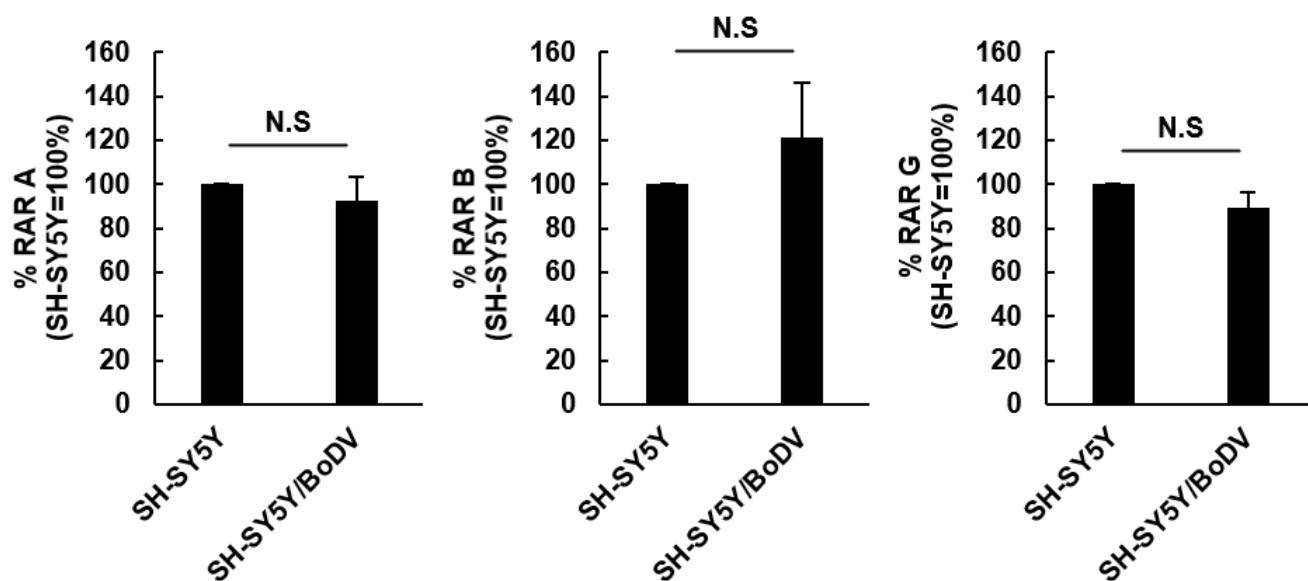
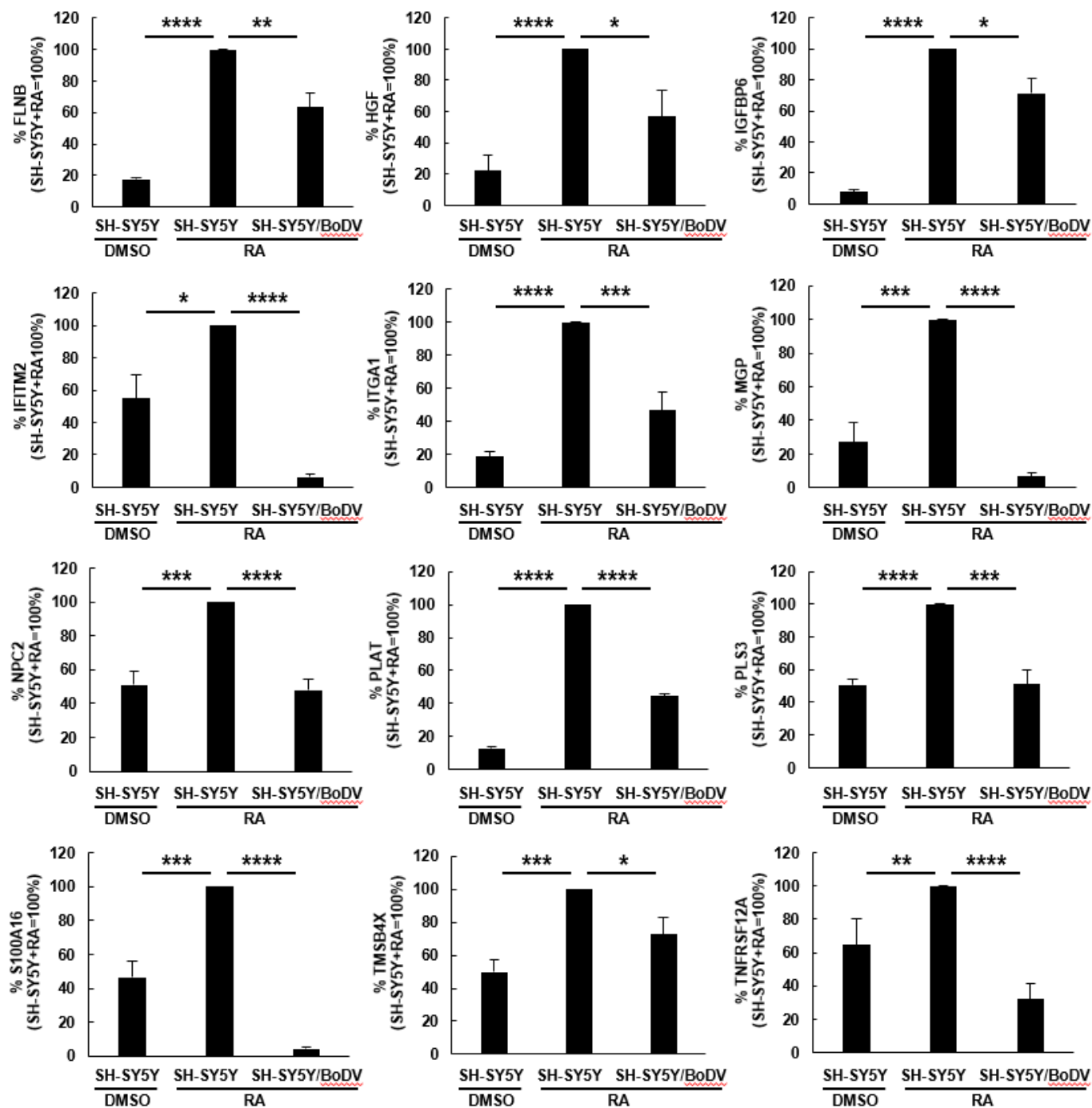


**Table S1.** List of primers and probes used in this study.

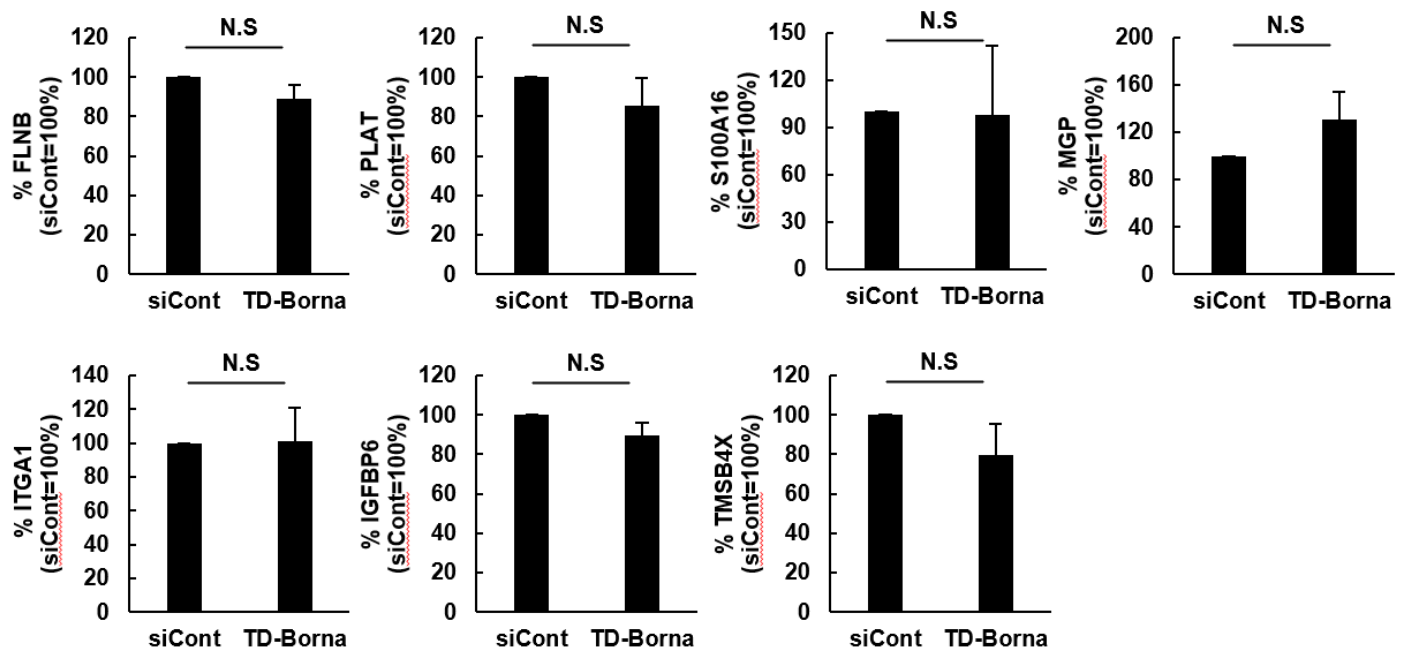
Primer	Sequence 5'to3'	Description
NCAM2-F	CTTATGTCTGCAGGGCCACA	Real-time RT-qPCR
NCAM2-R	TGCTGCCCTTTGACTTCGAT	Real-time RT-qPCR
RARA-F	CATTGAGACCCAGAGCAGCA	Real-time RT-qPCR
RARA-R	CCGGGTCACCTTGTTGATGA	Real-time RT-qPCR
RARB-F	GCACCAGGTATACCCAGCA	Real-time RT-qPCR
RARB-R	GTCCGTTCTCAAGGTCCTG	Real-time RT-qPCR
RARG-F	CAAGGTGACCAGGAATCGCT	Real-time RT-qPCR
RARG-R	GTCTCCTGATGGGCTTTGCT	Real-time RT-qPCR
MGP-F	GCCGCTTAGCGGTAGTAAC	Real-time RT-qPCR
MGP-R	AGCGTTCTCGGATCCTCTCT	Real-time RT-qPCR
HGF-F	ACCCTGGTGTTCACAAGCA	Real-time RT-qPCR
HGF-R	GCAAGAATTTGTGCCGGTGT	Real-time RT-qPCR
S100A16-F	GCAGTCATTGTCCTGGTGGA	Real-time RT-qPCR
S100A16-R	GATGAGCTTATCCGCAGCCT	Real-time RT-qPCR
NPC2-F	CTGGCAGCTACATTCTGCT	Real-time RT-qPCR
NPC2-R	CACGGCCTTGCTGCTTTTAG	Real-time RT-qPCR
ITGA1-F	GGCAGCACAATTCATGCACA	Real-time RT-qPCR
ITGA1-R	AAATGTACACAGCTCCCCCG	Real-time RT-qPCR
FLNB-F	GGAGTTGCTCGTGGAAGACA	Real-time RT-qPCR
FLNB-R	CCTGCAGCTTTGGTGTC AAC	Real-time RT-qPCR
TGM2-F	GGCGAACCACTGAACAAAC	Real-time RT-qPCR
TGM2-R	CAGGTACTTGGTGCCACACT	Real-time RT-qPCR
IGFBP6-F	AACCGCAGAGACCAACAGAG	Real-time RT-qPCR
IGFBP6-R	TGGTCACAATTGGGCACGTA	Real-time RT-qPCR
PLAT-F	TACAGCCAGCCTCAGTTTCG	Real-time RT-qPCR
PLAT-R	TTCTGCCCAAGATCACCGTC	Real-time RT-qPCR
PLS3-F	ACTCTGCTTCTGCCATTGGG	Real-time RT-qPCR
PLS3-R	TCGGAGTAAAGCAGCCAAGG	Real-time RT-qPCR
Primer	Sequence 5'to3'	Description
TMSB4X-F	CTTCGCTTTTCTCCGCAAC	Real-time RT-qPCR
TMSB4X-R	ATTTCAGTGTCTGCCACCC	Real-time RT-qPCR
IFITM2-F	AGCAGGAAGTGGCTATGCTG	Real-time RT-qPCR
IFITM2-R	CACGGAGTACGGAATGCTA	Real-time RT-qPCR
TNFRSF12A-F	CGGACCTGGACAAGTGCAT	Real-time RT-qPCR
TNFRSF12A-R	GTCTCTCTATGGGGGTGGT	Real-time RT-qPCR
GAPDH-F	AGCGAGATCCCTCCAAAATC	Real-time RT-qPCR
GAPDH-R	AAATGAGCCCCAGCCTTCTC	Real-time RT-qPCR
BoDV-1 PmRNA/gRNA probe	FAM-AGAACCCCTCCATGATCTCAGACCCAGA-TAMRA	Real-time RT-qPCR (Hayashi et al., 2009)
BoDV-1 gRNA-specific RT primer for huP2Br strain	TGTTGCGCTAACAACAAACCAATCAC	Real-time RT-qPCR (Hayashi et al., 2009)
BoDV-1 P mRNA/gRNA-forward primer for huP2Br strain	ATGCATTGACCCAACCAAGTC	Real-time RT-qPCR
BoDV-1 P mRNA/gRNA-reverse primer for huP2Br strain	ATCATTCGACAGCTGCTCCCTTC	Real-time RT-qPCR
BoDV-1 N-forward primer (OU277)	GGAGCCGAGCAGATCAAGAA	Real-time RT-qPCR
BoDV-1 N-reverse primer (OU278)	CACAAAGGAGCCTACCCAGG	Real-time RT-qPCR
BoDV-1 L probe	FAM-CGAGGCATCCGTGGTCAGCAGAT-TAMRA	Real-time RT-qPCR
BoDV-1 L-forward	GGAAGCGCCCCGTGTT	Real-time RT-qPCR
BoDV-1 L-reverse primer (OU283)	CCCCCAGAGTGATTCGCTTA	Real-time RT-qPCR



**Figure S1.** Comparable expression of receptors for RA in SH-SY5Y and SH-SY5Y/BoDV cells. The mRNA amounts of the RA receptors were evaluated by real-time RT-qPCR. We calculated the expression of each gene by the DDCT method. The expression of each gene was normalized with that of the *GAPDH* gene in each condition. Values are expressed as the mean percentage + standard error of mean (SEM) of three independent experiments. N.S, non-significant.



**Figure S2.** Confirmation of RNA-seq results using real-time RT-qPCR analyses. The mRNA amounts of differentiation-related genes that were downregulated by BoDV-1 infection in our RNA-seq analysis were evaluated by real-time RT-qPCR. We calculated the expression of each gene by the DDCT method. The expression of each gene was normalized with that of the *GAPDH* gene in each condition. Values are expressed as the mean percentage  $\pm$  SEM of at least three independent experiments. \*,  $P < 0.05$ ; \*\*,  $P < 0.01$ ; \*\*\*,  $P < 0.005$ ; \*\*\*\*,  $P < 0.001$ .



**Figure S3.** Effect of TD-Borna on BoDV-1-related transcriptomic changes in differentiated SH-SY5Y/BoDV cells. RA-treated SH-SY5Y/BoDV cells were treated with the scrambled siRNA (siCont) or TD-Borna. The mRNA amounts of differentiation-related genes that were downregulated by BoDV-1 infection were evaluated by real-time RT-qPCR. We calculated the expression of each gene by the DDCt method. The expression of each gene was normalized with that of the *GAPDH* gene in each condition. Values are expressed as the mean percentage + SEM of at least three independent experiments. N.S, non-significant.