
VA ID Number: 03-084

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Location: Miami, FL

Topic: Oncology

USPTO Issue Date: 10/18/2008

Patent Number: 7,452,865

The VA has a joint ownership interest with The Administrators of the Tulane Educational Fund

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Abstract There is provided a novel series of synthetic antagonistic analogs of hGH-RH(1-29)NH₂. These analogs inhibit the activity of endogenous hGH-RH on the pituitary GH-RH receptors, and therefore prevent the release of growth hormone. The analogs also inhibit the proliferation of human cancers through a direct effect on the cancer cells. The higher inhibitory potencies of the new analogs, as compared to previously described ones, results from replacement of various amino acids.