RESEARCH COMMUNICATION

Regulation of neuroblast competence: multiple temporal identity factors specify distinct neuronal fates within a single early competence window

Michael D. Cleary and Chris Q. Doe

University of Oregon Department of Neuroscience, Howard Hughes Medical Institute, Eugene, Oregon 97403, USA

Cellular competence is an essential but poorly understood aspect of development. Is competence a general property that affects multiple signaling pathways (e.g., chromatin state), or is competence specific for each signaling pathway (e.g., availability of cofactors)? Here we find that Drosophila neuroblast 7-1 [NB7-1] has a single early window of competence to respond to four different temporal identity genes [Hunchback, Krüppel, Pdm, and Castor]; that each of these factors specifies distinct motor neuron identities within this competence window but not outside it; and that progressive restriction to respond to Hunchback and Krüppel occurs within this window. Our work raises the possibility that multiple competence windows may allow the same factors to generate different cell types within the same lineage.

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Cellular diversity is often achieved through the generation of multiple cell types from a single progenitor. One of the most striking examples of this process occurs in the CNS, where vast numbers of neurons and glia with unique functions are derived from a limited number of progenitors. Regulated expression of intrinsic and extrinsic cues that confer spatial and temporal identity is one mechanism for controlling the generation of cellular diversity in the CNS [Pearson and Doe 2004]. However, a common feature of progenitor cells is that they do not remain competent to respond to such cues throughout development. Restriction of progenitor competence is an important mechanism ensuring that the correct progeny are made at the proper time and place.

The vertebrate CNS provides at least two examples of the role of progenitor competence in the generation of cellular diversity. The cerebral cortex is derived from progenitor cells whose progeny migrate to distinct layers depending on their time of birth [McConnell 1992]. Clonal analysis and heterochronous transplant experiments have shown that early cortical progenitors are multipotent, but the ability to make early cell types is gradually lost as progenitors age, in a process known as "progressive restriction" [Desai and McConnell 2000]. Both extrinsic and intrinsic cues appear to regulate cortical progenitor competence [Bohner et al. 1997; Hanashima et al. 2004]. Restriction of competence also occurs during vertebrate retinal development, but instead of a progressive restriction of competence, retinal progenitors pass through distinct, nonoverlapping competence states [Cepko et al. 1996]. While retinal progenitors at each stage can produce multiple cell types, once a competence state has been passed it is not possible to induce production of the earlier cell types. As with cortical progenitors, both extrinsic and intrinsic cues appear to regulate retinal progenitor competence [Waid and McLoon 1998; Belliveau and Cepko 1999; Rapaport et al. 2001].

While studies of cortical and retinal progenitors demonstrate the general significance of progenitor competence in generating cell diversity, little is known about the molecular basis of "competence" or "progressive restriction of competence". Is competence a general cellular state that allows changes in gene expression in response to multiple signaling pathways? Or is competence specific for each signaling pathway, perhaps reflecting the availability of a particular cofactor?

Here we investigate these questions using the well-characterized Drosophila neuroblast 7-1 [NB7-1] system [Fig. 1]. NB7-1 is a neural progenitor that buds off a series of ganglion mother cells [GMC-1, GMC-2, GMC-3, etc.; named by birth order], and each GMC makes two postmitotic neurons [U1/sib, U2/sib, U3/sib, etc.; named by birth order], the U1–U5 neurons can be identified by their position and staining for Even-skipped [Eve] protein [Isshiki et al. 2001; Pearson and Doe 2003]. Four different transcription factors—Hunchback [Hb], Krüppel [Kr], Pdm1/Pdm2 [Pdm], and Castor [Cas]—are sequentially expressed in NB7-1 [Isshiki et al. 2001; Pearson and Doe 2003; Grosskortenhaus et al. 2005], similar to other neuroblast lineages [Kambadur et al. 1998; Brody and Odenwald 2000; Isshiki et al. 2001; Novotny et al. 2001; Kanai et al. 2005]. NB7-1 has high Hb and low Kr during the first division [and maintained in the U1 neuron]; low Hb and low Kr during the second division [and maintained in the U2 neuron]; Kr alone is present during the third division [and maintained in the U3 neuron]; Pdm alone is present during the fourth division [and transiently detected in the U4 neuron], and both Pdm and Cas are present during the fifth division [with Cas maintained in the U5 neuron] [Fig. 1; Isshiki et al. 2001]. All four transcription factors specify temporal identity within the NB7-1 lineage. Hb is necessary and sufficient to specify U1 and U2 fates [high levels give U1, lower levels give U2] [Isshiki et al. 2001; Pearson and Doe 2003; Grosskortenhaus et al. 2005]; Kr is necessary and sufficient to specify U3 fate [Isshiki et al. 2001; this work], and Pdm specifies U4 cell fate, whereas Pdm/Cas together specify U5 cell fate [R. Grosskortenhaus and C.Q. Doe, unpubl.].

Most importantly for this work, the competence of NB7-1 to respond to Hb has been well-characterized.
NB7-1 has a single competence window for responding to Hunchback and Krüppel

To determine whether NB7-1 undergoes progressive restriction in competence to respond to Kr, similar to that observed for Hb [Pearson and Doe 2003], we generated pulses of Kr at progressively later points in the NB7-1 lineage. We used hsp70-Kr to induce expression of Kr within the NB7-1 lineage from the fourth division onward [Fig. 3A]. When we use one copy of UAS-Kr at 22°C, which provides relatively low levels of Kr, we observed only five to six Eve+ U neurons [Fig. 3C], mostly U1, U2, and three U3 neurons [91%], but also U1, U2, and four U3 neurons [9%]. Thus, NB7-1 loses competence to respond to prolonged Kr expression after five to six cell divisions, similar to results from the Kr pulse experiments described above. Prolonged expression of Hb using the same conditions (prospero-gal4, one copy of UAS-Hb, 22°C) also results in just five to six Eve+ U neurons [Pearson and Doe 2003]. We conclude that NB7-1 has a single competence window for generating U1–U3 neurons in response to Hb and Kr.

We next wanted to test whether the later-expressed temporal identity factors Pdm and Cas share the same early competence window with Kr, or if they have distinct competence windows. Pdm specifies the U4 neuronal identity, while Pdm/Cas together specify U5 neuronal identity [R. Grosskortenhaus and C.Q. Doe, unpubl.]. We used scabrous-gal4 to prolong Kr expression for a variable length of time within the NB7-1 lineage [two copies of UAS-Kr at 29°C], which delayed but did not prevent the sequential expression of Kr, Pdm, and Cas [Fig. 4]. This experiment allowed us to assay NB7-1 competence when presented with Kr, Pdm, or Cas at different times in its lineage.

We found that the scabrous-gal4 UAS-Kr embryos always had a total number of seven to eight Eve+ U neurons, although ectopic U3 neurons ranged from two to six in number [Fig. 5]. Interestingly, hemisegments with only two ectopic U3 neurons typically had U4/U5 neu-
rons; those with three ectopic U3 neurons had only a U4 neuron, and those with four or more ectopic U3 neurons lacked both U4/U5 neuronal fates (Fig. 5B,C). We interpret these data in the following way. In segments where Kr declines the fastest [fewest ectopic U3 neurons], there is time for Pdm to induce U4 fate and Pdm/Cas to induce U5 fates prior to loss of competence; however, in segments where Kr lasts the longest, both Pdm and Cas expression occur after the competence window and no U4/U5 fates are produced. Taken together, this experiment allows us to draw several conclusions (summarized in Fig. 5D): (1) Prolonged Kr expression can partially extend the neuroblast competence window (from five to six divisions to seven to eight divisions); (2) competence to respond to Kr, Pdm, and Cas is simultaneously lost at the end of this competence window, suggesting that there is a single competence window for responding to multiple temporal identity factors; and (3) each temporal identity factor specifies different U1–U5 motor neuron identities within the competence window, but not outside it. It is currently an open question as to how prolonged expression of one factor [Kr, this work; Hb, Grosskortenhaus et al. 2005] can extend the competence window to respond to three distinct factors [Kr, Pdm, and Cas].

Loss of neuroblast competence requires down-regulation of Hunchback but not Krüppel

Our previous experiment showed that prolonging Kr expression [scabrous-gal4 UAS-Kr] in NB7-1 lineage can only partially extend neuroblast competence. Interestingly, similar experiments prolonging Hb expression [scabrous-gal4 UAS-hb] revealed that the neuroblast maintains full competence for as long as Hb is expressed, in some cases over 15 divisions, with normal U3–U5 fates appearing after Hb levels decline (Grosskortenhaus et al. 2005). Thus, extended Hb expression—but not extended Kr expression—can maintain the neuroblast in a young, fully competent state. This raised the possibility that down-regulation of Hb is required for loss of neuroblast competence, alternatively, Hb may be more potent than Kr in maintaining neuroblast competence.

To distinguish these models, we compared the effect of high-level Hb or Kr expression beginning at the fourth neuroblast division (prospero-gal4, 2× UAS-hb or UAS-Kr, 29°C), which would allow Hb down-regulation and permit us to compare the efficacy of Hb versus Kr in extending neuroblast competence. Performing this experiment with Hb resulted in a partial extension of neuroblast competence and the production of an average of 9.1 Eve+ U neurons: U1–U3, 6.1 extra U1, and no U4/U5 (Pearson and Doe 2003). Performing the experiment with Kr resulted in an almost identical phenotype of 9.8 Eve+ U neurons: U1/U2, 7.8 U3s, and no U4/U5 (Fig. 3D). Thus, Hb and Kr appear equally efficient at extending neuroblast competence; this is supported by their equivalent effect when expressed under heat shock or lower level prospero-gal4 control [competence lost after five divisions] (Figs. 2, 3C; Pearson and Doe 2003). More importantly, a comparison of the scabrous-gal4 UAS-hb and prospero-gal4 UAS-hb experiments shows that Hb down-regulation is critical for loss of neuroblast competence. When Hb is maintained from the beginning of the lineage [scabrous-gal4 UAS-hb], competence persists for the length of Hb expression, in some cases over 15 divisions; when Hb down-regulation occurs followed by permanent Hb re-expression one division later [prospero-gal4 UAS-hb], then competence is lost after approximately nine divisions [Pearson and Doe 2003; Grosskortenhaus et al. 2005]. We conclude that down-regulation of Hb, but not Kr, initiates progressive restriction in neuroblast competence that is normally complete after five divisions.
Mitotic progenitors, but not post-mitotic neurons, are competent to respond to Krüppel

Thus far, we have been investigating how neuroblast competence changes over multiple rounds of cell division. Here, we consider how competence changes during neuronal differentiation. We expressed Kr in high levels in the newborn post-mitotic U1–U5 neurons (eve-gal4 UAS-Kr). In these embryos, Kr is first detected just as the U1–U5 neurons are born. Despite high levels of Kr protein, no change in U1–U5 fate was ever detected (Supplementary Fig. 2). Conversely, transient expression of Kr in NB7-1/GMCs can occasionally generate ectopic U3 neurons that do not maintain Kr expression (Supplementary Fig. 3), despite the ability of Kr to positively autoregulate within the CNS (Supplementary Fig. 4). Thus, mitotic progenitors but not post-mitotic neurons are competent to respond to Krüppel. Similar results have been observed for competence to respond to Hb [Pearson and Doe 2003].

Conclusions and future directions

Our experiments, combined with previous studies [Isshiki et al. 2001; Pearson and Doe 2003; Grosskortenhaus et al. 2005], allow us to make four major conclusions.

1. A single early competence window is used by multiple temporal identity factors. The molecular basis for the early competence window is unknown, but it must be general enough to modulate response to four distinct transcription factors rather than being factor specific. Perhaps loss of competence leads to restricted access of Hb, Kr, Pdm, and Cas to target loci involved in U1–U5 neuronal specification; other loci may remain unaffected, allowing these four transcription factors to induce different cell fates later in the neuroblast lineage (see below). Identifying Hb and Kr target genes, and in-
vestigating how or whether they undergo chromatin modifications during the process of progressive restriction will help resolve this question, and may provide insight into the mechanism of progressive restriction in mammalian neural progenitors.

(2) Each temporal identity factor specifies distinct motor neuron fates within the competence window, but not outside of it. Within the early competence window, each temporal identity factor specifies a unique U1–U5 neuronal identity: high Hb, U1; low Hb, U2, Kr, U3; Pdm, U4; Pdm/Cas, U5. The loss of competence to generate U1–U5 fates may allow each of these transcription factors to be “reused” later in the NB7–1 lineage to generate different subsets of neurons. This model is supported by the fact that we and others (T. Isshiki, pers. comm.) have observed a second round of Kr and Cas neuronblast expression later in embryonic development [Fig. 4]. These findings suggest that neuroblasts have the potential for cycling through distinct competence windows, and may provide a model for understanding how successive competency states are established (e.g., in vertebrate retinal progenitors).

(3) NB7–1 undergoes progressive restriction in competence to respond to both Hb and Kr. Competence to respond to both Hb and Kr is progressively restricted early in the lineage, then completely lost after five divisions of NB7–1. Progressive restriction may be regulated autonomously in the neuroblast or by changing environmental cues, such as inhibitory feedback from GMC or neuronal progeny. We favor a lineage-intrinsic mechanism because different neuroblasts lose competence to respond to Hb at different times (e.g., NB7–1 remains competent to respond to Hb for five divisions, whereas the adjacent NB1–1 is only competent to respond to Hb for two to three divisions) [Isshiki et al. 2001]. A feedback inhibition model would have parallels with vertebrate retinal progenitors, where differentiated amacrine cells send an inhibitory feedback signal to terminate amacrine cell production [Bellido and Cepko 1999]. In this case, the signal would likely depend on the number of progeny produced rather than the type of progeny, because loss of competence can occur without production of the last-born neurons in the competence window [U4/ U5] (see Figs. 3, 5).

(4) Down-regulation of Hb but not Kr initiates progressive restriction and loss of competence. Neuroblast competence is maintained if Hb is expressed from the beginning of the lineage [Grosskortenhaus et al. 2005]. However, neuroblast competence is not maintained when Kr is expressed from the beginning of the lineage [where Hb is down-regulated normally] [Fig. 5] or when Hb or Kr are expressed later in the lineage after normal Hb down-regulation [Figs. 2, 3; Pearson and Doe 2003]. We propose that Hb down-regulation initiates progressive restriction in neuroblast competence, ultimately leading to a complete loss of competence.

Materials and methods

Fly stocks

[1] hs70-Kr/Cyo hb-lacZ; [2] hs70-hb (Hb476.2)/CyO; hs70-hb [Hb476.1] TM3; [3] yw, UAS-Kr; UAS-Kr; [4] yw; UAS-Kr; [5] prospero-gal4 homozygous on chromosomes II and III; [6] eve-gal4;[1–3.5–4.3] homozygous on chromosomes II and III; [7] scabrous-gal4 homozygous on chromosome II; [8] engrailed-gal4 homozygous on chromosome II.

Unless otherwise noted, all Gal4 lines were crossed to yw; UAS-Kr; UAS-Kr Bies, and embryos were collected at 29°C.

Heat-shock experiments

Embryos were collected for 1 h, aged to the indicated time, and subjected to two cycles of 30 min at 37°C, then 30 min at 22°C, and allowed to develop to stage 16–17. This generates high levels of Hb and Kr without affecting control embryos. Before heat shock, some embryos were fixed and stained with Eve/Hb or Eve/Kr for developmental staging. When scoring the frequency of extra cells per hemisphere, no more than five hemisegments were scored per embryo (typically just one side of the CNS) so that, on average, at least 10 different embryos were used in the analysis of each timepoint.

Protein immunolocalization

Detection of Biological Sciences, National University of Singapore, Singapore); rabbit anti-Cas, 1:500, W. Odenwald [Neural Cell-Fate Determinants Section, National Institute of Neurological Disorders and Stroke, National Institutes of Health, Bethesda, MD]; guinea pig anti-Eve, 1:400, EADC; mouse anti-Eve monoclonal 288, 1:20, guinea pig anti-Runt, 1:40, EADC; rabbit anti-Runt, 1:500 of 1:10 preabsorbed, A. Brand [Wellcome Trust/Cancer Research UK Institute and Department of Genetics, University of Cambridge, Cambridge, UK]; rabbit anti-Zhid, 1:200, M. Lundell [Division of Life Sciences, University of Texas at San Antonio, San Antonio, TX] and sheep anti-Digoxigenin, 1:1000, Boehringer. Species-specific secondary antibodies were conjugated to Alexa 488, Alexa 555 (Molecular Probes), Cy5 (Jackson ImmunoResearch), or Biotin (Vector Labs) and were used at 1:200. The TSA-plus Fluorescein System [PerkinElmer] was used to detect the anti-Pdm1 antibody. All images were collected as confocal image stacks on a Leica TCS SP2 AOBS confocal microscope, processed in ImageJ (NIH), and shown as two-dimensional projections. In all experiments, analysis was restricted almost exclusively to the abdominal segments of the CNS.

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