cDNA and CRE-Luc DNA (a cAMP reporter). Methods: 24 hours after transfection, ALLO or LCA, lithocholic acid (a known TGR5 receptor ligand) was added for 24 hours. Stimulation increased cAMP synthesis, leading to an activation of CRE-Luc which causes Luciferase synthesis. Fluorescent light detection was used to measure Luciferase activity. Light was quantified in Relative Light Units (RLU). Results: ALLO stimulates the TGR5 receptor in a dose-dependent manner.](image)

**Possible ALLO pathways via TGR5 receptor:**

- ALLO → TGR5 receptor
- → activated TGR5 receptor
- → cAMP → ERK
- → Gα
- → IP₃, ERK
- → β-arrestin
- → CAMP → ERK
- → ERK

**Figure 1:** Effect of allopregnanolone and GABA on RAW 264.7 mouse macrophage BrdU incorporation. Methods: 5x10⁵ cells were plated. ALLO, GABA, or ALLO & GABA was added for 72 hours. BrdU was added in the last 24 hours. Cells were fixed and incubated with a BrdU antibody and a colorimetric substrate. Absorbance, a measure of cell proliferation, was read at 370 nm (y-axis). N = average of 12 wells/treatment ± SEM. Results: ALLO and GABA reduce RAW cell proliferation.

**Figure 2(a):** Effect of ALLO on 293 kidney cells that were transiently transfected with hTGR5 receptor cDNA and CRE-Luc DNA (a cAMP reporter). Methods: 24 hours after transfection, ALLO or LCA, lithocholic acid (a known TGR5 receptor ligand) was added for 24 hours. Stimulation increased cAMP synthesis, leading to an activation of CRE-Luc which causes Luciferase synthesis. Fluorescent light detection was used to measure Luciferase activity. Light was quantified in Relative Light Units (RLU). Results: ALLO stimulates the TGR5 receptor in a dose-dependent manner.

**Figure 2(b):** Western blot showing amount of ERK phosphorylation at various time points after ALLO stimulation of 293 cells transiently transfected with TGR5 receptor cDNA.

**Conclusions**

- ALLO reduces RAW cell proliferation
- ALLO appears to work like GABA in reducing RAW cell proliferation
- ALLO may work through a GABA₆ receptor to reduce neuroinflammation
- We have identified a new receptor for ALLO—the TGR5 receptor
- TGR5 is expressed in the brains of normal mice
- Binding of ALLO to TGR5 may contribute to the neuroprotective effects of ALLO that we have seen in mouse models of neurodegeneration.

**Future Directions**

- Create a plasmid containing TGR5 cDNA and the neomycin resistance gene to better select for transfected 293 cells and to further study signaling caused by ALLO stimulation
- Study the effect of ALLO on intracellular calcium levels (flow cytometry experiments)
- Study the effect of ALLO on lymphocyte NMDA receptors

**References & Acknowledgements**


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