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## Introduction

Understanding how brain complexity develops is one of the challenges of neurobiology. Neurons and glia mature in a succession of transitions between cell states. The orchestration of these processes for different cell types requires the tight regulation of both intrinsic and extrinsic factors. Although each cell type undergoes its own program of differentiation, the timing of differentiation must be coordinated between different cell types. Thus, during brain formation, the regulation of cell number and the precise timing of differentiation require the interplay between intrinsic programs of development and extrinsic factors.

The rodent cerebellum is an attractive model system for the study of cell differentiation, as it consists of a small number of neuronal types that have been morphologically and molecularly well characterized (for review see [Armengol and Sotelo, 1991](#B4) ; [Sotelo, 2004](#B128) ). Three major types of neurons with very different properties are found in the cerebellar cortex: Purkinje cells, granule cells, and GABAergic interneurons (basket, stellate, and Golgi neurons). They differ in their function, morphology, origin, migration routes, and differentiation timing. Neurogenesis in the cerebellum extends over a protracted period of time, between embryonic day 10. 5 and postnatal day 15 in mice, and parallels glial cell proliferation and differentiation. As a consequence, successive steps in the differentiation of several cerebellar cell types are spread over a long time period.

Purkinje cells, some of the largest neurons in the central nervous system, have a central place among the different cell populations of the cerebellar cortex. Purkinje cells are the only efferents of the cerebellar cortex, mainly sending inhibitory projections to the deep cerebellar nuclei, and thus constitute the sole output for all motor coordination and learning from the cerebellar cortex. Many fundamental concepts of modern neuroscience have been established by a focus on this spectacular cell type (see review [Sotelo, 2004](#B128) ). They are the targets of numerous naturally occurring neurological mutations ( [Dusart et al., 2006](#B36) ; [Sajan et al., 2010](#B122) ). 5-hydroxy-methyl-cytosine, a modified nucleotide which function remains enigmatic, was first discovered in Purkinje cell DNA ( [Kriaucionis and Heintz, 2009](#B78) ). Furthermore, Purkinje cells, being the only output of the cerebellar cortex, control the function of the cerebellum. Thus, when Purkinje cells are affected, it is very easy to detect through behavioral phenotypes. Furthermore, their typical location and morphology make them very easy to study. For these reasons, they have been more frequently studied than other neurons and consequently they have been more often detected as implicated in pathologies. The Purkinje cell constitutes thus a classical model to study a number of aspects of neuronal differentiation.

Purkinje cells are the first neuron of the cerebellar cortex to be generated; they are born during the early fetal development (embryonic days 11–13 in mice, [Miale and Sidman, 1961](#B95) ; [Carletti and Rossi, 2008](#B24) ); they end their phase of migration 2 or 3 days before birth and their axons reach their targets in the deep cerebellar nuclei by the end of the fetal period (embryonic day 17, [Eisenman et al., 1991](#B39) ). Between embryonic day 19 and the day of birth, Purkinje cells receive their first climbing fiber synapses ( [Mason et al., 1990](#B91) ; [Chedotal and Sotelo, 1992](#B25) ). However, the development of mouse Purkinje cells occurs largely during the first three postnatal weeks of life, during which Purkinje cells develop dendrites and establish synaptic connections ( [Sotelo and Dusart, 2009](#B130) ). Although it largely coincides with granule cell proliferation and maturation of GABAergic interneuron precursors in neighboring cerebellar layers, the postnatal Purkinje cell maturation process is probably also governed by an intrinsic genetic program.

The aim of this review is to focus on the events that occur during the transition between the first and the second postnatal week and to investigate to which extent these events can be driven by the peak of circulating thyroid hormone that occurs at the end of the first postnatal week in the mouse.

## The End of the First Postnatal Week: A Period of Transition in Purkinje Cell Differentiation

### Dendritic Differentiation

#### Morphological transition at the end of the first postnatal week

Shortly after birth, cerebellar Purkinje cells have a bipolar shape reminiscent of their migratory morphology, with a primary dendrite at their apical pole and an axon at the basal pole. Ramon y Cajal was the first to describe this stage, which he called “ phase of fusiform corpuscle” ( [Cajal, 1926](#B23) ). In 1991, Armengol and Sotelo described two types of fusiform stages in the rat: a simple fusiform corresponding to Cajal's description, and a complex fusiform, presenting a more elaborated dendritic tree in which some collateral branches develop from the primary dendrites ( [Armengol and Sotelo, 1991](#B4) ). Around postnatal day 3 in rats, the primary dendrite has regressed, in parallel with the emergence of numerous perisomatic dendritic processes. At this stage, the Purkinje cells are not polarized, and Cajal called this stage “ stellate.” Several different stellate stages have since been more precisely described ( [Armengol and Sotelo, 1991](#B4) ). At least two different morphological types can be distinguished: a true stellate form, in which the processes are thin, long, and without spines, and an atrophic stage, in which the processes are short. A third intermediate type has also been described in which few (two or three) dendrites emerge from the soma at the apical pole. Interestingly, the small dendrites of this later stage have spines. All these stages can be visualized in parallel during the first postnatal week, as the development of the cerebellar cortex is not synchronous: different developmental stages can coexist within a same lobule and among neighboring Purkinje cells ( [Armengol and Sotelo, 1991](#B4) ). Only video-microscopy would show whether all the Purkinje cells go through these different stages and in which order. However, the study of the proportions of cells in the different stages over time either *in vivo* in rats ( [Armengol and Sotelo, 1991](#B4) ) or *ex vivo* in organotypic cultures ( [Boukhtouche et al., 2006b](#B16) ; [Poulain et al., 2008](#B111) ) suggests that the Purkinje cells pass through these different stages in the described order ( [Sotelo and Dusart, 2009](#B130) ).

At the beginning of the second postnatal week, the Purkinje cells have a single stem segment at their apical pole. From this time, a “ cerebellist” can easily recognize the early form of the future mature Purkinje cell dendritic tree. One particularity of this dendritic tree is that the growth and the ramification occur in the sagittal plane ( [Kaneko et al., 2011](#B70) ). Thus, as described by Cajal, mature Purkinje cell dendritic tree resembles an “ espaliered” fruit tree ( [Cajal, 1911](#B22) ). Larramendi proposed that this transition from multiple dendritic trees to a single one could be the result of the sudden drop of the Purkinje cell nucleus toward the basal pole ( [Larramendi, 1969](#B79) ). In parallel with this transition, Purkinje cell somata merge from multiple irregular rows into a single layer. During the second postnatal week, and up to the end of the third postnatal week, the dendritic tree grows first wider and then taller ( [Berry and Bradley, 1976](#B9) ).

#### Morphological changes: cell-autonomous versus non-cell-autonomous processes

The first dendritic differentiation phases are likely to be driven by intrinsic Purkinje cell developmental programs. The very few purified newborn mouse Purkinje cells (0. 2%) that survive in dissociated culture have smaller dendrites after 21 days *in vitro* than after 4 days *in vitro* , suggesting that the regressive events occur *in vitro* ( [Baptista et al., 1994](#B5) ). In these culture conditions, Purkinje cells never acquire their typical dendritic form. In organotypic culture, Purkinje cells grown in the absence of climbing fiber present similar dendritic developmental phases as those described *in vivo* ( [Boukhtouche et al., 2006b](#B16) ; [Poulain et al., 2008](#B111) ), suggesting that climbing fibers are not necessary for the general sculpting of the dendritic trees. However, in the absence of climbing fibers, the size of the dendritic tree was reduced, due to a decrease in the total number of dendritic segments whereas individual segment lengths were largely unaltered ( [Bradley and Berry, 1976](#B20) ). In contrast, the study of experimental models or mutant mice in which the development of parallel fibers is impeded has revealed that parallel fibers are very important for the growth and planar arrangement of the mature dendritic tree (for review see [Sotelo and Dusart, 2009](#B130) ). Thus, although the first postnatal phases of Purkinje cell dendritic differentiation are likely to be intrinsic, the later phases occurring from the second postnatal week dependent on the environment.

An interesting example of the importance of intrinsic factors has been described for the nuclear receptor RORα, which is deleted in *staggerer* mouse ( [Hamilton et al., 1996](#B58) ). The effect of this mutation on Purkinje cells has been long known ( [Sidman et al., 1962](#B125) ; [Boukhtouche et al., 2006a](#B15) ; [Gold et al., 2007](#B55) ). More recently, the role of RORα in the first stages of Purkinje cell dendritic development has been studied using lentiviral RORα overexpression in organotypic culture of newborn cerebellar slices ( [Boukhtouche et al., 2006b](#B16) ). In this model, 58% of RORα transduced Purkinje cells are already in an atrophic stage after 3 days of culture, while 94% of control Purkinje cells are still in the fusiform stages. After 5 days in culture, 57% of the transduced Purkinje cells are already in a mature stage and present numerous spines. These results indicate that the overexpression of RORα first promotes the regression of the primary dendritic tree and then accelerates dendritic development ( [Boukhtouche et al., 2006b](#B16) ). Later over-expression does not alter Purkinje cell morphology, suggesting a restriction of the developmental function of RORα to early stages ( [Boukhtouche et al., 2006b](#B16) ). Although the growth of the characteristic form of Purkinje cells is dependent of the environment (for review see [Sotelo and Dusart, 2009](#B130) ), the factors that drive its specific form are still unknown.

### Transition of Synaptic Components at the End of the First Postnatal Week

#### General description of the development of synaptic connections on purkinje cells

Purkinje cells are at the center of the cerebellar neuronal circuit. Each Purkinje cell receives up to 200 000 synapses and transmits the integrated signal to the deep nuclei. The innervation of Purkinje cells undergoes profound modifications during the first two postnatal weeks. As it has been recently reviewed, cerebellar developing circuits typically differ substantially from their mature counterparts, which suggests that development may not simply involve synaptic refinement, but rather involves restructuring of key synaptic components and network connections, in a manner reminiscent of metamorphosis ( [van Welie et al., 2011](#B137) ).

Purkinje cells establish functional synapses with deep nuclear neurons between postnatal days 2 and 6 ( [Gardette et al., 1985](#B48) ) and at this time Purkinje cell axons grow many collaterals ( [Gianola et al., 2003](#B54) ). These recurrent axon collaterals underlie facilitating synapses between cerebellar Purkinje cells ( [Orduz and Llano, 2007](#B106) ). Interestingly, the Purkinje–Purkinje connection is asymmetric and provides a robust substrate for propagating waves of activity in the developing, but not adult, cerebellum ( [Watt et al., 2009](#B143) ).

During the first postnatal week, Purkinje cells are contacted by the presynaptic inputs of glutamatergic climbing fibers, the olivocerebellar afferents ( [Mason et al., 1990](#B91) ; [Chedotal and Sotelo, 1992](#B25) , [1993](#B26) ; [Morara et al., 2001](#B98) ). Interestingly during this period, Purkinje cells pass through a phase of climbing fibers multi-innervation ( [Crepel et al., 1976](#B30) ; [Mariani and Changeux, 1981](#B88) ; [Kano and Hashimoto, 2009](#B71) for review). Some mossy fibers that in adult innervate granule cells can also transiently innervate Purkinje cells ( [Mason and Gregory, 1984](#B92) ; [Takeda and Maekawa, 1989](#B134) ; [Kalinovsky et al., 2011](#B69) ). In parallel, some GABAergic axon terminals abut on Purkinje cell somata ( [Sotelo, 2008](#B129) ; [Ichikawa et al., 2011](#B67) ).

At the end of the first postnatal week, the supernumerary climbing fibers begin to be eliminated and the remaining one translocates and synapses onto the proximal dendritic compartment of PCs. The parallel fibers (the axons of granule cells) make synapses at the more distal part of Purkinje cell dendritic tree. In parallel, the different GABAergic interneurons start to innervate specific parts of the Purkinje cell (soma or dendritic tree) ( [Sotelo, 2008](#B129) ; [Ichikawa et al., 2011](#B67) ). Interestingly, during the second postnatal week, considerable fraction of Purkinje somatic spines is succeeded from glutamatergic climbing fibers to GABAergic Basket fibers, in parallel with the switching of postsynaptic receptor phenotypes ( [Ichikawa et al., 2011](#B67) ).

Thus, at the end of the first postnatal week, the development of Purkinje cells is marked by increase of spinogenesis, synaptogenesis with parallel fibers and GABAergic molecular interneurons ( [Sotelo and Dusart, 2009](#B130) ; [van Welie et al., 2011](#B137) ). In parallel with these events that will continue up to the third postnatal week, there is a transition between depolarizing and hyperpolarizing GABA, and the regression of climbing fiber multi-innervation occurs.

#### Transition between depolarizing and hyperpolarizing GABA

Based on two independent sets of experiments (calcium imaging using fura-2 loaded Purkinje cells and perforated-patch recordings) Eilers et al. demonstrated a depolarizing action of GABA on immature Purkinje cells ( [Eilers et al., 2001](#B38) ). They showed that the transition from depolarization to hyperpolarization occurs around postnatal day 6 (P6) in rats. Interestingly, GABA-mediated Ca 2+ signaling was never detected in Purkinje cells with more elaborate dendritic trees (aged P8/9). The depolarizing action of GABA has also been observed in Purkinje cells from 3 day-old mice ( [Rakotomamonjy et al., 2011](#B118) ). Thus Purkinje cells, like some other neuronal populations, exhibit GABA-mediated depolarization during early postnatal stages of life (for review see [Ben-Ari et al., 2007](#B7) ). It is thought that GABA depolarizes immature neurons because of a “ reversed” chloride gradient in a wide range of neuronal types and animal species ( [Ben-Ari et al., 2007](#B7) ). The chloride accumulation in immature neurons can be due either to the early expression of transporters such as the Na-K-Cl co-transporter (NKCC) which accumulates chloride within the cell and/or the lack of expression of co-transporters such as K-Cl transporter (KCC) that export the chloride out of the cell ( [Delpire, 2000](#B32) ). For the majority of the neurons, the expression of KCC2 increases indeed at the end of the first postnatal week in rodents (for review see [Ben-Ari et al., 2007](#B7) ). Surprisingly, Purkinje cells express KCC2 very early during development ( [Mikawa et al., 2002](#B96) ; [Takayama and Inoue, 2007](#B133) ), but the intracellular chloride concentration can be regulated by other factors, such as the expression of WNK family kinases ( [Rinehart et al., 2011](#B120) ). How the intracellular chloride concentration is regulated within immature Purkinje cells is still an open question, and it is, therefore, not understood how the transition between GABA depolarization to hyperpolarization would be triggered in this neuron.

#### Multi-innervation of purkinje cells by climbing fibers during the first postnatal week

In the adult, each Purkinje cell receives synapses from only one climbing fiber (mono-innervation). However, just after birth, around P3 in the rat, several climbing fibers converge and synapse onto the same Purkinje cell body, so that most Purkinje cells are shown to be initially innervated by multiple climbing fibers ( [Crepel et al., 1976](#B30) ; [Mariani and Changeux, 1981](#B88) ). The peak of multi-innervation is around P5, and the regression of the multi-innervation starts at the end of the first postnatal week. In parallel, there is a translocation of the climbing fibers from the soma to the emergent dendritic tree ( [Cajal, 1911](#B22) ) although these two events can be dissociated. From this time until the end of the third week, one climbing fiber input is strengthened while supernumerary climbing fibers are weakened and finally eliminated, resulting in mono-innervation of Purkinje cells in the mature system ( [Hashimoto and Kano, 2003](#B61) ; [Hashimoto et al., 2009](#B60) ). As reviewed recently, the phase of synaptic elimination can be divided in two phases: the first is between P7 and P12, and is independent of the parallel fibers; the second phase depends on parallel fibers [for review see ( [Kano and Hashimoto, 2009](#B71) )]. Interestingly, it has been shown using co-culture and grafting experiments that climbing fiber synapse elimination occurs only during a Purkinje-cell-dependent critical period ( [Gardette et al., 1990](#B47) ) and triggers indelible processes that prevent synapse competition in the mature system ( [Letellier et al., 2007](#B81) , [2009](#B82) ). Whereas numerous actors (such as mGluR1, IGF, BDNF etc.) have been shown to be involved in the second phase of climbing fiber elimination [for review see ( [Kano and Hashimoto, 2009](#B71) )], the mechanisms of the first phase at the transition between multi-innervation and regression of this multi-innervation are less understood. The progressive replacement of full-length TrkB by its truncated form on terminal climbing fibers at the end of the first postnatal week is likely to be involved in this process ( [Sherrard et al., 2009](#B124) ).

### Developmental Purkinje Cell Death Ends at the End of the First Postnatal Week

#### Evidence for developmental purkinje cell death

It took a long time for the importance of developmental cell death to be recognized ( [Ameisen, 2002](#B2) ), and even longer in the case of the Purkinje cell. During the intense phases of cell proliferation, cell death is a counterintuitive notion, and cell death by apoptosis is rapid (about 20-fold faster than proliferation) and difficult to observe directly. Thus, the best way to reveal neuronal cell death during development is to count a population of neurons at different time points. This implies the identification of a neuronal population with specific markers, which were lacking for early Purkinje cells ( [Madalosso et al., 2005](#B85) ; [Dusart et al., 2006](#B36) ). However, numerous early indirect measures suggested that Purkinje cells pass through a phase of programmed cell death during their early development. Some Purkinje cells present pycnotic characteristics in the cerebellar primordium from embryonic day 15–16 (in mouse or chicken; [Bertossi et al., 1986](#B10) ). A small number of TUNEL-positive or activated-Caspase3-positive Purkinje cells were observed in the P3–4 mouse cerebellum ( [Kitao et al., 2004](#B73) ; [Marin-Teva et al., 2004](#B89) ). [Jankowski et al. (2009)](#B68) more precisely described the temporal and spatial distribution of pyknotic Purkinje cells during postnatal mouse development, observing dying Purkinje cells during the first postnatal week with a peak at P3, and very few if any pyknotic Purkinje cells after P9.

Furthermore, overexpression of the anti-apoptotic *bcl-2* gene at various stages of mouse development, and knock-out of the pro-apoptotic *bax* gene provided indirect indications of the existence and periods of programed developmental Purkinje cell death. Counting numbers of adult Purkinje cells shows a 40% increase in the transgenic overexpressing bcl-2 either from embryonic day 13 (E13) and a 27% increase if bcl-2 is overexpressed from P0, as well as a 30% increase for the bax deficient mice compared to wild types ( [Zanjani et al., 1996](#B149) ; [Fan et al., 2001](#B40) ). In contrast, the expression of the human *Bcl-2* gene after P7 (using a *L7-HuBcl2* transgene, selectively expressed in Purkinje cells) did not change the total number of Purkinje cells, suggesting that the period of Purkinje developmental death ends before P7 ( [Goswami et al., 2005](#B56) ).

In organotypic culture, good survival is obtained when the cerebellum is explanted between E19 and P0, or after P10. By contrast, the great majority of the Purkinje cells die by apoptosis when the cultures are prepared from cerebellum between P1 and P8, with a maximum of death observed between P3 and P5 ( [Dusart et al., 1997](#B34) ; [Ghoumari et al., 2000](#B51) , [2002](#B52) ). Thus Purkinje cells are more vulnerable to the culture conditions between P1 and P8. Interestingly during this period, they are also more vulnerable to the noxious effects of alcohol ( [Pierce et al., 1999](#B109) ). This phase of high vulnerability has been proposed to reflect a period of programmed cell death ( [Dusart et al., 2005](#B35) ).

All these results suggest the existence of two periods of programmed Purkinje cell death, a first period during the embryonic life between E13–15, and a second period between P3 and P5 ( [Zanjani et al., 1996](#B149) ). Due to the identification of early Purkinje cell specific markers, it is likely that developmental Purkinje cell death will soon be re-evaluated. It is remarkable that the period of developmental cell death ends for mouse Purkinje cells at the end of the first postnatal week.

#### Purkinje cell survival factors

The program of cellular death is generally engaged by default when a mammalian cell is deprived of survival signals released by other cells ( [Raff et al., 1994](#B117) ). In contrast to other cell types, the vast majority of neurons are not renewed throughout the life of individuals. The neuronal periods of programmed cell death must be, therefore, tightly regulated. During development, neurons depend on trophic factors released by either their targets or their afferents for their survival ( [Oppenheim, 1991](#B104) ). According to the neurotrophic theory, this dependence allows the adjustment of numbers of neurons with their targets or afferent fibers.

During development, Purkinje cells, like other neurons, are dependent for their survival on signals produced by themselves and by other cellular types. The survival of dissociated and purified Purkinje cells *in vitro* increases by 14-fold in the presence of astrocytes, and by 32-fold in the presence of granule cells ( [Baptista et al., 1994](#B5) ). During their maturation, Purkinje cells express the neurotrophin receptors: LNGFR, TrkC, and TrkB ( [Yan and Johnson, 1988](#B148) ; [Cohen-Cory et al., 1989](#B28) ; [Lindholm et al., 1993](#B84) ; [Minichiello and Klein, 1996](#B97) ; [Velier et al., 1997](#B139) ). In parallel, the mRNA encoding neurotrophin-3 (NT-3) is abundant in rat granule cells between P5 and P20 and then replaced by brain-derived neurotrophic factor (BDNF) mRNA ( [Lindholm et al., 1993](#B84) ; [Rocamora et al., 1993](#B121) ; [Gao et al., 1995](#B45) ). In addition, Purkinje cells produce both insulin-like growth factor (IGF-1) and its receptor during their postnatal development ( [Bartlett et al., 1991](#B6) ; [Bondy et al., 1992](#B13) ; [Garcia-Segura et al., 1997](#B46) ). Interestingly, some of the IGF-1 present in the cerebellar cortex is transported by the climbing fibers from inferior olivary neurons to the Purkinje cells ( [Nieto-Bona et al., 1995](#B103) ). Furthermore, the deep nuclear neurons, the target of Purkinje cells, also express IGF-1 ( [Bondy et al., 1992](#B13) ). The mRNA and protein of Glial cell line-derived neurotrophic factor (GDNF) receptor are present in Purkinje cells during development ( [Burazin and Gundlach, 1999](#B21) ).

After the period of programmed cell death, neurons can survive in the absence of trophic factors or their targets: postnatal sympathetic neurons and septo-hippocampal cholinergic neurons gradually lose their dependency on Nerve Growth Factor or on their targets for survival ( [Lazarus et al., 1976](#B80) ; [Sofroniew et al., 1990](#B127) , [1993](#B126) ; [Svendsen et al., 1994](#B132) ; [Orike et al., 2001](#B107) ). Similarly, adult Purkinje cells survive for very long periods in the absence of connections with their main target (deep nuclear neurons) and afferents (climbing fibers) ( [Dusart and Sotelo, 1994](#B37) ; [Morel et al., 2002](#B99) ). From the pattern of expression of trophic factors in the developing cerebellum, it is difficult to propose a model to explain why Purkinje cells become independent from the presence of their targets at the end of the first postnatal week. It is likely that this target independence is the consequence of other processes than trophic factor availability.

Organotypic culture has been successfully used to unravel the role of different molecules ( [Ghoumari et al., 2000](#B51) , [2002](#B52) , [2003](#B49) , [2006](#B50) ; [Rakotomamonjy et al., 2011](#B118) ; [Repici et al., 2011](#B119) ) and of microglial cells ( [Marin-Teva et al., 2004](#B89) ) in developmental Purkinje cell death. During the first week of postnatal life, mouse Purkinje cells show high expression of Caspase-3 mRNA ( [de Bilbao et al., 1999](#B31) ), suggesting that they are competent to die. The role of Lifeguard in Purkinje cell survival has been recently underlined ( [Hurtado de Mendoza et al., 2011](#B64) ). The mechanisms responsible for closing the period of neuronal target dependence have been studied in depth at the level of the apoptotic pathway ( [Putcha et al., 2000](#B112) ; [Orike et al., 2001](#B107) ; [Wright and Deshmukh, 2006](#B146) ; [Wright et al., 2007](#B147) ; [Vaughn and Deshmukh, 2008](#B138) ; [Kole et al., 2011](#B76) ), but it is likely that for the moment, we have only seen the tip of the iceberg.

### Other Processes With a “ Transition Phase” at the End of the First Postnatal Week

#### Glial cell differentiation

The form and cytoskeletal content of Bergmann glia, a cerebellum-specific type of radial glia whose nuclei are in the Purkinje cell layer, change considerably during embryonic and postnatal development. Their content in glial fibrillary acidic protein increases at the end of the first postnatal week ( [Bovolenta et al., 1984](#B18) ).

Oligodendrocyte precursor cells (OPCs) are already present in the cerebellum at embryonic stages ( [Levine et al., 1993](#B83) ), but myelination in the mouse cerebellum begins only at the end of the first postnatal week ( [Foran and Peterson, 1992](#B44) ). Thus, for at least one week, the OPCs are in the presence of Purkinje cell axons but their differentiation is somehow inhibited. The synchronization of cell development is particularly important for cell types that have strong interactions, the case for oligodendrocytes and neurons. OPCs start to differentiate if a mitogenic stimulus is removed or a differentiation stimulus is added, and conversely this cell differentiation is inhibited in the presence of a mitogenic stimulus and the absence of a differentiation stimulus ( [Durand and Raff, 2000](#B33) ). This delayed differentiation period can be reproduced *in vitro* using organotypic culture ( [Bouslama-Oueghlani et al., 2003](#B17) ). However, how OPC and Purkinje cell differentiation are synchronized remains an open question.

#### Axon regeneration

Axonal regeneration in the mammalian CNS is a characteristic of immature neurons which is lost during development ( [Schwab and Bartholdi, 1996](#B123) ; [Dusart et al., 2005](#B35) ). Rodent Purkinje cells progressively lose their ability to regenerate their axons during the first postnatal week: axotomy in the early postnatal period is followed by axonal regeneration, but this capacity for regeneration is absent by the second postnatal week ( [Dusart et al., 1997](#B34) ; [Gianola and Rossi, 2001](#B53) ; [Ghoumari et al., 2002](#B52) ). Thus, Purkinje cells, like the vast majority of CNS neurons, lose their ability to regenerate their cut axons at the end of the first postnatal week.

## A Role for Thyroid Hormone in these Important Transition Processes?

### Effect of Thyroid Hormone on Purkinje Cell Differentiation

Thyroid hormones (TH, which includes both thyroxine, the inactive precursor and 3, 3′, 5-triiodo-L-thyronine (T3), the active deiodinated derivative) are required for proper neurodevelopment ( [Oppenheimer and Schwartz, 1997](#B105) ; [Koibuchi and Chin, 2000](#B74) ; [Bernal, 2007](#B8) ). Whereas the T3 level in serum is remarkably stable in adults, it rapidly increases at birth and peaks during the second week of mouse postnatal development ( [Hadj-Sahraoui et al., 2000](#B57) ). It should be noted however that the distribution of T3 to neurons is a highly regulated process ( [Heuer and Visser, 2009](#B63) ), and that the exact T3 concentration sensed by Purkinje cells is unknown ( [Quignodon et al., 2004](#B115) ). In chicken embryos, it has been proposed that Purkinje cells first gain the ability to convert thyroxine into T3, increasing local signaling at early stages, and only later express the type 3 deiodinase which catabolizes TH ( [Verhoelst et al., 2002](#B141) , [2005](#B140) ). Whether rodent Purkinje cells are also able to metabolize thyroxine and T3 is unknown. In rodents, T3 deficiency results in a number of histological alterations, mainly visible in cerebellum. Purkinje cell alignment is affected, their dendritic arborisations are drastically reduced, and they have fewer synapses. This cellular phenotype can be rescued only if the TH level is restored at an early stage.

T3 acts directly on gene transcription by binding to nuclear receptors (mainly TRα1 and TRβ1), which are both present in Purkinje cells. TRα1 and TRβ1 not only activate transcription upon T3 binding, but also repress gene expression in the absence of ligand. This explains why knock-out mice usually display a mild phenotype, while major neurological disorders result from point mutations making one isoform dominant negative ( [Flamant et al., 2002](#B43) ; [Morte et al., 2002](#B100) ). The genes activated by ligand-bound TR in Purkinje cells remain completely unknown, but some reports have described changes in Purkinje cell gene expression in hypothyroid animals. For example, *Pcp2* (L7) expression is sensitive to TH deficiency, but its down-regulation is only delayed, and reaches normal levels at a later stage, as is the case for several other markers ( [Strait et al., 1992](#B131) ). This observation is consistent with the possibility that T3 only sets the timing of Purkinje cell differentiation. In line with this theory, the elimination of multiple climbing fiber innervation of Purkinje cells occurs 2–3 days later in hypothyroid animals ( [Crepel et al., 1981](#B29) ). More recently, T3 treatment in organotypic cultures was found to accelerate the progression of the early steps of Purkinje cell dendritic differentiation ( [Boukhtouche et al., 2010](#B14) ).

Although this has not been addressed in great detail, it appears that the respective abundance of both T3 receptors changes in Purkinje cells during rodent postnatal development. TRα1 appears to be expressed at birth, and then TRβ1 expression gradually increases, becoming progressively predominant after several weeks ( [Mellstrom et al., 1991](#B94) ; [Bradley et al., 1992](#B19) ; [Wallis et al., 2010](#B142) ). Expressing a dominant-negative mutation of either TRα1 ( [Quignodon et al., 2007](#B116) ; [Fauquier et al., 2011](#B41) ) or TRβ1 ( [Hashimoto et al., 2001](#B59) ; [Portella et al., 2010](#B110) ) is sufficient to affect Purkinje cell differentiation in mice. As the two phenotypes are not identical, a cross-repression between the two receptors appears to be unlikely. The sequential expression of TRα1 and TRβ1 might be a more plausible explanation: in that case, one should be able to demonstrate that Purkinje cell differentiation is impaired at an earlier stage in TRα1 mutant. Interestingly, primary cultures showed that *in vitro* morphological changes promoted by T3 are dependent on TRα1, not TRβ1 ( [Heuer and Mason, 2003](#B62) ). In humans, many germline mutations have been reported for TRβ1 which lead to a complex syndrome without obvious cerebellar disorders, known as resistance to thyroid hormone ( [Weiss and Refetoff, 2000](#B144) ). The first case of a young patient with a TRα1 mutation was recently reported ( [Bochukova et al., 2012](#B12) ). The child's deficit was consistent with those seen in congenital hypothyroidism.

### Cell Autonomous and Non-Autonomous Effects

T3 also exerts an influence on other cerebellar neuronal and glial cell populations, and can thus influence Purkinje cells indirectly. For example, Bergmann glia promote synapse formation between Purkinje cells and GABAergic interneurons ( [Ango et al., 2008](#B3) ). As T3 deficiency affects the differentiation of both Bergmann glia ( [Manzano et al., 2007a](#B86) ) and GABAergic interneurons ( [Manzano et al., 2007b](#B87) ), it has the potential to explain the observed reduction in synaptogenesis ( [Nicholson and Altman, 1972](#B102) ; [Fauquier et al., 2011](#B41) ). T3 deficiency also impairs the production of neurotrophins by granular neurons, which stimulate Purkinje cell differentiation ( [Neveu and Arenas, 1996](#B101) ; [Koibuchi et al., 2001](#B75) ). As the production of several other neurotrophins and growth factors by Purkinje cells changes as they mature, this creates a situation of interdependence that is very difficult to unravel. Finally, oligodendrocyte precursor differentiation, at least *in vitro* , is strictly dependent on T3 ( [Ahlgren et al., 1997](#B1) ; [Durand and Raff, 2000](#B33) ). Myelin formation is retarded by T3 deficiency and accelerated by T3 excess ( [Ibarrola and Rodriguez-Pena, 1997](#B65) ; [Marta et al., 1998](#B90) ; [Billon et al., 2002](#B11) ). The temporal control exerted by T3 on the differentiation of both Purkinje cells and oligodendrocytes provides a simple hypothetical mechanism to ensure that myelination takes place soon after Purkinje axon outgrowth. Cre/loxP technology was used to address whether T3 activates early oligodendrocytes differentiation and myelin formation in a cell-autonomous manner or not. Whereas expression the dominant negative TRα1 mutation only in oligodendrocytes precursors had no visible effect on their differentiation, a delay was observed when the mutation was expressed in GABAergic neurons before P8, or when it was expressed in the astrocyte lineage ( [Picou et al., 2012](#B108) ). The ability of T3 to promote the secretion of several factors at early postnatal stage is thus likely to be determinant in the control exerted on the timing of oligodendrocytes differentiation. In that respect, T3 deficiency could be regarded mainly as a desynchronization of interdependent differentiation processes, whose consequences become rapidly irreversible.

## Conclusions—Perspectives

At the end of the first postnatal week, rodent Purkinje cells undergo profound morphological and electrophysiological changes. They also deeply modify their contacts with their other cellular partners. From a cellular point of view, since these transitions occur when TH levels in serum increase ( [Hadj-Sahraoui et al., 2000](#B57) ), it is tempting to propose that Purkinje cells undergo a metamorphosis: these transitions are reminiscent of amphibian metamorphosis ( [Tata, 2006](#B135) ) and suggest an interesting analogy ( [Kress et al., 2009](#B77) ). The recent concern that some chemicals might have neurotoxic effects due to interference with T3 signaling should bring new attention to this process ( [Ibhazehiebo et al., 2011](#B66) ). Indeed, 1, 2, 5, 6, 9, 10-αHexabromocyclododecane (HBCD) impairs thyroid hormone induced dendrite arborization of Purkinje cells ( [Ibhazehiebo et al., 2011](#B66) ).

With our present knowledge, it is very difficult, however, to decipher whether all the processes occurring at this transition period are only concomitant or are truly related. Furthermore, these processes could also be completely independent of TH. It is not known, for example, whether or not axon regeneration or the factors that contribute to chloride concentration within Purkinje cells are driven by TH. During this transition period, it is important to remember that Purkinje cells synthesize other hormones such as progesterone and estradiol ( [Tsutsui, 2008](#B136) ). Both progesterone and estradiol promote dendritic growth, spinogenesis, and synaptogenesis via their nuclear receptors in developing Purkinje cells ( [Tsutsui, 2008](#B136) ). Indeed, a large fraction of the 50 members of nuclear receptor family are expressed in Purkinje cells ( [Qin et al., 2007](#B113) ) suggesting that several of the small ligand molecules that bind these transcription factors are required locally. We have already described the effect of TRα, TRβ, and RORα mutations on Purkinje cells, but rev-erbα and COUP-TFII also affect either the survival or the development of Purkinje cells ( [Chomez et al., 2000](#B27) ; [Kim et al., 2009](#B72) ). Furthermore, T3 action on early Purkinje cell dendritic differentiation requires the presence of functional RORα ( [Boukhtouche et al., 2010](#B14) ). This raises interesting questions concerning interactions between the TRα and RORα signaling pathways ( [Qiu et al., 2009](#B114) ). Furthermore, the role of RORα in later stages of development is unknown.

Finally, the direct target genes of all these nuclear receptors in Purkinje cells, and how these target genes are connected to build or maintain a functioning Purkinje cells, are currently unknown. A major innovation is the systematic development of models with somatic mutations, mainly based on CRE/loxP technology, which allow the analysis of cell autonomous consequences of these mutations ( [Winter et al., 2009](#B145) ; [Fauquier et al., 2011](#B41) ). It will greatly improve our understanding of transcription factors, like RORα and TRα1, which are already well known to be required for Purkinje cell maturation and/or survival but are expressed in many other cell types as well. In such mutant mice, combining Purkinje cell sorting, sequence-based transcriptome analysis and electrophysiology should lead to a better characterization of the Purkinje cell status and pave the way to a deeper understanding of the molecular mechanisms at work. Whether the molecular mechanisms governing Purkinje cell differentiation and maturation will be transposable to other cell types is difficult to predict. Whatever the extent of their originality, it is likely that this neuron will continue to occupy the minds of many researchers in the future and bring new important results for developmental neurobiology.

Another open question remains: what could be the role of such a metamorphosis in the rodents? In precocial birds (such as chicken), thyroid function is already well developed during the latter part of incubation and hatchings exhibit relatively mature sensory and locomotor capabilities: these birds are able to walk just after hatching. In contrast in altricial birds (such as dove), thyroid function shows little maturation until after hatch as also is the case for sensory, and motor functions: these birds remains in the nest for a while after hatching ( [McNabb, 2006](#B93) ). Interestingly, in contrast to rodents, the increase of TH circulation occurs before birth in sheep, an animal mature enough at birth to walk ( [Fisher et al., 1994](#B42) ). It remains to demonstrate which components of the metamorphosis of the developing cerebellar microcircuit are indeed triggered by thyroid hormone. However, it is tempting to speculate that in cerebellum as well as in other brain parts the burst availability of high levels of T3 might trigger a general process allowing the animal to switch from a developmental program to a mature one adapted to its final environment.

## Conflict of Interest Statement

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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