

***GRIN3A* and *MAPT* stimulate nerve overgrowth in macrodactyly**

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Abstract. As an uncommon and congenital condition, macrodactyly is characterized by an increase in the size of all the elements or structures of the digits or toes; however, the underlying pathogenesis remains to be fully elucidated. In the present study, the gene expression profiles of abnormal nerves were examined in three patients with macrodactyly using microarray analysis to identify potential genes contributing to nerve overgrowth. Gene expression profiling in the nerve tissue samples were scanned using the microarray and the differentially expressed genes were verified at the transcription level using reverse transcription-quantitative polymerase chain reaction analysis. Western blot analysis was used to determine the expression of target genes at the translational level. To confirm the upregulated genes during the process of nerve proliferation, SH-SY5Y cells were induced to differentiate into a neuronal-like phenotype using retinoic acid. A total of 165 genes showed significant changes (≥ 5 -fold) in gene expression, which may be associated with the development of limbs in macrodactyly. Glutamate ionotropic receptor NMDA 3A (*GRIN3A*) and microtubule-associated protein tau (*MAPT*) were identified as important contributors in promoting nerve overgrowth. Furthermore, it was identified that *GRIN3A* and *MAPT* were regulated by the cAMP-protein kinase A and extracellular signal-regulated kinase 1/2 pathways, respectively. The identification of genes expressed at high levels in macrodactyly may reveal potential factors, which contribute

to abnormal nerve proliferation and underpin the pathogenesis of macrodactyly, and provide potential application targets in nerve tissue regeneration engineering.

Introduction

Macrodactyly is an uncommon congenital condition characterized by an increase in the size of all the elements or structures of the digits or toes, including phalanges, tendons, vessels, subcutaneous fat and finger nails. The malformation often occurs unilaterally or asymmetrically and affects more than one digit or toe. The clinical conditions associated with this deformity are carpal tunnel syndrome, syndactyly, neurofibromatosis type 1 cafe au lait spots, lipoma and nevi (1-3).

The malformations in the fingers or toes are considered to be associated with other syndromes of macrodactyly, including vascular malformations, multiple enchondromatosis, maffucci syndrome, tuberous sclerosis and neurofibromatosis type 1 (4,5). The exact pathogenesis underlying this condition and, in particular, the role of nerve growth stimulators in macrodactyly, remains to be fully elucidated. The present study aimed to determine nerve overgrowth stimuli using gene expression profiling was performed in malformed enlarged nerve tissues from the digits or toes of patients with macrodactyly. A total of six overexpressed (>10 -fold) genes, including creatine kinase, mitochondrial 2 (*CKMT2*), vasoactive intestinal peptide (*VIP*), FXYD domain-containing ion transport regulator 3 (*FXYD3*), Glutamate ionotropic receptor NMDA 3A (*GRIN3A*), *GSTT1* and microtubule-associated protein tau (*MAPT*) were identified for their potential contribution to abnormal nerve overgrowth. In addition, *MAPT* and *GRIN3A* were identified as key regulators of nerve outgrowth. These nerve growth stimulators may contribute to nerve regeneration and reconstruction following nerve injury.

Materials and methods

Patients. The present study was reviewed and approved by the Ethics Committee of the First Hospital of Jilin University, (Changchun, China). Three male patients between the ages of 17 and 25 were recruited from the First Hospital of Jilin University between May 2011 and September 2013. All enrolled patients underwent surgical management for isolated nonsyndromic macrodactyly. Normal nerve tissue samples, which served as healthy controls, were obtained from five

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Table I. Primer sequences used in reverse transcription-polymerase chain reaction analysis.

Gene	Accession no.	Forward primer	Reverse primer
<i>CKMT2</i>	NM_001825.2	5'-GCCGCAATGCTTCTCTG-3'	5'-GGCCATCCCAGCACAT-3'
<i>VIP</i>	NM_003381.3	5'-TCCTTGTGCTCCTGACTC-3'	5'-CTGCTCCTCTTTCCATTC-3'
<i>FXYP3</i>	NM_005971.3	5'-GTGACCCTGGGCCTGCT-3'	5'-CAGCTTTGGGCTGAGCCT-3'
<i>GRIN3A</i>	NM_133445.2	5'-GGAGGTAGATGATGAAGGC-3'	5'-AAACAAGAGGGCATAACAG-3'
<i>GSTT1</i>	NM_000853.2	5'-ATCTTTGCCAAGAAGAACG-3'	5'-TGTGAGGACCAGTAAGGAAG-3'
<i>MAPT</i>	NM_016835.4	5'-GCCAAAGGGCAGGATG-3'	5'-TTCGGGAAGTGACAGAAGAG-3'
<i>GAPDH</i>	NM_002046	5'-GGAGTCAACGGATTGGTC-3'	5'-CCCCAGCCTTCTCCAT-3'

CKMT2, creatine kinase, mitochondrial 2; *VIP*, vasoactive intestinal peptide; *FXYP3*, FXYP domain-containing ion transport regulator 3; *GRIN3A*, glutamate ionotropic receptor NMDA 3A; *GSTT1*, glutathione S-transferase θ 1; *MAPT*, microtubule-associated protein tau.

patients undergoing elective surgery for unrelated reasons, including road traffic accidents or fires. Written informed consent was obtained from guardians on the behalf of all participants.

Tissue treatment and reverse transcription-quantitative polymerase chain reaction (RT-qPCR) analysis. A sample (~10 mg) of enlarged nerve tissue from a digit or toe was obtained from each of the patients with macrodactyly, which were sectioned into smaller sections (~3 mm³), snap frozen in liquid nitrogen and stored at -80°C until further use. The frozen tissues were homogenized in cold normal saline (0.9% NaCl), using a ULTRA-TURRAX Tube Drive Workstation (IKA Werke GmbH & Co. KG, Staufen, Germany). Next, RNA was extracted from the nerve tissue samples using TRIzol (Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA USA) according to the manufacturer's protocol. A total of 500 ng RNA was used for reverse transcription was performed using the GoScript Reverse Transcription system (Promega Corporation, Madison, WI, USA). Next, the products were amplified using Power SYBR Green Master Mix (containing SYBR Green I Dye, AmpliTaq Gold® DNA Polymerase, dNTPs, passive reference and optimized buffer) on an ABI 7300 (Applied Biosystems; Thermo Fisher Scientific, Inc.). RT-qPCR was performed with an initial 10 min at 95°C followed by 40 cycles of 95°C for 15 sec, and 60°C for 1 min. All experiments were repeated three times. Relative gene expression was calculated with the $2^{-\Delta\Delta C_q}$ method (6) following normalization to the expression of GAPDH. The primers used for PCR are listed in Table I.

Cell culture. The SH-SY5Y cell line was obtained from the Cell Bank of the Chinese Academy of Sciences (Shanghai, China), and cultured in Dulbecco's modified Eagle's medium supplemented with 10% fetal bovine serum (all from Gibco; Thermo Fisher Scientific, Inc.) and 1% penicillin/streptomycin (Sigma-Aldrich; Merck Millipore, Darmstadt, Germany) at 37°C with 5% CO₂. The cells were sub-cultured when they reached 85-90% confluence and then seeded at a density of 5x10³ cells/cm² into 12-well plates (Nest Scientific USA, Rahway, NJ, USA) for 24 h at 37°C. The cells were then induced to differentiate into neuronal cells by treating the cells with 10 μ M retinoic acid for 72 h at 37°C (7). Subsequently,

the differentiated cells were treated with protein kinase A (PKA) inhibitor H89 or extracellular signal-regulated kinase (ERK)1/2 inhibitor PD98059 (Beyotime Institute of Biotechnology, Haimen, China) for up to 4 h at 37°C.

Western blot analysis. The cells were collected and treated with lysis buffer (Beyotime Institute of Biotechnology) supplemented with 1% protease inhibitor mixture (Sigma-Aldrich; Merck Millipore). Subsequently, cell lysates were centrifuged at 10,000 x g for 15 min at 4°C. Protein quantification was performed using BCA Protein Quantification kit (Thermo Fisher Scientific, Inc.) and 30 μ g protein per lane were separated on 10% SDS PAGE gel and then transferred onto PVDF membranes (Invitrogen; Thermo Fisher Scientific, Inc.). The membranes were incubated with 5% skimmed dry milk for 30 min and washed three times with TBST (10 mM Tris, 150 mM NaCl and 0.1% Tween-20). Next, the membranes were incubated with rabbit anti-human GRIN3A polyclonal antibody (cat. no. sc-98986; 1:5,000) and rabbit anti-human MAPT polyclonal antibody (cat. no. sc-32828; 1:1,000) at 4°C overnight. Then the goat anti-rabbit IgG-HRP secondary antibody (cat. no. sc-2302; 1:1,000) was used to incubate the membranes for 2 h at 37°C. All antibodies were purchased from Santa Cruz Biotechnology, Inc. (Dallas, TX, USA). Densitometry scores were determined using Quantity One software, version 4.6.9 (Bio-Rad Laboratories, Inc., CA, USA).

Microarray. The RNA from the nerve tissue samples was hybridized and scanned using the Agilent Microarray Scanner (Agilent Technologies, Inc., Santa Clara, CA, USA) at the National Engineering Center for Biochip (Shanghai, China), according to the manufacturer's protocol. The raw data were obtained using Feature Extraction 10.7 software (Agilent Technologies, Inc.) with default settings and normalized using the Quantile algorithm in Gene Spring 11.0 software (Agilent Technologies, Inc.).

Spring 11.0 software. Gene Ontology (GO) functional annotation clustering analysis was used to process the microarray data. The GO project database (geneontology.org/) was used following the criteria of the Database for Annotation, Visualization and Integrated Discovery (david.abcc.ncifcrf.gov/)

Table II. Downregulated genes (>10-fold) in the macrodactyly tissue samples.

Gene symbol	Fold change	Description
<i>ABL2</i>	0.0901	Similar to <i>ABL1</i> ; associated with cytoskeletal rearrangement
<i>CCL20</i>	0.0571	Chemotactic factor; associated with inflammation
<i>CDC14B</i>	0.0523	Serine/threonine phosphatase; dephosphorylate p53
<i>LAMA1</i>	0.0674	α subunit of laminin
<i>IL1B</i>	0.0740	Inflammatory cytokine
<i>LIF</i>	0.0613	Inhibits embryonic stem cell differentiation; promotes germ cell proliferation
<i>MBP</i>	0.0945	Myelin sheath component
<i>MFSD2</i>	0.0872	Mediates syncytin-2-dependent cell-cell fusion
<i>NR4A1</i>	0.0764	Nuclear transcriptional activator
<i>OSM</i>	0.0558	Inhibits tumor proliferation; regulates cytokines
<i>PLAUR</i>	0.0825	Degrades the extracellular matrix
<i>SPATA13</i>	0.0836	Activates Rho-associated pathway; spermatogenesis-associated
<i>SOD2</i>	0.0947	Superoxide dismutase;
<i>SOX11</i>	0.0828	Antigen in B lymphomas; tumor suppressor

ABL2, ALB proto-oncogene 2, non-receptor tyrosine kinase; *CCL20*, chemokine (C-C motif) ligand 20; *CDC14B*, cell division cycle 14 B; *LAMA1*, laminin subunit α 1; *IL1B*, interleukin 1 β ; *LIF*, leukemia inhibitory factor; *MBP*, myelin basic protein; *MFSD2*, major facilitator superfamily domain-containing 2; *NR4A1*, nuclear receptor subfamily 4, group A, member 1; *OSM*, oncostatin M; *PLAUR*, plasminogen activator, urokinase receptor; *SPATA13*, spermatogenesis-associated 13; *SOD*, superoxide dismutase.

Table III. Upregulated genes (>10-fold) in the macrodactyly tissue samples.

Gene symbol	Fold change	Description
<i>CKMT2</i>	10.4	Creatine kinase
<i>VIP</i>	12.6	Neuropeptide, belongs to a glucagon/secretin superfamily
<i>FXYP3</i>	11.2	Na, K-ATPase ion channel regulator
<i>GRIN3A</i>	10.8	Ionotropic glutamate receptor
<i>GSTT1</i>	13.4	Catalyzes the conjugation of reduced glutathione to electrophilic moieties of xenobiotics or endogenous compounds
<i>MAPT</i>	12.7	Microtubule-associated protein tau

CKMT2, creatine kinase, mitochondrial 2; *VIP*, vasoactive intestinal peptide; *FXYP3*, FXYP domain-containing ion transport regulator 3; *GRIN3A*, glutamate ionotropic receptor NMDA 3A; *GSTT1*, glutathione S-transferase θ 1; *MAPT*, microtubule-associated protein tau.

for grouping genes according with the relevant biological processes, molecular functions and cellular component categories.

Statistical analysis. Statistical analyses were performed using GraphPad software, version 5.0 (GraphPad, Inc., La Jolla, CA, USA). All data are expressed as the mean \pm standard deviation. Student's *t*-test was performed to analyze the results of the gene expression profiling assays. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

Gene expression profile analysis in macrodactyly. The present study first examined the gene expression profile in nerve tissue samples from three patients with macrodactyly. Of the 29,378 genes analyzed, 22 genes were upregulated (≥ 5 -fold)

and 143 genes were downregulated (≥ 5 -fold) in the macrodactyly samples, compared with genes in the normal control samples (Fig. 1A). In particular, 14 genes were downregulated (Table II) and six genes were upregulated >10-fold in the macrodactyly samples (Table III). The genes upregulated in the samples from the patients with macrodactyly were considered as possible targets, which may promote nerve overgrowth.

The differentially expressed genes were grouped according to their biological process using Gene Ontology annotation (Fig. 1B). In the patients with macrodactyly, the majority of the differentially expressed genes were involved in metabolic process (27.5%), response to stimulus (20.1%) and immune system process (16.9%).

Confirmation of genes, which may contribute to nerve overgrowth. The upregulated genes identified in the present study (≥ 10 -fold) may promote nerve overgrowth, therefore, these

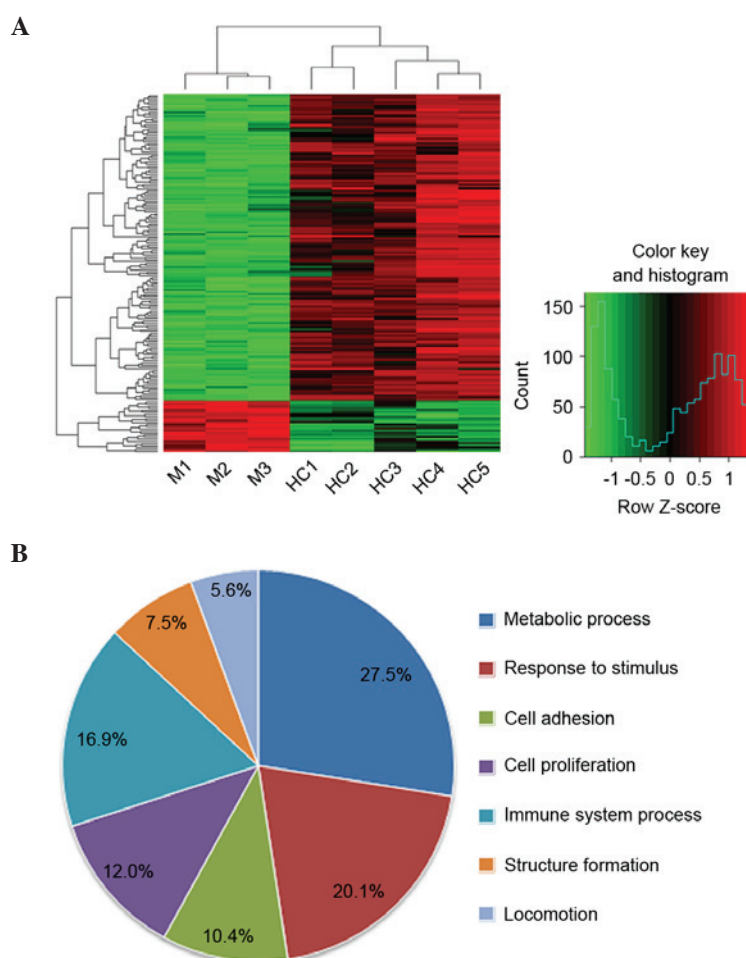


Figure 1. Gene expression profile analysis. (A) Hierarchical clustering of differentially expressed genes in macrodactyly. Dendrogram indicates gene expression profile using microarray analysis of three patients with macrodactyly (M1, M2 and M3) and five healthy controls (HC1, HC2, HC3, HC4 and HC5). Red represents upregulated genes and green represents downregulated genes. The heatmap shows 165 transcripts filtered with a fold change threshold ≥ 5 , including 22 upregulated and 143 downregulated genes associated with nerve overgrowth. (B) GO term analysis of differentially expressed genes in macrodactyly. Expression data were obtained from microarray analysis of transcripts with ≥ 5 -fold expression change and $P < 0.05$, and grouped into GO terms. The pie chart shows that 27.5% of all GO terms identified were broadly grouped into metabolic process, 20.1% were involved in the response to the stimulus and 16.9% were involved in immune system process. GO, Gene Ontology.

target genes were further confirmed at the transcriptional level. The results showed that, of the six candidate genes, *VIP*, *FXRD3*, *GRIN3A* and *MAPT* were upregulated, which was consistent with the microarray data (Fig. 2A). To further confirm the upregulation of the above genes, retinoic acid was used to induce the neuronal differentiation of SH-SY5Y cells (Fig. 2B), and the mRNA levels of *VIP*, *FXRD3*, *GRIN3A* and *MAPT* in the SH-SY5Y cells were measured using RT-qPCR analysis. As presented in Fig. 2C and D, the transcriptional expressions of *GRIN3A* and *MAPT* were increased along with the induction. Subsequently, the translational levels of these two genes (Fig. 2E and F) were determined and the results also confirmed the result that *GRIN3A* and *MAPT* were upregulated along with the process of nerve axon growth during differentiation of the SH-SY5Y cells.

Signaling pathways responsible for the upregulation of target genes. To investigate the signaling pathways and events, which may stimulate nerve cell proliferation, the SH-SY5Y cells were induced by retinoic acid for 72 h. Retinoic acid has been shown to induce cell differentiation

by sustained phosphorylation of ERK1/2 or cAMP response element binding protein (8-10). Therefore, PD98059 and H89 were added to the cells to inhibit the ERK1/2 and cAMP/PKA pathways, respectively. The results showed that at transcriptional and translational level, the expression of *GRIN3A* was regulated by the cAMP/PKA pathway, whereas the expression of *MAPT* was affected by the ERK1/2 pathway (Fig. 3A-D).

Discussion

Macrodactyly is an uncommon congenital nervous system disease, and is characterized by the proliferation of nerve fibers, vessels, subcutaneous fat and other tissues (1-3). The pathogenesis of macrodactyly remains to be fully elucidated, therefore, the identification of genes expressed at high levels in macrodactyly may reveal factors, which promote nerve overgrowth and contribute to the pathogenesis of this condition.

In the present study, the gene expression profiles were compared between patients with macrodactyly and healthy

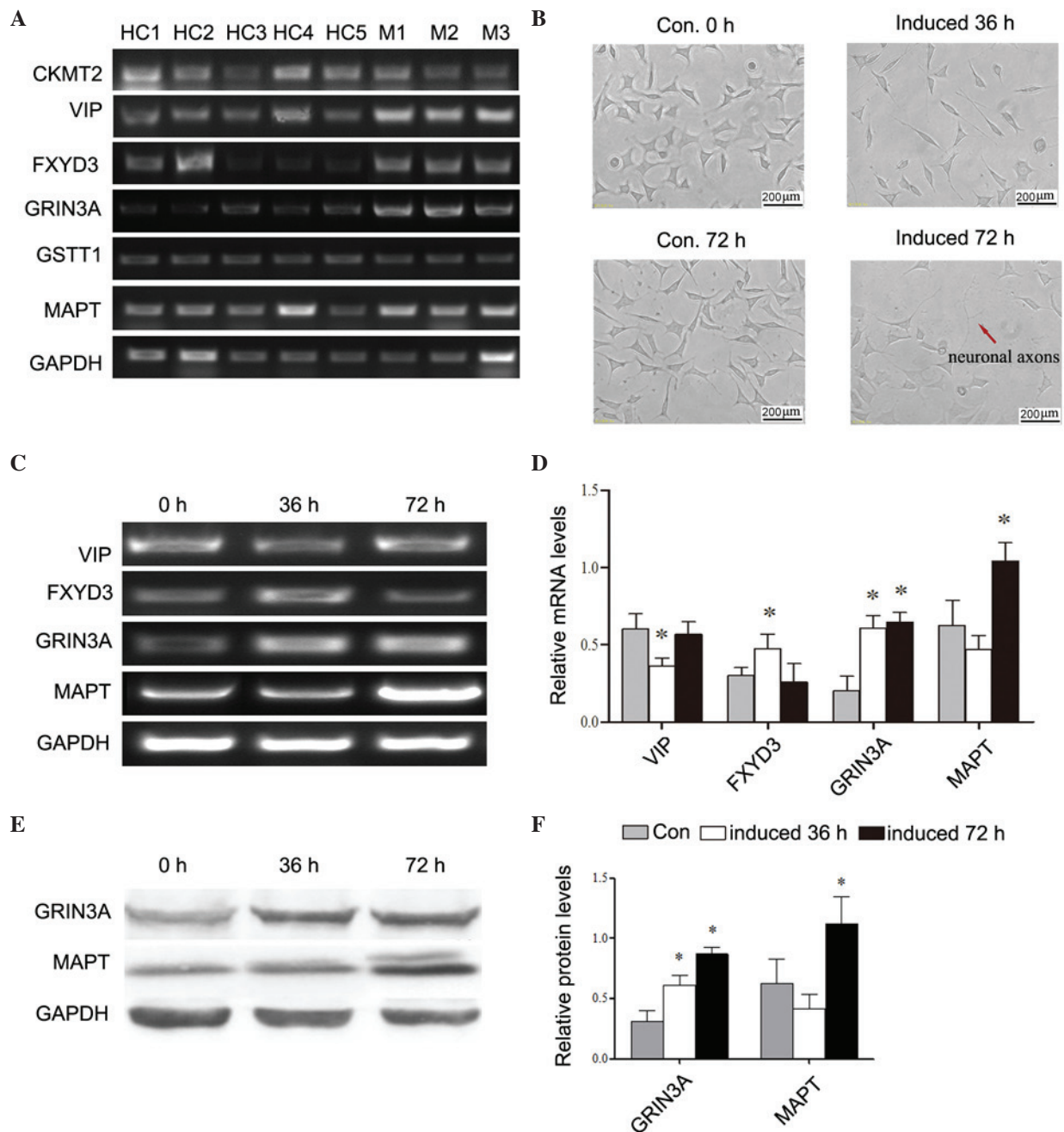


Figure 2. Analysis of mRNA levels of candidate genes associated with nerve outgrowth. mRNA levels were measured to confirm the expression profile of the six candidate genes (*CKMT2*, *VIP*, *FXYD3*, *GRIN3A*, *GSTT1* and *MAPT*). (A) mRNA levels of *CKMT2*, *VIP*, *FXYD3*, *GRIN3A*, *GSTT1* and *MAPT* were determined with *GAPDH* as the reference. (B) Differentiation of SH-SY5Y cells induced by retinoic acid for 72 h. Scale bar=200 μ m. (C) Confirmation of selected genes following cell differentiation. (D) Histogram showing results of the analysis of relative mRNA levels of target genes. (E) Western blot analysis of translational levels following differentiation. (F) Quantification of western blotting. * $P < 0.05$, vs. Con. *CKMT2*, creatine kinase, mitochondrial 2; *VIP*, vasoactive intestinal peptide; *FXYD3*, FXYD domain-containing ion transport regulator 3; *GRIN3A*, glutamate ionotropic receptor NMDA 3A; *GSTT1*, glutathione S-transferase $\theta 1$; *MAPT*, microtubule-associated protein tau; Con, control.

controls. It was found that a significant number of genes (143; 87%) were downregulated (≥ 5 -fold), whereas only 22 (13%) genes were upregulated (≥ 5 -fold). Among the downregulated genes, apoptosis-associated factors, including B cell lymphoma 2, cell division cycle 14 B and pentraxin 3, and microtubule formation associated promoters, including *MAP1*, *MAPT* and sorting nexin 22, were downregulated, as expected. Of note, pro-inflammatory cytokines or associated receptors, including interleukin (*IL*) 1β , *IL1* receptor (*R*), *IL7R* and *IL8*, and the regulators of cell-cell or cell-extracellular

matrix interactions, including chemokine (C-C motif) ligand 20, C-X-C chemokine receptor type 4, integrin α X, integrin β 8, major facilitator superfamily domain-containing 2, plasminogen activator, urokinase and selectin L, were significantly downregulated. It is known that inflammatory cytokines and cell adhesion molecules are frequently over-expressed in human cancer, and are involved in oncogenesis and metastasis by improving extracellular matrix degradation (11-15). The results of the present study indicated that cell invasion and metastasis may be restrained in macrodactyly,

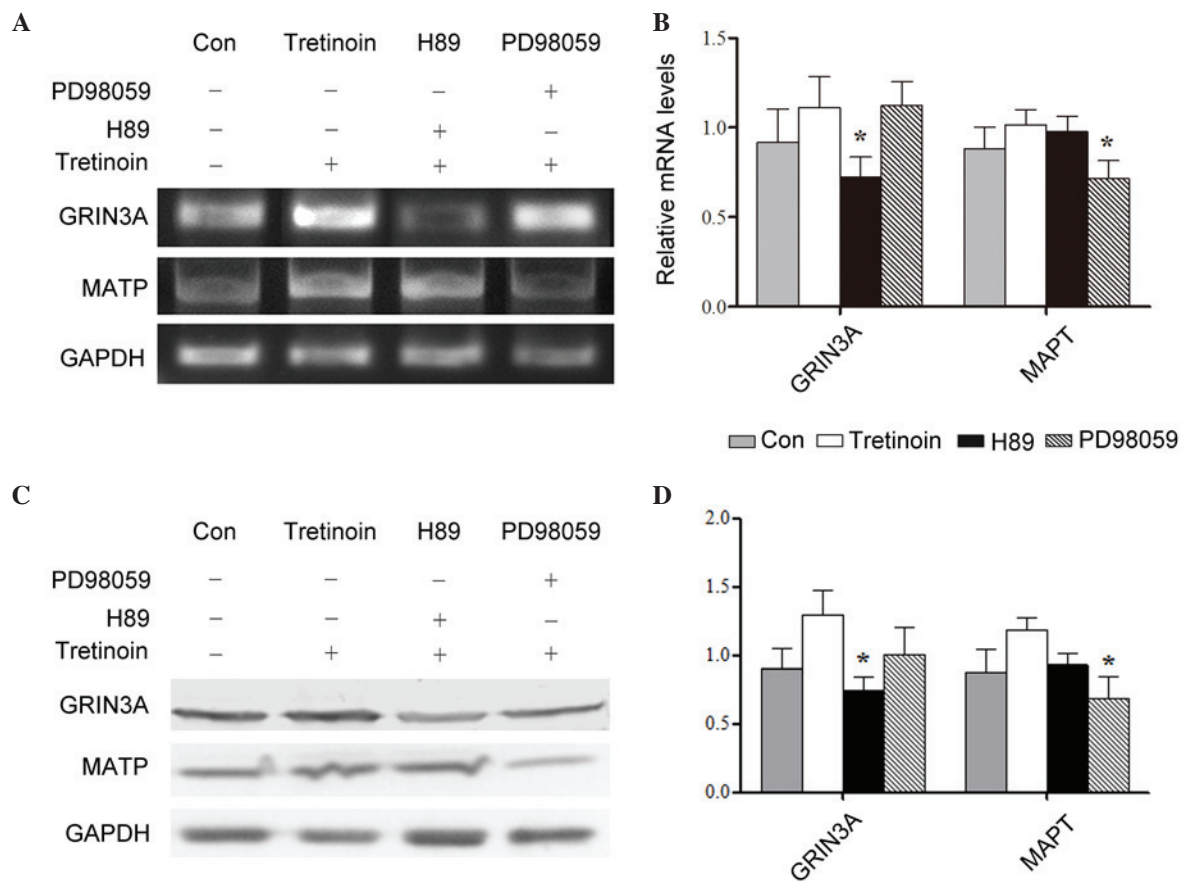


Figure 3. Signaling pathways relevant to nerve overgrowth. SH-SY5Y cells were induced by retinoic acid, and treated with PD98059 and H89 to inhibit extracellular signal-regulated kinase 1/2 and cAMP/protein kinase A pathways, respectively. Expression levels of *GRIN3A* and *MAPT* were detected using (A and B) reverse transcription-quantitative polymerase chain reaction and (C and D) western blotting analyses. * $P < 0.05$, vs. retinoic acid (tretinoin) and Con groups. *GRIN3A*, glutamate ionotropic receptor NMDA 3A; *MAPT*, microtubule-associated protein tau; Con, control.

and this may explain why macrodactyly seldom develops to malignancy.

In addition, the present study found that angiogenesis-associated positive regulators, including vascular endothelial growth factor (*VEGF*)A/B/C, *Notch*, δ -like 4 and macrophage migration inhibitory factor, were not expressed at high levels, and a number even showed decreased expression, including fibroblast growth factor receptor 1, *microRNA 2* (*mir21*) and *mir221*. It is well known that vessel overgrowth is common in tumors, particularly in malignancy (16,17). Therefore, the present study hypothesized that the abnormal vascular proliferation observed in macrodactyly is not due to hyperplastic nerve tissue directly.

The genes, which identified as being upregulated in the patients with macrodactyly were considered to be involved in promoting nerve overgrowth. The present study found 14 down-regulated (>10 -fold) genes and six upregulated (>10 -fold) genes (*CKMT2*, *VIP*, *FXRD3*, *GRIN3A*, *GSTT1* and *MAPT*) in the macrodactyly samples. Subsequently, the upregulated levels of *VIP*, *FXRD3*, *GRIN3A* and *MAPT* were confirmed using RT-qPCR analysis. *GRIN3A* and *MAPT* have been shown to be involved in stimulating nerve growth. *MAPT*, is involved in improving nerve proliferation, and is expressed at high levels in multiple neurodegenerative disorders, including progressive supranuclear palsy, Parkinson's disease and Alzheimer's disease (18,19). As an ionotropic glutamate receptor, *GRIN3A* is involved in regulating nerve signal transduction (20,21).

Nerve growth is a result of the combined effects of multiple genes and signaling networks, thus selecting positive regulators from a nerve outgrowth model may be a promising strategy for identifying key factors and signaling pathways. To investigate the signaling pathways responsible for promoting nerve growth, retinoic acid was used to induce the differentiation of SH-SY5Y cells (7). By inhibiting the ERK1/2 pathway and cAMP-PKA pathway, which are involved in neurite proliferation, it was found that *GRIN3A* showed a significant correlation with the cAMP-PKA pathway, whereas *MAPT* was affected by the ERK1/2 pathway.

Increased expression levels of *MAPT* and *GRIN3A* may lead to the abnormal pathologic condition of nerve tissue in macrodactyly and, to a certain extent, may have potential applications in nerve regeneration. It is known that axon continuity is always interrupted during nerve damage, and the distal axonal tract finally undergoes degeneration. Thus, identifying techniques to reconstruct the structure of nerve tissue, and restore its function following nerve injury and repair has been one of the key areas of investigation in tissue engineering. Gene therapy is one potential approach in the treatment of traumatic nerve injury. Various genes are being considered as candidates for promoting nerve regeneration, including *VEGF* (22,23). Based on the findings from the present study, *MAPT* and *GRIN3A* warrant further examination as two potential factors, which may contribute to nerve regeneration and reconstruction following nerve injury.

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