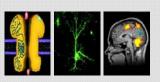


SFB 874 / IGSN

CONFERENCE



Cortical and subcortical representation of sensory and cognitive memory

April 28 - 29, 2015, Ruhr University Bochum

Wednesday April 29, afternoon (13:15 – 16:15)

Session 4: Cognition relevant information processing

at the subcortical level

OXANA ESCHENKO

Max Planck Institute for Biological Cybernetics, Tuebingen, Germany

The role of Locus Coeruleus for sensory processing within mesocortical dopaminergic pathway

Salient events evoke burst-like responses of noradrenergic (NE) neurons of the Locus Coeruleus (LC) and dopaminergic (DA) neurons of the ventral tegmental area (VTA). The associated NE and DA release modulates information processing in the projection targets of LC and VTA. In the rat, terminal fields of both LC-NE and VTA-DA neurons converge in the medial prefrontal cortex (mPFC), a cortical area controlling many cognitive capacities. We investigated the role of LC phasic activation for sensory responses in VTA and mPFC. Under urhetaine anestesia, noxious stimulation (foot shock, FS) produces a robust short-latency (~20 ms) excitation of LC-NE neurons. In VTA and mPFC, the firing rate modulation induced by FS was present in ~30% of neurons. We classified FS-induced responses of VTA neurons according to latency (early: ~40 ms or late: ~150 ms) and duration (phasic: < 300 ms or sustained: > 300 ms). Similarly, the mPFC single-unit responses differed by latency and/or duration. Supression of LC ongoing and FS-evoked activity by iontophoretic injection of clonidine, an alpha2-adrenergic receptor agonist, reduced responsiveness in both VTA and mPFC. Population of initially 'non-responsive' mPFC neurons showed 'gating-effect'. Spontaneous discharge of substantial proportion of VTA and mPFC neurons was bidirectionally modulated. These results suggest that depending on the motivational valence of a salient event, LC phasic activation and associated NE release may selectively enhance or supress signalling within different and, possibly competing mesolimbic and mesocortical pathways. The behavioral data supporting this hypothesis will be presented.



