The Intracellular Domain of Teneurin-1 Induces the Activity of Microphthalmia-associated Transcription Factor (MITF) by Binding to Transcriptional Repressor HINT1

Teneurins are large type II transmembrane proteins that are necessary for the normal development of the CNS. Although many studies highlight the significance of teneurins, especially during development, there is only limited information known about the molecular mechanisms of function. Previous studies have shown that the N-terminal intracellular domain (ICD) of teneurins can be cleaved at the membrane and subsequently translocates to the nucleus, where it can influence gene transcription. Because teneurin ICDs do not contain any intrinsic DNA binding sequences, interaction partners are required to affect transcription. Here, we identified histidine triad nucletide binding protein 1 (HINT1) as a human teneurin-1 ICD interaction partner in a yeast two-hybrid screen. This interaction was confirmed in human cells, where HINT1 is known to inhibit the transcription of target genes by directly binding to transcription factors at the promoter. In a whole transcriptome analysis of BS149 glioblastoma cells overexpressing the teneurin-1 ICD, several microphthalmia-associated transcription factor (MITF) target genes were found to be up-regulated. Directly comparing the transcriptomes of MITF versus TENI-ICD-over-expressing BS149 cells revealed 42 co-regulated genes, including glycoprotein non-metastatic b (GPNMB). Using real-time quantitative PCR to detect endogenous GPNMB expression upon overexpression of MITF and HINT1 as well as promoter reporter assays using GPNMB promoter constructs, we could demonstrate that the teneurin-1 ICD binds HINT1, thus switching on MITF-dependent transcription of GPNMB.

Significance:

This is a novel mechanism for a teneurin-mediated transcriptional control.

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1 The data reported in this paper have been deposited in the Gene Expression Omnibus (GEO) database, www.ncbi.nlm.nih.gov/geo (accession nos. 61704 and 61705). 

2 The abbreviations used are: ECD, extracellular domain; ICD, intracellular domain; qPCR, quantitative PCR; IPA, ingenuity pathway analysis; GBD, glucocorticoid binding domain; Dox, doxycycline; Dex, dexamethasone; SEAP, secreted embryonic alkaline phosphatase; tet, tetracycline; MITF, microphthalmia-associated transcription factor.

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tion sites, and polyproline-rich regions shown to bind SH3 domain-containing proteins like CAP/Ponsin (29, 30).

Teneurins were first identified as pair-rule genes (2, 3), which was curious because pair-rule genes were known to be transcription factors rather than transmembrane proteins (5). Although this evidence was recently refuted (31), it still paved the way to implicate the teneurin ICDs in transcriptional regulation. Regulated intramembrane proteolysis by either site-2 protease, signal peptide peptidase, or signal peptide peptidase-like proteases is predicted to release the ICD (24) before it translocates to the nucleus, as shown by several in vitro and in vivo studies (14, 29, 32). In the nucleus, the teneurin-2 ICD affects zic-mediated transcription and is localized to PML bodies, which are so-called transcriptional hotspots (32). The teneurin-1 ICD can bind to MBD1, a transcriptional repressor (29). Interaction partners are crucial for the ICDs to influence transcriptional regulation because they do not contain any intrinsic DNA binding sequences.

Although the teneurin ICDs are strongly implicated in transcriptional regulation, target genes as well as mechanisms have yet to be elucidated. In this study, we are investigating the transcriptional activity of the human teneurin-1 ICD, which will be referred to as simply TEN1-ICD hereafter. We performed two unbiased screens to identify 1) interacting proteins and 2) target genes involved in the transcriptional control of these target genes. We will show how the ICD can influence transcription via HINT1, a transcriptional repressor that directly binds to transcription factors at the promoters of their target genes (33, 34).

**EXPERIMENTAL PROCEDURES**

Cloning—All constructs were prepared by classical cloning procedures and verified by sequencing. TEN1-ICD was cloned from human adult brain cDNA (20), HINT1 from pPR3-N-HINT1 (Dualsystems), eGFP-His from pcDNA3-EgFP (Addgene), RFP-HA from pQCXIX-RFP (Clontech), CFP-MYC from pECFP1-C1 (Addgene), and MITF from pCMV6-MITF variant 4 (Origene). The following constructs were prepared in pcDNA3 (Invitrogen): HINT1-MYC, TEN1-ICD-HA, HINT1-CFP-MYC, RFP-HA, and MITF-RFP-HA (note that all tags are C-terminal). For inducible overexpression studies with the TEN1-ICD, we used the highly predictable and tightly regulated gene expression system described by Anastassiadis et al. (35) using the following plasmids: pirtetR-GBD as tetracycline (tet) activator plasmid and eGFP-His and TEN1-ICD-eGFP-His in the tet promoter plasmid ptetO as described previously (35). For the yeast two-hybrid screen, TEN1-ICD was cloned into pDHBl (Dualsystems). For promoter reporter studies, the GPNMB and GPNMB ΔM-box promoters were cloned into pSEAP2basic (Clontech). Promoter sequences GPNMB (1096 bp) and GPNMB ΔM-box (144 bp) as described (36) were cloned from HT1080 cell whole genomic DNA and extracted with the DNeasy tissue kit (Qiagen) using the following primers: ACTAGCTAGGCGCAACTTTTTCTGCATACTCTG (forward) and ACTACTCGAGCATCTGGTGTGCCTCCCTC (reverse) and ATGCTAGGAACTTGGAGAACAGATCGGAC (forward) and ATCTCGAGCAGTCTCCCTGCCATCTGGTGTGCCTCCCTC (reverse), respectively.

**Cell Culture**—BS149, SH-SY5Y, U373, LN229, MO95S, U343MG, and T98G cells (kind gift of the Hemmings laboratory at the Friedrich Miescher Institute for Biomedical Research) and COS-7 (ATCC) cells were cultured in DMEM containing 10% fetal bovine serum (FBS), 100 mg/liter penicillin, and 100 mg/liter streptomycin. Genes were either overexpressed by transient transfection or by a modified tet system kindly provided by Gerrit Fischedick of the Schöler laboratory at the Max Planck Institute for Molecular Biomedicine (35). Cells were transfected with jetPEI (Polyplus). BS149 cells were first transfected with pirtetR-GBD and made stable by 0.5 μg/μl puromycin selection and then, after further transfection with either ptetO-eGFP-His or ptetO-TEN1-ICD-eGFP-His, by 150 μg/μl hygromycin selection. The tet system was induced by adding 10−7 M dexamethasone (Dex) 100% ethanol and 1 μg/ml doxycycline (Dox) in Milli-Q water to the medium.

**Yeast Two-hybrid Screen**—The yeast two-hybrid screen was performed using the DUALHunter starter kit (Dualsystems) and following its manual. Prey proteins were fished with the bait protein TEN1-ICD, out of the normalized human fetal brain cDNA (NubG-X) library (Dualsystems).

Proximity Ligation Assay and Immunocytochemistry—The proximity ligation assay was performed using the DUOl ink system (Sigma-Aldrich) and following the appropriate protocol provided by the manufacturer. COS-7 cells were transfected with either HINT1-MYC as negative control or both HINT1-MYC and TEN1-ICD-HA to identify an interaction. For detection, mouse monoclonal α-HA (42F13, used as hybridoma supernatant) and rabbit polyclonal α-MYC (Abcam) antibodies were used. For immunocytochemistry, parallel cultures were stained with the same primary antibodies and Alexa Fluor 488 goat anti-rabbit and Alexa Fluor 568 goat anti-mouse (both from Life Technologies) as secondary antibodies. Cells were viewed with an Axioskop 50 microscope (Zeiss), and pictures were taken with an ORCA-ER digital camera (Hamamatsu).

**Real-time qPCR**—To determine endogenous transcript levels of teneurin-1 in normal tissues, brain total RNA, fetal brain RNA, and cerebellum RNA were acquired (Clontech). For the glioblastoma cell line screen of endogenous teneurin-1 expression, 1 × 106 cells were used for RNA extraction. Experiments were performed in biological triplicates, where for each cell line RNAs were extracted from independent cultures on three consecutive days. All overexpressing cells were FACS-sorted directly into RLT lysis buffer (Qiagen) at a 3:1 volume ratio of lysis buffer to cells in PBS, 24 h after induction or transfection. Total RNA was extracted with an RNeasy Mini kit and QIASHredder columns (both from Qiagen) and reverse-transcribed into cDNA, using random hexamer primers with the high capacity cDNA reverse transcription kit (Applied Biosystems).

For real-time qPCR, gene-specific primers (Table 1) were designed, and all data were normalized to TBP, using Platinum SYBR Green qPCR SuperMix-UDG with ROX (Invitrogen) on a StepOnePlus real-time PCR system (Applied Biosystems). Endogenous levels were calculated by using the comparative Ct method (37).

Western Blot—BS149 cells were induced or kept uninduced as a negative control at 70% confluence on a 3.5-cm plate and...
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TABLE 1
Real-time qPCR primers
shown are human primers for all real-time qPCR experiments, using Platinum SYBR Green qPCR SuperMix-UDG with ROX.

| Target       | Forward primer | Reverse primer |
|--------------|----------------|----------------|
| Teneurin-1 ICD | CTACCTGCGTTTCTCCAAACCC | CCAGAATTGCTTCTTCCAGGA |
| Teneurin-1 ECD | GCCATATTCTGTGTTGCTCAA | TCTCAATCTCTGAGTGAC |
| Teneurin-1 ICD-L | GAACACACTTTCTCCGATCC | CGAGACCCCTGCACTTCTG |
| GPNMB         | AAGTGAGAGGGTCTCATCTGGAAGG | TGGCGATGCGTCTGCTTCTG |
| SCARB1        | ATTAAGCCTCAGCTGCGAACGC | GCCCTCACTGATCTCACCC |
| EDNRB         | CTGCTGCGCTTCTGCGTAC | GCTCCGATTGCCCATCTCT |
| SLC1A4        | CAGGCCCTTCCCTCTCTTGA | GCCGGCAATGGGAGAATAC |
| SEMA6A        | ACATCTGCTGCTAGAGCATCA | TCTCAGTGCTGCTACGTGCCC |
| ERBB3         | GCTCGTGTGACCCACTGAACT | GGGTGCGACGAAAGCATTT |
| CHL1          | ACCACATTTTTCTGGAGTCAAGG | TCAATGCGATATCCTGGGGTA |
| TRP           | TGGACAGGGGCCAGAAGGAG | CGACTCCAGTCTCCACAC |

harvested after 24 h directly in sample buffer, containing β-mercaptoethanol. Samples were run on a 10% polyacrylamide gel and blotted on a PVDF membrane. Protein bands were detected by a mouse monoclonal α-GFP antibody (Roche Applied Science) in 5% skim milk in 1× TBS-Tween 20 (0.05%), a mouse monoclonal α-vinculin antibody (Sigma-Aldrich) as a loading control, and a goat α-mouse secondary horseradish peroxidase-conjugated antibody (Invitrogen) with SuperSignal West Dura Chemiluminescent Substrate (Thermo Scientific).

Microarray—For the whole transcriptome analysis of TEN1-ICD overexpression, the stable BS149 cell lines with pirtetR-GBD/petoto-eGFP-His or pirtetR-GBD/petoto-TEN1-ICD-eGFP-His were split into three 10-cm Petri dishes each. The triplicate cell lines were cultured once before induction with Dex and Dox. For the whole transcriptome analysis of MITF, cells were transiently transfected in triplicates with either pcDNA3-RFP-HA or pcDNA3-MITF-RFP-HA. The overexpressing cells were then FACS-sorted directly into RLT lysis buffer (Qiagen) at a 3:1 volume ratio of lysis buffer to cells in PBS, 24 h after induction or transfection. Total RNA was extracted with an RNeasy Mini kit and QIAshredder columns in PBS, 24 h after induction or transfection. SEAP activity measurements were taken 24 h post-transfection by using the chemiluminescent SEAP reagent (Clontech). All values were measured by the Mithras LB940 Luminometer (Berthold Technologies).

Ingenuity Pathway Analysis (IPA)—Normalized unscaled expression values for the genes passing -fold change cut-off = 1.5 and a Benjamini-Hochberg corrected p value = 0.05 filter were up-loaded into IPA (Qiagen) and run through the pathway analysis. The upstream analysis was filtered for transcriptional regulators and sorted by activation z-score. The MITF network was created via the My Pathways option, only allowing direct connections related to transcriptional regulation.

SEAP Reporter Assays—Promoter reporter assays were performed by cotransfecting BS149 cells with 1 μg of pSEAP2basic-promoter constructs and either 0.5 μg of empty pcDNA3 vector, 0.5 μg of MITF, 0.05 μg of MITF plus 0.45 μg of empty pcDNA3 vector, or 0.05 μg MITF plus 0.45 μg of TEN1-ICD and 150 ng of the Metridia luciferase pMetLuc reporter vector (Clontech) for the normalization of the transfection efficiency. SEAP activity measurements were taken 24 h post-transfection by using the chemiluminescent SEAP reporter gene assay (Roche Applied Science). The Metridia luciferase assay for normalization was performed with the Ready-to-Glow secreted luciferase reporter assay (Clontech). All values were measured by the Mithras LB940 Luminometer (Berthold Technologies).

Statistical Analysis—All grouped data are means ± S.D. Statistical analysis was completed using GraphPad InStat version 3.05. Differences between two groups were evaluated using a two-tailed Student’s t test for parametric data or a Mann-Whitney U test for nonparametric data. Multiple comparisons were performed using one-way analysis of variance. Values of p < 0.05 were considered statistically significant.

RESULTS
Identification of TEN1-ICD Interaction Partners—To identify new interaction partners, we performed a yeast two-hybrid screen using the TEN1-ICD as a membrane-anchored bait protein fishing for prey encoded by a human fetal brain cDNA library. The benefit of this method is that baits that might activate transcription independently of a prey are prevented from enterin the nucleus, making them nevertheless usable for a screen. Furthermore, the readout is based on the split ubiquitin method, in which an interaction of the bait and prey proteins also joins the N- and C-terminal parts of split ubiquitin. A ubiquitin-specific protease then releases the artificial transcription factor LexA-VP16 to switch on the two reporters HIS3 and LacZ (Fig. 1A). The yeast strain NY51 can thus grow on a defined minimal medium lacking histidine only when an inter-
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Several MITF Target Genes Are Up-regulated in TEN1-ICD-overexpressing Cells—In parallel, we wanted to identify a system to study how the overexpression of the TEN1-ICD influences transcription, assuming that cells with endogenous teneurin-1 expression also have the machinery to respond to teneurin-1 signaling. To get started, we performed a qPCR screen identifying glioblastoma cell lines that express endogenous teneurin-1 (Fig. 2A, left). We chose the BS149 cell line for our further studies because it has an endogenous expression of teneurin-1 and excellent transfection efficiency. We also compared the levels of TEN-1 transcripts in BS149 cells to tissues known to express teneurins, such as the brain. Whereas total adult brain RNA contained about 18.3-fold more teneurin-1 transcripts, the levels in fetal brain and adult cerebellum were 50.5- and 181.7-fold higher, respectively, than in BS149 cells (Fig. 2A, right).

To study the influence of teneurin-1 on transcriptional regulation in BS149 cells, we needed an inducible method to overexpress the TEN1-ICD transiently. Thus, we used a modified and improved tet system, which eliminates the leakiness of the promoter (35). Two separate vectors are required to make the cell line stable (Fig. 2B). One contains the tet-activating domain fused to a glucocorticoid binding domain (GBD). The other contains the tetO operator sequences directly upstream of a CMV promoter and the gene to be overexpressed. The tet-activating domain is continuously expressed and bound to HSP90 in the cytosol via the GBD. Adding Dex releases the fusion protein from HSP90 and subsequently binds the tetO-CMV promoter, switching on gene expression due to the addition of Dox. This system allows for well controlled induction of transcripts in a physiological range. First, we were able to show the functionality of the system by Western blot of cell extracts of the stable BS149 cells induced by Dex/Dox addition. TEN1-ICD-eGFP-His and eGFP-His fusion proteins were detected only when the modified tet system was induced (Fig. 2C). qPCR confirmed the specific overexpression of the TEN1-ICD-eGFP-His fusion construct, resulting in a 214.9-fold increase, compared with endogenous teneurin-1 expression, on a transcript level (Fig. 2D). Although this might appear as very strong overexpression, this is not more than is observed in total RNA of adult cerebellum. Thus, at least on a transcript level, the overexpressed TEN1-ICD is in a physiological range.

To show the effects on transcriptional regulation in BS149 cells, we overexpressed the TEN1-ICD-eGFP-His construct by induction of the tet system and compared it with the overexpression of the eGFP-His construct in a whole transcriptome microarray analysis. For more rigorous results, we compared biological triplicates and separately induced and FACS-sorted the cells for GFP expression before extraction of their total RNA. With a -fold change cut-off excluding values below 1.5, we obtained a list of 430 differentially regulated genes. An upstream pathway analysis by IPA was used to identify the top 20 transcriptional regulators with a significant number of downstream targets in the data set (Table 3). The list was sorted based on positive activation z-scores (i.e. these transcriptional regulators are all activated). One of the transcriptional regulators in the list, MITF, is a well described basic helix-loop-helix leucine zipper transcription factor that has previously been shown to be directly inhibited by the histidine triad nucleotide-binding protein HINT1 (33, 34). Interestingly, HINT1 was one of the interaction partners found in the yeast two-hybrid screen described above, and we considered that this interaction may be involved in the induction of MITF target genes by the TEN1-ICD.
The relationships of MITF to its downstream targets are shown in an IPA My Pathways network. Seven of the eight target genes present in this database are consistently up- or down-regulated in our microarray experiment, with only the exception of NGFR (Fig. 3). A literature search revealed three further positively regulated MITF target genes in our microarray data, ERBB3, SLC1A4, and CHL1 (38). All target genes are marked in a volcano plot (Fig. 4A), summarizing the microarray data. Consistent up- or down-regulation of the MITF target genes in the triplicates of the microarray is shown in Fig. 4B and was confirmed for some of the target genes by qPCR (Fig. 4C). To exclude the possibility that these target genes are a result of the particular stable cell line used for the transcript profiling, we created a second set of stable BS149 cell lines using the same vectors of the modified tet system. Again, we confirmed the particular stable cell line used for the transcript profiling, as in the nucleus (Fig. 5A). Using a proximity ligation assay, the HINT1-MYC and TEN1-ICD-HA co-transfection confirms an interaction, as seen by the fluorescent red dots in the cells that are overexpressing both proteins (Fig. 5B, left and center image). As a negative control, the same primary and secondary antibodies were used, but only HINT1-MYC was overexpressed in COS-7 cells (Fig. 5B, right image). Therefore, we concluded that the TEN1-ICD indeed binds to HINT1, and we hypothesized that the TEN-1 ICD might act by sequestering HINT1, thereby compromising its repression of MITF-mediated transcription.

The proteins interact when expressed in COS-7 cells. We co-transfected vectors encoding HINT1-MYC and TEN1-ICD-HA; if the two proteins are located in close proximity to each other, the oligonucleotides present on the secondary antibodies used to detect the two proteins can hybridize and be ligated and amplified using fluorescence-labeled nucleotides. Immunocytochemistry images of co-transfected cells show that many COS-7 cells overexpress both proteins in the cytoplasm as well as in the nucleus (Fig. 5A). Using a proximity ligation assay, the HINT1-MYC and TEN1-ICD-HA co-transfection confirms an interaction, as seen by the fluorescent red dots in the cells that are overexpressing both proteins (Fig. 5B, left and center image). As a negative control, the same primary and secondary antibodies were used, but only HINT1-MYC was overexpressed in COS-7 cells (Fig. 5B, right image). Therefore, we concluded that the TEN1-ICD indeed binds to HINT1, and we hypothesized that the TEN-1 ICD might act by sequestering HINT1, thereby compromising its repression of MITF-mediated transcription.

### TABLE 2
**Teneurin-1 ICD as a Transcriptional Regulator**

| Gene       | Protein name                                      | Information                                                                 |
|------------|---------------------------------------------------|-----------------------------------------------------------------------------|
| CSNKB      | Casein kinase II subunit β                        | Regulatory subunit of casein kinase II (CK2); not required for catalytic activity; recruits substrates/ regulators; CK2 is a pleiotropic kinase involved in e.g. cell proliferation, transcription, etc. (59) |
| POLR2J     | DNA-directed RNA polymerase II subunit RPBI1-a     | Subunit of the RNA polymerase II complex; forms heterodimer with POLR2C, which is part of the core enzyme; involved in the termination of transcription; binds to SATB1 and Che-1 (60–62) |
| APBB1      | Amyloid β A4 precursor protein-binding family B member 1 | Adapter protein; aids in processing of amyloid precursor protein (APP) by binding to it; this interaction also influences production of amyloid-β peptides that are found in Alzheimer patients; promotes neurite outgrowth (63) |
| BEX1       | Brain-expressed X-linked protein 1                | Small adapter protein; involved in NGFR signaling and transcriptional regulation of cell cycle arrest genes; implicated in promoting survival of neurons and neurite outgrowth (51, 52) |
| NIIBP1     | Nucleotide binding protein 1                      | MRPI/MnD-type P-loop NTPa; interacts with motor protein KIF5A; involved in regulation of centriole duplication; implicated in assembly of cytosolic iron-sulfur proteins (64) |
| HINT1      | Histidine triad nucleotide-binding protein 1       | Member of the evolutionarily conserved HIT superfamily; tumor suppressor gene; inhibits transcription factors like MITF and β-catenin by directly binding them at the promoters of their target genes (34) |
| MACF1      | Microtubule-actin cross-linking factor 1          | Spectraplakin and TIP protein; conserved SH3 domain; role in cross-linking microtubules to actin cytoskeleton; regulates growth of neuronal microtubules, and subsequently filopodia formation and axon extension (53, 54) |
| PTPN12     | Tyrosine-protein phosphatase non-receptor type 12  | Part of non-receptor PTP subfamily; ubiquitously expressed phosphatase; important functions in early embryogenesis, like development of mesenchyme; involved in cell spreading and migration (65) |
| FIBP       | Acidic fibroblast growth factor intracellular binding protein | Selectively binds aFGF; implicated in autocrine action of aFGF; up-regulated in some cancers; involved in angiogenesis of tumors (66, 67) |
| DICALM     | Phosphatidylinositol-binding clathrin assembly protein | Ubiquitously expressed adapter protein; strongest expression in neurons, especially in pre- and postsynaptic structures; risk factor for Alzheimer disease; plays a role in clathrin-mediated endocytosis (68, 69) |
| GNB2L1     | Guanine nucleotide-binding protein                 | Scaffold protein; originally identified as anchoring protein for protein kinase C; also recruits and binds other proteins like integrins; altered expression levels in many cancers (70) |

Information about all interaction partners of the TEN1-ICD, as identified in the yeast two-hybrid screen.

| Gene       | Protein name                                      | Information                                                                 |
|------------|---------------------------------------------------|-----------------------------------------------------------------------------|
| POLR2J     | DNA-directed RNA polymerase II subunit RPBI1-a     | Subunit of the RNA polymerase II complex; forms heterodimer with POLR2C, which is part of the core enzyme; involved in the termination of transcription; binds to SATB1 and Che-1 (60–62) |
| HINT1      | Histidine triad nucleotide-binding protein 1       | Member of the evolutionarily conserved HIT superfamily; tumor suppressor gene; inhibits transcription factors like MITF and β-catenin by directly binding them at the promoters of their target genes (34) |
| MACF1      | Microtubule-actin cross-linking factor 1          | Spectraplakin and TIP protein; conserved SH3 domain; role in cross-linking microtubules to actin cytoskeleton; regulates growth of neuronal microtubules, and subsequently filopodia formation and axon extension (53, 54) |
| PTPN12     | Tyrosine-protein phosphatase non-receptor type 12  | Part of non-receptor PTP subfamily; ubiquitously expressed phosphatase; important functions in early embryogenesis, like development of mesenchyme; involved in cell spreading and migration (65) |
| FIBP       | Acidic fibroblast growth factor intracellular binding protein | Selectively binds aFGF; implicated in autocrine action of aFGF; up-regulated in some cancers; involved in angiogenesis of tumors (66, 67) |
| DICALM     | Phosphatidylinositol-binding clathrin assembly protein | Ubiquitously expressed adapter protein; strongest expression in neurons, especially in pre- and postsynaptic structures; risk factor for Alzheimer disease; plays a role in clathrin-mediated endocytosis (68, 69) |
| GNB2L1     | Guanine nucleotide-binding protein                 | Scaffold protein; originally identified as anchoring protein for protein kinase C; also recruits and binds other proteins like integrins; altered expression levels in many cancers (70) |
in BS149 cells, MITF overexpression should result in the induction of genes in common with TEN1-ICD induction. To determine the overlap of differentially regulated genes, the MITF-RFP-HA construct was transiently overexpressed in BS149 cells and compared with RFP-HA in a whole transcriptome microarray analysis. Again, we compared biological triplicates by separately transfecting and sorting the cells and extracting their total RNA. With a -fold change cut-off (excluding values below 1.5) and an adjusted \( p \) value cut-off (excluding values above

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**FIGURE 2.** Overexpression of the teneurin-1 ICD through a modified tet system in BS149 cells, a cell line expressing endogenous teneurin-1. A, left, quantitative RT-PCR screen of teneurin-1 expression in glioblastoma cell lines. Values are normalized to TBP, using the comparative \( C_r \) method. Right, quantitative RT-PCR results of teneurin-1 expression comparing the glioblastoma cell line BS149 with different brain tissues. Values are normalized to TBP, using the comparative \( C_r \) method. Note that teneurin-1 transcript levels in cerebellum are 181.8-fold higher than in BS149 cells. B, scheme of the modified tet system. Through the addition of Dex and Dox, the tet activator construct (irtTA/VP16/GBD) is released from HSP90 in the cytosol and induces expression of the ICD through the tet-CMV promoter. C, Western blot with anti-GFP showing the expression of the control construct GFP-His and the teneurin-1 ICD-GFP-His 24 h after the addition of Dex and Dox. Anti-vinculin is the internal control for equal loading. D, quantitative RT-PCR results show increased levels of the ICD-GFP fusion construct (using teneurin-1 ICD-GFP primers) compared with endogenous teneurin-1 levels (using teneurin-1 ECD primers) in BS149 cells overexpressing the ICD-GFP construct by a modified tet system, using cDNA prepared from the RNA used in the microarray. Values were normalized to TBP, using the comparative \( C_r \) method. Note that teneurin-1 ICD transcript levels are 214.9 times higher than endogenous teneurin-1, which is in the range of the endogenous teneurin-1 present in cerebellum. Error bars, S.D.

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**TABLE 3**

Predicted activated transcriptional regulators

| Transcriptional regulator | Activation \( z \)-score | \( p \) value of overlap | Number of downstream targets in data set |
|---------------------------|--------------------------|--------------------------|----------------------------------------|
| SREBF1                    | 3.872                    | 5.60E-12                 | 23                                      |
| SREBF2                    | 3.342                    | 1.85E-13                 | 17                                      |
| IRF5                      | 2.415                    | 1.49E-03                 | 6                                       |
| PPARC1                    | 2.209                    | 4.97E-06                 | 8                                       |
| IRF3                      | 2.122                    | 3.51E-02                 | 2                                        |
| MYOD1                     | 2.000                    | 3.62E-02                 | 8                                       |
| KLF2                      | 1.977                    | 8.46E-03                 | 8                                       |
| CEBP1                     | 1.964                    | 5.16E-04                 | 8                                       |
| MITF                      | 1.913                    | 1.45E-02                 | 2                                        |
| BCL6                      | 1.849                    | 3.44E-04                 | 10                                      |
| MYCN                      | 1.830                    | 2.75E-01                 | 7                                       |
| IRF1                      | 1.807                    | 6.12E-05                 | 12                                      |
| IRF7                      | 1.719                    | 1.40E-05                 | 13                                      |
| STAT1                     | 1.677                    | 4.88E-09                 | 22                                      |
| BRCA1                     | 1.660                    | 5.48E-07                 | 15                                      |
| EGR2                      | 1.633                    | 6.83E-04                 | 10                                      |
| NFATC2                    | 1.510                    | 7.29E-04                 | 10                                      |

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**FIGURE 3.** Teneurin-1 ICD overexpression affects MITF target genes. IPA identified all of its eight MITF target genes in our microarray to be differentially regulated, six up-regulated (red) and two down-regulated (green). The arrows indicate that all genes are directly activated or inhibited by MITF. The regulation of these target genes is consistent with the database for all genes except for NGFR, which according to IPA is up-regulated by MITF but is down-regulated in our microarray.
we obtained a list of 497 differentially regulated genes. Between the 430 genes differentially regulated by the TEN1-ICD and the 497 genes affected by MITF, there is an overlap of 42 genes that were either up- or down-regulated in both microarrays (Fig. 6A).

One of the overlapping genes is **GPNMB**, a target gene of MITF with a well described promoter that is directly bound by MITF (36). Therefore, we further investigated the effect of the TEN1-ICD on the expression of this particular gene. First, we could confirm by qPCR that MITF also regulates **GPNMB** in BS149 cells, showing a 4.14-fold increase in expression (Fig. 6B). To show a link of the TEN1-ICD and MITF via HINT1, HINT1 would have to influence MITF-regulated transcription. We therefore compared **GPNMB** transcript levels after co-

![FIGURE 4. Microarray identifies nine up-regulated MITF target genes due to teneurin-1 ICD overexpression. A, volcano plot showing the 430 differentially regulated genes when the teneurin-1 ICD is overexpressed in BS149 cells compared with BS149 cells with overexpression of GFP only (-fold change >1.5; p < 0.05). MITF target genes are marked in the plot. B, detailed view of the expression levels of the 11 MITF target genes in a heat map, where bright green means little or no expression and bright red means high expression. Nine genes are up-regulated, and two genes are down-regulated, with the -fold change indicated. C, quantitative RT-PCR confirmed the up-regulation of some of the MITF target genes. Values are normalized to TBP, and -fold change values compare overexpression of teneurin-1 ICD-GFP-His with GFP-His. Error bars, S.D.](image-url)

![FIGURE 5. Proximity ligation assay confirms the interaction between HINT1 and the TEN1-ICD. A, immunocytochemistry (ICC) images of COS-7 cells co-transfected with TEN1ICD-HA and HINT1-MYC stained with anti-HA (red) and anti-MYC (green) and the merged channels showing co-expression in yellow. B, proximity ligation assay images of COS-7 cells co-transfected with TEN1ICD-HA and HINT1-MYC (left and center images) and COS-7 cells transfected only with HINT1-MYC as a negative control (right image); nuclei are stained with DAPI; white bar, 50 μm.](image-url)
transfection of cells with the MITF-RFP-HA and HINT1-CFP-MYC constructs. To make sure to analyze cells that over-express both proteins, cells were FACS-sorted for expression of RFP and CFP before RNA isolation. Indeed, we saw a significant up-regulation of GPNMB transcripts when we transiently over-expressed the MITF-RFP-HA together with the empty control plasmid CFP-MYC, and this induction was attenuated by co-transfection with HINT1-CFP (Fig. 6C).

Finally, we wanted to investigate whether MITF directly influences GPNMB transcription in BS149 cells using promoter reporter experiments and whether this was affected by TEN1-ICD. To do this, we used two different promoter constructs of GPNMB in SEAP reporter gene assays: the full promoter and one missing the crucial MITF binding site M-box (GPNMB ΔM-box) as described previously (36) (Fig. 7A). Overexpressed MITF in BS149 cells strongly induced the full GPNMB promoter, with a 24.15-fold increase compared with the control, whereas the GPNMB ΔM-box promoter could not be induced by MITF (Fig. 7B). Thus, MITF directly binds to the GPNMB promoter and influences its transcription in BS149 cells. Next, we tested the influence of the TEN1-ICD on MITF-induced transcriptional activation of the GPNMB promoter reporter.

This experiment showed that the TEN1-ICD further increases MITF-dependent transcription by 1.59-fold.

In summary, our results show that the transcriptional repressor HINT1 is a previously unknown interaction partner of teneurin-1, through which the TEN-1 ICD can influence MITF-dependent transcription, as depicted in the model presented in Fig. 8.

**DISCUSSION**

We propose a function for the ICD of teneurin-1 that is very similar to the Notch signaling pathway. Notch is a well studied type I transmembrane protein with a single-spanning transmembrane domain and an ICD that is much smaller in size than the ECD. The ICD can regulate transcription of several target genes by replacing repressors with other positive regulators in a transcription complex once it translocates to the nucleus (39).

In this study, we show that the teneurin-1 ICD can influence MITF-dependent transcription of GPNMB by binding to repressor HINT1. Although the ICDs of teneurins have been
implicated in transcriptional activity in the past (29, 32), here we are elucidating a new molecular mechanism to explain how a teneurin ICD can influence transcriptional activity. In our model, the ECD domain is released following homo- or heterophilic interaction, which is required for subsequently releasing the ICD.

Following its release, the ICD can translocate to the nucleus due to its predicted nuclear localization signal, as has previously been demonstrated in experimental studies (14, 29, 32). Here we have shown that the ICD can bind HINT1, thus switching on MITF-dependent transcription (Fig. 7). We showed this by using the MITF target gene GPNMB as an example, although there were 41 other genes differentially regulated by MITF and the TEN1-ICD. Most importantly, this highlights the potential of how teneurins can influence transcriptional regulation.

The mechanism of action of the TEN-1 ICD may be slightly different from Notch ICDs. Rather than replacing transcriptional repressors, recruiting positive regulators, and taking part in a transcriptional complex, the teneurin-1 ICD might regulate transcription by binding the transcriptional repressor and either releasing it from the transcription factor MITF or competing for binding to MITF. We are not ruling out the possibility that the TEN-1 ICD or the ICDs of the other teneurins have other modes of action, more like the Notch ICD. As we will discuss below, there are other interesting genes that might be regulated by the teneurin-1 ICD independent of MITF.

Although MITF-dependent transcription has mostly been studied in a melanocyte-specific context, other sites of expression like the retinal pigment epithelium have also been identified (40). Additionally, according to the Allen Mouse Brain Atlas (41), there is a weak MITF expression in the mouse olfactory bulb. However, MITF expression seems to have been rarely studied in the context of the central nervous system.

The expression of teneurin-1, HINT1, and GPNMB in the CNS has been studied more extensively. Unrelated studies suggest at least partially overlapping expression patterns of the three genes in the olfactory bulb, hippocampus, and cerebral cortex (14, 17, 42, 43). It will be very interesting to further explore the overlapping expression patterns of these genes in the CNS, including MITF, and to determine the function of the teneurin-1 ICD regulating MITF target genes like GPNMB. In the real-time qPCR screen, all tested glioblastoma cell lines expressed endogenous teneurin-1. In a recent review discussing teneurins in human tumorigenesis and malignancy, teneurin-2 and -4 were described as potential tumor suppressors or oncogenes (44). Teneurin-1 was not mentioned in the review, which may be due to a lack of data about the expression of this gene in tumors. GPNMB not only promotes invasiveness of glioma cells; it is also elevated in malignant glioblastomas. Taken together, this could be a first link of teneurin-1 to cancer and would make it a potential target in glioblastomas (45).

GPNMB is a type I transmembrane protein originally identified in human melanoma cells (45). Since then, its expression has also been identified in chondrogenesis (46), in differentiating osteoclasts and osteoblasts (46), and in normal and diseased brain tissue (43, 47). The function in the CNS is mostly unknown, although it is suggested to play a role in the stability of neurons and the immune/inflammatory response in the CNS (43). Interestingly, GPNMB also contains an RGD sequence in its ECD, which is likely to bind integrins (45). Integrins have also been studied in the context of axon guidance and neuronal connectivity (48). This leaves us with several potential functions of the teneurin-1 ICD regulating GPNMB expression. For one, GPNMB could regulate the stability of neurons once they have found their postsynaptic partner due to teneurin-1 homophilic binding. On the other hand, the capability of GPNMB to bind integrins could further specify or stabilize an interaction of the presynaptic neuron with its postsynaptic partner. Finally, we cannot exclude the possibility that our identified mechanism takes place in a non-neuronal context. Recent papers have identified teneurin-3 and -4 as important players in chondrogenic differentiation (49, 50). Although teneurin-1 has not been studied in chondrogenesis, the likely role of GPNMB as a key regulator in chondrogenesis could implicate a function of both proteins in this event.
Teneurin-1 ICD as a Transcriptional Regulator

Here we elucidated a novel mechanism of how the teneurin ICD can influence transcription. It is likely that additional mechanisms will be identified in the future, as this is a very diverse protein family with a wide range of important functions.

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