

Neuroscience in China 2010–2011

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Over the past two years, there have been greater numbers of contributions made by Chinese scientists in various areas of neuroscience compared with the last 10 years, indicating the unprecedented rapid development of Chinese science. The research has related to various areas, including cellular and molecular neuroscience, developmental neuroscience, systems and computational neuroscience, mechanisms of neural disorders, and neuropharmacology in animal models.

Developmental neuroscience is always a “hot topic”, where progress is helpful not only to the understanding of the formation of neural networks, but also towards the reconstruction of a deficient nervous system. The developmental regulation and isolation of neural stem/progenitor cells (NSPCs) and their functions in neurogenesis have been reviewed [1]. Unlike fetal brain, the persistence of neuroblasts in adult brain is controversial. Recently, evidence from adult monkey and human brains suggests that neuroblasts may be generated in the subventricular zone (SVZ) and then continuously migrate to the rostral migratory stream (RMS) [2]. Furthermore, Cheng et al. [3] found that newborn neurons in the dentate gyrus (DG) migrated to the inner part of the granule cell layer using a mouse model based on the Cre/loxp system, providing an efficient way to selectively label and manipulate newborn neurons in the adult mouse DG. However, the integration of newborn neurons into the existing circuit is difficult because of their poor survival, insufficient proliferation and differentiation abilities. A ciliary neurotrophic factor (CNTF)-collagen gel-controlled delivery system was found to provide a physical scaffold for the transplanted NSPCs to adhere to and migrate, as well as facilitating NSPCs survival, growth and proliferation, and simultaneously reducing the consumption of expensive growth factors [4]. Treatment with salvianolic acid B can also increase the number of neural

stem cells (NSCs) and their derivative neurospheres, and promote the neurite outgrowth of NSCs and their differentiation into neurons [5].

Formation of the nervous system requires correct neuron functions, including axon guidance, neural migration [6], synaptic formation and remodeling [7], neural plasticity [8], glial cells [9], and neurite outgrowth. However, the precise neuroanatomical architecture remains unknown. A new tool for the mapping of neurite circuits at the mesoscale level, named micro-optical sectioning tomography, has been developed. It can provide a map in which the morphology and spatial locations of neurons and traces of neurites can be clearly distinguished, and was used to develop a 3D structural data set of a Golgi-stained whole mouse brain at the neurite level [10]. The strength of excitatory synaptic inputs in layer V pyramidal neurons of the medial prefrontal cortex were found to regulate social hierarchy in mice [11]. Myelination is another important part of neuronal development. In mice, the *Rheb1* transgene in neural progenitor cells increased mTORC1 activity and promoted myelination in the brain, indicating a crucial role for *Rheb1* [12]. Glial cell line-derived neurotrophic factor (GDNF) activation of neurite outgrowth has been widely reported. Its negative regulation has been studied and Rap1GAP is reported to interact with RET to suppress GDNF-induced neurite outgrowth [13]. Lg11 activation of Rab10 promotes axonal membrane trafficking required for neuronal polarization [14]. *N*-cadherin-dependent interaction between neurons is required for maintaining the activity-induced growth of dendrites [15].

The importance of signal transduction and signaling pathways in the nervous system are self-evident. However, mechanisms involved in the release of neurotransmitters, transport of receptors and interactions between receptors are still unclear. Recently, one study found cross-talk between NMDA and GABA_A receptors as well as Ca²⁺/calmodulin-dependent protein kinases II in cultured neurons of the rat inferior colliculus, a central auditory system [16]. Another

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study observed large dense-core vesicles (LDCVs) transfer various GPCRs, ion channels and peptides in the dorsal spinal cord, enabling a rapid, activity-dependent modulation of neuronal sensitivity [17]. Interaction between μ - and δ -opioid receptors in nociceptive afferent neurons is involved in morphine analgesic tolerance [18,19]. Moreover, follistatin-like 1 secreted from nociceptive afferent terminals acts as an activator of presynaptic sodium pump and suppresses excitatory neurotransmission [20]. Furthermore, the formation, function and plasticity of sensory circuits are studied in the *Drosophila* by several groups [21–23]. Interestingly, neuregulin-ErbB4 signaling is found to contribute to the mechanism of epilepsy [24,25]. Neurodegenerative diseases have been widely studied in terms of the features described above [26]. Recent data has shown that the concentration of copper ions and formaldehyde are highly related to the pathology of Alzheimer's disease (AD) [27,28], and the investigation of ion channels in neuronal survival during the past 10 years in China was recently reviewed [29]. More evidence indicates that formaldehyde can not only promote Tau to form pore-like aggregates, but also induce hyperphosphorylation of Tau and disrupt its protection of DNA [30]. Furthermore, Su et al. [31] modified 2,4-dinitrophenylhydrazine (2,4-DNPH) method to analyze endogenous formaldehyde with higher sensitivity and lower relative standard deviations using UV-HPLC. All these studies improve the research about pathology of AD. In Huntington's disease, an important domain of huntingtin protein containing Htt4-17 was shown to regulate the cytoplasmic localization of this protein and impact on aggregate formation [32]. The mechanisms of PD related proteins, α -synuclein and Parkin for example, have been summarized [33]. Biomarkers for neural diseases remain to be improved. Currently, there is no specific biomarker for early-stage AD in the cerebrospinal fluid or blood [34]. Although recent studies indicated thiobarbituric acid reactive substances, nitric oxide, catalase (CAT) and glutathione peroxidase (GP) (molecules related to oxidative stress) may be biomarkers for schizophrenia, further study and confirmation is required [35]. Both attention deficit hyperactivity disorder (ADHD) and schizophrenia are related to the dysfunction of midbrain dopamine neurons. A study in mice demonstrated that the GC-C/PKG signaling pathway may be responsible [36]. Non-coding RNA (ncRNA) has recently received more attention and given plenty of research results. The role of ncRNA in pathophysiological mechanism in several neurodegenerative diseases has been reviewed [37]. Some other research studied the formation and process of memory and cognition [38–41], which may help us to understand the deflection of cognition and memory in psychotic. More and more emphases are being laid on the studies of endophenotypes in neurodegenerative diseases [42–47].

Other studies have focused on drug usage and pharmacology especially in the central nervous system. The effective treatment of central nervous system disorders is limited

by the blood brain barrier (BBB). A simple diffusion system proposed by Han et al. [48] has been validated and can deliver drugs effectively through bypassing the BBB. Fu et al. [49] identified a selective GABA_B agonist, Baclofen, which can differentially modify cortical electroencephalography activity, elucidating the relationship between GABA_B receptor-mediated neurotransmission and aging. Drug addiction is common in pain treatment. Recent studies of the mechanism of pain is popular and reviewed by Zhang [50]. A new study by Chen et al. [51] showed that endogenous enkephalin may not be involved in the antihyperalgesic effects of gabapentin. The inflammation-induced afferent neurotransmission can be inhibited by activating B-Type natriuretic peptide signal pathway [52]. Moreover, intrathecal treatment of follistatin-like 1 could reduce chronic pain induced by nerve injury [53].

Computational neuroscience is an exciting new field that has developed over the last ten years, with multidisciplinary interactions including experimental neuroscience, robotics and computer vision [54]. The current review reports on the activity of rat pre-frontal cortex (PFC) neurons controlled by a one-dimensional (1D) machine using an encoding method to quench thirst [55]. This may benefit disabled patients. However, mature technology such as imageology plays important roles in other areas. For instance, high-resolution magic-angle spinning proton nuclear magnetic resonance spectroscopy and pattern recognition (HRMAS ¹H NMRS) are used in grade classification of neuroepithelial tumors according to metabolic fingerprints [56]. Based on number series completion, a typical data-driven scientific discovery task, the resulting data obtained by functional magnetic resonance imaging (fMRI) revealed the left dorso-lateral prefrontal cortex (DLPFC) is related to rule identification, while the left anterior prefrontal cortex (APFC) may be involved in mental set maintenance needed during rule identification and extrapolation [57]. Furthermore, using fMRI-BOLD signals and ERP, Zhang et al. [58] indicated a bottom-up saliency map is created in V1, challenging the dominant view that the saliency map is generated in the parietal cortex.

In neuroscience studies, greater attention is being directed towards the development of animal models. Since the well-known genetic background and measurable cognitive behaviors, *Drosophila* has been widely used as an animal model for research in visual cognition and information processing in the perspective of genes-brain-behavior [59,60]. To understand the regulatory mechanisms of dendritic development, retinal ganglion cells (RGCs) of mice have been used [61]. Furthermore, the rat was also used in behavior studies such as the influence of social isolation [62]. Interestingly, an animal model for social hierarchy has been developed and used to analyze the neural circuits controlling the social behavior [11]. In addition to sleep and emotion, orexin plays a role in central vestibular motor control [63].

Despite amazing progress over the past few decades, the

nature of the nervous system remains unresolved. For example, the generation and survival of NSPCs, neuronal migration and plasticity, synaptic formation and shaping and the neural circuits underlying functions still require further study. Elucidation of these mysteries requires multidisciplinary combination at different levels. We look forward to greater contributions from Chinese neuroscientists in these areas.

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