Steroids as external temporal codes act via microRNAs and cooperate with cytokines in differential neurogenesis

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The generation of neuronal cell diversity is controlled by interdependent mechanisms, including cell intrinsic programs and environmental cues. During development, the astonishing variety of neurons is originated according to a precise timetable that is managed by a complex network of genes specifying individual types of neurons. Different neurons express specific sets of transcription factors, and they can be recognized by morphological characteristics and spatial localization, but, most importantly, they connect to each other and form functional units in a stereotyped fashion. This connectivity depends, mostly, on selective cell adhesion that is strictly regulated. While intrinsic factors specifying neuronal temporal identity have been extensively studied, an extrinsic temporal factor controlling neuronal temporal identity switch has not been shown. Our data demonstrate that pulses of steroid hormone act as a temporal cue to fine-tune neuronal cell differentiation. Here we also provide evidence that extrinsic JAK/STAT cytokine signaling acts as a spatial code in the process. Particularly, in *Drosophila* mushroom bodies, neuronal identity transition is controlled by steroid-dependent microRNAs that regulate spatially distributed cytokine-dependent signaling factors that in turn modulate cell adhesion. A new era of neuronal plasticity assessment via manipulating external temporal cues such as hormones and cytokines that specify individual types of neurons might open new possibilities for brain regenerative therapeutics.

How Multiplicity of Neuron Types is Generated

The development of multiple compartments of the brain is a highly orchestrated process, where commitment of certain types of neurons to specific zones, layers and compartments is linked to the developmental stage, at which neurons are generated. During the last few years, significant progress has been made in the discovery of genes that identify and control development of different neuronal subtypes (reviewed in refs. 3–5). A subsequent series of intrinsic signaling programs are described in invertebrate and vertebrate organisms where neuronal progenitors in a time-dependent manner progressively acquire specific identity via expression of unique sets of genes that coordinate the generation of the multiple projection neuron subtypes.

Like in vertebrates, neuronal stem cells in *Drosophila* produce different types of neurons depending on embryonic anterior-posterior and dorsal-ventral polarity that establish gradients of morphogens and induce expression of gap, pair-rule and Hox genes that subsequently assemble a set of differentially expressed transcription factors. Following the lineage specification, the neuronal stem cell generates a characteristic set of neuron subtypes. The exact birthdate of specialized neurons suggests an interaction between temporal cues and neuron-intrinsic cell fate factors. Despite the broad data about existence of these intrinsic programs, it is important to note that the extrinsic temporal determinants of differential morphogenesis have not been revealed in any organism. We
Steroid Hormone Regulates Chronological Neurogenesis in Drosophila

As a model to study extended neurogenesis we use Drosophila learning center or mushroom body (MB) neurons that are responsible for olfactory learning and memory. MB neuron subtypes are generated in the same lineages by type I neuroblasts and specified in a birth-order-dependent fashion. MB γ and α'/β' neurons are produced during larval stages, while α/β neurons are born from the same neuronal precursors after transition from larval to pupal stages. This MB neuron diversification is coincident with key developmental time periods (Fig. 1). In Drosophila there are two systemic developmental timers—steroid ecdysone and juvenile hormone—that synchronize the genetic, morphological and behavioral changes associated with developmental transitions. Pulses of the steroid hormone ecdysone trigger major postembryonic developmental transitions, including molting and metamorphosis. Ecdysone interacts with a heterodimer of Ecdysone Receptor (EcR) and Ultraspiracle (Usp) — two members of nuclear receptor superfamily. This complex directly induces expression of primary-response targets, which in turn multiply hormonal signal by regulation of secondary-response gene transcription. These mechanisms determine stage- and tissue-specific responses to each developmentally regulated ecdysone pulse. Moreover, ecdysone signaling is patterned spatially as well as temporally; depending on the cell type and the developmental stage, the ecdysone receptor complex binds different co-activators or co-repressors that can have other binding partners, regulated by additional signaling pathways. For example, the putative transcription factor Abrupt attenuates ecdysone signaling by binding to its co-activator Taiman, and we showed that this interaction plays an important role in cell non-autonomous regulation of early germline progeny differentiation. Moreover, other signaling pathways discovered that in Drosophila steroid hormones regulate the chronological neuronal identity switch that is executed by steroid-dependent microRNAs (miRNAs) (Fig. 1).

Figure 1. Model of differential neurogenesis regulation by cooperation of developmentally controlled temporal systemic signaling and intrinsic spatiotemporal codes. Scheme represents the chronologically regulated signaling cascade controlling α'/β' to α/β neuronal identity switch in the Drosophila MB that takes place at the larva-to-pupa developmental transition. Amount of ecdysone at different stages of development is represented as relative levels (scheme adopted from ref. 15). Developmentally regulated pulse of the steroid hormone ecdysone acts as an extrinsic temporal signaling code to activate expression of miRNAs from the let-7 complex in the differentiating MB neurons. Temporally induced miRNAs let-7 and miR-125 are intrinsic spatiotemporal codes that downregulate at least two BTB domain containing transcription factors Abrupt and Chinmo, which allows for the α'/β' to α/β neuronal cell fate transition. Cell adhesion molecule FasII is downstream of let-7/Abrupt signaling. During larval stages Abrupt suppresses FasII expression allowing for early-born lobes to be formed, while at the pupal stage downregulation of Abrupt allows FasII expression and promote α/β neuronal differentiation.
Hormones and MicroRNAs

Development of the living organism is organized into discrete temporal stages, each of which is characterized by a unique program of gene expression that controls tissue formation and differentiation. miRNAs were first found because of their role in the regulation of developmental staging of the nematode *C. elegans*. Multiple studies in insects also suggest an important role for miRNAs in the coordination of the developmental transitions; depletion of Dicer-1 (protein required for miRNAs biogenesis) in *B. germanica* and mutations in *Drosophila* miRNAs *let-7* and *miR-125* impair regulation of metamorphic processes. The temporal regulation of these and many other miRNAs expression is mediated by developmentally controlled hormonal signals. For example, in *Drosophila*, the upregulation of *miR-100, miR-125*, and *let-7* encoded by the miRNA *let-7*-C locus and downregulation of *miR-34*, *miR-14*, and *miR-8* require the steroid hormone ecdysone. Recent work from Chawla and Sokol identified and mapped three Ecdysone Response Elements within the *let-7*-C locus, proving that miRNAs can be first-response targets of the hormonal signaling. Importantly, not only do hormones regulate miRNA expression but also miRNAs can affect the strength of systemic signaling. For example, *miR-14* has been identified to mediate a positive autoregulatory loop of EcR that amplifies ecdysone response, while miRNA *bantam* activity in ecdysone-producing cells represses hormone production and thereby promotes systemic growth. A number of studies in vertebrate models and cell cultures also show relationships between hormones and miRNAs. Glucocorticoids influence a variety of physiological processes in vertebrates, including adaptation to stress, metabolism, immunity and neuronal development. Kawashima et al. show that glucocorticosteroids regulate levels of brain-derived neurotrophic factor (BDNF) via suppression of *miR-132* expression, which possibly contributes to the regulation of synaptic plasticity in the brain. On the other hand, *miRs-18* and *-124a* can regulate levels of corticosteroid receptor and therefore modulate downstream effectors of this hormonal signaling. Recent work from Huang et al. demonstrates that the *miR-21* promoter has a thyroid hormone response element that allows miRNA to be activated in response to hormonal stimuli. Thyroid hormone in vertebrates is an important regulator of development, differentiation and growth. Overactivation of *miR-21* promotes hepatoma cell migration and invasion, analogous of that observed with thyroid hormone stimulation. In breast cancer, the estrogen receptor α (ERα) binds the *miR-221/222* transcription start site and recruits co-repressors to suppress their transcriptional activity, while miRNAs *miR-191* and *miR-425* are upregulated via estrogen-mediated activation. Another study shows that *miR-221/222* acts as a negative regulator for ERα supporting the idea for the existence of negative regulatory loop involving miRNAs and hormonal receptors.

Together, these data confirm that hormones and miRNAs are prone to work together in regulation of multiple processes. On one hand, cell-specific miRNAs can be used as additional factors that fine-tune the specificity of cellular responses to global hormonal signaling; on the other hand, miRNAs are also involved in feedforward and feedback loops to readjust the precision of this systemic signaling in a given cell type.

MicroRNAs in the Brain

Biogenesis of miRNAs exhibits specific temporal and spatial profiles in different types of cells and tissues and, therefore, affects a wide range of biological functions. Conditional knockout of Dicer has been extensively used to address the collective role for miRNAs in specific tissues and cell types in mice. The essential functions for the miRNA pathway have been uncovered in the brain: miRNAs regulate neuronal development and synaptic plasticity, oligodendroglia differentiation and myelin formation and are implicated in brain tumor development and in the regulation of neurodevelopmental and neurodegenerative disorders. The role of specific miRNAs in the regulation of embryonic and adult neurogenesis, particularly in the proliferation and differentiation of neural stem cells, is emerging. Recent work from Parsons et al. provided a genome-scale profiling of miRNA differential expression patterns in human embryonic stem cell neuronal lineages. This allowed identifying molecular miRNA signatures for human embryonic neurogenesis: the in vitro neuroectoderm-originated human neuronal cells acquire their identity by downregulation of pluripotency-associated miRNAs (such as *hsa-miR-302* family). In addition, induction of high levels of expression of miRNAs required for regulation of human central nervous system development (such as *hsa-miR-10* and *let-7*) occurs in a stage-specific manner. In a similar study Stappert et al. demonstrated that time-controlled modulation of specific miRNA activities not only regulates human neural stem cell self-renewal and differentiation but also contributes to the development of defined neuronal subtypes; hence *miR-125b* and *miR-181* promote and *miR-181a* inhibits generation of dopaminergic fate neurons. Boissart et al. found that *miR-125* potentiates early neural specification of human embryonic stem cells by regulating SMAD4, a key factor for pluripotent stem cell lineage commitment. Using primary cultures derived from P1 rat cortex, neuron-enriched (*miR-376a* and *miR-434*) and glia-enriched (*miR-223, miR-146a, miR-19* and *miR-32*) miRNAs were identified. MiRNAs have been also found to direct development of specific brain regions during embryogenesis. Nowakowski et al. showed that *miR-92b* is involved in the regulation of a number
of intermediate progenitors populations in mice brain that give rise to the cerebral cortical neurons.

A number of studies in vertebrates reveal the role for miRNAs in the regulation of adult neurogenesis that is largely restricted to two major brain regions: subventricular zones of the ateral ventricle and of the dentate gyrus in the hippocampus. MiRNAs let-7b, miR-9, miR-106b-25 cluster, miR-137, miR-184, miR-124, and their specific targets were identified to regulate neural cell proliferation and/or neuronal differentiation during adulthood. Latest studies from Liu et al. uncovered the molecular mechanism by which miR-17-92 cluster regulates ischemia-induced neural progenitor cell proliferation which stimulates adult neurogenesis after injury. It has been discovered that stroke substantially upregulates miR-17-92 cluster expression in neural progenitor cells of the adult mouse. Overexpression of miR-17-92 cluster in the cell culture and in vivo significantly increased cell proliferation, whereas inhibitions of individual members of miR-17-92 cluster, miR-18a and miR-19a suppressed cell proliferation and increased cell death. Subventricular zone neuronal fate is determined by miR-124: in vivo inhibition of miR-124 causes a block in neurogenesis and leads to an accumulation of ectopic cells with astrocyte characteristics (neural stem cells) in the olfactory bulb, while upon miR-124 overexpression neural stem cells are not maintained in the subventricular zone of mouse brain and neurogenesis is lost.

Studies from Drosophila revealed that this evolutionary ancient miR-124 controls neural stem cells proliferation by targeting anachronism—an inhibitor of neuroblast proliferation. Studies from Drosophila mutant lacking miR-124 shows reduced proliferative activity of neuronal progenitor cells and decreased production of adult postmitotic neurons. We showed that ecdysteroid signaling induces expression of let-7-C in Drosophila brain, which is required for proper differentiation of the last-born MB neurons. let-7 deficiency or ecdysone signaling defect leads to MB morphological defects that result in learning and memory disabilities.

Involvement of miRNAs in regulation of neuronal development, plasticity and maintenance provides a new additional layer of gene regulation, which has an effect on nervous system functions and contributes to therapeutic approaches toward neurological diseases. These new findings also propose miRNAs as possible candidates for innovative brain therapies. However, since the general role for miRNAs is the transcriptional repression of their targets, upcoming studies should be focused on finding functional miRNA-target pairs that are also defined at the spatiotemporal level.

### BTB Transcription Factors as Temporal Codes

We established a spatiotemporal connection between the ecdysteroid-induced miRNA let-7 and its target, the BTB transcription factor Abrupt in the developing brain. BTB/POZ zinc finger factors are a class of nuclear DNA-binding proteins containing the BTB domain, which was first identified as a conserved element in the developmentally regulated Drosophila proteins Broad-complex, Tramtrack and Bric-a-brac. Afterwards, the BTB protein-protein interaction motif was found in hundreds of different proteins virtually in all organisms, ranging from yeast to humans. It is involved in the regulation of gene expression through the local control of chromatin conformation and the recruitment of degradation targets to E3 ubiquitin ligase complexes.

Interestingly, the BTB domain can form dimers and mediate interactions with non-BTB domain containing proteins and can establish both stable and transient interactions. This explains the ability of BTB containing proteins to participate in multiple processes and implies that management of their proper levels is of a particular significance.

BTB/POZ domain zinc finger factors were linked to broad range of developmental processes in vertebrates and invertebrates: chromatin remodeling, cancer development and intriguingly, regulation of cell fate specification in the nervous system. For example, the BTB/POZ zinc-finger transcription factor-encoded by gene Rp58 is required for the correct differentiation of neural progenitors into neurons, since its neural-specific deletion results in severe cerebellar hypoplasia and developmental failure of several neuronal types. By coherently repressing multiple proneurogenic genes in a timely manner this BTB protein supports neuronal differentiation and brain growth. During embryonic development of the murine cerebral cortex another mammalian BTB factor, HOF is specifically expressed in immature non-dividing cells and is downregulated in differentiated cells of the hippocampus; importantly, it is one of the factors that might be involved in early definition of hippocampal compartment within the neocortex.

Similarly, in the Drosophila nervous system several BTB/POZ domain zinc finger transcription factors have been implicated in specifying neuronal and glial cell lineages. For example, Tramtrack proteins transcriptionally repress genes that promote transformation of neuronal support cells into neurons, while Lola, Fruitless, Abrupt, and Chinmo are intrinsically required for development of different subsets of neurons. Such data provide evidence that BTB/POZ zinc-finger proteins play an important role in the transcriptional program that controls differentiation of progenitors into neurons. Since the growth and organization of the brain is tightly correlated with the speed of the whole organism development, it implies that neuron differentiation should be responsive to external temporal cues. Interestingly, the neuronal temporal identity of Drosophila MB neurons is governed by two BTB transcription factors, Chinmo and Abrupt and both of them are subjects to miRNA-mediated regulation. We found that this regulation is chronologically induced by systemic steroid signaling that controls the major larva-to-pupa transition during Drosophila development, which also coincides with the time-point when the last-born neurons are generated. This demonstrated for the first time that differential neurogenesis is hierarchically regulated by extrinsic systemic signaling, which, in chronological manner, adjusts programs of intrinsic temporal determinants of neuronal cell fate and that BTB transcription factors play a role as temporal codes in the process.

Next, we aimed to understand whether intercellular environmental signaling,
such as extrinsic cell-to-cell signaling would also cooperate to fine-tune the outcome of differential neurogenesis.

**Concerted Action of Cytokines and Steroids in Differential Neurogenesis**

Interestingly, let-7 target Abrupt that is expressed in MBs is associated with the evolutionary conserved JAK/STAT signaling pathway, which plays key roles in multiple developmental and physiological processes in the brain, ranging from the regulation of neurogenesis and stem cell fate to memory formation. In the adult brain, endogenous cytokine levels are very low under normal physiological conditions; however, various types of injuries, including trauma, seizures and ischemia induce an increase of cytokine ligand levels, which in turn promotes neuronal stem cell self-renewal. In the developing brain, some neuroepithelial cells become neuroblasts and generate the neuronal and glial cells, and in the *Drosophila* optic lobe, the timing of this transition is negatively regulated by JAK/STAT signaling. Secretion of the JAK/STAT ligand Unpaired (Upd) shapes an activity gradient in the neuroepithelium and negatively regulates the progression of the proneural wave. JAK/STAT signaling is further integrated with the Notch and EGFR signals to balance neuroblast self-renewal and neuron differentiation. Since the BTB transcription factor Abrupt has been shown previously to be negatively regulated by the JAK/STAT signaling pathway in ovaries, we evaluated whether JAK/STAT plays a role in Abrupt regulation during MB development.

We used a 10xSTAT-GFP reporter line (Fig. 2A and C) and antibodies against STAT92E, the *Drosophila* homolog of mammalian STAT (signal transducer and activator of transcription) proteins (Fig. 2D) to visualize JAK/STAT signaling activity in the developing brain. At the larval stage, JAK/STAT activity was predominantly observed in neuroblasts (Miranda positive cells, arrows) and in glia (Repo positive cells) (Fig. 2A). Mushroom body neuroblasts (MBNs) are the only neuronal stem cells that continue to divide during later stages; interestingly, in the pupal and pharate brains, apart from glial cells, GFP signal indicating JAK/STAT activity was restricted to these mitotically active neuronal stem cells (Fig. 2C). Similar pattern of JAK/STAT signaling activity was detected with STAT92E antibodies (Fig. 2D). This expression analysis shows that the JAK/STAT signaling pathway is active in all postembryonic neuronal stem cells regardless of the developmental stage or ecdyson signaling activity.

Previously, we found that Abrupt is expressed in early-born γ, α/β’ neurons and miRNA let-7 in the late-born α/β neurons and this temporally induced let-7 expression is necessary to downregulate Ab, which is critical for proper specification of the last-born neurons. Abrupt is a very potent cell fate regulator, since its misexpression is sufficient to even induce homeotic transformation. Therefore, we hypothesized the possibility that spatially distributed cytokine signaling would repress Abrupt expression in the MB neuronal stem cells (Fig. 2B).

To test this we analyzed different JAK/STAT pathway mutants (see Materials and Methods) and found that downregulation of JAK/STAT signaling via expression of dominant negative form of dome specifically in the neuroblasts resulted in changed Abrupt expression pattern in the MB cell body clusters and in the appearance of ectopic Abrupt protein in some of the neuroblasts (Fig. 2E and F). Next, we wanted to test if this misexpression would affect the neuronal stem cell progeny differentiation. MB neuroblasts are continuously dividing to give rise to MB neurons (Kenyon cells) that based on their birthdate and cell adhesion molecule expression, are clustered into three types of MB lobes (γ, α/β’ and α/β) with distinct axonal projection patterns. We used FasII antibodies as a molecular marker for γ and α/β MB axons to evaluate whether downregulation of JAK/STAT signaling or overactivation of the transcription factor Abrupt in the MBNs affect overall MB morphology. We observed that downregulation of JAK/STAT activity via overexpression of a dominant negative form of dome or STAT RNAi using pan-neuronal and neuroblast-specific driver lines (insGal4 and wortGal4, respectively) indeed caused morphological changes in the adult mushroom bodies; MBs with slim α/β lobes and fused β-lobes (Fig. 2G–H and 2J–K; Table 1) were observed. Importantly, similar MB morphological defects were identified upon overexpression of Abrupt in the MB neuroblasts (Fig. 2I–J and 2L; Table 1). This evidence supports the hypothesis that spatially distributed JAK/STAT signaling represses the transcription factor Abrupt in neuronal stem cells and this downregulation is critical for proper neurogenesis.

Since previously we found that ecdyson signaling also targets this BTB transcription factor via let-7 miRNA, we conclude that two extrinsic signaling pathways, global hormonal and local cytokine, collaborate to regulate extended neurogenesis during *Drosophila* MB development.

Interestingly, another BTB-zinc finger protein Chinmo that has been found to control stem cell self-renewal and direct neuroblast temporal identity also depends on JAK/STAT activity and can be targeted by miRNA let-7 and miR-125. This implies that regulation of expression of JAK/STAT dependent BTB factors Abrupt and Chinmo should be under strict developmental control to guarantee faithful cell fate determination. Our current and previous data provide evidence that in the developing brain, the temporally induced by ecdyson miRNA let-7 negatively regulates Ab, which is additionally targeted by the local JAK/STAT cytokine signaling pathway to ensure proper MB development. The interaction between global developmental and local tissue-specific signaling results in formation of a robust spatio-temporal pattern to fine-tune the fidelity of neuronal cell differentiation, which is essential for proper brain morphogenesis (Fig. 3).

**Cell Adhesion as a Final Outcome of Differential Neurogenesis**

The complexity of the brain is generated by multiple types of neurons that connect to each other in a specialized manner, which often depends on selective cell adhesion. Neurons expressing similar cell adhesion proteins not only cluster together to organize brain compartments that have distinct functions, selective cell adhesion is also used for establishment of
Figure 2. For figure legend, see page 179.
synaptic connections that allow neurons to communicate and transfer information. Significant alterations in the brain structure and functions are generated even by moderate changes in the quantities of adhesion molecules on the neuronal cell surfaces. Therefore, differential cell adhesion is the final aftermath of differential neurogenesis, suggesting that timing and levels of cell adhesion protein expression must be precisely regulated (Fig. 3).

Among the most important cell adhesion molecules (CAMs) involved in the development of the nervous system, synaptic plasticity and cognition and memory are neural cell adhesion molecules (NCAMs) that belong to the immunoglobulin superfAMILY. Previous data show that levels of human NCAM2 that is primarily expressed in the brain to stimulate neurite outgrowth and facilitate dendritic and axonal compartmentalization are essential for normal brain development. For example, the increased expression of NCAM2 as a result of trisomy 21 may cause dosage-related detrimental effects in Down syndrome; also, in genomewide association studies, NCAM2 was suggested as a candidate gene for the development of autism and Alzheimer’s disease, and multiple NCAM1 proteins are differentially altered in bipolar disorder and schizophrenia. Furthermore, NCAMs play a critical role in plasticity of the nervous system and in mechanisms controlling learning and memory and their expression levels are known to be highly susceptible to modulation by stress. Moreover, NCAM is involved in some of the bidirectional effects of stress on memory processes, where its increased expression is facilitating stress actions while its decreased expression is impairing effects of stress on memory consolidation. All these data imply that regulation of NCAM expression is a prerequisite for proper brain development and function. However, the question remains: What genetic machinery regulates precise expression of adhesion molecules in the brain?

Ample sets of regulatory elements are required for spatiotemporally restricted expression pattern of a given gene; however, it is not well-defined which set of transcriptional factors regulates differential expression of appropriate cell adhesion proteins that modulate the degrees to which various neurons adhere to each other to make synapses. In the Drosophila MBs, the ortholog of NCAMs, Fasciclin 2 (Fas2) displays specific temporal patterns of expression that plays a significant role in the spatial segregation of MB neurons. Low levels of Fas2 are detected in the γ
MB neurons makes their axons to project into the places, where the later-born α'/β' neurons would send their axons. Since distinct MB neurons have different functions in Drosophila behavior regulation, it would be interesting to analyze whether this alteration in the cell adhesive characteristic would change fly cognition. Importantly, we also show that miRNAs are mediators between extrinsic temporal cues and intrinsic spatiotemporal codes that determine the precision of neuronal adhesiveness during brain development. It would be important in the future to explore the role of these factors in the adult brain plasticity. Interestingly, it has been proposed that the increased stickiness of human neurons might explain the accelerated evolution of the human brain beyond the brains of primates. Another differential cell adhesion as a fundamental mechanism of neuronal cell differentiation that controls the finest aspects of neuronal specification (Fig. 3). Once a specific neuron is born, it must recognize and join other neurons of the correct type to assemble into a specific brain compartment that normally is determined and maintained by the system of preferential cell affinities. Even more, neurons send out axons and dendrites that via differential cell adhesion make synapses with other neurons. However, neurons do not simply reside inertly stuck together; instead, the new synapses are established and actively maintained by selective adhesion created and gradually adjusted by neurons; thus, contributing to the nervous system plasticity. We found that misexpression of Fas2 in the early-born α'/β' lobes. Fas2 provides specific adhesive codes among MB neurons preventing them from intermingling and assuring formation of distinct MB lobes. We showed that the transcription factor Abrupt suppresses Fas2 expression in the earlier-born neurons, while steroid-induced miRNA let-7 via downregulation of Abrupt allows this critical adhesion molecule to be highly expressed in the late-born α/β neurons. Thus, the precise Fas2 expression is essential for proper MB morphology and function (Fig. 1). Together, these data show that NCAMs are multifunctional proteins involved in neurogenesis and neurodevelopment and their expression levels are critical for dendritic and axonal compartmentalization and synaptic plasticity. This makes MB neurons makes their axons to project into the places, where the later-born α/β neurons would send their axons. Since distinct MB neurons have different functions in Drosophila behavior regulation, it would be interesting to analyze whether this alteration in the cell adhesive characteristic would change fly cognition. Importantly, we also show that miRNAs are mediators between extrinsic temporal cues and intrinsic spatiotemporal codes that determine the precision of neuronal adhesiveness during brain development. It would be important in the future to explore the role of these factors in the adult brain plasticity. Interestingly, it has been proposed that the increased stickiness of human neurons might explain the accelerated evolution of the human brain beyond the brains of primates. Another
factor that distinguishes humans from other primates is that developmental profiles of miRNAs, as well as their target genes, show the fastest rates of human-specific evolutionary change, which allows for the faster evolutionary rate in divergence of developmental patterns. One of the key features of the miRNA function is that miRNAs normally do not turn their target genes on and off, but just modulate their expression. This allows building novel networks between newly originated genes and miRNAs softly, not necessarily causing the lethality. Analysis of recently originated brain genes in Drosophila showed that numerous newly evolved genes are expressed in the brain and all of the MB-positive new genes are expressed in the α/β, but not in more ancestral γ and α/β’ lobes. Since miRNAs and newly evolved genes are frequently co-expressed in the brain, the hypothesis can be put forward that the establishment of novel sets of spatiotemporal codes for differential neurogenesis that are generally fine-tuned by miRNAs is a common mechanism that might contribute to the phenotypic evolution of behavior and individual plasticity of the nervous system. Management of genetic programs that temporally specify individual subtypes of neurons could help to evaluate the true limits of progenitor plasticity within the developing and adult brain and initiate a new phase of plasticity assessment.

Materials and Methods

Fly strains and genetics

We used wortGal4 and insGal4 (BDSC) driver lines crossed to a dominant negative form of Dome (UAS-domeDδ or UAS-domeDN) and STAT RNAi transgenic line (UAS-STAT2ERα, VDRC) to downregulate JAK/STAT signaling; and UAS-Abrupt (BDSC) to overexpress Abrupt in the neuroblasts. Oregon R animals were used as a wild-type control. To visualize active JAK/STAT signaling 10xSTAT-GFP reporter was used. All crosses were maintained at 25°C on standard medium.

Immunohistochemistry

Brains were dissected in PBS and fixed in 4% formaldehyde (Polysciences, Inc.), adult and pupal for 30 min, larvae for 15 min. Staining was performed as described. The following antibodies were used: mouse anti-Fas II 1:20 (marker for γ and α/β lobes) and mouse anti-Repo 1:20 (glia marker) (DSHB), rabbit anti-Abrupt 1:500,9 rabbit anti-STAT2ERα 1:500,8 guinea pig anti-Miranda (gift from A Wodarz), Alexa 488, 568, or 633 goat anti-mouse, anti-rabbit, anti-guinea pig (1:500, Molecular Probes). Images were obtained with a confocal laser-scanning microscope Zeiss LSM700 and processed with ZEN 2010 and Adobe Photoshop software.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

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