Rab15 expression correlates with retinoic acid-induced differentiation of neuroblastoma cells

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Abstract. Neuroblastoma is the most common extracranial solid tumor in children and accounts for 15% of pediatric cancer deaths. Although retinoic acid (RA) is currently used to treat high-risk neuroblastoma patients in the clinic, RA-responsiveness is variable and unpredictable. Since no alterations in the RA-signaling pathway have been found in neuroblastoma cells, molecules correlated with RA-induced differentiation will provide predictive markers of RA-responsiveness for clinical use. The Rab family of small G proteins are key regulators of membrane traffic and play a critical role in cell differentiation and cancer progression. Although an increasing number of cancer-associated alternative splicing events have been identified, alternative splicing of Rab proteins remains to be characterized in neuroblastoma. In the present study, we focused on Rab15 that was originally identified as a brain-specific Rab protein and regulates the endocytic recycling pathway. We identified alternatively spliced Rab15 isoforms designated as Rab15CN and Rab15SN in neuroblastoma cells. Rab15CN was composed of 7 exons encoding 212 amino acids and showed brain-specific expression. Alternative splicing of exon 4 generated Rab15SN that was predicted to encode 208 amino acids and was predominantly expressed in testis. RA-induced neuronal differentiation of neuroblastoma BE(2)-C cells and specifically up-regulated Rab15SN expression. Reciprocally, RA-induced differentiation was observed in Rab15CN-expressing BE(2)-C cells in preference to Rab15SN-expressing BE(2)-C cells. Furthermore, Rab15CN expression was also specifically up-regulated during RA-induced differentiation of newly established neuroblastoma cells from high-risk patients. These results suggest that Rab15 expression correlates with RA-induced differentiation of neuroblastoma cells.

Introduction

Neuroblastoma is the most common extracranial solid tumor in children and accounts for 15% of pediatric cancer deaths. More than 50% of high-risk neuroblastoma patients develop early or late relapse (1-3). This is mainly due to the difficulty in eradicating the minimal residual disease (MRD) that resides in the bone marrow and peripheral blood even after achieving the complete remission. With the aim of inducing the differentiation of MRD, the vitamin A metabolite retinoic acid (RA) is currently used in the clinic for high-risk neuroblastoma patients. Although RA can let neuroblastoma cells cease proliferation, differentiate into neuronal cells or undergo apoptosis, RA-responsiveness of high-risk neuroblastoma patients is variable and unpredictable. Consequently, less than 40% of high-risk neuroblastoma patients can expect a long-term cure (4,5). Since no alteration in the RA-signaling pathway is found in neuroblastoma cells, a molecule correlated with RA-induced differentiation will give insight into a predictive marker of RA-responsiveness for clinical use.

Phenotypic characterization of a large number of neuroblastoma cells identifies three major cell types, designated as N (neuroblastic), S (substrate-adherent and non-neuronal) and I (intermediate) (6,7). N-type neuroblastoma cells have small and rounded cell bodies with neurite-like processes and attach poorly to the substrate. In contrast, S-type cells have large and flattened cell bodies without neurite-like processes and grow in culture as an adherent monolayer. I-type cells have an intermediate morphology between N-type and S-type, and can differentiate into either N-type or S-type when induced by specific agents. BE(2)-C cells have a typical I-type phenotype and are frequently used as a model for RA-induced differentiation of neuroblastoma cells (8).

Rab family small G proteins are central in ensuring the spatiotemporal regulation of membrane traffic. Over 60
different Rab proteins are identified in mammalian cells, and each Rab protein recognizes distinct subsets of intracellular membranes and regulates the specific membrane transport pathway (9-11). Accumulating evidence reveals that aberrant expressions of several Rab proteins are associated with cancer progression (12). Increased expression of brain-specific Rab3A and Rab23 is implicated in brain tumor, neuroendocrine tumors, hepatocellular carcinoma and gastric cancer (13-15). Rab25 and its effector Rab-coupling protein (RCP) control the endocytic recycling pathway and are identified as drivers of genomic amplicon in ovarian and breast cancer (16,17).

Alternative splicing represents an important molecular mechanism of gene regulation in normal cell growth and differentiation. In the case of hyaluronan synthase 1 (HAS1) and receptor for hyaluronan-mediated motility (RHAMM), their aberrant splicing are associated with multiple myeloma, and their isoform balance is shown to be a prognostic marker (18,19). Although an increasing number of cancer-associated alternative splicing events are identified, alternative splicing of Rab proteins remains to be characterized in neuroblastoma (20,21).

In the present study, we focused on Rab15 that was originally identified as a brain-specific Rab protein and regulated the endocytic recycling pathway (22,23). We identified and characterized alternatively spliced Rab15 isoforms in neuroblastoma cells.

**Materials and methods**

**Cell culture, transfection and neuronal differentiation.** BE(2)-C cells were obtained from ATCC (Manassas, VA), cultured at 37°C (5% CO₂ and 95% air) in Dulbecco’s modified Eagle’s medium (DMEM)/Ham’s F12 (Wako Pure Chemical, Osaka, Japan) with 10% fetal bovine serum (FBS; Invitrogen, Carlsbad, CA), and transfected using Lipofectamine 2000 transfection reagent (Invitrogen) according to the manufacturer’s instructions. For neuronal differentiation, BE(2)-C cells were obtained from BE(2)-C cell line, and transfected using lipofectamine 2000 transfection reagent (Invitrogen) according to the manufacturer’s instructions. For neuronal differentiation, BE(2)-C cells were obtained from BE(2)-C cell line, and transfected using lipofectamine 2000 transfection reagent (Invitrogen) according to the manufacturer’s instructions.

**Quantitative real-time RT-PCR.** Human tissue total RNA was purchased from Clontech (Palo Alto, CA). Total RNA from neuroblastoma cells was isolated with RNasy Mini kit (Qiagen, Valencia, CA) and reverse transcribed using Quantitect Reverse Transcription kit (Qiagen). Real-time PCR analysis was performed with an ABI 7500 Fast real-time PCR System (Applied Biosystems, Foster City, CA) using FastStart Universal SYBR-Green Master (Roche) according to the manufacturer’s instructions. Each sample was analyzed in triplicate. Relative mRNA expression of Rab15 to PGK1 and Rab15 isoform balance were calculated by the comparative Cₚ method. Primers for Rab15cn (ENST00000436278) were forward 5'-AGAGATACAGACACATACA-3' (sense) and reverse 5'-TTCTTGTCGATCCTCACACGC-3' (anti-sense). Rab15an (ENST00000267512) were forward 5'-TACAGATCTGGGACACTGCA-3' (sense) and reverse 5'-CAGGCAGTGAATGGCATCT-3' (anti-sense), and phosphoglycerate kinase 1 (PGK1, NM_000291) were forward 5'-GAAGACCTCCGTCTTTC-3' (sense) and reverse 5'-GGCTCGGCTTTAAC-3' (anti-sense).

**Plasmid construction.** The full-length cDNAs of human Rab15cn (ENST00000436278) and Rab15an (ENST00000267512) were cloned by RT-PCR from BE(2)-C cell cDNA, and cloned into the bicistronic expression vector pCMV6-AC-ires-GFP (Origene, Rockville, MD). The resulting pCMV6-Rabl5cn-ires-GFP and pCMV6-Rabl5an-ires-GFP plasmids were sequenced using an ABI Prism 3100 Genetic Analyzer (Applied Biosystems).

**Phase-contrast and immunofluorescence microscopy.** Phase-contrast images of neuroblastoma cells were acquired using a BZ-9000E fluorescence microscope (Keyence, Osaka, Japan). Neuroblastoma cells grown on Collagen I-coated cover glass (Iwaki, Tokyo, Japan) were fixed with 2% formaldehyde in PBS for 15 min at room temperature. After permeabilization with 0.1% Triton X-100 in PBS for 5 min and blocking with 5% goat serum (Sigma) in PBS (GS/PBS) for 60 min at room temperature, cells were incubated with primary antibodies in GS/PBS for 60 min, with Alexa 568-conjugated secondary antibodies (Invitrogen) in GS/PBS for 60 min, and with 1 µg/ml 4,6-diamidino-2-phenylindole (DAPI; Sigma) in PBS for 5 min at room temperature. Fluorescent images were acquired using a BZ-9000E fluorescence microscope (Keyence).

**Newly established neuroblastoma cells from high-risk patients.** Bone marrow and tumor samples were obtained from patients with high-risk neuroblastoma who were admitted to Hyogo Children’s Hospital and gave the written informed consent approved by Hyogo Children's Hospital Ethics Board. These samples were handled in accordance with the Guidelines for Clinical Research of Kobe University Graduate School of Medicine. Bone marrow samples were

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**Table I. Summary of clinical samples.**

<table>
<thead>
<tr>
<th>Sample name</th>
<th>Age (years)</th>
<th>Gender</th>
<th>Tumor risk group</th>
<th>Specimen type</th>
</tr>
</thead>
<tbody>
<tr>
<td>NBBM2</td>
<td>6</td>
<td>Male</td>
<td>High</td>
<td>Bone marrow</td>
</tr>
<tr>
<td>NBBM3</td>
<td>3</td>
<td>Female</td>
<td>High</td>
<td>Bone marrow</td>
</tr>
<tr>
<td>NBT2D</td>
<td>2</td>
<td>Male</td>
<td>High</td>
<td>Adrenal gland</td>
</tr>
</tbody>
</table>

Tumor risk group was based on the Children’s Oncology Group Neuroblastoma Risk Stratification System (30).
separated by Mono-Poly Resolving Medium (DS Pharma Biomedical, Osaka, Japan) and mono- and poly-nucleated cells were collected. The resulting bone marrow cells were washed with PBS, cultured at 37°C (5% CO₂ and 95% air) in DMEM/Ham's F12 containing 10% FBS, and designated as NBTT2D cells (Table I).

Antibodies and other reagents: Mouse anti-β-1, 3-glucuronyltransferase 1 (B3GAT1)/CD57/HNK-1 antibody was purchased from BD and mouse anti-tyrosine hydroxylase (TH) from Millipore (Temecula, CA). Other reagents were obtained from commercial sources.

Results

Identification and tissue distribution of alternatively spliced Rab15 isoforms. Rab15 cDNA was originally isolated from a rat brain cDNA library and contained a 639 bp open reading frame encoding 212 amino acids (22). In human, Rab15 gene is composed of 7 exons and predicted to have an alternatively spliced exon 4 (Fig. 1A). We first cloned the full-length cDNAs for two human Rab15 transcripts (ENST00000436278 and ENST00000267512) by RT-PCR from neuroblastoma BE(2)-C cells and confirmed their DNA sequences. Here, we designated ENST00000436278 and ENST00000267512 as canonical Rab15 (Rab15cn) and alternative Rab15 (Rab15An), respectively. Rab15cn was a human counterpart of rat Rab15 and contained a 78-bp exon 4, resulting in a 639 bp open reading frame encoding 212 amino acids. Rab15An was an alternatively spliced isoform of human Rab15 and had a 131-bp insertion (exon 4b) between exon 4 and exon 5, resulting in a 627 bp open reading frame encoding 208 amino acids (Fig. 1A). Amino acid sequence analysis revealed that exon 4b insertion resulted in a frameshift in Rab15An and generated a unique carboxy-terminal tail of 100 amino acids lacking the conserved residues for gDp/gtp-binding (g2 and g3) and geranylgeranylation (CxC) in Rab proteins (24). Next we examined the expression of Rab15cn and Rab15An in various human tissues by a quantitative real-time RT-PCR using the Rab15 isoform-specific primers (Fig. 1A). In accordance with previous report (22), Rab15cn mRNA was highly expressed in brain, and at low levels in other tissues and testis (Fig. 1B). In contrast, testis showed the highest level of Rab15An expression, and other tissues and brain showed only slight expression (Fig. 1C).

Differential expression of Rab15cn and Rab15An during RA-induced differentiation of BE(2)-C cells. As reported previously (8), BE(2)-C cells responded to RA and extended neurite-like processes (Fig. 2A). We then analyzed the expression of Rab15cn and Rab15An to PGK1 (C) in various human tissues was analyzed by quantitative real-time RT-PCR. The mean expression in skeletal muscle was set to 1. The data shown are the means ± SD of three independent experiments.

Distinct RA-responsiveness of Rab15cn-expressed and Rab15An-expressed BE(2)-C cells. To gain insight into the function of Rab15cn and Rab15An, we then examined the effect of Rab15cn and Rab15An expression on the RA-induced
Characterization of newly established neuroblastoma cells from high-risk patients. Because neuroblastoma cells displayed considerable heterogeneity in response to RA (25), we have established three neuroblastoma cells designated as NBBM2, NBBM3, and NBTT2D from high-risk patients (Table I). The morphology of NBBM2 and NBTT2D cells were characterized by a round and flattened cell body with little neurite-like processes, and closely resembled I-type BE(2)-C cells (Fig. 3). In contrast, NBBM3 cells displayed a spindle-shaped cell body with longer neurite-like processes that is typical for N-type neuroblastoma cells (Fig. 5A). To confirm the origin of NBBM2, NBBM3, and NBTT2D cells, we determined whether these cells indeed expressed neuroblastoma markers, β-1,3-glucuronyltransferase 1 (B3GAT1, also known as CD57 and HNK-1) and tyrosine hydroxylase (TH), by an immunofluorescence microscopy. B3GAT1 is a marker for histologically immature neuroblastoma cells and neural crest stem cells (26), while TH is a catecholamine biosynthetic pathway enzyme used as a clinical neuroblastoma marker (27). Although the percentage of positive cells was varied, NBBM2, NBBM3, and NBTT2D cells indeed expressed both B3GAT1 and TH (Fig. 4).

Differential expression of Rab15CN and Rab15AN during RA-induced differentiation of newly established neuroblastoma cells from high-risk patients. To characterize newly established neuroblastoma cells, we first examined their RA-responsiveness. I-type NBBM2 and NBTT2D cells extended neurite-like processes and displayed a typical n-type morphology in response to RA (Fig. 5A). N-type NBBM3 cells were also responded to RA and elongated the length of neurite-like processes (Fig. 5A). Next, we analyzed the
expression of Rab15CN and Rab15AN by a quantitative real-time RT-PCR using the Rab15 isoform-specific primers. Upon RA-treatment, Rab15CN expression was increased up to 1.9-, 1.8-, and 2.0-fold in NBBM2, NBBM3, and NBTT2D cells, respectively (Fig. 5B). In contrast, Rab15AN expression in NBBM2, NBBM3, and NBTT2D cells was changed 1.3-, 1.4-, and 1.2-fold, respectively (Fig. 5C). These results indicate that RA induces differential expression of Rab15CN and Rab15AN in NBBM2, NBBM3, and NBTT2D cells as well as BE(2)-C cells.

Correlation between Rab15 expression and RA-induced differentiation of neuroblastoma cells. As the isoform balance of some aberrantly spliced genes in cancer cells was reported to be a prognostic marker (18,19), we first determined Rab15 isoform balance measured by the Rab15CN/Rab15AN ratio in various human tissues. It was considerably varied, ranging from 0.6 in testis to 11.5 in brain (Fig. 6A). We then determined Rab15 isoform balance in BE(2)-C, NBBM2, NBBM3, and NBTT2D cells treated with DMSO and RA. It was significantly increased upon RA-treatment in all neuroblastoma cells (Fig. 6B). These results suggest that Rab15 expression correlates with RA-induced differentiation of neuroblastoma cells.

Discussion

In the present study, we have identified alternatively spliced Rab15 isoforms designated as Rab15CN and Rab15AN in neuroblastoma cells and obtained two novel findings. First, Rab15AN is predominantly expressed in testis, while Rab15CN shows a brain-specific expression. Second, Rab15 expression correlates with RA-induced differentiation of neuroblastoma cells.

Neuroblastoma cells are widely assumed to originate from neural crest cells that had a cancerous change during their sympathoadrenal differentiation (1-3). In response to RA, neuroblastoma cells can differentiate into neuronal cells or undergo apoptosis. Although the heterogeneity in RA-responsiveness is widely recognized (25), all neuroblastoma cells treated with RA in the present study extended neuron-like processes and up-regulated Rab15CN expression (Fig. 5). Rab15CN expression in BE(2)-C cells was not sufficient to induce the extension of neuron-like processes (Fig. 3). Although Rab15 up-regulation likely corresponds to the increased demand for the endocytic recycling pathway to remodel cell membranes and induce neuronal differentiation (23), the exact way by which Rab15 up-regulation could affect the RA-induced differentiation is an open question for future study.
An increasing number of genes including HAS1 and RHAMM are shown to predict clinical outcome depending on the differential expression of alternatively spliced isoforms in cancer patients (18-21). Although alternative splicing events of Rab proteins remain to be characterized in neuroblastoma, Rab6A and Rab28 are reported to contain alternatively spliced isoforms. Rab6A isoforms named Rab6Aα and Rab6Aβ are ubiquitously expressed at similar levels, whereas Rab28 isoforms named Rab28S and Rab28L show differential expression: ubiquitous Rab28S and testis-specific Rab28L (28,29). In the case of Rab15, it is not clear whether alternatively spliced isoform Rab15α has actual cellular function, or whether Rab15α is a consequence of aberrant mRNA splicing. We are now trying to elucidate the exact function of Rab15α mRNA and its encoded protein in neuroblastoma cells. As RA-responsiveness of MRD in high-risk neuroblastoma patients is variable and unpredictable, Rab15 expression will give insight into a predictive marker of RA-responsiveness for clinical use (4,5).

In summary, we identified alternatively spliced Rab15 isoforms, brain-specific Rab15β and testis-specific Rab15α, in neuroblastoma cells, and we revealed that Rab15 expression correlated with RA-induced differentiation of neuroblastoma cells.

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