Special Focus: Synapse-Glia Interactions

More than just glue

The diverse roles of cell adhesion molecules in the Drosophila nervous system

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Key words: drosophila, cell adhesion, nervous system, glia, axon, synapse

Cell adhesion is the fundamental driving force that establishes complex cellular architectures, with the nervous system offering a striking, sophisticated case study. Developing neurons adhere to neighboring neurons, their synaptic partners, and to glial cells. These adhesive interactions are required in a diverse array of contexts, including cell migration, axon guidance and targeting, as well as synapse formation and physiology. Forward and reverse genetic screens in the fruit fly Drosophila have uncovered several adhesion molecules that are required for neural development, and detailed cell biological analyses are beginning to unravel how these factors shape nervous system connectivity. Here we review our current understanding of the most prominent of these adhesion factors and their modes of action.

Introduction

Cell adhesion is the main force that sorts cells into distinct functional groups, a prerequisite for the establishment and maintenance of tissues and organs. In this context, adhesion plays many roles, affecting the survival and proliferation of cells through interactions with the substrate, as well as controlling the morphogenesis and assembly of the more complex cellular arrangements seen in all organs. Intriguingly, relative to our knowledge of the plethora of factors and their modes of action.

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in the Drosophila oocyte and the eye.6,7 These latter studies also demonstrated that by restricting N-cadherin expression to a specific cell type, differential adhesive interactions can occur amongst subsets of cells within a larger structure. This specificity allows cells expressing the same cadherins to arrange themselves by minimizing their mutual surface tensions in a highly localized way, within the in vivo environment.7 Similar interactions may play important roles in shaping the fine structure of the nervous system and we anticipate that as adhesion factors are identified and characterized in more detail, many of their actions will be attributable to differential adhesion. The challenge that we are only beginning to confront is how all of these adhesive cues might be integrated by developing neurites into “decisions” about where to project.

**Adhesive Interactions Amongst Neurons in the Developing Nervous System**

Many genetic screens for mutations affecting the establishment of neuronal connectivity have identified cell adhesion molecules of many different molecular families. Foremost among these are members of the cadherin superfamily, Immunoglobulin (Ig) domain containing proteins and members of the leucine-rich repeat (LRR) family. We focus on three contexts in which these molecules have been studied extensively, examining the formation of specific connections in the adult visual system and the neuromuscular junction, and discussing the use of cell adhesion in neuronal self-recognition. We further narrow our discussion to molecules that have been shown to directly mediate adhesion, without excluding the possibility that they might act as signaling molecules as well. However, at present, the signaling pathways that might be regulated by these molecules have not been studied in flies. Finally, it is important to recognize that drawing a distinction between adhesion and signaling molecules becomes somewhat arbitrary as one begins to consider that cell signaling pathways can regulate cell adhesion and vice versa.

The structure and development of the adult visual system. The visual system has provided a facile system to examine the molecular mechanisms by which developing axons choose appropriate synaptic partners. These choices are informed by adhesive interactions between axons and their targets, and surprisingly, also amongst afferent axons themselves. Here we briefly review the anatomy of the system, and then describe these interactions, and their molecular mechanisms.

The Drosophila visual system comprises the retina and four optic ganglia, the lamina, medulla, lobula and lobula plate (Fig. 1A). The retina consists of approximately 800 subunits, ommatidia, each containing eight distinct photoreceptor neurons, designated R1–R8. Photoreceptor axons from each ommatidium form a fascicle that innervates the optic lobe in a retinotopic pattern. R1–R6 cells project to the most peripheral optic neuropil, the lamina, while the R7 and R8 cells send their projections through the lamina into the medulla (Fig. 1A). During development, R1–R6 cells initially stop between two layers of glial cells in the lamina. Subsequently, each R cell growth cone defasciculates, and extends away from the ommatidial bundle along a unique trajectory to join a neighboring column of post-synaptic targets (Fig. 1A). R1–R6 axons, together with their post-synaptic targets, then form a new fascicle, called a cartridge, and initiate synapse formation. By contrast, R7 and R8 target within distinct layers in the medulla. These targeting choices emerge in two steps: initially both R7 and R8 axons target to relatively shallow, temporary layers.9 Then, after additional medulla layers assemble, both R7 and R8 axons extend deeper to their final target layers where they form stable connections with higher order neurons.8 Thus, the adult visual system captures at least two distinct types of synaptic specificity: cartridge formation in the lamina, and layer-specific targeting in the medulla.

Adhesive interactions in the developing visual system. The molecular mechanisms that underlie the formation of these connections are incompletely understood. However, three different cell adhesion molecules, the classical cadherin N-cadherin, the leucinerich repeat adhesion molecule Capricious, and the protocadherin Flamingo play important roles.10-12 N-cadherin is required both for cartridge formation and layer-specific targeting while Capricious regulates layer-specific targeting, and Flamingo cartridge formation. Moreover, N-cadherin and Capricious mediate interactions between R cell axons and their targets, whereas Flamingo mediates interactions only amongst R cell axons. Here we consider the functions of each one in turn.

N-cadherin is broadly expressed in the nervous system, and homozygous mutant animals display pleiotropic defects in axon fasciculation and guidance.13 On the other hand, mosaic studies demonstrated specific roles for N-cadherin in particular aspects of R cell axon targeting.10,14 (Fig. 1B). These studies revealed that N-cadherin was required for R1–R6 axons to reach their appropriate lamina cartridges, as well as for R7 axons (but not R8 axons) to reach the correct layer of the medulla. Subsequent analyses revealed that N-cadherin is also required in lamina and medulla neurons for R cell axon targeting.15,16 Finally, these studies demonstrated that N-cadherin mediates homophilic, attractive interactions between R cell axons and their targets, providing the first functional evidence for such an N-cadherin mediated interaction in any context.15

N-cadherin activity is subject to sophisticated, dynamic and cell-type specific regulation of its expression level and sub-cellular localization. In the context of R1–R6 axon targeting in the lamina, N-cadherin adhesion appears to be asymmetrically regulated within the growth cone to allow for defasciculation and lateral movement towards the correct targets.17 This lateral extension is dependent upon two additional proteins, the receptor tyrosine phosphatase LAR, and the synaptic scaffolding molecule Liprin-α, both of which are required in R cells for target selection, raising the possibility that they could interact with N-cadherin.17 Additional complexity has been revealed through studies of N-cadherin in other visual neurons. In particular, N-cadherin functions in lamina neurons, controlling the targeting of their axons to specific layers in the medulla.18 Individual lamina neurons have complex and varied N-cadherin requirements, with some requiring N-cadherin cell-autonomously while others necessitate N-cadherin function in their neighbors. At least part of this complexity is achieved by dynamic regulation of N-cadherin localization to specific neuronal processes. These data support a model where cell- and stage-specific modulation of N-cadherin controls discrete targeting steps. Finally, it is also likely that N-cadherin insertion into the plasma membrane is regulated in developing R cell axons, perhaps through the actions of the exocyst complex.19 These studies underscore that molecules such as N-cadherin do not simply act as “on and off” cellular adhesives, but rather that they are under complex regulation at multiple levels.
The roles of cell adhesion in the nervous system

The roles of cell adhesion in the nervous system are both necessary and sufficient to specify targeting to a single layer, implying that other layer-specific cues remain to be discovered. Flamingo encodes an unusual, evolutionarily conserved member of the cadherin superfamily, containing both cadherin repeats and a seven trans-membrane domain. Initial studies of Flamingo focused on its roles in epithelial planar cell polarity and dendrite patterning in the peripheral nervous system of the embryo and larva.20-23 Flamingo also plays at least three distinct roles in photoreceptor target selection. First, Flamingo regulates the spacing of R cell axon bundles, establishing the local topography of cartridges and columns in the lamina and the medulla, respectively.12,24,25 Second, Flamingo appears to be required for layer-specific targeting. In flamingo mutants, R8 axons (but not R7 cells) fail to reach their target layer and instead terminate in more superficial layers of the medulla.24 Finally, Flamingo mediates homophilic interactions among R1–R6.

Future studies will undoubtedly uncover both additional regulatory mechanisms as well as provide molecular details regarding the known mechanisms.

The adhesive leucine-rich repeat protein Capricious plays an instructive role in R8 cell layer-specific targeting in the medulla.11 In particular, capricious loss-of-function mutations cause targeting errors in R8 axons, with many extending laterally and others terminating in inappropriate medulla layers (Fig. 1C). In contrast, the layer-specific targeting of R7 axons is unaffected by capricious mutations. Consistent with a function restricted to R8, Capricious protein is expressed in R8 axons, and their target layer, but not in R7 cells, or their recipient layer. Remarkably, Capricious is sufficient to specify axonal targeting, as ectopic expression of Capricious in R7 cells redirects them to the Capricious-positive R8 layer. This implies that by mediating afferent-target interactions, Capricious provides a cue that is both necessary and sufficient to specify targeting to a single layer, implying that other layer-specific cues remain to be discovered.

Flamingo encodes an unusual, evolutionarily conserved member of the cadherin superfamily, containing both cadherin repeats and a seven trans-membrane domain. Initial studies of Flamingo focused on its roles in epithelial planar cell polarity and dendrite patterning in the peripheral nervous system of the embryo and larva.20-23 Flamingo also plays at least three distinct roles in photoreceptor target selection. First, Flamingo regulates the spacing of R cell axon bundles, establishing the local topography of cartridges and columns in the lamina and the medulla, respectively. Second, Flamingo appears to be required for layer-specific targeting. In flamingo mutants, R8 axons (but not R7 cells) fail to reach their target layer and instead terminate in more superficial layers of the medulla. Finally, Flamingo mediates homophilic interactions among R1–R6.
axons that guide these axons to their appropriate target cartridges. Detailed mosaic analyses have provided new insight into Flamingo’s mechanism of action. In particular, Flamingo acts exclusively in a cell non-autonomous fashion, mediating afferent-afferent interactions that take place prior to growth cone extension. These studies demonstrated that while Flamingo function in any single growth cone is dispensable for normal target selection, individual growth cones are sensitive to differences in Flamingo activity between themselves and their neighbors. In effect, individual growth cones balance Flamingo-mediated interactions with their immediate neighbors, creating a form of “opponency” where the strength of the interaction with one neighbor is opposed by an interaction with another neighbor. This simple model provides a mechanism by which differences in adhesivity between cells, perhaps reflecting differences in cell fate, can be translated into changes in growth cone orientation necessary for appropriate target selection. While Flamingo’s molecular structure, containing a seven pass trans-membrane domain similar to those of G-protein coupled receptors, suggests that it functions as more than simply an adhesive factor, there is no evidence in any experimental context that it acts as a signaling receptor. In summary, the cellular opponency model that emerged from studies of Flamingo in Drosophila provides a powerful framework for considering homophilic, adhesive interactions more broadly.

**The structure and development of the neuromuscular junction.** Adhesive interactions also play critical roles in patterning the connections between motor neurons and their muscle targets in the developing embryo and larva. The abdominal body wall of embryos and larvae is segmented, with each segment containing more than 30 bilaterally symmetrical muscle fibers, targeted in a precise pattern by approximately 40 identified motor neurons (Fig. 1E). Each fiber is unique, and can be defined by morphological criteria. Synapses between motor neurons and muscles, the neuromuscular junctions (NMJ), form during embryogenesis and reflect a developmental sequence of axon guidance decisions made at a series of so-called “choice points.” These decisions culminate in the arrival of each motor neuron at a specific muscle target, and the formation of a NMJ of unique morphology. Thus, studies of cell adhesion in this context have focused both on the roles of adhesive interactions in shaping axon guidance decisions, and regulating the strength and patterning of the NMJ.

**Adhesive interactions at the neuromuscular junction.** As in the visual system, our understanding of the molecular factors that shape the NMJ are only partly understood. Here we focus on just two adhesion molecules where both gain and loss-of-function studies demonstrate critical developmental roles, namely Flamingo and Capricious.

Recent work has linked Flamingo to target selection and synaptogenesis at the NMJ. Homozygous mutant animals display increased numbers of ectopic synapses, normal “en passant” synapses distal to the axon terminal, as well as pre-synaptic varicosities within normally asynaptic axonal segments (Fig. 1F). These observations suggest that Flamingo suppresses synapse formation in this context. Intriguingly, this study also described a function for Flamingo in the maintenance of axons and synapses as homozygous mutant animals display age-dependent axon degeneration and loss of muscle innervation. Thus, Flamingo acts differently at the NMJ in comparison to the visual system, specifically affecting synapse formation in one case, while affecting target selection in the other.

Capricious acts as a molecular matchmaker between pre- and post-synaptic cells at the neuromuscular junction, a function similar to its role in the visual system. In the embryo, Capricious is expressed in a small number of muscles and neurons. In one well-studied case, Capricious is expressed in both a single muscle and the motor neurons that innervate it. In capricious mutants, motor neurons form ectopic synaptic contacts with muscles that neighbor the appropriate target, arguing that Capricious functions to restrict neuronal innervation to a single target (Fig. 1G). Conversely, expression of Capricious in all muscles causes the formation of many ectopic synapses. Intriguingly, the intracellular domain of Capricious is required in the muscle but not the innervating neuron, suggesting an asymmetric signaling role. Taken together, these studies argue that Capricious mediates specific adhesive interactions between a particular muscle and its neuronal targets, and that this interaction promotes synapse formation with the target while inhibiting synapse formation on neighboring cells.

**DSCAM: A new paradigm for self-recognition in the nervous system.** Many neurons display complex axonal and dendritic architectures whose organization reflects, in part, interactions that discriminate between processes elaborated by the same cell, versus those elaborated by neighbors. The Ig superfamily member Down Syndrome cell adhesion molecule (DSCAM) plays a central role in these interactions. Mutations in DSCAM cause defects in the development of a variety of neurons, affecting the outgrowth patterns of simple branched axons in the mushroom body, the elaboration of complex dendritic fields in the larva, and the connectivity of the olfactory system, among others. Additional analysis has provided evidence for a novel, DSCAM-mediated mechanism that supports neuronal self-recognition. DSCAM displays astonishing molecular diversity, having more than 38,000 alternatively spliced isoforms that differ in their extracellular and transmembrane domains. Individual neurons express multiple isoforms, with single cells of the same type expressing distinct, but overlapping subsets of the genomic repertoire. Homophilic adhesive interactions are isoform-specific, with negligible binding of different splice variants to each other. Finally, diversity per se, rather than expression of particular isoforms in specific cells, is central to DSCAM function. Taken together, these data support a model in which neuronal processes can distinguish self from non-self because only neurites from the same cell will contain exactly the same complement of DSCAM isoforms. Through unknown signaling pathways, these interactions then alter neurite behavior to prevent processes from the same cell from crossing one another.

**Glial-Neuron Interactions in Development**

The preceding studies have focused on adhesive interactions between neurons and their targets. However, this restricted view ignores an array of significant interactions between neurons and the non-neuronal constituents of the nervous system, namely glia. Glial cells form elaborate shapes in order to ensheath neuronal cell bodies and axons, and to become intimately associated with synapses (Fig. 2A). Although our knowledge about the functions of glia is still fragmentary, neuron-glial interactions are required during glial migration, axon guidance and targeting, as well as synapse formation and function. For many of these processes, contact-mediated, potentially adhesive interactions are particularly important and it is apparent
that we have only just begun to identify the critical molecular players. Here we focus on instances where neuron-glia interactions influence migration, axon targeting, synapse formation and physiology.

The roles of neuron-glial interactions in cell migration and axon extension. Neuron-glial interactions influence both glial migration and axon extension during visual system development (Fig. 2B). During late larval stages, R cells differentiate in a wave that spreads from posterior to anterior. As they differentiate, R cell axons extend basally and exit the eye disc at the posterior, through the optic stalk. Concomitantly, retinal basal glia migrate through the optic stalk, into the eye disc, moving in the opposite direction, while staying in close contact with R cell axons.44,45 This migration is induced by a retinal signal: in mutants that lack R cells, glia do not enter the eye disc and instead accumulate in the optic stalk.46 Surprisingly, when R cells are present, but fail to extend axons, glial migration is unaffected. Thus, R cell axons are not a required substrate for glial migration. However, retinal basal glia will migrate towards large ectopic patches of differentiating R cells, suggesting that R cells provide an attractant that is sufficient for migration, but not necessary.45 Intriguingly, although glia do not need R cell axons to direct their migration, R cell axons do require glial cells as guideposts. When glial migration into eye disc is inhibited, R cell axons frequently stall and fail to exit the eye disc into the optic stalk.45,47 Similarly, in gilgamesh mutants, glia migrate too far into the eye disc, causing R cell axons to extend anteriorly rather than posteriorly.47 A related guidepost role can be found in the embryonic nervous system. There, the formation of the main axon tracts depends on longitudinal glia: when these glia are ablated, the axons of pioneer neurons, which form the first longitudinal pathways, frequently stall or extend inappropriately.43,48,49

Glia can also shape neuronal architecture by serving as intermediate targets. In the visual system, glial cells are the first to colonize the lamina. By arriving before R cell axons reach the brain, and before lamina neuron differentiation occurs, glial cells play a critical role in directing R cell axon targeting. Glial cells establish two distinct layers, designated the epithelial layer and the marginal layer, and provide a short-range signal that directs R cell axons to stop between them (Fig. 2B). When glia fail to migrate into the lamina, R1–R6 axons fail to stop, and instead project into the medulla.50,51 However, while this stop signal is very likely to be a local, possibly adhesive signal, its molecular identity is unknown.

Neuroglian mediates neuron-glial interactions. While many neuron-glial interactions are local, the molecular identities of the factors involved are largely unknown. The only known adhesion molecule that appears to mediate homophilic interactions between neurons and glia during axon guidance is Neuroglian. Neuroglian mutant animals display defects in both neuronal and glial morphogenesis: sensory axons form ectopic branches and glia fail to ensheath them.52 This ectopic branching phenotype could only be rescued when Neuroglian was expressed in both neurons and glia, suggesting that Neuroglian acts as a homophilic adhesion molecule. These axon branching defects were not caused by defects in glial ensheathment, suggesting that Neuroglian plays distinct roles in neurons and glia.
Intriguingly, Neuroglian is differentially spliced, with one form being restricted to neurons, while another form is expressed by neurons, glia and epithelial cells. These splice variants have identical extracellular but different intracellular domains, suggesting that Neuroglian-mediated interactions could activate distinct signaling pathways in neurons and glia. Finally, the role of Neuroglian as a mediator between axons and glia appears to be conserved, as its vertebrate homolog, Neurofascin 155, mediates neuron-glia contacts in myelinated axons.

Gliaal functions in synapse development. The best understood requirement for neuron-glial interactions at the developing synapse occurs during axon pruning, a synaptic remodeling process that takes place during metamorphosis. In one prominent brain structure, the mushroom body, γ neurons initially project two branches, one medially and one dorsally (Fig. 2C). Both branches degenerate during early metamorphosis; at later stages only the medial branch re-extends. This axonal degeneration is triggered by the steroid hormone Ecdysone and relies on interactions between γ neurons and glia. Indeed, pruning is implemented by glial processes that invade the mushroom body neuropil to phagocytose degenerating axonal debris. These events are mediated by the scavenger receptor Draper. Blocking glial phagocytosis either by ectopic expression of a dominant-negative Dynamin, or by removal of Draper, results in failure of axon degeneration and persistence of ectopic axon branches. Finally, while Ecdysone signaling within γ neurons triggers both cytoskeletal degeneration in the axon, and induces glial invagination, it is also required in the glia themselves to upregulate Draper expression. Thus, one signal activates complementary pathways in neurons and glia, which then act in synchrony to prune axons.

Gliaal activities at mature synapses. Anatomical studies of the NMJ and the visual system demonstrate that glial processes are in close proximity to synapses, forming highly specialized contacts that must surely be regulated by adhesion. In addition, glial cells regulate synaptic function by establishing and maintaining extracellular homeostasis. For example, glia form the blood-brain barrier to protect neurons from the high potassium and glutamate concentration of the hemolymph.

Glia also express ion channels required for neurotransmitter recycling, such as the cysteine-glutamate exchanger Gendrblind and the glutamate transporter dEAAT1, both of which remove excess glutamate from the synapse.

Moreover, glia modify biogenic amines using the β-aminergic amine synthase Ebony. This function is best understood at R cell synapses, where Ebony conjugates the neurotransmitter histamine to β-alanine, generating carcinine. Carcinine is thought to be transported back into R cell terminals, where another enzyme, Tan, hydrolyses it back to histamine. This observation suggests a tightly coupled neuron-glial biosynthetic cycle. At this synapse, the close association of neurons and glia is morphologically apparent, as glial processes, called capitate projections, invaginate into the pre-synaptic terminal. These structures are thought to be highly dynamic and show differential localization of Dynamin and Endophilin, suggesting that they are the sites of vesicle endocytosis and recycling.

The Ig-domain containing molecule Basigin regulates capitate projection formation and basigin mutants display reduced numbers of capitate projections and defective synaptic transmission. However, the causal relationship between defects in capitate projections and synaptic transmission are unclear. Finally, although Basigin is expressed both by R cells and glia, it binds the Laminin receptor Integrin. Thus while morphological and functional data demonstrate intimate neuron-glial interactions, the molecules that actually hold these cells together remain largely unknown.

Concluding Remarks These studies highlight three fundamental themes. First, it is striking how diverse the functions of adhesion molecules truly are; in no sense can the actions of these proteins be attributed to something akin to a simple “glue” acting to hold cell surfaces together. While it is unclear whether this functional diversity reflects differences between the molecules themselves, or between the cellular contexts in which they act, understanding how it arises presents a critical challenge. Second, it is apparent that both the timing and precise levels of expression of each adhesion molecule are critical for their function: manipulations that moderately alter expression often cause dramatic phenotypes. Thus we need to understand the regulatory mechanisms that control the activities of adhesion molecules, and have better models that define how cells read and interpret differences in adhesivity. Finally, while we have made some progress in identifying and characterizing the adhesive molecules that play critical roles during development, our understanding of adhesive interactions amongst neurons, and between neurons and glia in the adult nervous system remains incomplete. A systematic study of the mechanisms responsible for the long-term maintenance of brain architecture will likely lead us to new insights into cell adhesion.

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www.landesbioscience.com Cell Adhesion & Migration 41
The roles of cell adhesion in the nervous system

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