

Thu 9. May 2019
Time: 16:00 h

University of Zürich -
Irchel, Winterthurerstr.
190, 8057 Zürich
Lecture Hall Y03-G-85

Everybody is welcome

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This lecture is hosted by
Prof. Raimund Dutzler
(UZH).

NCCR TransCure Lecture in Biology by Cristina Paulino

The KdpFABC Complex: What Happens When a P-type ATPase Hijacks an Ion Channel

P-type ATPases ubiquitously pump cations across biological membranes to maintain vital ion gradients. Among those, the chimeric K^+ uptake system KdpFABC is unique. While ATP hydrolysis is accomplished by the P-type ATPase subunit KdpB, K^+ has been assumed to be transported by the channel-like subunit KdpA. A first crystal structure uncovered its overall topology, suggesting such a spatial separation of energizing and transporting units. The use of single particle cryo-EM allowed us to determine two additional structures of the 157 kDa, asymmetric KdpFABC complex in an E1 and E2 state, at 3.7 Å and 4.0 Å resolution, respectively. Unexpectedly, the new structures suggest a so far unprecedented transport mechanism through two half-channels along KdpA and KdpB, uniting the alternating-access mechanism of actively pumping P-type ATPases with the high affinity and selectivity of K^+ -channels. This way, KdpFABC functions as a true chimeric complex, synergizing the best features of otherwise separately evolved transport mechanisms.