

## "Sulfated Gallic Acids and Uses Thereof"

VCU #12-96

## **Applications**

- Therapy for thromboembolic disease and cardiovascular disease
- Anticoagulant
- Cardiovascular disease
- Possible use as a therapeutic agent for cancers

## **Advantages**

- Allosteric mechanism facilitating high specificity
- Aromatic architecture enhances selectivity and potency
- Limited off-target interactions including intracranial bleeding and fetal toxicity
- Minimal risk of bleeding
- Easy high yield preparation

### **Inventors**

Umesh R. Desai, Ph.D. Rami A. Al-Horani, Ph.D.

#### Contact

Magdalena K. Morgan, Ph.D. Licensing Associate mkmorgan@vcu.edu Direct 804-827-6095

#### **Market Need**

Thromboembolic diseases are amongst the most frequent causes of death around the globe. Anticoagulants are the main means of treatment and prevention of thromboembolic disorders. Clinically, heparins have been extensively used as an anticoagulant both for treating thrombotic disorders and for many patients undergoing surgery. The use of heparins come with a number disadvantages. Heparins usually consist of a heterogeneous mixture, and thus contain many additional ingredients which result in a variety of side effects. Additionally, heparins are highly sulfated molecules, and thus highly hydrophilic and result in poor bioavailability. Some of the side effects observed with heparin usage are thrombocytopenia, osteoporosis, as well as others. There is a clinical need for a new anticoagulant with heparin-like activity without the many side effects currently observed with its use.

# **Technology Summary**

The Technology is a group of direct, allosteric, small molecule Factor XIa inhibitors. Factor XIa is part of the intrinsic pathway of the coagulation cascade. By targeting through this pathway, there is a lower chance of bleeding. There are several novel molecules that have been developed in this new class. These molecules are the first to use an allosteric mechanism to inhibit this coagulation factor. Their aromatic architecture enhances selectivity and potency by promoting hydrophobic interactions. Off-target interactions including crossing the blood brain barrier and fetal toxicity are avoided by limiting the number of sulfate groups as compared to heparins. Off-target interactions are also limited by the high selectivity of the molecules for Factor XIa. These molecules are good for commercial applications due to their easy, high yield preparation.

## **Technology Status**

In vitro testing has been performed on the proposed molecules in clinically relevant settings.

Publications: Al-Horani et al 2013

Patent Pending: US and Foreign Rights available.

This technology is available for licensing to industry for further development and commercialization.