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**Spatially patterned gradients of synaptic connectivity are established early in the developing retina**

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How do appropriate synaptic patterns of connectivity emerge during development? What is the relationship between a neuron’s dendritic architecture and its synaptic connectivity? These questions have puzzled developmental neurobiologists for many years, but only recently, with the advent of novel neuronal labeling and imaging techniques, have answers to these questions been obtained in the living organism. In a recent article in *Neural Development*, Morgan et al. [1] investigate the developmental mechanisms by which excitatory synaptic inputs become distributed across retinal neurons so that reliable visual information can be transferred to the brain. This study demonstrates that synaptic inputs to retinal projection neurons are spatially patterned from early stages of dendritic development and that the density of inputs is maintained as constant, even as the retinal circuits remodel and mature.

The organized laminated structure of the vertebrate retina (Figure 1a) provides an excellent model in which to study how synaptic circuits are established during development and to what extent intrinsic versus extrinsic signals contribute to this process. Synaptic circuits in the retina transform visual information that is collected by photoreceptors into electrical and chemical signals, which are then transferred to retinal ganglion cells (RGCs), the output neurons of the retina. RGCs relay visual information to the brain through their long projecting axons. Those RGCs, whose cell bodies reside in the ganglion cell layer, receive synaptic input onto their dendrites in the inner plexiform layer in the form of excitatory and inhibitory synapses from bipolar cells and amacrine cells. The inhibitory synapses signal using $\gamma$-aminobutyric acid (GABA), whereas the excitatory synapses signal using glutamate. The dendrites of RGCs are remodeled extensively during development: initially as the first inhibitory GABAergic synapses between amacrine cells and RGCs are formed, and then in response to the first excitatory glutamatergic inputs from bipolar cells to RGCs [2,3]. This dynamic remodeling of dendritic arbors [4] works together with molecular cues [5] to organize inputs into different sublaminae in the inner plexiform layer in response to visual signals. In this way, the distinct types of RGCs attain their characteristic dendritic lamination, architecture and synaptic connectivity.

Evidence of dynamic mechanisms of synaptogenesis and their relation to dendritic arbor structure has been obtained in recent studies that have expressed fluorescently tagged postsynaptic components in individual neurons in live fish,
dendrites that stratify or branch in both sublaminae and receive inputs from both ON and OFF visual pathways.

Confirming previous observations that chimeric PSD-95 expression localizes to ultrastructurally identified synapses in vivo [7], the authors [1] found that expression of PSD95-YFP on RGC dendrites localized to sites where bipolar cell synapses form. Synaptic sites were found to be evenly distributed along the RGC dendritic arbor (which was visualized by the expression of the red fluorescent protein td-Tomato) before synaptic glutamate responses could be recorded, and the patterned distribution of synaptic sites on individual branches was maintained as the dendritic arbor stratified and remodeled. A centro-peripheral gradient of synaptic number, with more synapses closer to the cell body, emerges early, when bipolar cells are forming synapses onto RGCs, and this gradient is maintained despite ongoing remodeling and synaptogenesis. It is likely that centro-peripheral synapse gradients are established by competition between bipolar inputs at the borders of the dendritic arbor, where the border of an RGC receptive field locates. Live imaging of synaptic sites on distal branches [3] could now be used to demonstrate the existence of a dynamic competitive process at receptive field borders.

Using a tool that measures the lamination index of the dendritic branches and of synaptic sites, Morgan et al. [1] also revealed that the dendritic arbors of RGCs begin to stratify both their branches and their synapses at the time when synaptic glutamate neurotransmission begins (postnatal days 7-12 in the mouse). During this period, when the dendritic arbor actively enlarges and also refines by pruning back dendrites, the number of synaptic contacts per unit area of the bipolar cell surface is maintained by the increase in the density of synapses on those branches that are retained. For visual clarity, synaptic sites are illustrated only in a portion of the dendritic arbor.

frog and mouse embryos [6-8]. The postsynaptic density protein PSD-95 is a scaffolding protein that participates in synapse maturation and has served as a marker for glutamatergic postsynaptic sites in vivo [9]. In their new study, Morgan and colleagues [1] show a new correlation of the emergence of glutamatergic synaptic inputs on RGCs with their dendritic arbor structure by analyzing RGCs expressing PSD-95 tagged with yellow fluorescent protein (YFP) at key stages in retinal circuit development. They developed a set of elegant measuring tools to examine the density and distribution of putative glutamatergic synaptic sites on RGC dendrites (bipolar cell inputs) in explanted mouse retinas, from postnatal day 5, before functional glutamatergic responses are recorded, until the first postnatal month, when functional circuits are mature. They focused on monostitified and ON and OFF bistratified RGCs to determine whether the distinct spatial patterns of bipolar cell inputs are established at the onset of synaptogenesis or whether they emerge through a remodeling process. Monostitified RGCs (ON-center or OFF-center) position their dendrites in either one of two sublaminae to receive functional input from ON or OFF visual pathways. Bistratified RGCs have
Morgan et al. [1] also suggests that branches that are eliminated or pruned back during arbor refinement bear synaptic contacts, and these branches can be those that do make contact with presynaptic bipolar cells. This notion is in agreement with observations that have been made in studies that have followed pre- and postsynaptic components in individual branches by time-lapse imaging of axons and dendrites in vivo [6,11-13]. These studies indicate that RGCs, like other central neurons, transition from an exploratory state to a mature state by removing transient contacts made by excess dendritic branches. Dynamic mechanisms of synaptic and dendritic arbor remodeling seem to be similar for RGCs that maintain their stratification order (dendrites of ON and OFF bistratified RGCs) and those that transition from an immature bistratified to a monostratified dendritic arbor. Time-lapse imaging studies could now be used to reveal the sequence of events by which spatially patterned gradients of synaptic connectivity are established and refined in the retinal circuit, and to ascertain the contribution of early inhibitory and late excitatory inputs to this dynamic process.

In summary, the elegant study by Morgan and colleagues [1] provides new insights into the mechanisms that shape the functional receptive field of an RGC and this contributes to our understanding of the cellular and molecular mechanisms that control synaptic connectivity in the developing retina. It is known that intercellular communication between RGCs and their presynaptic neurons (amacrine and bipolar cells), in the form of activity-dependent as well as molecular signals, is responsible for the remodeling and shape of dendritic arbors. Stratification and laminar refinement are mediated by interactions between transmembrane recognition molecules that are conserved through evolution and that guide the branching patterns of distinct neuronal subtypes [5]. Laminar and synaptic refinement is also modulated by activity-dependent signals, which in turn may control the expression and function of important proteins, such as brain-derived neurotrophic factor (BDNF), within the local retinal circuitry. BDNF contributes to RGC dendritic branching and laminar refinement, acting both locally within the retina and also through retrograde mechanisms acting at the axonal target(s) [14-17]. To what extent these and other local molecular signals contribute to the patterned distribution of excitatory synaptic inputs in the retinal circuit, and how this pattern is modified by activity and potential retrograde signals from the brain, is now open to new investigation. Studies that employ both static and dynamic analyses of synaptic components in living neurons in their natural setting are beginning to provide a long-sought window into the developing brain.