

# ECE 2023

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## AZP-3813, a Bicyclic, 16-Amino Acid Peptide Antagonist of the Human Growth Hormone Receptor, Effectively Suppresses IGF1 in Beagle Dogs

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# CONFLICT OF INTEREST

Michael D. Culler

**X I have the following potential conflicts of interest to report:**

☐ Research Contracts

☐ Consulting

**X Employment in the Industry (Employee of Amolyt Pharma)**

**X Stockholder of a healthcare company**

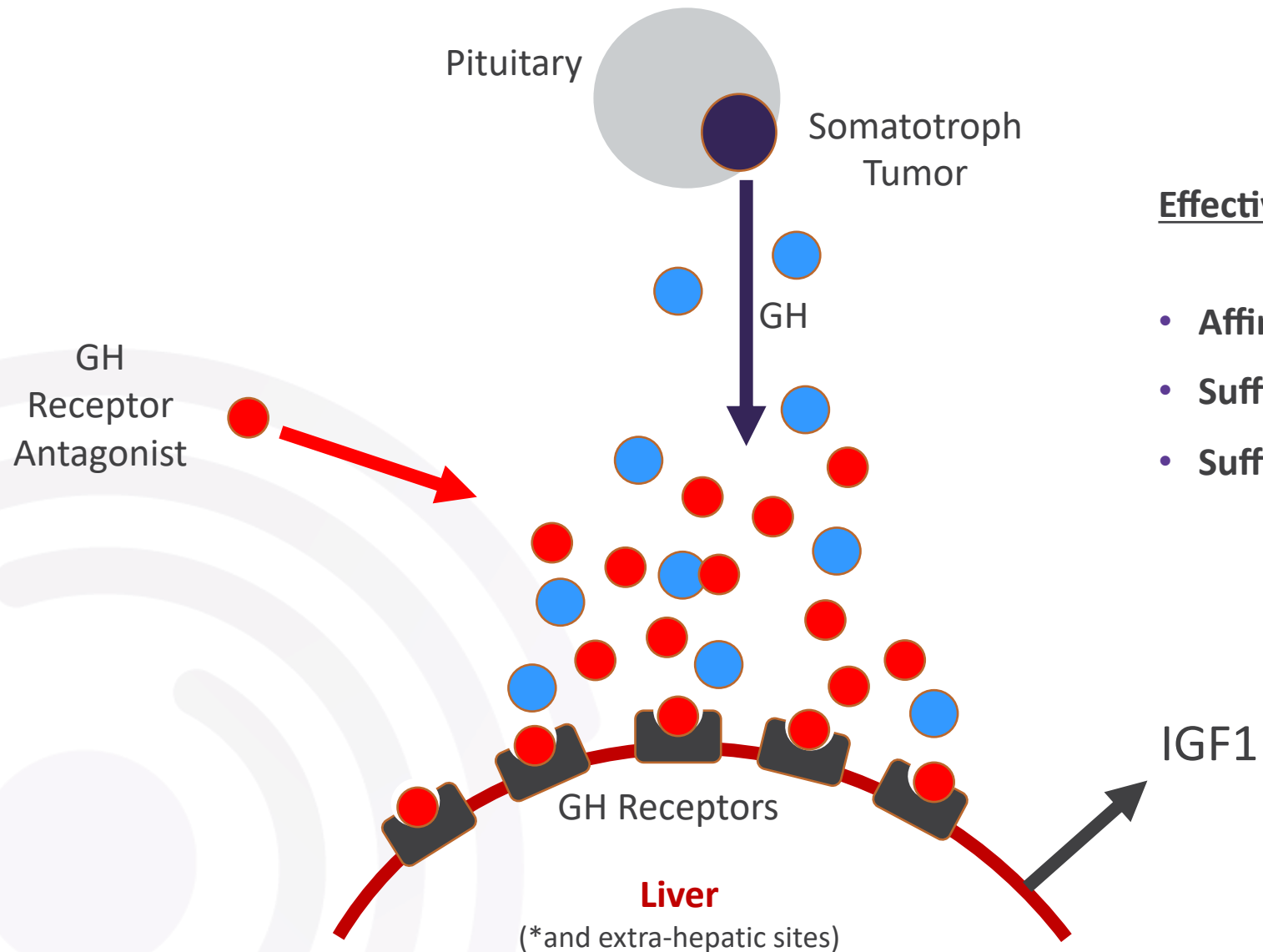
☐ Owner of a healthcare company

☐ Other(s) – *please include details*

*No commercial logos or product names to be included please.*

☐ I declare that I have no potential conflict of interest.

# GH Receptor Antagonist to Suppress Excess IGF1 in Acromegaly



## Effectiveness Depends On:

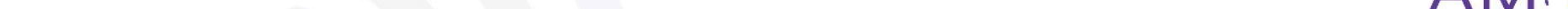
- Affinity for GH receptor
- Sufficient concentration
- Sufficient half-life

# AZP-3813 – GH Receptor Antagonist

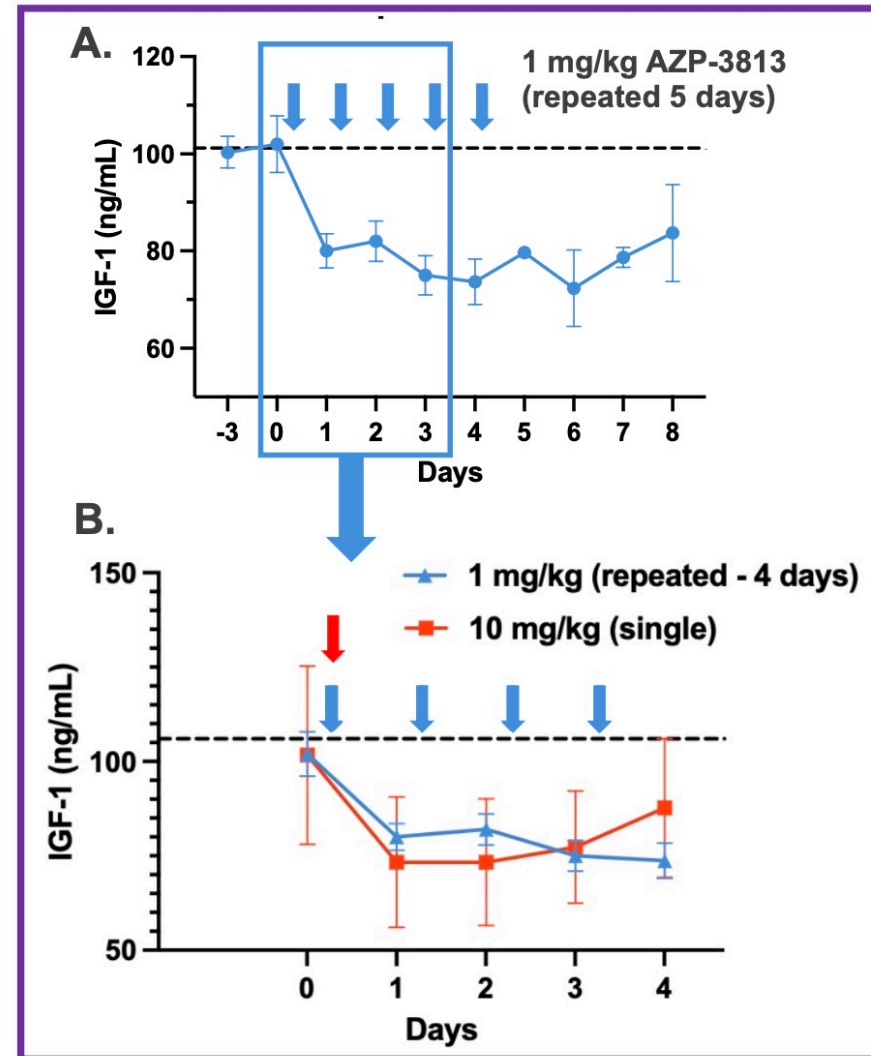
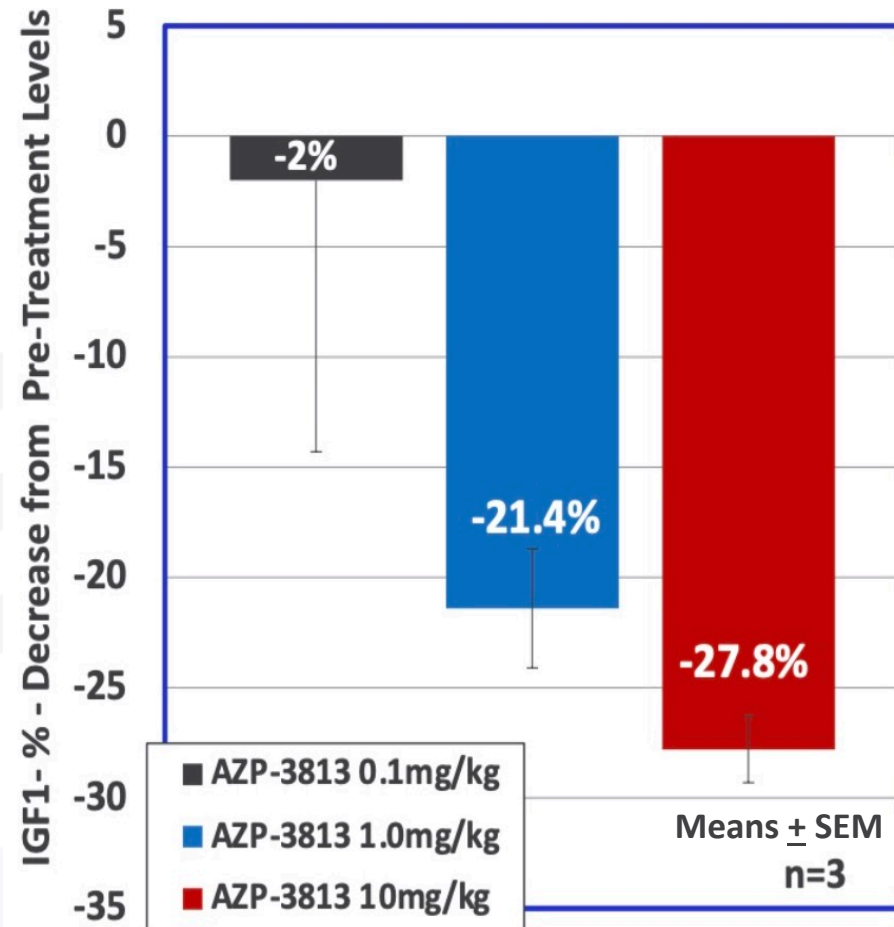
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## AZP-3813: 16 Amino Acid, Bi-Cyclic Peptide

- MW = 2479.9
- hGH-R affinity ( $K_D$ ) = 2.9nM
- hGH-R antagonism ( $IC_{50}$ ) = 9.9nM



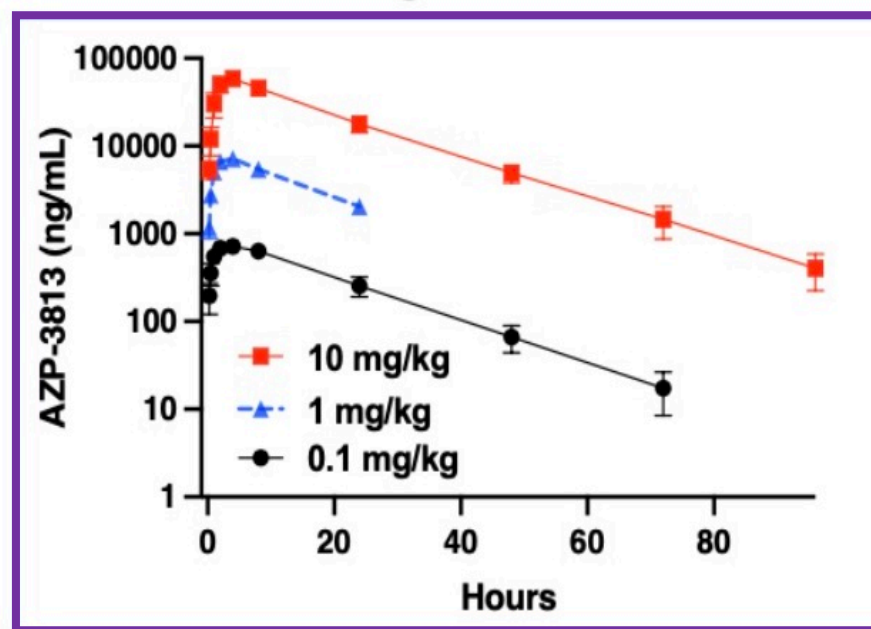
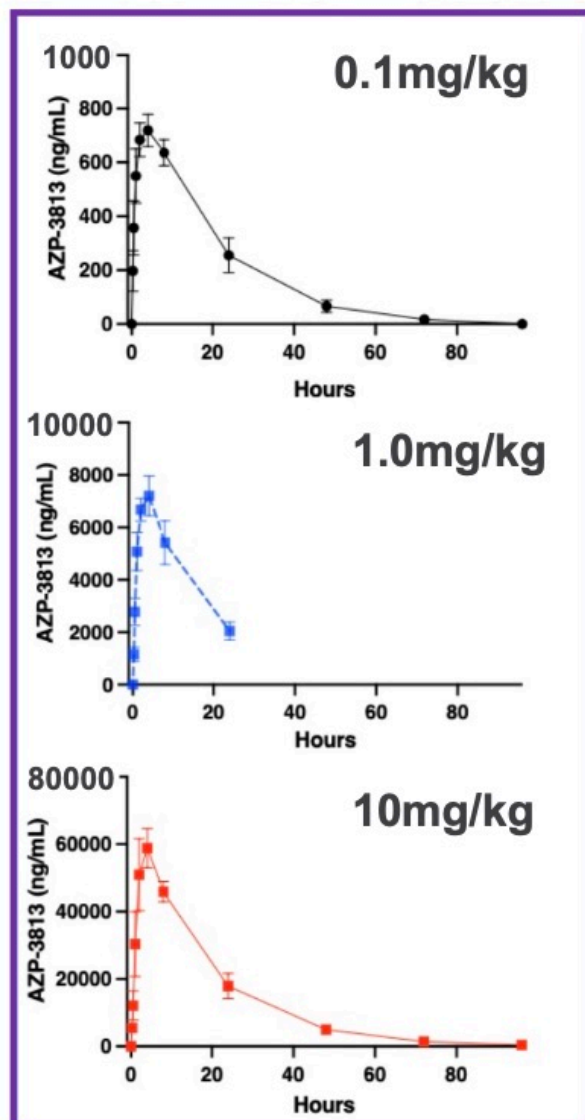
# Dose-Related and Sustained Suppression of IGF1 in Beagle Dogs





# AZP-3813 Pharmacokinetics in Beagle Dogs

Increasing doses of AZP-3813 produce linear increases in blood concentration



Increasing doses of AZP-3813 produce parallel elimination curves

Dose	Half-Life (h)
0.1mg/kg	15.1
1.0mg/kg	13.2
10mg/kg	14.3

Mean circulating half-life =  $14.2 \pm 0.47$  hours

# Conclusion

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The results of the current study demonstrate that the potent GH receptor antagonist activity exhibited by AZP-3813 translates to effective, sustained in vivo suppression of IGF1 levels in normal Beagle dogs, and further support the development of AZP-3813 as a potential therapy for acromegaly

For more detail, please visit our poster: [RC 7.3](#)

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Thank you!