NDRG2 Promotes GATA-1 Expression through Regulation of the JAK2/STAT Pathway in PMA-stimulated U937 Cells

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Background: N-myc downstream-regulated gene 2 (NDRG2), a member of a newly described family of differentiation-related genes, has been characterized as a regulator of dendritic cells. However, the role of NDRG2 on the expression and activation of transcription factors in blood cells remains poorly understood. In this study, we investigated the effects of NDRG2 overexpression on GATA-1 expression in PMA-stimulated U937 cells. Methods: We generated NDRG2-overexpressing U937 cell line (U937-NDRG2) and treated the cells with PMA to investigate the role of NDRG2 on GATA-1 expression. Results: NDRG2 overexpression in U937 cells significantly induced GATA-1 expression in response to PMA stimulation. Interestingly, JAK2/STAT and BMP-4/Smad pathways associated with the induction of GATA-1 were activated in PMA-stimulated U937-NDRG2 cells. We found that the inhibition of JAK2 activation, but not of BMP-4/Smad signaling, can elicit a decrease of PMA-induced GATA-1 expression in U937-NDRG2 cells. Conclusion: The results reveal that NDRG2 promotes the expression of GATA-1 through activation of the JAK2/STAT pathway, but not through the regulation of the BMP-4/Smad pathway in U937 cells. Our findings further suggest that NDRG2 may play a role as a regulator of erythrocyte and megakaryocyte differentiation during hematopoiesis.

[Immune Network 2011;11(6):348-357]

INTRODUCTION

Hematopoiesis is controlled by a number of cell type-specific transcription factors that induce differentiation of cells to fulfill particular functions (1). These factors give rise to a specific lineage commitment via a coordinated regulation of expression and activation. However, little is known about how transcription factors regulate hematopoiesis. To this end, it is important to gain an understanding of how transcription factors are modulated during the differentiation of hematopoietic cells.

Globin transcription factor 1 (GATA binding protein 1, GATA-1) is a transcription factor known to be involved in the development of various hematopoietic lineages (2). GATA-1 is a C4 zinc finger transcription factor that recognizes WGATAR DNA motifs and is essential for erythrocyte, megakaryocyte, mast cell, and eosinophil differentiation. GATA-1 has a reciprocal interaction with PU.1, another transcription factor that promotes macrophage and dendritic cell development (3). It has been reported that there are various genes associated with the modulation of GATA-1 expression. A gain of function JAK2 (JAK2V617F) mutation leads to enhanced GATA-1 expression, whereas morpholino knock-down of jak2a or a specific JAK2 inhibitor (TG101209) significantly suppresses GATA-1 expression in zebrafish embryos (4,5). Rab7b-induced IL-6 production and STAT3 activation promote GATA-1 activity in K562 cells (6). GATA-1 expression is also increased by treatment with recombinant BMP-4, but is reduced by Smad5 knockdown in the embryoid body (EB) (7,8). Moreover, dorsomorphin, a selective inhibitor of BMP-induced Smad activation, decreases expression of GATA-1 during embryonic stem (ES) cell differentiation (9,10). Ectopic expression of erythroid differentiation-associated gene (EDAG) induces GATA-1 expression in 32D cells (11), whereas HSP27 promotes ubiquitination of GATA-1 in K562 cells (12).
N-myc downstream-regulated gene 2 (NDRG2) belongs to the NDRG family, a new family of differentiation-related genes composed of four members which share 57~65% amino acid identity (13). NDRG proteins have common structural features, including an NDR-domain and an α/β hydrolase fold, which show high homology among NDRG members (13). Of the NDRG family members, NDRG2 is highly expressed in the adult brain, salivary glands, and skeletal muscle (13). It has been characterized as a regulator of dendritic cell differentiation from monocytes, CD34+ progenitor cells, and myelomonocytic leukemia cell line (14,15). NDRG2 has also been shown to regulate cell growth, apoptosis, and neurodegeneration (16-19). Recently, it has been proposed to be a novel intrinsic factor for the modulation of IL-10 production in myeloid cells (20). However, the role of NDRG2 in the expression and activation of transcription factors in blood cells has remained poorly understood.

Interestingly, NDRG2 overexpression induces a significant decrease of PU.1 expression in U937 cells. We previously showed that NDRG2 overexpression activates the STAT3 pathway in PMA-treated U937 cells (21) and also induces BMP-4 production in MDA-MB-231 cells (17). Given that STAT3 and BMP-4 are involved in GATA-1 expression, and that NDRG2 inhibits PU.1 expression, we hypothesized that NDRG2 increases GATA-1 expression through regulation of either the JAK2/STAT or BMP-4/Smad pathway. To test this hypothesis, we investigated whether NDRG2 promotes expression of GATA-1 in PMA-stimulated U937 cells. GATA-1 expression was substantially increased in NDRG2-overexpressing U937 cells in response to PMA. In addition, NDRG2 expression induced activation of the JAK2/STAT and BMP-4/Smad pathway in PMA-stimulated U937 cells. Inhibition of JAK2 decreased PMA-induced GATA-1 expression in U937-NDRG2 cells. However, inhibition of the BMP-4/Smad pathway did not suppress PMA-induced GATA-1 expression in U937-NDRG2 cells. Taken together, these data indicate that NDRG2 expression promotes the GATA-1 expression through regulation of the JAK2/STAT pathway and not through the BMP-4/Smad pathway in U937 cells.

**MATERIALS AND METHODS**

**Cell lines and reagents**

U937 cells were cultured in RPMI 1640 (Gibco/Invitrogen, Carlsbad, CA) supplemented with 10% heat-inactivated FBS (Gibco), Phorbol 12-myristate 13-acetate (PMA), AG490, and dorsomorphin (6-[(4R,5R)-2-piperidin-1-yl-ethoxy)phenyl]-3-pyridin-4-yl-pyrazol-1,5-d]-pyrimidine) were purchased from Sigma-Aldrich (St. Louis, MO). Recombinant human noggin was obtained from Peprotech, Inc. (Rocky Hill, NJ).

**Construction and transfection of U937 cells**

Cells expressing NDRG2 were previously generated in our laboratory (20). Among the established clones, clone #61 was used as a stable NDRG2-expressing U937 cell line.

**RT-PCR**

Total RNA from harvested cells was isolated using the TRI reagent (Molecular Research Center, Cincinnati, OH). cDNA was synthesized from 1 μg of total RNA and amplified by PCR for 35~40 cycles in a 20 μl reaction mixture containing 1 μl cDNA, 10 pmol of each primer, 10 mM dNTP, and 0.5 U Taq DNA polymerase (Bioneer, Daejeon, Republic of Korea). β-actin was amplified for 22 cycles and used as a loading control. PCR products (10 μl) were electrophoresed on a 1% agarose gel and visualized by ethidium bromide staining. The primer sequences used for RT-PCR are summarized in Table I.

**Western blot analysis**

The cells were lysed in a protein extraction solution (iNtRON Biotechnology, Seongnam, Korea). The proteins were separated on an SDS-polyacrylamide gel and transferred onto a PVDF membrane (Amersham Biosciences, Burkes, UK), which was then blocked with Tris-buffered saline and 0.05% Tween-20 (TBST) containing 5% skim milk. The membranes were incubated with specific antibodies and washed with TBST. The antibodies for phospho-STAT1, phospho-STAT3, phospho-STAT5, phospho-Smad1/5/8, and Smad5 were purchased from Cell Signaling Technology Inc. (Beverly, MA). Antibodies against GATA-1, SOCS1, SOCS2, SOCS3, NDRG2, Lamin A/C, actin, and α-actinin were purchased from Santa Cruz Biotechnology (Santa Cruz, CA). The phospho-JAK2 antibody was purchased from Millipore (Billerica, MA), and the antibody against BMP-4 was purchased from R&D Systems (Minneapolis, MN). After incubating the membrane with diluted secondary antibody coupled to horseradish peroxidase, the antigen-antibody complexes were detected by enhanced chemiluminescence and exposed using LAS 3000 imaging system (FUJIFILM Corporation, Tokyo, Japan).
Table I. PCR primers

| Gene     | Sense primer                     | Antisense primer                     |
|----------|----------------------------------|--------------------------------------|
| PU.1     | 5'-GCA GGC CCT TCT ATA AAA TC-3' | 5'-GCG TTT GGC GTT GTG ATA GA-3'     |
| CSF-1R   | 5'-AGT GCA GTG GCC ACA CTG AT-3' | 5'-AGC TCC GAG GTT GTG ACC AA-3'     |
| GATA-1   | 5'-TTT CCA GTA CCT TCT TTT CTC CC-3' | 5'-ACA GTG GAG TGG CTG TTG CTC-3' |
| SOCS1    | 5'-AGA CCC CTC ACC TCT TG-3'     | 5'-CTG CAC AGC AGA AAA TAA AGC-3'    |
| SOCS2    | 5'-GGA TGG TAC TGG GGA AGT ATG ACT-3' | 5'-AGT GGA TCA GAT GCA CCA CAT TAC-3' |
| SOCS3    | 5'-CAG TTT TGC CTT TCC TCC TCC CAC C-3' | 5'-AAG TGT CCC CTG TTA GGA GC-3' |
| SOCS4    | 5'-AGA GAA GGT CCT ATG ACT GG-3' | 5'-GGG CAT TCC TCC TGG ACC GA-3'     |
| SOCS5    | 5'-CAG TTC TGC AAT TCC ACA AG-3'  | 5'-GTC TTA GTC TCG TCC TCA ACA CT-3' |
| BMPRIA   | 5'-GCA TAA CTA ATG GAC ATT CCT-3' | 5'-TAG AGT TCC TCC TCC CAT GA-3'     |
| BMPRIB   | 5'-GCA GCA CAG ACG GAT ATT GT-3'  | 5'-TCT GCG TCC TCA TCA ACA CT-3'     |
| BMPRII   | 5'-AGG GGA GAG AAG AGC AC-3'     | 5'-CTG TGC TCT TCT CAG ACT GA-3'     |
| ActRI    | 5'-GCA TTC CCA GAG CA CAA-3'     | 5'-GCC TGG TCT TGG CCA CC-3'         |
| ActRII   | 5'-ATG GCT AGA GGA TGT GCA TAT T-3' | 5'-TCT GCA ATG GTC TCA CAG AGC A-3' |
| ActRIIB  | 5'-ACA GGA AGC CAC GAG AGG GT-3'  | 5'-GTA GTG CGG TGG ACC CAC GT-3'     |
| PC1/3    | 5'-TCC ACA CTG GCC ACC TCT TA-3'  | 5'-GCC TGC TCC ATT CTT TTC CC-3'     |
| PC2      | 5'-CAA GCA GCA GCT GGA GAG AG-3' | 5'-CAG TGG AGG CGG TTG GAG ATA GT-3' |
| PC4      | 5'-CTG GGC GTA CTG GAA AGG G-3'   | 5'-CAG CCT TGT GTC GGG AAA GA-3'     |
| PC5/6    | 5'-CAG AAG GCA TGG TGA TGG AG-3' | 5'-AGA GGG CGA GGT CAG TGA A-3'      |
| PC7      | 5'-GCT GTT CAG AGA GGT GTG TG-3'  | 5'-AGG GCT AGA GGT GGT GGT-3'        |
| PACE4    | 5'-TCA AGA GCA GCA GGA AGT GC-3' | 5'-CCT TCG ACA CAC TCT GTG GG-3'     |
| Furin    | 5'-CCA CAG ACC CCA AGT TCT CT-3'  | 5'-GTG CCT CCA CTG CAT GTG C-3'      |
| GATA-2   | 5'-GCG CTG TGG CCT CTA CTA CA-3'  | 5'-GCC TCT GTT CCC TAG CCC AT-3'     |
| c-Myb    | 5'-CAC TGG GAA GGG GAG ACT CAT-3' | 5'-GTG TCA CAT GAC CAG CGT CC-3'     |
| FOG-1    | 5'-CTG TGA AGA AGG ACG CAG C-3'   | 5'-GAC AGG CAG ATG AGC AC             |
| RUNX1    | 5'-AGA AAT CAT TGA GTC CCC CG-3'  | 5'-AGC AGG GAG CAG AAG AGT TT-3'     |
| LMO2     | 5'-ATC GAA AGG AAG AGC CAG C-3'   | 5'-CAG GGG TCC TAT ATG TCA CT TC-3'  |
| NF-E2    | 5'-CAA CTT ACT GCC CCT GCT CA-3'  | 5'-CAA GGA GTG GGG CAT GAG TG-3'     |
| EKLF     | 5'-TGG CCT CTC AGT CTG GTG AG-3'  | 5'-CGG CCG TCT GAA GAC ACC AC-3'     |
| NDRG2    | 5'-GGA TTA GTC ATG GGG GAG GTG G-3' | 5'-GAA TTC ACA GGA GAC CTC CAT GGT-3' |

§Number indicates the reference.

Isolation of cytoplasmic and nuclear fraction

The cells were suspended with 100 μl cold buffer A (25 mM Tris, pH 8.0, 10 mM KCl, 1 mM DTT, 0.5 mM PMSF) and incubated on ice for 15 min. Then, 25 μl 10% NP-40 was added to the sample and mixed by vortexing for 10 sec. After centrifugation for 2 min at 4°C, supernatants were collected as the cytoplasmic fraction. The pellet was suspended in 50 μl buffer C (50 mM Tris, pH 8.0, 400 mM NaCl, 1 mM DTT, 1 mM PMSF) and incubated for 10 min at 4°C on a shaking platform. After centrifugation for 5 min at 4°C, supernatants were collected as the nuclear fraction.

Trypan blue exclusion assay

The cells were cultured in a 24-well plate in RPMI 1640 medium containing 10% FBS and treated with 0.1 μg/ml PMA for 2 days. The cells were harvested every 1 day, stained with 0.4% trypan blue, and live cells were counted using a hemocytometer.

CCK-8 assay

The cells were seeded at a density of 1×10^4 cells/150 μl in 96-well plates in RPMI-1640 medium containing 10% FBS and treated with 0.1 μg/ml PMA for 2 days. Then, 10 μl of CCK-8 solution (Dojindo, Gaithersburg, MD) was added to each well. After incubation for 2~3 h at 37°C, the absorbance at 450 nm was recorded using a VICTOR3™ plate reader (PerkinElmer, Waltham, MA).
Statistical analysis
All the experiments were performed more than three times, The Student’s t-test was used for statistical analysis. The values are represented as the mean±SD. A value of p<0.05 was considered significant.

RESULTS
NDRG2 overexpression prevents cell growth arrest in response to PMA
U937 cells differentiate into monocyte-like cells upon treatment with phorbol 12-myristate 13-acetate (PMA), and their growth is arrested by enhanced expression of PU.1-dependent genes (22). To identify the effect of NDRG2 expression during PMA-induced differentiation in U937 cells, U937-mock and U937-NDRG2 cells were treated with PMA for 48 h. Ectopic expression of NDRG2 was detected only in the cytoplasm of U937 cells (Fig. 1A). After treatment with PMA for 48 h, U937-mock cells attached to culture plate, but U937-NDRG2 cells showed comparatively reduced attachment (data not shown). A trypan blue exclusion assay revealed that both U937-mock and U937-NDRG2 cells showed no significant differences in cell proliferation. However, U937-mock cells did not proliferate more in response to PMA. Interestingly, U937-NDRG2 cells continued to proliferate during PMA-induced differentiation into macrophages (Fig. 1B). In addition, it was also confirmed using a CCK-8 assay that U937-NDRG2 cells maintained a higher level of proliferation than U937-mock cells during PMA-induced differentiation (Fig. 1C). Interestingly, NDRG2 expression significantly inhibited PU.1 expression in U937 cells although PU.1 levels were also weakly increased in U937-NDRG2 cells after treatment with PMA (Fig. 1D). Consistent with these results, CSF-1R expression, which is regulated by PU.1, was not observed in U937-NDRG2 cells after treatment with PMA. Accordingly, these data demonstrate that NDRG2 hinders the differentiation of U937 precursors into macrophages by PMA stimulation and promotes continued cell growth through the suppression of PU.1 expression.

GATA-1 expression is induced by NDRG2 overexpression
PU.1, which is suppressed by NDRG2 expression in U937 cells, has a reciprocal interaction with GATA-1 during hematopoiesis. Thus, we examined whether NDRG2 induces GATA-1 expression in U937 cells. As shown in Fig. 2A and

Figure 1. Overexpression of NDRG2 in U937 cells. (A) Cell lysates were fractionated into cytoplasmic and nuclear components. Western blotting using an anti-NDRG2 antibody was performed for each fraction. Lamin A/C and α-actinin were used as control for protein loading as well as marker proteins for nuclear and cytoplasmic fraction, respectively. (B, C) U937-mock and U937-NDRG2 cells were treated with 0.1 μg/ml PMA for 48 h. Cell viability was determined using either the trypan blue exclusion assay (B) or the CCK-8 assay (C). The results represent the means±SD of duplicates. *p<0.05, **p<0.01. (D) PU.1, CSF-1R, and NDRG2 mRNA levels were measured by RT-PCR. β-actin was used as a loading control.
B, when U937-mock and U937-NDRG2 cells were treated with PMA for 48 h, GATA-1 expression at both the mRNA and the protein levels was only detected in PMA-treated U937-NDRG2 cells. These data suggest that although PU.1 levels in U937-NDRG2 cells are weakly induced after stimulation with PMA, GATA-1 expression is remarkably enhanced by NDRG2 expression.

GATA-1 induction by NDRG2 correlates with the gene expression patterns marking differentiation into erythrocytes/megakaryocytes

Because GATA-1, which is increased by NDRG2 overexpression in response to PMA, is essential for differentiation of erythrocytes and megakaryocytes, we explored whether NDRG2 controls the expression levels of other genes that regulate erythroid/megakaryocyte differentiation processes. It is well known that a number of factors help to promote the differentiation of erythrocytes and megakaryocytes, including FOG-1, NF-E2, RUNX1, SCL/tal, LMO2, EKLF, and c-Myb (23,24). U937-mock and U937-NDRG2 cells were treated with 0.1 μg/ml PMA and gene expression was analyzed after 48 h using RT-PCR. Interestingly, the expression of these factors was strongly inhibited by treatment with PMA in U937-mock cells but was maintained in U937-NDRG2 cells. Moreover, EKLF expression was newly induced in U937-NDRG2 cells. Additionally, GATA-2, which is often replaced by GATA-1 during hematopoiesis, disappeared in U937-NDRG2 cells. Collectively, these results indicate that NDRG2 can help to maintain the expression of other genes involved in promoting the differentiation of erythrocytes and megakaryocytes.

JAK2/STAT signaling is activated by NDRG2 expression

To better define the mechanism that regulates GATA-1 expression by NDRG2, we investigated whether NDRG2 can control the activation of the JAK2/STAT pathway, which is involved in GATA-1 expression. After a 48 h treatment with 0.1 μg/ml PMA, protein production and mRNA expression of related genes were analyzed in U937-mock and U937-NDRG2 cells. As shown in Fig. 3A, phosphorylation of JAK2 and STAT1 was markedly increased by NDRG2 expression and PMA stimulation. The phosphorylation of STAT3 and STAT5 was severely impaired by PMA treatment in U937-mock cells, whereas phosphorylation was maintained in U937-NDRG2 cells, SOCS family proteins serve as negative regulators of the...
JAK/STAT pathway. U937-mock cells showed an increased expression of SOCS3 after treatment with PMA, whereas U937-NDRG2 cells did not (Fig. 3B and C). Other SOCS family proteins did not show any significant alterations by NDRG2 expression. Together, our findings suggest that NDRG2 activates JAK2/STAT signaling, which can mediate GATA-1 expression in U937 cells in response to PMA.

BMP-4/Smad signaling is activated by NDRG2

To evaluate other mechanisms by which NDRG2-induced GATA-1 expression is regulated, we further examined whether NDRG2 can control the activation of the BMP-4/Smad pathway that is associated with the induction of GATA-1 expression. After 48 h of treatment with 0.1 μg/ml PMA, Western blot analysis was performed. NDRG2 expression promoted the production of active BMP-4 (37 kDa) in PMA-treated U937 cells (Fig. 4A). Consistent with BMP-4 production, phosphor-
regulation of Smad1/5/8, which can be induced by active BMP-4, was also detected only in PMA-treated U937-NDRG2 cells. These results demonstrate that NDRG2 can activate the BMP-4/Smad pathway when cells are stimulated with PMA.

**NDRG2 regulates the expression of PC and BMPR to modulate the BMP-4/Smad signaling pathway**

BMP-4, similar to other TGF-β family members, binds to type I and type II serine/threonine kinase receptors. Type I receptors are activated by phosphorylated type II receptors to initiate intracellular signals through the activation of Smad proteins (25-27). We therefore evaluated the effects of NDRG2 on BMP receptor (BMPR) expression in response to PMA. Type I receptors include BMPRIA, BMPRIB, and ActRI, and Type II receptors include BMPRII, ActRII, ActRIIB. As shown in Fig. 4B, BMPRIA expression was inhibited by treatment with PMA in U937-mock cells but increased in U937-NDRG2 cells. The expression of ActRII and ActRII was enhanced by NDRG2 expression. Although the expression of ActRI and ActRIII was detected in U937-mock cells in response to PMA, it was still higher in U937-NDRG2 cells, Expression of BMPRIA, BMPRII, and ActRIIB was highly induced in PMA-treated U937-NDRG2 cells. These data show that NDRG2 expression generally increases expression of BMPR in U937 cells and also suggest that phosphorylation of Smad1/5/8 may be induced by the increased BMPR activation in PMA-treated U937-NDRG2 cells.

The activity of BMP-4 is regulated by pro-protein convertase (PC) family members. Nine members of the PC family, including PC1/3, PC2, PC4, PC5/6, PC7, PACE4, Furin, SKI-1/S1P, and PCSK9, have been identified (28). The seven PCs, except for SKI-1/S1P and PCSK9, cleave pro-proteins at single or pairs of basic residues in the Golgi, secretory granules, cell surface, or endosomes. In particular, PC5/6, PC7, PACE4, and Furin are known to cleave pro-BMP-4 (29-31). As shown in Fig. 4C, expression of PC2, PC4, and PC5/6 increased with NDRG2 expression. Although the expression of PC2 and PC4 was detected in U937-mock cells in response to PMA, their expression was still higher in U937-NDRG2 cells. Furin was expressed only in PMA-treated U937-NDRG2 cells. These data suggest that NDRG2 expression increases the expression of PC to promote production of active BMP-4.

**NDRG2 modulates GATA-1 expression through regulation of the JAK2/STAT pathway but not through regulation of the BMP-4/Smad pathway**

To identify which pathway regulates NDRG2-mediated GATA-1 expression, U937-NDRG2 cells were pre-treated with a JAK2 inhibitor (AG490), a BMP-4 antagonist (Noggin), or a Smad1/5/8 inhibitor (dorsomorphin). As shown in Fig. 5A, phosphorylation of JAK2 was suppressed by AG490 treatment in U937-NDRG2 cells. Phosphorylation of Smad1/5/8 was not significantly altered by JAK2 inhibition in U937-NDRG2 cells. Importantly, PMA-induced GATA-1 expression was inhibited by pre-treatment with AG490 in U937-NDRG2 cells, PMA-induced Smad1/5/8 phosphorylation was decreased by pre-treatment with dorsomorphin (DM) or noggin (Fig. 5B and C). Phosphorylation of JAK2 was not regulated by treatment with dorsomorphin or noggin. Interestingly, PMA-induced GATA-1 expression was not inhibited by pre-treatment with DM or

![Figure 5](image-url)
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noggin in U937-NDRG2 cells. Together, these data suggest that NDRG2 increases GATA-1 expression through the regulation of the JAK2/STAT pathway, but not through regulation of the BMP-4/Smad1/5/8 pathway, in response to PMA.

DISCUSSION

NDRG2 belongs to the NDRG family, a new characterized group of differentiation-related genes composed of four family members (13). NDRG2 has been characterized as a regulator of dendritic cell differentiation from monocytes, CD34+ progenitor cells, and myelomonocytic leukemic cell line (14,15). Recently, it was suggested to be a novel intrinsic factor for the modulation of IL-10 production in myeloid cells (20). However, the role of NDRG2 in the expression and activation of transcription factors in hematopoietic cells remains poorly understood.

Since Friend et al. (32) first demonstrated an induction of murine virus-induced erythroleukemic cell (MEL) differentiation in vitro in the early 1970s, several groups have used this system to study hematopoiesis. We previously reported that NDRG2 induces BMP-4 production in MDA-MB-231 breast tumor cells (17). Thus, we hypothesized that NDRG2 expression may regulate the expression of other genes that promote the differentiation of erythrocytes and megakaryocytes. Therefore, further studies of how NDRG2 controls gene expression involved in hematopoiesis dependent or independent of GATA-1 are warranted.

It has been suggested that various genes are involved in GATA-1 expression during embryonic developmental processes and hematopoiesis. We previously reported that NDRG2 expression activates the STAT3 pathway upon treatment with PMA for up to 60 min in U937 cells (21). Furthermore, NDRG2 increased BMP-4 production in MDA-MB-231 breast tumor cells (17). Thus, we hypothesized that NDRG2 expression increases GATA-1 expression through the regulation of the JAK2/STAT or BMP-4/Smad signaling pathway. While U937-mock cells did not express GATA-1, U937-NDRG2 cells showed an increased GATA-1 expression in response to PMA after 48 h (Fig. 2). NDRG2 did not appear to form a complex with GATA-1, which was induced by NDRG2 overexpression in response to PMA (data not shown). It has previously been reported that NDRG2 is downregulated by myc (13). GATA-1 is able to promote repression of the c-myc proto-oncogene during erythroid maturation (33). Thus, it is conceivable that a positive feedback loop exists between GATA-1 and NDRG2.

There are many proteins capable of controlling the differentiation of erythroid and megakaryocytic cells. Most of these genes, including NF-E2, RUNX1, LMO2, c-Myb, and FOG-1, were expressed in both U937-mock and U937-NDRG2 cells. However, their expression was inhibited by treatment with PMA in U937-mock cells but maintained in U937-NDRG2 cells. Moreover, the EKLF gene, which is specific for erythroid lineages, was only expressed in U937-NDRG2 cells, and RUNX1, which is specific for megakaryocyte lineages, was stably expressed even after treatment with PMA. Because both EKLF and RUNX1 were modulated by NDRG2 in U937 cells, the type of lineage specifically regulated by NDRG2 still remains unclear, GATA-2, which promotes proliferation of progenitor cells during early hematopoiesis and is suppressed by GATA-1 directly during hematopoiesis (34), was detected in U937-mock cells but not in U937-NDRG2 cells (Fig. 3). Based on these results, we speculate that NDRG2 expression can regulate the expression of other genes that promote the differentiation of erythrocytes and megakaryocytes. Therefore, further studies of how NDRG2 controls gene expression involved in hematopoiesis dependent or independent of GATA-1 are warranted.

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late cleavage of pro-BMP-4 to active BMP-4, increased due to NDRG2 expression (Fig. 4C). Together, these findings suggest that NDRG2 expression can induce PC expression, produce active BMP-4, and phosphorylates Smad1/5/8 through increased BMPR activity in U937 cells.

To identify the signal transduction pathways by which NDRG2 induces GATA-1 expression, we analyzed JAK-STAT and BMP4-SMAD pathways using specific inhibitors, PMA-induced GATA-1 expression was suppressed by pre-treatment with AG490, a JAK2 inhibitor, in U937-NDRG2 cells (Fig. 5A). However, dorsomorphin (DM), a Smad1/5/8 inhibitor, and noggin, a BMP-4 antagonist, did not have any inhibitory effect on PMA-induced GATA-1 expression in U937-NDRG2 cells (Fig. 5B and C). Recently, the cross-talk between the BMP and JAK/STAT signaling pathways has been a focus of research to better understand hematopoiesis, embryonic development, and leukemia development (5). When the relationship between the BMP-4/Smad and JAK/STAT pathways was evaluated using specific inhibitors in PMA-treated U937-NDRG2 cells, there was no significant effect on activation of either pathway (Fig. 5), raising the possibility that these systems act independently.

In summary, we report in this study that NDRG2 induces GATA-1 expression through activation of the JAK2/STAT signaling pathway, but not through BMP-4/Smad signaling, in PMA-treated U937 cells. These results suggest that NDRG2 may serve as a regulator in erythrocyte and megakaryocyte differentiation during hematopoiesis. Further study is required to identify the effects of NDRG2 expression on the differentiation of erythroid/megakaryocytic lineages in primary hematopoietic stem cells and in vivo. Moreover, study of the effects of NDRG2 expression on myeloid cell differentiation through the activation of the BMP-4/Smad pathway also requires further elucidation.

CONFLICTS OF INTEREST

The authors have no financial conflict of interest.

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