Thyroid Hormone Induces Cerebellar Purkinje Cell Dendritic Development via the Thyroid Hormone Receptor α1

Heike Heuer and Carol Ann Mason
Department of Pathology, Center for Neurobiology and Behavior, Columbia University, College of Physicians and Surgeons, New York, New York 10032

The thyroid hormone L-3,3′,5-triiodothyronine (T3) plays an important role during cerebellar development. Perinatal T3 deficiency leads to severe cellular perturbations, among them a striking reduction in the growth and branching of Purkinje cell dendritic arborization. The molecular mechanisms underlying these effects are poorly understood. Despite the well documented broad expression of thyroid hormone receptors (TRs), analysis of different TR-deficient mice has failed to provide detailed information about the function of distinct TRs during neuronal development. The cerebellar cell culture systems offer an excellent model by which to study the effects of T3, because differentiation of cerebellar neurons in mixed and purified cultures proceeds in the absence of serum that contains T3. Addition of T3 to cerebellar cultures causes a dramatic increase in Purkinje cell dendrite branching and caliber in a dose- and time-dependent manner. Furthermore, we demonstrate for the first time that T3 acts on Purkinje cells directly through TRα1 expressed on the Purkinje cell and not on the granule cell, the presynaptic partner of Purkinje cells. In contrast, TRβ isoforms are not involved, because Purkinje cells derived from TRβ−/− mice show the same T3 responsiveness as wild-type cells. T3-promoted Purkinje cell differentiation was not mediated via neurotrophins, as suggested previously, because dendritogenesis of Purkinje cells from BDNF−/− mice could be effectively stimulated in vitro by T3 treatment. Furthermore, the effects of T3 observed were not abolished by tyrosine kinase receptor B (TrkB)–IgG, TrkC–IgG, or K252a, agents known to block the actions of neurotrophin. These results indicate that T3 directly affects Purkinje cell differentiation through activation of the TRα1.

Key words: thyroid hormone; T3; TR; Purkinje cell; neurotrophins; BDNF; NT3; K252a; dendrite development; cerebellum

Introduction

The thyroid hormone L-3,3′,5-triiodothyronine (T3) is essential for normal CNS development (Oppenheimer and Schwartz, 1997). T3 deficiency during fetal and neonatal periods causes striking abnormalities in neuronal migration and differentiation, outgrowth of neuronal processes, synaptogenesis, myelination, glial cell proliferation, and neuronal cell death (Legrand, 1984; Anderson, 2001). In humans, if thyroid hormone replacement is not instituted within a critical time window after birth, T3 deficiency leads to the syndrome termed cretinism (Porterfield and Hendrich, 1993).

The development of the murine cerebellum in hypothyroid animals has been extensively examined, especially with regard to Purkinje cell differentiation. Previous studies on hypothyroid rats have revealed abnormal organization of the trunk of the Purkinje cell dendritic tree, persistent hypoplasia of the dendritic field, and a reduction in spine number (Vincent et al., 1982; Legrand, 1984). Proliferation, migration, and differentiation of granule cells, however, appear to be transiently retarded, although parallel fibers of granule cells are shorter and have fewer synaptic contacts with Purkinje cells (Nicholson and Altman, 1972; Lauder, 1977, 1978). Despite the detailed description of cerebellar development under T3 deficiency, the molecular mechanisms underlying T3-induced Purkinje cell morphogenesis are essentially unknown.

The actions of thyroid hormone are primarily mediated by thyroid hormone receptors (TRs), which belong to the family of nuclear hormone receptors containing a DNA-binding and a ligand-binding domain (Mangelsdorf et al., 1995). By interacting with response elements on DNA [thyroid hormone responsive elements (TREs)] and recruiting corepressor or coactivator proteins, TRs can mediate transcriptional control of target genes (Wu and Koenig, 2000; Zhang and Lazar, 2000). Two genes (α and β) have been identified, and these give rise to different splice variants. Among these are the T3 receptors TRα1 and TRβ1, both reported to be expressed during rat cerebellar development (Mellström et al., 1991; Bradley et al., 1992). However, the functions of these receptors with regard to Purkinje cell development are not clear, because obvious cerebellar abnormalities have not been noted in thyroid hormone receptor-deficient mice (Forrest and Vennstrom, 2000; Forrest et al., 2002).

In this study, we used cerebellar cell culture models to analyze the effects of T3 on Purkinje cell dendritic development and to define the receptor involved in this action. Our previous studies indicated that granule cells are potent regulators of Purkinje cell...
development (Baptista et al., 1994; Morrison and Mason, 1998; Shimada et al., 1998). The coculture of purified granule and Purkinje cells allowed us to ask on which cell the key receptors must be expressed: on the Purkinje cell, thus a direct route for T3 stimulation of Purkinje cell dendritic development, or on the granule cell, its presynaptic partner, which would release growth-promoting factors in response to T3 and indirectly mediate the effects of T3 on Purkinje cell differentiation. In addition, we studied the role of BDNF and neurotrophin-3 (NT3) signaling in these cell culture systems, because these neurotrophins have been implicated in collaborations with T3 during cerebellar development (Lindholm et al., 1993; Neveu and Arenas, 1996; Koibuchi et al., 1999).

Materials and Methods

Animals. Experiments were performed either with C57BL/6J mice or with transgenic mice from a timed pregnancy breeding colony under our direction. Breeding pairs of TRβ−/− mice (Forrest et al., 1996), TRα1−/−/− mice (Wikström et al., 1998), and BDNF−/−/− mice (Ernfors et al., 1994) were obtained from The Jackson Laboratory (Bar Harbor, ME). Genotypes of animals were determined using protocols provided by The Jackson Laboratory. For any single Purkinje cell purification experiment, 30 pups at postnatal day 0 (P0)→P1 were used. Granule neurons were purified from 4-d-old pups. Mixed cerebellar primary cultures were prepared as described previously (Morrison and Mason, 1998). Briefly, newborn C57BL/6J mice were killed by decapitation. Cerebella were dissected and resuspended in horse serum, penicillin, streptomycin, and 10% horse serum (Invitrogen). Serum-free medium was composed of Eagle’s basal medium (BME; Invitrogen, Gaithersburg, MD) supplemented with bovine serum albumin (10 mg/ml; Sigma), insulin (5 μg/ml), transferrin (5 ng/ml), sodium selenite (30 ng/ml; Sigma), penicillin-streptomycin (25 μU/ml; Invitrogen), and Sigma 1-8844 supplement (1:100 dilution, resulting in final concentrations of 5 μg/ml insulin, 5 μg/ml transferrin, and 10 μg/ml sodium selenite). Serum-containing medium was composed of BME, gluta mine, penicillin—streptomycin, and 10% horse serum (Invitrogen).

Cerebellar cultures. Mixed cerebellar cultures were prepared as published previously (Baptista et al., 1994; Hatten et al., 1998). Briefly, newborn C57BL/6J mice were killed by decapitation. Cerebella were dissected in PBS, and meninges were removed. Tissue was treated with trypsin (1% in PBS; Worthington, Freehold, NJ) for 5 min at room temperature (RT). After replacing trypsin with Dnase (0.05% in BME; Worthington), cerebella were triturated successively with three fire-polished Pasteur pipettes of decreasing bore sizes. Cells were centrifuged and resuspended in PBS with Dnase, and the cell slurry was passed through a 33 μm nylon mesh filter. Cells were resuspended in horse serum, diluted 1:8 in serum-free medium, and plated at a density of 11 × 10^5 cells/cm^2 in Lab-Tek (Nunc, Naperville, IL) 7-mm-diameter wells (300,000 cells/well in 0.2 ml). Plated cells were allowed to attach overnight, and then the medium was changed to complete serum-free medium that included reagents of interest. Thereafter, medium (including reagents of interest) was replaced every 3–4 d over a 14 d culture period.

Purkinje cell–granule cell cocultures. Cerebellar Purkinje cells and granule neurons were purified as described previously (Baptista et al., 1994; Hatten et al., 1998; Morrison and Mason, 1998). For granule cell purification, dissociated cells were subjected to a Percoll gradient. The dense cell fraction at the interface between the 35 and 60% Percoll phases was collected, and non-neuronal cells were removed by two sequential platings on Petri dishes precoated with 100 μg/ml poly-β-lysine. Nonadherent neuronal cells were collected and plated under serum-free conditions at a density of 11 × 10^5 cells/cm^2 onto poly-β-lysine-coated Lab-Tek wells (Nunc) (300,000 cells per well). These cultures typically consisted of at least 95% granule cells and <5% GFAP-positive cells. Cells with the morphological characteristics of oligodendroglia could not be detected (Hatten, 1985; Baptista et al., 1994).

Purkinje cell–enriched preparations were obtained by passing papain-dissociated cerebellar cells over a 35% Percoll cushion. Non-neuronal cells as well as the majority of containing granule cells were removed by negative immunopanning using anti-GD3 supernatant (HB-8448, from the R24 hybridoma; American Type Culture Collection, Manassas, VA). Purkinje cells were selected by positive immunopanning using anti-Thyl.1.2 (Leinco Technologies, St. Louis, MO) and released from the surface with trypsin. After sedimentation, Purkinje cells were resuspended in serum-free medium and plated at a density of 1 × 10^5 cells/cm^2 (30,000 cells per well in 0.2 ml) on granule cell monolayers prepared 1 d before. Cultures prepared this way were composed of at least 90% calbindin-positive Purkinje cells and <5% GFAP-positive glial cells (Baptista et al., 1994; Morrison and Mason, 1998).

Immunocytochemical analysis of Purkinje cell morphology. Purkinje cells were visualized as published previously (Morrison and Mason, 1998). Briefly, cultures were fixed with 4% paraformaldehyde at room temperature for 60 min and immunostained with a rabbit polyclonal antibody against calcitonin gene–related peptide (CGRP, 50 ng/ml; Swant, Bellinzona, Switzerland) followed by incubation with a cyanine 3 (Cy3)-labeled goat anti-rabbit antibody (1:400; Jackson ImmunoResearch, West Grove, PA). Calbindin-positive Purkinje cells were photographed with an AxioCam digital camera (Zeiss, Oberkochen, Germany) on an Axiohot microscope (63× objective). All experiments were performed at least three times. To quantify dendritic outgrowth, the total area covered by the somatic and dendritic tree on randomly selected Purkinje cells (n = 20) in each experiment was determined by tracing the outlines of the cell and dendritic branches and computing the area using NIH Image software. The values are expressed as mean ± SEM, and results from one experiment are shown graphically. Statistical significance was determined by an unpaired Student’s t test and defined as *p < 0.01 or as **p < 0.001.

In situ hybridization combined with immunocytochemistry. Digoxigenin (Dig)-labeled RNA probes for the detection of TRα1 and TRβ1 transcripts were generated according to standard protocols (Ehrchen et al., 2001). cDNA fragments corresponding to nucleotides 1115–1446 of mouse thyroid hormone receptor α1 (GenBank accession number X51983) and nucleotides 271–495 of mTRβ1 (NM_009380) subcloned in pGEM-T easy vector (Promega, Mannheim, Germany) were kindly provided by S. Friedrichsen (MPI, Hannover, Germany) and used as templates for in vitro transcription. Probes were diluted to a final concentration of 5 μg/ml in hybridization buffer consisting of 50% formamide, 10 mM dithiothreitol, 10% dextran sulfate, 0.6 μM NaCl, 10 mM Tris–HCl, pH 7.4, 1× Denhardt’s solution, 0.1 mg/ml tRNA, 100 μg/ml sonicated salmon sperm DNA, 1 mM EDTA–di–Na, and 10 mM dithiothreitol. In situ hybridization was performed as described previously (Ehrchen et al., 2001) on mixed cerebellar cultures grown on Lab-Tek slides (Nunc) for 14 d in the presence or absence of 1 μM T3. Cultures were fixed in 4% buffered paraformaldehyde for 1 h, treated with 0.1% Triton X-100 in PBS for 10 min, and acetylated (0.25% acetic anhydride in 0.1 M triethanolamine, pH 8.0) for 10 min to reduce nonspecific labeling. After applying hybridization mix, Lab-Tek slides (Nunc) were coveredslipped and incubated in a humid chamber at 37°C for 16 hr. After hybridization, coverslips were removed in 2× SSC (0.3 M NaCl and 0.03 M sodium citrate, pH 7.0), and the cells were treated with RNase A (20 μg/ml) and RNase T1 (1 U/ml) at 37°C for 30 min. Successive washes followed at RT in 1×, 0.5×, and 0.2× SSC for 20 min each and in 0.2× SSC at 65°C for 1 hr. For detection of Dig-labeled probes, cells were exposed overnight to alkaline phosphatase-conjugated anti-digoxigenin antibody (final dilution, 1:5000; Roche Diagnostics, Mannheim, Germany) at 4°C for 16 hr. Phosphatase activity was visualized as described previously (Ehrchen et al., 2001).
mg/ml Hoechst 33258 (Molecular Probes) for 5 min at room temperature. The cultures were examined by bright-field illumination as well as epifluorescence using an AX70 microscope (100× objective; Olympus Optical, Tokyo, Japan) equipped with the appropriate filters and photographed using a DP50 digital camera (Olympus Optical), followed by image manipulation with Adobe Photoshop (Adobe Systems, San Jose, CA).

Results
T3 induces Purkinje cell dendritic development in a dose- and time-dependent manner
As an initial approach to explore the involvement of T3 in Purkinje cell dendrite development, mixed cerebellar cell cultures were prepared; aside from Purkinje cells and granule cells, these cultures contain other interneurons as well as astroglia. They were kept under serum-free conditions with or without addition of T3 at different concentrations. After 14 d in vitro, cultures were fixed and immunostained with an anti-calbindin antibody to visualize Purkinje cells. These neurons respond to rising T3 concentrations with an increase in dendritic length and caliber as well as an increase in spine numbers. In the absence of T3, dendrites can form, but there is a greater preponderance of cells with multipolar dendritic trees, and the branches, including the dendritic stems, are thinner. With T3 added, numerous sculpted spines covered the dendrites, in contrast to the more elongated and filopodial spines on Purkinje cells without T3 (Fig. 1A). To quantify these effects, Purkinje cells from two different wells were randomly selected, and their shape was determined by measuring the area covered by their soma and dendritic tree using NIH Image software. As shown in Figure 1B, dendritic extent increases as T3 concentration rises, reaching an optimal factor of 3 at 3 nM T3.

To determine the temporal response of T3-induced Purkinje cell arborization, cultures were treated with 1 nM T3, a concentration equal to T3 serum levels in adult mice (Friedrichsen et al., 2003), only for a certain period of time. In the first set of experiments, serum-free medium was supplemented with 1 nM T3 at day 10, 7, 5, 3, or 1 in vitro (Fig. 1C, left column). In a complementary set of experiments, cells were plated in the presence of 1 nM T3, but T3 was withdrawn by changing to serum-free medium at day 1, 3, 5, 7, or 10 in vitro (Fig. 1C, right column). All cultures were then analyzed after 14 d in vitro. Quantification of Purkinje cell morphology revealed that continuous T3 treatment during the entire culture period is required for optimal dendritic growth (Fig. 1D), in that even if T3 is withheld for a few days, dendrites did not reach the full extent of differentiation. Likewise, if T3 was withdrawn, dendrites progressed no further than the stage of development at which T3 was removed.

TRα1 and TRβ1 are expressed in Purkinje cells in vitro
Analysis of TR expression pattern on rat brain sections has revealed that TRα1 is highly expressed in granule as well as in Purkinje cells (Bradley et al., 1992). In addition, TRβ1 has been
reported to be expressed in Purkinje cells exactly during a critical time window of dendritic outgrowth and synaptogenesis in vivo (Strait et al., 1991). To determine whether a similar expression pattern occurs in our in vitro cell culture system, we performed colocalization studies by combining nonradioactive in situ hybridization histochemistry for detecting TRα1 and TRβ1 isoforms and immunofluorescence labeling for detecting Purkinje cells. In situ hybridization experiments revealed strong labeling for TRα1 and TRβ1 mRNA in calbindin-positive Purkinje cells (Fig. 2). Therefore, we conclude that Purkinje cells are capable of expressing both TR isoforms in vitro as well as in vivo.

TRα1 but not TRβ isoforms mediate T3 action on Purkinje cell growth

To identify the thyroid hormone receptor responsible for the T3-induced dendritic growth, we took advantage of the availability of mice deficient in specific thyroid hormone receptor isoforms. First, we analyzed Purkinje cell differentiation in mixed cerebellar cultures of mice deficient in all known splice variants of the TRβ gene. Treatment of the TRβ⁻/⁻ mixed cultures with increasing concentrations of T3 led to the same morphological changes as those observed in wild-type Purkinje cells (Fig. 3A). Therefore, we conclude that TRβ receptor isoforms are not involved in mediating T3-promoted dendritogenesis. Next, we examined transgenic mice deficient in the TRα1 receptor isoform but that still express the non-ligand-binding splice variants TRα2 as well as TRΔ-α-2 and TRΔ-α-1. Although the TRα1⁻/⁻-derived Purkinje cells show more extensive dendritic outgrowth in T3-free conditions than TRβ⁻/⁻-deficient Purkinje cells (Fig. 3B) or wild-type controls with no T3 present, TRα1⁻/⁻-derived Purkinje cells do not respond to T3 treatment by further increasing dendritic area, as shown in Figure 3C. This insensitivity to T3 implies that TRα1 is important for mediating the actions of T3 on Purkinje cell dendritic differentiation.

T3 acts directly via TRα1 in Purkinje cells

Because TRα1 has been shown to be widely expressed during cerebellar development, with highest transcript levels in the granule cell layers (Bradley et al., 1989, 1992), it remains to be examined whether T3 influences Purkinje cell dendritic development directly via these receptors on Purkinje cells. Alternatively, granule cells could be the primary target of T3 action, which then in turn would affect Purkinje cell development (e.g., by the synthesis and release of other growth-promoting factors). To address this key question, we took advantage of the cerebellar coculture system developed previously in our laboratory (Baptista et al., 1994). Addition of T3 to cocultures of purified Purkinje cells and purified granule cells led to morphological changes in Purkinje cell dendritic parameters that were the same as those observed in mixed cerebellar cultures (Fig. 4A), including a robust increase in dendritic caliber and spine density. The same effect was observed if granule cells derived from TRα1⁻/⁻ mice were cocultured with wild-type Purkinje cells (Fig. 4B). However, if Purkinje cells were purified from TRα1⁻/⁻ mice and cocultured with either wild-type or TRα1⁻/⁻-derived granule cells, T3 stimulation of Purkinje cell dendritic outgrowth did not occur (Fig. 4C,D). Nonetheless, as already observed in mixed cerebellar cultures, TRα1⁻/⁻-derived Purkinje cells cultured under T3-free conditions showed a greater degree of dendritic differentiation than wild-type Purkinje cells (Fig. 4E). These experiments strongly suggest that T3 acts directly on TRα1 expressed in Purkinje cells to promote dendritic outgrowth.

Figure 2. Both TRα1 and TRβ1 transcripts are present in Purkinje cells in culture. Mixed cerebellar cultures were treated with or without 1 nM T3 for 14 d. In situ hybridization was performed to detect TRα1 (A, left) and TRβ1 (B, left) mRNA. Subsequently, the cultures were immunostained against calbindin (red) and GFAP (green) (right). Hoechst staining revealed localization of cell nuclei (blue). For both receptor isoforms, strong hybridization signals were found in cells with large cell bodies that are calbindin-positive and GFAP-negative, indicating that Purkinje cells (but not astrocytes) clearly express both receptor isoforms. Hybridization experiments with the respective sense probe revealed no specific labeling above background. Scale bars, 25 μm.
Neurotrophin signaling cascades do not interfere with T3-induced dendrite growth

Despite the well acknowledged importance of thyroid hormone for proper brain development, little is known about the genes that are regulated by thyroid hormone receptor interactions. Only a few genes containing a TRE have been identified so far, among them members of the neurotrophin family (Koibuchi and Chin, 2000). In the hypothyroid rat model, although transcript levels of BDNF and NT3 are suppressed during cerebellar development, grafting of cell lines expressing NT3 or BDNF into the rat brain

Figure 3. TRα1 but not TRβ isoform mediates the effects of T3 on Purkinje cell growth. A, Purkinje cells in mixed cerebellar cultures derived from TRβ−/− mice show a dose response to T3 treatment that is similar to that seen for wild-type Purkinje cells. B, With increasing T3 concentrations, Purkinje cell area increases up to threefold. C, Purkinje cells in mixed cultures prepared from TRα1−/− mice do not respond to T3 treatment by increasing dendritic dimensions. Under T3-free conditions, however, they exhibit a higher degree of arborization compared with those from TRβ−/− mice. Scale bar, 50 μm.

Figure 4. T3 acts directly via TRα1 expressed in Purkinje cells. To distinguish which thyroid hormone receptor mediates the effects of T3 on Purkinje cells and whether the required receptors must be on Purkinje cells (direct effect) or on granule cells (indirect effect), coculture of purified cells was performed. Purkinje cells purified from wild-type (WT) or TRα1−/− mice were cultured with granule cells purified from TRα1−/− or wild-type mice and treated with or without 1 nM T3 for 14 d. A, C, Wild-type Purkinje cells showed T3-promoted dendrite growth even when cultured with TRα1−/−-derived granule cells. B, D, However, Purkinje cells derived from TRα1−/− mice did not respond to T3 treatment at all, even in the presence of wild-type granule cells. These observations imply that T3 acts directly via TRα1 in Purkinje cells. E, Morphological analysis clearly indicates that TRα1−/−-derived Purkinje cells cultured under T3-free conditions showed a higher degree of dendritic differentiation than wild-type Purkinje cells. Data were derived from 20 randomly selected Purkinje cells in each group. Each column and error bar represents the mean ± SEM. *p < 0.01; **p < 0.001 (by Student’s t test). PC, Purkinje cells; GC, granule cells. Scale bar, 40 μm.

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ameliorates cerebellar defects caused by hypothyroidism (Neveu and Arenas, 1996). Because these findings implicate neurotrophins as a direct target of thyroid hormone action, we studied the putative role of neurotrophins with regard to T3-induced Purkinje cell dendritogenesis. In the first set of experiments, mixed cerebellar cultures were treated with BDNF (10 ng/ml) for 14 d in the presence or absence of 1 nM T3. After incubation for 14 d, Purkinje cell dendritogenesis was analyzed after immunostaining with anti-calbindin. Addition of exogenous BDNF did not further enhance T3-dependent dendritic arborization. BDNF+/- mice showed the same T3-induced increase in dendritic parameters as Purkinje cells prepared from BDNF +/- mice. To block the actions of neurotrophin, mixed cerebellar cultures were incubated with 1 nM T3 in the presence of 25 μg/ml TrkB–Fc or TrkC–Fc or with Fc alone (control). None of these agents perturbed T3-promoted dendritogenesis in Purkinje cells. Data were derived from 20 randomly selected Purkinje cells in each group. Each column and error bar represents the mean ± SEM.

In another approach, we treated mixed cerebellar cultures for 14 d with inhibitors of neurotrophin signaling. Soluble TrkB–Fc or TrkC–Fc fusion proteins (25 μg/ml) that consist of TrkB or TrkC extracellular domain fused to the human IgG heavy chain act as scavengers for released neurotrophins (Shelton et al., 1995). However, neither of these agents prevented T3-induced dendritic growth of Purkinje cells (Fig. 5C). To block the actions of neurotrophin, mixed cerebellar cultures were also treated for 14 d with K252a, a Trk-specific inhibitor. Control experiments were performed in the presence of vehicle with or without 1 nM T3. Addition of K252a (20 nM) did not abolish the effects of T3 on Purkinje cells. However, Purkinje cells develop long, thin, and highly branched dendrites. This effect could be quantified by determining the perimeter as well as the area of these cells. Compared with cultures (0 nM T3) that were only treated with vehicle (DMSO), the area of Purkinje cells increased 2.5-fold after T3 treatment. The perimeter, however, increased up to 4.6-fold if both agents (20 nM K252a and 1 nM T3) were present, thus indicating that dendrites indeed were longer and thinner compared with cells exposed to T3 alone. Adding 2 mM K252a or 20 nM K252b, a nonspecific inhibitor, had no effect on Purkinje cell morphology. Scale bar, 80 μm.
neurotrophin signaling, suggesting that, at least in our

in vitro experiment, neurotrophin signaling does not collaborate with

brain development. Moreover, changes could be observed in the presence of the protein kinase C blocker Gö6976 (Metzger and Kapfhammer, 2000) (data not shown), we propose that this effect of K252a is not attributable to a specific interaction with Trk receptors. In conclusion, we were not able to inhibit T3-induced dendrite expansion by blocking neurotrophin signaling, suggesting that, at least in our in vitro culture system, neurotrophin signaling does not collaborate with thyroid hormone signaling to stimulate Purkinje cell dendritic growth.

Discussion

Dendritic development is thought to be regulated primarily by the presence of afferents, because timing of afferent innervation and synapse formation coincides with the period of maximum growth and dendritic arborization. In addition, dendritic morphology is also thought to be determined by the number and type of synaptic inputs that a neuron receives. Granule cells and their parallel fiber axons, more than climbing fiber axons, play a key role in instigating Purkinje cell differentiation, studied most precisely in purified cocultures (Baptista et al., 1994). However, in vivo, when deprived of parallel fibers, Purkinje cells are capable of developing dendrites and spines, indicating that intrinsic as well as nonaffferent derived extrinsic factors contribute to the tightly regulated process of dendritogenesis (Scott and Luo, 2001; Whitford et al., 2002; Wong and Ghosh, 2002)

As an extrinsic factor, thyroid hormone strongly affects dendritogenesis of Purkinje cells. The absence of thyroid hormone during the first postnatal weeks causes profound Purkinje cell hypoplasia and also delays granule cell differentiation. As a consequence, humans as well as rodents suffering from congenital hypothyroidism develop neurological symptoms such as ataxia and poor fine motor movement, emphasizing the importance of proper thyroid hormone supply during cerebellar development. The mechanisms underlying these thyroid hormone effects, however, remain poorly understood. One question is whether thyroid hormone acts on Purkinje cell development directly or instead acts on granule cells, and thus indirectly on Purkinje cells. The generation of mice deficient in thyroid hormone receptors, singly and in combination, has not provided any additional insight with regard to cerebellar action of T3 (O’Shea and Williams, 2002). Neither Trb-/- animals that are deficient in all isoforms derived from the Trb locus (Forrest et al., 1996) nor mice lacking the Trα1 receptor isoform (Wikström et al., 1998) display any gross cerebellar abnormalities. Surprisingly, even Purkinje cells in Trα1-/-/β-/- mice appear to be quite unaffected (Göthe et al., 1999). However, in all of these receptor-deficient mouse models, the thyroid status is imbalanced and shifted to either hypothyroidism (e.g., in the case of Trα1-/-) or hyperthyroidism (e.g., in the case of Trb-/-). Even tissue-specific hyper/hypothyroidism can be observed as in the case of Trα2-/- mice, which overexpress Trα1 (Saltó et al., 2001). Because the thyroid axis is regulated in such a complex manner, compensatory pathways may take place in these TR mouse models that mask the function of each receptor for proper brain development.

Our approach to studying the effects of T3 in primary cerebellar cell cultures offers the opportunity to unravel specific pathways of T3 action, because the thyroid status can be exactly defined. Furthermore, cells from different thyroid hormone receptor-deficient mouse models can be combined with wild-type cells in vitro and checked for thyroid hormone response. Even direct and indirect effects of T3 can be distinguished, because coculture experiments with purified neuronal cell types allow identification of important receptor isoforms on cellular levels.

By taking advantage of this in vitro system, we focused on the effects of T3 with regard to Purkinje cell dendritogenesis. Addition of T3 to otherwise complete athyroid mixed cerebellar cell cultures revealed a dramatic increase in dendritic outgrowth of Purkinje cells in a dose-dependent manner, an observation that is in agreement with results published recently (Kimura-Kuroda et al., 2002). Purkinje cell soma and dendritic tree dimensions increase up to threefold within 14 d in vitro, reaching a plateau at 3 nM T3. This potent effect is observed only during continuous T3 exposure, indicating that the function of T3 is not restricted to serving as a molecular switch to initiate Purkinje cell maturation.

Which receptor isoform is involved in this process? Theoretically, both receptors could participate, because both receptors are expressed during cerebellar development. From our results, however, we can conclude that TRβ1, although expressed in Purkinje cells during the peak of dendrite formation (Strait et al., 1991), is not essential. Our data emphasize a pivotal role for Trα1, a receptor isoform highly expressed in granule as well as in Purkinje cells (Bradley et al., 1992). For T3-promoted dendritogenesis, however, Trα1 must be expressed in Purkinje cells, because the absence of Trα1 in granule cells does not interfere with the Purkinje cell response. Therefore, we can exclude an indirect effect by which T3 acts on granule cells, which, in turn, would release growth factors to control Purkinje cell dendritogenesis. In contrast, our results clearly demonstrate for the first time that Purkinje cells are a direct target of T3 action.

Why is only Trα1 but not Trβ1 essential for promoting T3-induced dendritogenesis? At first glance, this finding is surprising, because both receptors are closely related in their DNA- and T3-binding domain and can transactivate through similar TREs. However, differences in the conformation of Trα1 and Trβ could result in differential recognition of TREs or interaction with cofactors (Forrest et al., 2002). Indeed, several studies have postulated isoform-specific functions. In the kidney, T3-dependent regulation of deiodinase type 1 expression relies solely on Trβ despite the expression of both TRs (Amma et al., 2001). Alternatively, it has been proposed that TRs are essentially interchangeable, and that it is the total level of TR expression in a cell that determines its response to TH (Yen et al., 2003). In line with this argument, one has to assume that Trα1 is the major functional isoform in Purkinje cells, because Trβ cannot compensate for the absence of Trα1.

Signaling of Trα1 seems to be complex and involves at least two different mechanisms. Binding of T3 to Trα1 can obviously promote dendritic arborization, most likely by inducing expression of genes involved in this process. Without T3, however, unliganded Trα1 can also bind to DNA and act as a repressor of gene transcription. In our cell culture system, unliganded Trα1 appears to repress dendrite formation, because Purkinje cells lacking Trα1 show a higher degree of dendritic arborization than wild-type Purkinje cells under T3-free culture conditions. This in vitro observation is in agreement with studies by Morte et al. (2003), who showed an insensitivity of Purkinje cells in Trα1-/- animals to drug-induced hypothyroidism. Additionally, Purkinje cells are abnormally developed in mice expressing dominant-
negative forms of TR that lack T3 binding capacity (Hashimoto et al., 2001; Kaneshige et al., 2001). Together, these findings imply that thyroid hormone receptors can regulate transcription by both T3-dependent and T3-independent means. Furthermore, absence of T3-induced derepression of gene transcription might be the primary reason for the severe cerebellar abnormalities found under hypothyroid conditions (Göthe et al., 1999).

Which genes are positively or negatively regulated by TRα1/−/− action and mediate T3-induced dendrite formation? Neurotrophins appeared to be attractive candidates, because their expression patterns have been shown to be tightly regulated by T3 and in the hypothyroid cerebellum (e.g., BDNF mRNA levels are suppressed) (Koibuchi et al., 1999). Moreover, grafting cell lines expressing BDNF or NT3 into the fourth ventricle in part, but not completely, prevents hypothyroidism-induced abnormal cerebellar development (Neveu and Arenas, 1996). In addition, BDNF has been shown to regulate the growth and branching of cortical pyramidal neuron dendrites (McAllister et al., 1995; Miller and Kaplan, 2003). Thus, a similar role of BDNF in Purkinje cell development was an obvious assumption supported by the fact that BDNF−/− mice exhibit a decreased arborization of Purkinje cell dendrites and a delayed migration of granule cells (Schwartz et al., 1997), a phenotype similar to that in hypothyroid rodents. For these reasons, we analyzed the role of neurotrophin signaling with regard to T3-induced Purkinje cell dendritogenesis in our cell culture system. Unexpectedly, Purkinje cells derived from BDNF−/− mice displayed the same T3 responsiveness as wild-type cells. In addition, blocking the actions of neurotrophin or adding BDNF did not affect T3-induced dendritogenesis, particularly in terms of the extent and caliber of dendritic pattern, similar to our findings on the role of BDNF in Purkinje cell dendritogenesis (Morrison and Mason, 1998). Blocking BDNF, however, did affect spine morphology, again, as we found previously (Shimada et al., 1998). Therefore, we conclude that neurotrophin signaling, although crucial for granule cell development (Borghesani et al., 2002), does not directly participate in T3-induced Purkinje cell dendritogenesis.

The challenge still ahead is to unravel the mechanisms by which T3 promotes dendrite formation. The experiments presented here using Purkinje cells deficient in the TRα1 isoform indicate that T3 exerts its powerful effect directly on Purkinje cells, whereas the actions of T3 on granule cells appear to have only minor consequences for Purkinje cell dendritogenesis. We cannot exclude that granule cells release other growth factors or signaling molecules in response to T3 that support Purkinje cell differentiation. However, to boost dendritogenesis, T3 signaling must be activated within the Purkinje cell.

In conclusion, understanding dendritic development requires knowledge of the mechanisms that coordinate the overall growth and detailed patterning of Purkinje cell arborization. Elucidating the actions of thyroid hormone during Purkinje cell dendrite formation provides an excellent paradigm to explore the intracellular signaling pathways that mediate this process. Here we have shown that the interaction of T3 with TRα1 in Purkinje cells defines one crucial step in proper dendrite formation. Based on this information, future attention should be focused on identifying downstream genes regulated by TRα1.

References


