

## **ADDRESS COLLOQUIA**

Venue: Medical University Vienna, Center for Physiology and Pharmacology,  
Institute of Pharmacology, Conference room, 3<sup>rd</sup> floor  
Währingerstrasse 13a, 1090 Vienna,  
(Harald H. Sitte, Tel.: (01) 40160 31323, [harald.sitte@meduniwien.ac.at](mailto:harald.sitte@meduniwien.ac.at))

---

Friday      17.3.2017   12.00 s.t.      Walter Sandtner (host: H. Sitte)

Medical University of Vienna  
Center of Physiology and Pharmacology  
Institute of Pharmacology  
Währingerstraße 13a  
1090 Vienna

### ***"Drugs with atypical properties targeting monoamine transporters."***

---

**Walter Sandtner** (walter.sandtner@meduniwien.ac.at)

---

Abstract:

The transporters for the monoamines serotonin, dopamine and norepinephrine (SERT, DAT and NET) are important drug targets. Ligands that bind to these transporters are used for treating a wide range of disorders such as depression, attention deficit hyperactivity disorder, neuropathic pain etc. Drugs which affect monoamine transporters usually fall into two categories : they are either inhibitors of monoamine uptake or they are transported as exogenous substrates. Here I will show that ligands exist that can be both: exogenous substrates and inhibitors. Moreover, I will demonstrate that the mode of action of these ligands is governed by the surrounding ionic composition which is sensed by the transporters.