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A cell-intrinsic timer that operates during oligodendrocyte development

Béatrice Durand^{1*} and Martin Raff²

Summary

Multicellular organisms develop on a predictable schedule that depends on both cell-intrinsic timers and sequential cell-cell interactions mediated by extracellular signals. The interplay between intracellular timers and extracellular signals is well illustrated by the development of oligodendrocytes, the cells that make the myelin in the vertebrate central nervous system. An intrinsic timing mechanism operates in each oligodendrocyte precursor cell to limit the length of time the cell divides before terminally differentiating. This mechanism consists of two components, a timing component, which depends on the mitogen platelet-derived growth factor (PDGF) and measures elapsed time, and an effector component, which depends on thyroid hormone and stops cell division and initiates differentiation at the appropriate time. The cell-cycle inhibitor p27/Kip1 accumulates in the precursor cells as they proliferate and is part of both components of the timer. It seems likely that similar timing mechanisms operate in other cell lineages. *BioEssays* 22:64–71, 2000.

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Philosophers and physicists have long debated whether time is linear or circular, directional or nondirectional, reversible or irreversible.⁽¹⁾ For biologists, time is unequivocally linear, directional, and irreversible: species evolve, and individual organisms develop on a predictable schedule and invariably senesce and die; there is no going back. An important challenge is to understand the mechanisms that underlie these directional processes.

Besides these linear processes, both individual cells and multicellular systems display various types of cycles, or rhythms, which could, in principle, be involved in timing the progress of development and senescence. Circadian clocks are an example. They control a variety of physiological and behavioural functions in most organisms.⁽²⁾ It seems unlikely, however, that circadian clocks are used as timing devices in development or senescence, at least in animals, as muta-

tions that disable them in flies or mice have little effect on either of these processes. Cells and organisms have clearly developed other ways of keeping time.

Timing in animal development, however, does seem to depend on intracellular timers (reviewed in Refs. 3 and 4), as well as on sequential cell-cell interactions. In principle, an intracellular timer could operate in a number of ways. It could involve noncircadian intracellular oscillators, such as those that control the cell division cycle,⁽⁵⁾ or those that generate periodic ion fluxes,⁽⁶⁾ or repeating bursts of specific gene expression.⁽⁷⁾ Such oscillations seem to be used, like the ticks of a clock, to time some developmental processes, but how this is achieved is unknown. An intracellular timer could work through the sequential activation of a series of genes encoding gene regulatory proteins, with each transcriptional activator activating the next in the sequence, until a final gene product triggers a developmental event. Alternatively, a timer could operate like an hourglass, where the progressive accumulation or loss of a molecule involved in a rate-limiting process serves as a measure of time, with the molecule triggering a developmental event when a threshold level is reached.

Some examples of intracellular developmental timing mechanisms

An example of an intracellular developmental timing mechanism that makes use of the cell division cycle is seen in the early *Xenopus* embryo, where zygotic gene transcription begins at a specific time point called the midblastula transition (MBT). The timing of MBT depends on repeated cycles of DNA replication and cell division. Because the cells divide without cell growth, the ratio of nuclear DNA to cytoplasm in each cell increases with each division cycle, until it reaches a critical value that triggers zygotic gene transcription.^(8,9) Another example where timing depends on a form of cell-division counting is in the replicative senescence of cultured human fibroblasts, which stop dividing after a limited number of divisions when stimulated by mitogens. In this case, telomeres shorten with each cell division, until they become sufficiently short that they activate a checkpoint mechanism that arrests the cell cycle.⁽¹⁰⁾ Most of the mechanisms that control the timing of development, however, do not seem to depend on cell-division counting. Cells in a developing embryo, for example, frequently differentiate on close to a normal schedule even when cell division has been blocked

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by drug treatment.^(11,12) Moreover, many changes that occur in developing cells over time often occur after the cells have permanently withdrawn from the cell cycle.

Some developmental timers depend on other intracellular cycles. A dramatic example is seen during the development of vertebrate somites, which form every 30 min in early chick embryos. Remarkably, the expression of specific genes in the mesoderm cells that will form the somites in these embryos oscillates with a 30 min periodicity, suggesting that these oscillations somehow determine when these cells assemble into somites.^(3,7,13) It remains to be determined how the oscillations are controlled and how they are translated into developmental time.

Some developmental timers do not appear to depend on gene transcription or protein synthesis at all. In *Xenopus*

embryo development, for example, a number of precisely timed early events occur normally even when DNA, RNA, and protein synthesis are inhibited by drug treatment.^(11,14) These timers presumably depend on post-translational controls, but their nature remains a mystery.

As mentioned earlier, the timing of development in multicellular organisms depends on more than intracellular timing mechanisms described above. It also depends on cell-cell interactions, which are mainly mediated by extracellular signalling molecules. This interplay between intracellular timers and extracellular signals is well illustrated in the development of oligodendrocytes, which are the cells that make myelin in the vertebrate central nervous system and are the main focus of this review.

Timing of glial cell development in cultures of embryonic brain cells

The vertebrate brain contains a variety of types of glial cells, each of which develops on a predictable schedule. Remarkably, if cells are dissociated from an embryonic rat brain and grown in culture, the major classes of glial cells develop on the same schedule as they do in the developing brain.⁽¹⁵⁾ These findings suggest that the mechanisms that control the timing of glial cell development in vivo also operate in a culture dish, where they are accessible to analysis. The complexity of the brain cell cultures, however, makes such an analysis difficult, which is why we turned to the optic nerve, which contains no neurons and is therefore far simpler.

Glial cell development in the rat optic nerve

The optic nerve contains the axons of retinal ganglion neurons and two types of glial cells: astrocytes, which support

the axons in the nerve structurally and functionally, and oligodendrocytes, which extend processes that wrap around the axons to form an insulating myelin sheath (Fig. 1). The astrocytes develop from the neuroepithelial cells of the optic stalk, the primordium of the nerve, while the oligodendrocytes develop from precursor cells that migrate into the developing nerve from the brain beginning just before birth.⁽¹⁶⁾ The oligodendrocyte precursor cells divide a limited number of times before they stop and terminally differentiate into postmitotic oligodendrocytes. The first oligodendrocytes appear in the nerve around the day of birth, and their numbers then progressively increase over the next 6 weeks.^(17,18)

The astrocytes play an important part in oligodendrocyte development. For example, they secrete platelet-derived growth factor (PDGF), the main mitogen that drives proliferation of oligodendrocyte precursor cells.^(19,20) In the absence of PDGF, the precursors divide very little in culture, and few oligodendrocytes develop in vivo.^(21–23)

Oligodendrocyte development in cultures of optic nerve cells

The normal timing of oligodendrocyte development can be reconstituted in cultures of dissociated embryonic optic nerves,⁽²⁴⁾ as long as the oligodendrocytes are stimulated to divide by PDGF.⁽²⁵⁾ The reason the precursors stop dividing and differentiate in such cultures is not because PDGF becomes limiting or because the PDGF receptors on these cells disappear or are no longer activated by PDGF. Adding excess PDGF does not alter the timing of oligodendrocyte

differentiation,⁽²⁵⁾ and newly formed oligodendrocytes still express large numbers of PDGF receptors,⁽²⁶⁾ stimulation of which induces both an increase in cytosolic Ca²⁺⁽²⁷⁾ and immediate-early gene expression.⁽²⁸⁾ These findings raised the possibility that the precursor cells have a cell-intrinsic mechanism that limits their proliferative capacity and controls when they withdraw from the cell cycle and undergo terminal differentiation.

A cell-intrinsic timer in the oligodendrocyte precursor cell

When individual or purified oligodendrocyte precursor cells isolated from postnatal day 7 (P7) rat optic nerve are stimulated to proliferate in culture by either astrocytes or saturating amounts of PDGF, the cells divide up to eight times before they stop and differentiate.^(29,30) The progeny of an individual precursor cell stop dividing and differentiate at about the same time.^(29,30) If the two daughter cells of an individual precursor are separated and cultured on astrocyte monolayers in separate microwells, they divide the same number of times before they differentiate, indicating that an intrinsic mechanism operates in the precursor cells to limit the number of times, or the period of time, that the cell normally divides.⁽²⁹⁾

Does the counting mechanism count cell divisions, or does it measure time in some other way? Oligodendrocyte precursor cells cultured at 33°C divide more slowly, but stop dividing and differentiate after fewer divisions compared to cells cultured at 37°C. This suggests that the intrinsic mechanism does not operate by counting cell divisions but, instead, measures elapsed time in some other way.⁽³¹⁾ We therefore refer to it as a timer.

Two functional components of the timer

Although the timer is intrinsic to the precursor cell, its activity depends on extracellular PDGF, which is normally produced by other cells, such as astrocytes in the optic nerve. If a single precursor cell is cultured on its own in the absence of PDGF, it stops dividing and differentiates within 1–2 days.⁽³²⁾ Similarly, in the absence of PDGF, purified precursor cells differentiate prematurely, no matter how high the density of the cells in the culture dish.⁽¹⁸⁾ Thus, the timer only works if the cells are continuously stimulated to proliferate by PDGF.

In addition to PDGF, the timer also needs hydrophobic signals, such as thyroid hormone (TH) or retinoic acid (RA), to function normally. Precursor cells cultured in the presence of mitogens but in the absence of TH or RA keep dividing and do not differentiate⁽³⁰⁾ (Fig. 2). If TH is added to such a culture after 8 days, when most precursor cells would have differentiated had TH been present, the precursor cells soon stop dividing and differentiate.⁽³⁰⁾ Thus, in the presence of mitogens and the absence of hydrophobic signals, the cells can still keep track of time, but they seem unable to stop dividing and differentiate when the appropriate time is reached.

Therefore, the timer consists of at least two functional components: (1) a timing component, which depends on PDGF but not TH and measures elapsed time, and (2) an effector component, which is regulated by TH and stops cell division and initiates differentiation when the timing component indicates it is time.⁽³³⁾ A similar conclusion was reached by Böglér and Noble,⁽³⁴⁾ who used a combination of PDGF and fibroblast growth factor (FGF) to keep the precursor cells dividing beyond their normal limit.

All of the studies investigating the intrinsic timing mechanism of oligodendrocyte precursor cells have used cells in culture. Does this timer also operate in the developing nervous system to determine when cells stop dividing and differentiate? Two lines of evidence suggest that it does. First, exposing the P8 rat optic nerve to excess PDGF does not increase the proliferation of oligodendrocyte precursor cells in the nerve.⁽¹⁸⁾ This suggests that precursor cells do not normally stop dividing and start to differentiate because of PDGF deprivation, at least at this stage of development. Second, a cell lineage study in the developing rat brain found that clones of oligodendrocyte lineage cells contained either precursors or oligodendrocytes, but not both,⁽³⁵⁾

which suggests that the progeny of an individual precursor cell differentiate synchronously.

The role of p27/kip1

It is not certain whether the timer in oligodendrocyte precursor cells primarily controls the onset of differentiation, the cessation of proliferation, or both. Because the precursor cells stop dividing immediately and differentiate when deprived of PDGF,^(25,29) we suspect that the timer primarily controls cell proliferation. If this is the case, the timer must, at some point, interact with the cell-cycle control system that regulates progress through the cell division cycle. Since the cells stop dividing and differentiate in the G₁ phase of the cell cycle, it is likely that the relevant interactions occur within the control system that operates in G₁ (Fig. 3).

The eucaryotic cell cycle is controlled by a family of cyclin-dependent protein kinases (Cdks), which are cyclically activated to trigger the different phases of the cell cycle at the right time and in the right sequence.^(36,37) These Cdks are regulated by at least three types of proteins: cyclins activate and direct them to their substrates; kinases and phosphatases activate or inhibit their activity; and cyclin-dependent kinase inhibitors block the assembly or activity of the cyclin-Cdk complexes.^(36–38) Two families of Cdk inhibitors have been identified in mammalian cells, the Cip/Kip family, which includes p21Cip1, p27Kip1, and p57Ink4b, and the Ink4 family, which includes p16Ink4a, p15Ink4b, p18Ink4c, and p19Ink4d.⁽³⁹⁾ When over-expressed in cells, each of these inhibitors arrest the cell cycle in G₁ (Fig. 3). Moreover, some cdk inhibitors are up-regulated when developing cells stop dividing and differentiate, so that they could well play a part in arresting the cell cycle at the appropriate time.⁽⁴⁰⁾

In principle, the timer in oligodendrocyte precursor cells might depend on a decrease in a positive intracellular regulator such as a cyclin or Cdk that promotes cell-cycle progression, an increase in a negative intracellular regulator such as a Cdk inhibitor that blocks cell-cycle progression, or both types of mechanism. It would be advantageous to use both mechanisms, as this would make the timer more accurate, and, if one mechanism were defective, the timer would still work, although inaccurately.

Several lines of evidence suggest that the Cdk inhibitor p27/Kip1 (p27) is part of the timer. First, p27 levels are high in oligodendrocytes, both in vivo and in vitro, whereas they are variable in precursor cells.⁽⁴¹⁾ Second, p27 levels increase rapidly when precursor cells in culture are deprived of PDGF; the rate of increase is similar to the rate at which the precursor cells commit to cell-cycle withdrawal and differentiation, suggesting that p27 is part of the effector mechanism.⁽⁴¹⁾ Third, p27 levels progressively increase when P7 precursor cells are cultured in the presence of PDGF but in the absence of TH,⁽⁴¹⁾ conditions where the

timing component continues to operate, but the effector component is defective.⁽³⁰⁾ p27 levels plateau after about 8 days in these cultures, when most of the cells would have differentiated had TH been present (Fig. 4). In the absence of TH, however, despite high levels of p27, cells continue to divide, indicating that high levels of p27 alone are insufficient to stop the cell cycle. Fourth, p27 levels rise faster at 33°C than at 37°C, as would be expected if p27 is part of the timer, since the timer runs faster at the lower temperature.⁽³¹⁾ Fifth, and most importantly, oligodendrocyte precursor cells isolated from p27-deficient mice divide with a normal cell-cycle time, but many of them go through one or two more divisions compared to wild-type cells before they differentiate.⁽⁴²⁾ Detailed analysis of p27-deficient cells suggests that they are defective in both the timing⁽⁴²⁾ and the TH-dependent effector^(42,43) components of the timer.

The generality of the p27-dependent timing mechanism

It seems likely that p27 plays a similar role in limiting cell proliferation and timing the onset of differentiation during normal development in many cell lineages. p27-deficient mice that lack both copies of the *p27* gene grow approximately one-third larger than normal and contain more cells in all of their organs, apparently as a result of an increase in cell division rather than a decrease in cell death.^(44–46) Mice with one inactive copy of the *p27* gene are one-sixth larger than normal,^(44–46) suggesting that the precise concentration of p27 is important, as would be expected if the accumulation of p27 helps determine when cells stop dividing.

It seems clear, however, that p27 is only a minor component of the timer that operates in precursor cells. p27-deficient oligodendrocyte precursor cells, for example, do stop dividing and differentiate in culture in the presence of saturating concentrations of PDGF and TH, just not at precisely the correct time. What, then, are the other components of the timer? The phenotype of mice deficient in another Cdk inhibitor, p18/Ink4c (p18), is very similar to that of p27-deficient mice, suggesting that p18 may be another component.⁽⁴⁷⁾ In mice deficient in both p27 and p18, some organs are larger than in mice deficient in either protein alone, but, even in these doubly deficient mice, precursor cells still stop dividing and differentiate during development, suggesting that there are additional components that stop the cell cycle in these mice.⁽⁴⁷⁾ It seems likely that some of these other components will turn out to be intracellular cell-cycle promoters, such as cyclins or Cdks, whose levels may decrease to help arrest the cell cycle and initiate differentiation.⁽⁴⁸⁾

Genetic studies indicate that p27-like proteins are also present in *C. elegans*⁽⁴⁹⁾ and *Drosophila*.^(49–51) If the genes encoding these proteins are inactivated by mutation, developing cells in multiple lineages go through one or more extra

divisions before withdrawing from the cell cycle.^(49–51) It seems, therefore, that Cdk inhibitors are involved in stopping the cell cycle at the appropriate time during development of all animals.

The role of thyroid hormone (TH)

There is abundant evidence that TH normally plays an important part in timing oligodendrocyte development. Myelination, for example, is delayed in hypothyroid animals^(52,53) and accelerated in hyperthyroid animals.⁽⁵⁴⁾ Moreover, perinatal hypothyroidism decreases the number of oligodendrocytes that develop in the rat⁽⁵⁵⁾ and mouse⁽⁵⁶⁾ optic nerve, and TH increases the number of oligodendrocytes that develop in cultures of brain cells.^(57,58) While it is clear that TH activates the effector component of the timer in oligodendrocyte precursor cells and so stops division and initiates differentiation, it is not clear how this is brought about. It is likely that TH effects are mediated by the $\beta 1$ isoform of the TH receptor (TR $\beta 1$), as TR $\beta 1$ levels increase in cultured precursor cells in parallel with their sensitivity to the triggering effects of TH, whereas levels of TR α are high from the start, before the cells become sensitive to these effects of TH.⁽⁵⁹⁾ Interestingly, TR $\beta 1$ levels increase with the same kinetics as p27, and, like p27, the levels increase faster at 33°C than at 37°C, suggesting that it is also part of the timer in these cells.⁽⁵⁹⁾

Because TH influences the timing of differentiation in various cell lineages, including muscle, neuronal, and erythrocyte lineages, it is likely that it plays a part in coordinating differentiation in different tissues during development. The rodent thyroid gland becomes active around the time of birth, and TH levels progressively increase during the first weeks of postnatal life.^(52,60,61) It is well suited therefore to coordinate events, such as oligodendrocyte differentiation, that occur during this period. It has been shown recently, for instance, that TH triggers the onset of myelin gene expression in both the Schwann cells and oligodendrocytes that myelinate the peripheral and central branches, respectively, of the VIIIth nerve in neonatal rats, which might help to ensure that nerve conduction rates are high when hearing begins.⁽⁶²⁾ Thus, TH may help coordinate some of the events of postnatal mammalian development, just as it orchestrates the events of amphibian metamorphosis.^(63,64)

An intrinsic maturation program in precursor cells

Visualisation of oligodendrocyte precursor cells in the rat optic nerve at different developmental times shows that they become progressively more complex as the rats age.⁽⁶⁵⁾ Moreover, when these precursor cells are purified at different ages and studied in culture, embryonic precursor cells have a simpler morphology, migrate and divide faster, and divide more times before they differentiate compared to

postnatal precursor cells. If purified embryonic cells are cultured for 10 days or so, however, so that they become equivalent in age to postnatal cells; the two are indistinguishable.⁽⁶⁶⁾ These findings suggest that the precursor cells have an intrinsic maturation program that plays out over time, even when the cells are maintained in a relatively constant environment and in the virtual absence of other cell types. It seems likely, therefore, that the intrinsic timer that controls when these cells stop dividing and differentiate is only one part of a more complex maturation program. It remains a mystery how such an intracellular program is controlled in a way that remains predictable over such a long period of time.

Other types of precursor cells also have an intrinsic maturation program that alters their behaviour and responses to extracellular signals as development proceeds.^(67–70) A dramatic example is seen in multipotential progenitor cells isolated from the rodent embryonic cerebral cortex. When cultured on their own in microwells, these cells display stereotyped division patterns, which appear to be cell autonomous and to change with development.^(69,71) Similar division patterns are observed during neural development in nematodes⁽⁷²⁾ and insects,⁽⁷³⁾ where they reflect the action of both cell-intrinsic developmental programs and cell-cell interactions, which cooperate to generate the diverse cell types that make up the nervous system.

Heterochronic genes

The identification of some of the components of the timer in oligodendrocyte precursor cells does not indicate how the timer works. One needs to understand how the levels of p27 and TRβ1 are controlled so they increase with time in the dividing cells in order to do this.

A powerful approach to determine the molecular basis of developmental timers is to identify genes that are involved specifically in regulating the timing of development, these are the so-called heterochronic genes. This approach, for example, has identified heterochronic genes in the nematode, *C. elegans*.^(74,75) After hatching from the egg, the worm develops through four larval stages, separated by molts, to become a sexually mature adult. During these larval stages, cell division and differentiation occur at specific times in invariant patterns.⁽⁷²⁾ Mutations in heterochronic genes cause stage-specific transformations in larval development, such that cells adopt fates that are characteristic of the lineage but from an earlier or later stage of development.^(74,75) Studies of these transformations suggest that heterochronic genes normally determine the timing of cell fate choice.⁽⁷⁴⁾

Two of these genes *Lin-14* and *Lin-28* encode intracellular proteins that play a part in cell fate determination. Both proteins decrease progressively during larval development,

and this decrease is required for the normal timing of development. The decrease depends on the progressive increase of two small RNAs encoded by a third heterochronic gene *Lin-4*. The small RNAs bind to the mRNAs encoded by *Lin-14* and *Lin-28* and inhibit their translation.^(76–78) It is unclear, however, what controls the increase in the small RNAs during development. Heterochronic genes have been identified in plants, slime moulds, and fungi, but they have yet to be described in mammals.

Conclusion

Most of the timing mechanisms that operate during animal development depend on both intracellular timers and cell-cell interactions. Although there has been a great deal of progress in understanding the cell-cell interactions that regulate development, much less attention has been paid to the intracellular timers, which therefore remain largely mysterious. Understanding how these timers operate is one of the great challenges for developmental biologists in the future.

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