



## ***12th Symposium on Glycosaminoglycans***

**September 23th-25th, 2004**

**Organizers:**

**Job Harenberg**

**4<sup>th</sup> Department of Medicine Faculty of Clinical Medicine  
Mannheim**

**Benito Casu**

**„G. Ronzoni“ Institute for Chemical and Biochemical Research,  
Milan**

## **Welcome to Villa Vigoni 2004**

As for the past eleven editions, the 12<sup>th</sup> Glycosaminoglycan Symposium will be hosted in Villa Vigoni (Lovenno, Lake Como). Started as Italian-German meetings (the villa is a property of the German Government), the symposia have become more and more international, with eminent specialists from all over the world invited to contribute to discussion of topics of actual or emerging interest in the field of chemistry, biochemistry, biology, pharmacology, and clinical applications of heparin and other glycosaminoglycans. Both basic and applied aspects will be covered and discussed, with about twenty percent of participants being associated with industrial companies. Participation in the symposium (only by invitation) is limited to a maximum of sixty participants.

Topics selected for discussions this year include the sequence-dependence of biological activity of glycosaminoglycans; differentiation of low-molecular weight heparins; heparanase inhibitors as potential antiangiogenic and antimetastatic drugs; clinical assessment of heparins and non-GAG antithrombotic agents, and (in a minisymposium devoted to new therapeutic targets) GAGs in Alzheimer disease. Keynote speakers will outline the state-of-the art in each field and present novel results. A sufficient time will be allotted to in-depth discussion. The informal, workshop-like character of the symposia and the pleasant surroundings of the villa traditionally stimulate also after-session interactions among participants.

The 12<sup>th</sup> Glycosaminoglycan Symposium will start in the early afternoon of Thursday, September 23 and end the early afternoon of Saturday, September 25. Accommodation for most of the participants will be arranged in Villa Vigoni and in its surroundings. Accommodation can be arranged also in hotels in Menaggio. Meals will be served in the villa.

Invited participants will receive further information in due course.

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## **Main topics**

### *Anticoagulants*

argatroban, dermatansulfate, glycosaminoglycans, heparins, hirudins, low-molecular-weight heparins, pentasaccharide, new antithrombotic drugs, thrombin inhibitors

### *Characterization of Glycosaminoglycans*

Anticoagulant assays, nuclear magnetic resonance, new biological assays, standardization and harmonization of methods

### *Structure-function relationship*

Drug interactions of anticoagulants, glycosaminoglycans and platelet factor 4, glycosaminoglycans and growth factors, glycosaminoglycans and inflammation, heparin-induced thrombocytopenia, protein-glycosaminoglycan interactions, glycosaminoglycans and Alzheimer's disease

### *Polysaccharides and biomaterials*

Coating of stents, coating of biological materials

### *Clinical studies*

Deep venous thrombosis and pulmonary embolism, glycosaminoglycans, ischemic stroke, myocardial infarction, new indications, prolonged prophylaxis of venous thromboembolism, pentasaccharides, thrombin inhibitors

## **Scientific board**

Breddin HK, Casu B, Harenberg J, Palumbo M, Cornelli U.

### **Sponsors:**

Aventis Glycochemistry, Vitry Sur Seine, France  
Aventis Pharma Deutschland GmbH, Bad Soden, Germany  
Aventis Pharma, US  
BioTie Therapies, Turku, Finland  
Celltech Pharma GmbH & Co.KG, Essen, Germany  
Cornelli Consulting Srl, Milan, Italy  
DFG (Deutsche Forschungsgemeinschaft, Bonn, Germany  
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LabService Analytica, Bologna, Italy  
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Laboratorio Derivati Organici (LDO), Trino Vercellese, Italy  
Momenta Pharmaceuticals, Boston  
Opocrin Biofarmaci, Corlo di Formigine, Italy  
Pentapharm Ltd, Basel, Switzerland  
Sanofi-Synthelabo, Germany  
Sigma-Tau, Pomezia, Italy

Thursday, September 23<sup>rd</sup>, 2004

14.30-14.40

**Harenberg J, Mannheim and Casu B, Milan**  
Welcome address

<b>Session I</b>	<b>Heparanase - Biology and Pharmacology</b> <b>Moderators: Lindahl U, Uppsala, Vlodavsky I, Jerusalem</b>
<b>14.40-15.00</b>	<b>Steinkühler C, Pomezia</b> Investigating structure and function of human heparanase by protein engineering
<b>15.00-15.20</b>	<b>J.-p. L, Uppsala:</b> Minimal heparin/heparansulfate sequences cleaved by heparanase
<b>15.20-15.40</b>	<b>Vlodavsky I, Jerusalem</b> Biological activities of heparanase
<b>15.40-16.00</b>	<b>Coffee and tea</b>
<b>16.00-16.20</b>	<b>L. Borsig, Zürich</b> Mechanisms for antimetastatic activity of polysulfates
<b>16.20-16.40</b>	<b>Casu B, Milan</b> Strategies for developing heparan sulfate mimics as heparanase inhibitors
<b>16.40-17.10</b>	<b>Discussion</b>
<b>17.10-17.30</b>	<b>Wenzel Thea, Vienna</b> <i>Sculpia Vienne Austriae MDCCXXXVII Lorenzo Matielli F – Lorenzo Matielli in Loveno and Vienna, the villa of the Mylius family in Loveno</i>

Friday, September 24<sup>th</sup>, 2004

<b>Session II</b>	<b>Novel biological implications of glycosaminoglycans</b> <b>Moderators:</b> <b>Petitou M, Toulouse</b> and <b>Salmivirta, M, Turku</b>
<b>9.00-9.20</b>	<b>Lindahl U, Uppsala</b> Structure of heparan sulfate – how regulated does it need to be?
<b>9.20-9.40</b>	<b>Gallagher J, Manchester</b> Heparin motifs for the assembly of growth factor signalling complexes
<b>9.40-10.10</b>	<b>Baici A, Zürich</b> Paradoxical effects in glycosaminoglycan-protein interactions
<b>10.10-10.30</b>	<b>Nowak G, Jena</b> Molecular weight extended direct antithrombotic drugs with anticancer activity
<b>10.30-10.50</b>	<b>Coffee, tea</b>
<b>10.50-11. 10</b>	<b>Sasisekharan R, Boston</b> Promoters of cellular uptake of heparin and cancer cell death
<b>11.10-11.30</b>	<b>Yates EA, Liverpool</b> Engineered heparins: a strategy for optimizing biological activities of heparin
<b>11.30-11.50</b>	<b>Venkataraman G, Boston</b> <i>To be defined</i>
<b>11.50-12.15</b>	<b>Discussion</b>

<b>Session III</b>	<b>Advances in low- and very low-molecular weight heparins</b>
	<b>Moderators: Fareed J, Maywood, and Nowak G, Jena</b>
<b>14.00-14.20</b>	<b>Viskov C, Vitry sur Seine</b> Process-dependent generation of heparin oligosaccharides with different location of antithrombin-binding sequences
<b>14.20-14.40</b>	<b>Laux V, Frankfurt</b> Octaparin, a new well-characterised ultra low-molecular-weight heparin
<b>14.40-15.00</b>	<b>Bianchini, P, Modena</b> Coagulation and osteoporosis. Deligoparin inhibits arterial calcification in "Villaverde" rats
<b>15.00-15.15</b>	<b>Guerrini M, Milano</b> Structural characterization of low-molecular-weight heparins by two-dimensional NMR spectroscopy
<b>15.15-15.35</b>	<b>Alban S, Kiel</b> From the production to the clinics – differences between LMWHs
<b>15.35-15.50</b>	<b>Bisio A, Milan</b> Characterization of ultra-low-molecular weight heparins obtained by physical methods
<b>15.50-16.00</b>	<b>Discussion</b>
<b>16.00-16.15</b>	<b>Coffee, tea</b>

<b>Session IV</b>	<b>Clinical assessment of glycosaminoglycans. Comparison with other antithrombotics</b>
	<p><b>Moderators: Palumbo M, Padova, Kirchmaier C, Wiesbaden</b></p> <p><b>16.15-16.30 Harenberg J, Mannheim</b> Anticoagulant effects by inhalation of low-molecular-weight heparin certoparin by new therapeutic targets</p> <p><b>16.30-16.45 Sasisekharan R, Boston</b> Novel results on differential adsorption of heparin oligosaccharide administered by non-invasive (nasal) route</p> <p><b>16.45-17.00 Markwardt F, Erfurt</b> Historical perspectives in the development of thrombin inhibitors as antithrombotic drugs</p> <p><b>17.00-17.15 Stemberger A, Munich</b> Antithrombotic and anti-infective coating of biomaterials: both reduce thrombogenicity</p> <p><b>17.15-17.30 Fenyvesi T, Mannheim</b> Inhibition of thrombin generation and platelet aggregation by factor Xa and factor IIa inhibitors</p> <p><b>17.30-17.45 Calatzis A, Munich</b> Monitoring of Heparins, Heparinoids and direct inhibitors: How can we cope with standardization and calibrations issues in clinical practice?</p>
<b>17.45-18.00</b>	Discussion

Saturday, September 25<sup>th</sup>, 2004

<b>Session V</b>	<b>Minisymposium</b> (organized by U. Cornelli):  <b>New therapeutic targets: Glycosaminoglycans and Alzheimer's disease</b>
<b>9.00-9.20</b>	<b>Moderator: Cornelli U, Milano</b>  <b>Lindah U, Uppsala</b> Heparan sulfate and amyloid disease
<b>9.20-9.40</b>	<b>Turnbull JE, Liverpool</b> Heparin turns down the BACE: novel heparan sulfate-protein interactions in Alzheimer's disease
<b>9.40-10.00</b>	<b>Bergamaschini L, Milan</b> Peripheral treatment with a low-molecular-weight heparin reduces plaques and beta-amyloid accumulation in a mouse model of Alzheimer's disease
<b>10.00-10.20</b>	<b>Coffee, tea</b>
<b>10.20-10.40</b>	<b>Fareed J, Maywood</b> Differential action of heparin-derived oligosaccharides and LMWHs and relationships with Alzheimer disease
<b>10.40-11.00</b>	<b>Lee J, Maywood</b> Molecular-weight dependence in the prevention of amyloid toxicity in brain by LMWHs
<b>11.00-11.20</b>	<b>Hanin I, Maywood</b> Activity of different glycosaminoglycans in the protection of cholinotoxin AF64A injected into the brain
<b>11.20-11.40</b>	<b>Discussion</b>
<b>11.40-12.00</b>	<b>Casu B, Milan/Harenberg J, Mannheim</b> Closing remarks



## **General Information**

### **Social Program**

#### **Thursday, September 23, 2004**

**13.00 -14.00:**      *Lunch in Villa Vigoni*  
**19.00:**              *Dinner in Villa Vigoni*

#### **Friday, September 24, 2004**

**8.00-9.00:**         *Breakfast*  
**13.00 -14.00:**      *Lunch in Villa Vigoni*  
**19.00:**              *Dinner in Villa Vigoni*

#### **Saturday, September 25, 2004**

**8.00-9.00:**         *Breakfast*  
**12.30-13.30:**      *Lunch in Villa Vigoni*

### **Location of the Symposium:**

"Villa Vigoni e.V."  
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Lovenò di Menaggio  
I-22017 Menaggio (Como)  
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**Registration:**      Thursday, September 23, 2004

### **Organisers:**

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