



## The novel $\alpha_{IIb}\beta_3$ antagonist TMV-7 and its derivative RR prevent thrombosis without increasing bleeding risk

**PI :** Prof. Tur-Fu, Huang

Department of Pharmacology, College of Medicine, National Taiwan University

### **Experience:**

1985~ 1991 Associate Professor, Department of Pharmacology, College of Medicine, NTU

1989~ 2001 Outstanding Research Award, National Science Council, Taiwan.

2001 Scientific Technology Award, Executive Yen, ROC

2002~2008 Chair, Department of Pharmacology, College of Medicine, NTU

2009 Outstanding Innovative Research Award, National Taiwan University

1991~present Professor, Department of Pharmacology, College of Medicine, NTU

Publications: <http://ppt.cc/sjpEM>

### **Market Needs:**

Use of antithrombotic therapy, including antiplatelet and anticoagulant agents, is a vital element in reducing the overall morbidity and mortality in patients with cardiovascular disease. Improving the problem that clinical anti-thrombotics have significant bleeding risk.

### **Our Technology:**

Disintegrin TMV-7 and its derivative RR prevented thrombosis without increasing bleeding risk by selectively inhibiting  $G\alpha_{13}$ -binding without affecting talin-binding to  $\beta_3$  in human and mouse thrombin-activated platelets and causing agonist-induced PAC1-binding to  $\alpha_{IIb}\beta_3$  and clot retraction, processes of hemostasis driven by talin. At efficacious antithrombotic doses, both TMV-7 and RR had no effect on tail-bleeding time even given at higher dose (i.e., 2.5  $\mu\text{g/g}$ , 20-fold higher), indicating that they are efficacious antithrombotic  $\alpha_{IIb}\beta_3$  antagonists with a greater safety profile than the current  $\alpha_{IIb}\beta_3$  antagonists. Furthermore, TMV-7 is also a potent antiarrhythmic agent with cardioprotective properties in rats with myocardial ischemia-reperfusion injury.

### **Strength:**

We have designed the optimal anti-thrombotic agents with better safety profile. We are constructing a new RR derivative using a specific N-terminal PEGylation technique to enhancing its pharmaceutical advantages and pharmacological activities.

### **Competing Products:**

Tirofiban, eptifibatide, and abciximab

### **Intellectual Properties:**

Provisional Patent (14/126,808)

### **Contact (do not need to fill out):**

Center for Industry-Academia Cooperation, NTU

Tel: 02-3366-9945, E-mail: [ntuciac@ntu.edu.tw](mailto:ntuciac@ntu.edu.tw)

This information herein is intended for potential license of NTU technology only. Other usage of all or portion of this information in whatever form or means is strictly prohibited. Kindly contact us and we will help to achieve your goal the best we can.