OUTLOOK

Creating cell type-specific mutants by enhancer mutagenesis

Stephen Crews

Department of Biochemistry and Biophysics, School of Medicine, The University of North Carolina at Chapel Hill, Chapel Hill, North Carolina 27599, USA

Cell signaling plays an essential role in development, and knowledge of the identities of the cells sending the signal is critical. This can be a challenge, since signaling pathways and ligands are commonly used at multiple times and in multiple cell types during development. One solution to this problem is to create cell type-specific mutants using CRISPR/Cas9 to mutate enhancers that control different patterns of expression. In this issue of *Genes & Development*, Rogers and colleagues (pp. 634–638) provide the first use of this method in *Drosophila* to solve a long-standing issue in patterning of the embryonic central nervous system.

One of the surprises to emerge from studying the molecular genetics of development is that a relatively small number of signaling pathways are used repeatedly in different cell types and at different times of development. This has been a challenge to developmental biologists, who generally need to know which cell type is doing the signaling. Ideally, mutants would exist that reduce function only in a specific cell type. These mutants most likely would reside in enhancers that control cell type-specific transcription. However, mutants of this type are rare. Fortunately, as demonstrated in the present study (Rogers et al. 2017), the advent of CRISPR/Cas9 mutagenesis and the relative ease in identifying and determining the cellular expression of enhancers now promise to make the creation of cell type-specific mutant strains routine.

The issue addressed by Rogers et al. (2017) concerns patterning of the *Drosophila* embryonic central nervous system (CNS). Early in development, the neuroectoderm is defined by a discrete set of CNS midline cells flanked by three bilaterally symmetrical rows of neuroectodermal cells: the ventral, intermediate, and dorsal neuroectoderm. The *intermediate neuroblasts defective* (*ind*) gene is expressed specifically in the intermediate neuroectoderm and promotes intermediate column neuroblast

formation (Fig. 1A, blue). It was demonstrated that the EGFR signaling pathway is critical for the formation of intermediate column neuroblasts and the expression of *ind*. Furthermore, EGFR signaling is dependent on two ligands: Spitz (Spi) and Vein (Vn). Generation of functional Spi requires processing by the Rhomboid (Rho) membrane protease, and expression of *rho* generally indicates a cellular source of Spi signaling. However, in this case, *rho* is expressed in two nearby cell types: the CNS midline cells and the ventral neuroectoderm (Fig. 1A; red). The CNS midline cells are known to be a prominent source of developmental signals in *Drosophila* (as well as in crustaceans, fish, and mammals) and are an attractive candidate as the source of Spi and Vn signaling. However, the closer spatial proximity of the ventral neuroectodermal expression of *spi* and *vn* to the *ind* expression domain makes this set of cells a reasonable alternative.

To directly address this issue, the investigators took advantage of detailed *rho* gene expression studies that had previously identified distinct enhancers: one for neuroectodermal expression and two for midline expression (Fig. 1B). Using CRISPR/Cas9 mutagenesis to create homozygous enhancer deletion mutants, they created an endogenous *rho* gene that lacked neuroectodermal expression yet retained midline expression (Fig. 1C, *rho NEE* knockout). Since the Vn ligand is likely to act redundantly with Spi, a *rho NEE* knockout, *vn* double mutant was assayed for embryonic *ind* expression. Only sporadic *ind* expression was observed, demonstrating that midline *rho* expression was not sufficient for normal *ind* expression (Fig. 1C). In contrast, deletion of the two *rho* midline enhancers (*rho SHA MLE* knockout) resulted in the absence of *rho* midline expression, but *ind* expression was normal (Fig. 1D). However, the observed *ind* expression could be due to Vn acting as a redundant EGFR ligand. This was resolved by examining a *single-minded* (*sim*) *vn* double mutant, which has greatly reduced midline *rho* expression and lacks *vn* expression yet produces a normal *ind* pattern

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Corresponding author: steve_crews@unc.edu

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Figure 1. (A) Schematic of a *Drosophila* embryo showing the expression of *rho* (red) in the CNS midline cells (ml) and ventral neuroectoderm (v-nec). The *ind* gene is expressed (blue) in the intermediate neuroectoderm (i-nec). (B) Schematic of the *rho* gene (not to scale) showing the two midline enhancers (SHA and MLE) and the neuroectodermal enhancer (NEE). Corresponding embryonic expression of *rho* and *ind* is at the right. (C) Deletion of the neuroectodermal enhancer resulted in an incomplete pattern of *ind* expression. (D,E) Removal of the two midline enhancers (D) and analysis of a *sim* *vn* double mutant (E) showed relatively normal *ind* expression.

(Fig. 1E). These results demonstrated that the source of Spi signaling is the neuroectoderm and not the midline.

This CRISPR/Cas9-mediated approach should prove generally useful for providing new insights into a variety of developmental and physiological questions regarding any experimental organism. One major advantage of cell type-specific mutants is avoiding pleiotropy. In many cases, this will make interpretation of phenotypes significantly clearer compared with examining typical loss-of-function mutants. It is worth noting that enhancers commonly control temporal expression, and enhancer mutants that affect only specific time windows of expression may be invaluable and help compensate for a general lack of effective temperature-sensitive mutants. Creating enhancer mutants has other benefits in understanding how genes are regulated and function. Although great progress has been made in understanding enhancer function and transcriptional regulation, the biological function of individual enhancers is not as well known. Understanding how individual enhancers function to generate patterns of gene expression is particularly important given that human disease loci commonly map to noncoding sequences and likely influence transcript levels (Maurano et al. 2012). For example, in a review on eukaryotic transcriptional autoregulation (Crews and Pearson 2009), it was noted that while the phenomenon of autoregulation was widespread, there was relatively little data regarding its biological significance—mutation of autoregulatory enhancers is a solution. Many genes have multiple enhanc-

ers that can drive similar patterns of expression (shadow enhancers). That enhancer mutations can provide novel insights into biological function was demonstrated by the removal of shadow enhancers, which revealed that they confer developmental robustness to varying environmental and genetic conditions (Frankel et al. 2010; Perry et al. 2010). In an extreme case, the *Drosophila* *sim* gene has potentially eight CNS midline enhancers (Pearson et al. 2016). What is the biological contribution of each enhancer toward gene function? Analyzing enhancer mutations in vivo will provide important new insights into the control and evolution of patterns of gene expression.

Using CRISPR/Cas9 to create cell type-specific mutants by enhancer deletion requires that relevant enhancers be identified and characterized. In fact, Rogers et al. (2017) made good use of extensive enhancer analyses of the *rho* gene begun 25 years ago. While identifying enhancers using in vivo transgenic enhancer testing approaches is generally straightforward and successful, it can be time-consuming. Fortunately, there are genomic advances that will accelerate progress. For *Drosophila*, there is the existence of large-scale Gal4 transgenic projects that have assayed thousands of genomic fragments from hundreds of genes for their in vivo transcriptional activity (Manning et al. 2012; Kvon et al. 2014) as well as an annotated enhancer database (Gallo et al. 2011). These data are a useful starting point for further deletion analyses to reduce a fragment to a manageable size for enhancer mutations. Another approach should also prove particularly valuable. This involves carrying out genome-wide surveys of open chromatin regions using ATAC-seq (assay for transposase-accessible chromatin [ATAC] with high-throughput sequencing) and FAIRE-seq (formaldehyde-assisted isolation of regulatory elements [FAIRE] coupled with sequencing) (McKay and Lieb 2013). These regions of open chromatin generally correspond to enhancers and promoters. With these data, one can delete these open chromatin regions individually and assay expression and phenotypes, thus bypassing the traditional transgenic method of enhancer analysis. Most useful will be identifying open chromatin regions from purified cells or nuclei corresponding to the cells of interest. This will help identify relevant cell type-specific enhancers. Recently, a comparison was made of FAIRE-seq data from purified *Drosophila* CNS midline cells with whole-embryo FAIRE-seq data (Pearson et al. 2016). It was demonstrated that peaks of open chromatin that are strong in the midline sample and weak in the whole-embryo sample had a high probability of being enhancers driving in vivo-relevant midline expression. With this type of information, one can go directly to using CRISPR/Cas9 for making cell type-specific enhancer mutants in vivo.

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