



OPEN TSC complex decrease the expression of mTOR by regulated miR-199b-3p

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The TSC complex formed by TSC1 and TSC2 is the most important upstream negative regulator of mTORC1. Genetic variations in either *TSC1* or *TSC2* cause tuberous sclerosis complex (TSC) disease which is a rare autosomal dominant disorder resulting in impairment of multiple organ systems. In this study, besides a reported variation, c.2509_2512del (p.Asn837Valfs*11, p.N837fs) in *TSC1*, we found a *de novo* *TSC2* variation c.1113delG (p.Gln371Hisfs*18, p.Q371fs), which these two mutation influence the formation of TSC complex. We found that the decrease of TSC complex with the appearance of the decreased miR-199b-3p expression. At the same time, the reduction of miR-199b-3p increased the expression of mTOR and the activation of mTORC1 and mTORC2, the additional miR-199b-3p caused the decrease the expression of mTOR and the activation of mTORC1 and mTORC2. In brief, our results may illustrate a novel mechanism of TSC caused by variations in either *TSC1* or *TSC2*, and a new mTOR expression regulator, miR-199b-3p.

Keywords TSC1, TSC2, MTOR, MiR-199b-3p, TSC

TSC (Tuberous sclerosis complex) (OMIM#191100#613254), with a morbidity of near to 1:6,000–1:10,000 in the world population, is an autosomal dominant disorder that multiple organ systems affected including skin, kidneys, brain, lung and heart¹. The prevalence of TSC is independent of nations, ethnicity and sex, however, some studies found that female patients with TSC have a higher incidence of renal disease and lymphangioleiomyomatosis burden than male patients². Among patients with TSC, more than 80% patients develop multiple and bilateral angiomyolipomas in the kidneys which are the leading cause of adult deaths with this disease³. TSC is usually caused by germline loss-of-function variations in either *TSC1* or *TSC2*, encoding hamartin (TSC1) and tuberlin (TSC2), respectively^{4,5}. The complex of TSC1 and TSC2 inactivates the small GTP-binding protein Rheb to negatively regulate mammalian target of rapamycin complex 1 (mTORC1), an important regulator of cellular biosynthesis⁶, which is now recognized as a classical model of TSC pathogenesis⁷. Once activated, mTORC1 phosphorylates the downstream effectors, such as p70S6K and eukaryotic initiation factor 4E-binding protein 1 (4EBP1), to promote translation initiation⁸. mTOR is a component of two functionally and structurally distinct complexes, one is rapamycin-sensitive mTOR complex 1 (mTORC1), the other is rapamycin-insensitive mTOR complex 2 (mTORC2)⁹. mTORC2 is known to control cell survival and proliferation⁸.

miRNAs are small, non-coding RNA molecules about 19 to 25 nucleotides which involve in the regulation of gene expression during post-transcriptional stages by pairing to the target mRNA, thereby promoting the degradation of target mRNA or decreasing its translational efficiency¹⁰. Many miRNAs are also involved in the pathogenesis of various malignancies that have been evidenced by many studies¹¹. It has been demonstrated that miR-199b is associated with the expression of mTOR^{12,13}, but the molecular mechanism is unclear. *Barbara et al.* reported that TSC2 might regulate the biogenesis of microRNA in mouse embryonic fibroblasts (MEF), which indicates that the microRNA may play an important role in the pathogenic mechanism of TSC¹⁴.

In this study, we reported two variations, include a *de novo* mutation (c.1113delG) in *TSC2* and one reported mutation (c.2509_2512del) in *TSC1* that caused TSC in two independent families. Our results showed that TSC2-p.Q371fs influenced the cell proliferation and apoptosis in fibroblast from the patient. Furthermore, the TSC complex may regulate the protein level and activity of mTOR through the manipulation of the level of intracellular miR-199b-3p, which may a novel mechanism of TSC caused by mutation either in *TSC1* or *TSC2* and expands our understanding of TSC pathogenesis.

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Materials and methods

Patients

This study was approved by the Ethics Committees of Shanxi University and the ethics number is SXULL20190049, written informed consent was obtained from all patients and healthy volunteers involved in this study.

Sequencing analysis

Whole-exome sequencing and validation by PCR were performed as described in our previous studies¹⁵.

Cell culture

Skin tissue samples were isolated from a TSC patient (22 years old, female) with the variation c.1113delG in *TSC2* and two healthy volunteers (female, 23 years old and female, 25 years old). Fibroblasts of skin tissues separated and then cultured in DMEM (Boster) with 10% FBS (Gibco), 100 U/mL penicillin and 0.1 mg/mL streptomycin and grown in 5% CO₂ at 37 °C as described before¹⁶.

HEK293T cells were maintained in DMEM (Boster) supplemented with 10% FBS (BI). Cells were transfected with plasmids in this study using Polythermide (PEI) (PolyScience, Cat.No.23966-2) as described before¹⁷.

The plasmids for overexpression and shRNA

The plasmids pcDNA3-Flag-TSC2-WT and pcDNA3-Myc3-hTSC1-WT were purchased from Addgene. The plasmids pcDNA3-Flag-TSC2-MT containing mutant *TSC2* (c.1113delG) and pcDNA3-Myc3-hTSC1-MT containing mutant *TSC1* (c.2509_2512del) were constructed using site-specific mutagenesis. Lentiviral plasmids expressing shRNA specific to either *TSC1* (NM_000368.5) or *TSC2* (NM_00548.5) were constructed using the targeting sequences 5'-AAAGAAGAAGCTGCAATATCT-3'¹⁸ and 5'-GCTCATCAACAGGCAGTTC TA-3'¹⁹ respectively.

Lentiviruses were harvested in HEK293T cells 72 h post co-transfection. The co-transfection of lentiviral plasmid vectors, psPAX and pMD2.G was performed at the ratio of 5:5:1. The medium containing lentiviruses was collected and then purified by centrifugation at 1000×g and filtration through 0.22 µm filter. Cells were transformed with the medium containing the lentiviruses and fresh medium at the ratio of 3:1 and cultured for 72 h, then drug selection with puromycin (1 mg/ml) was performed for 7–10 days to get stable transformed cell lines.

qRT-PCR analysis

The total RNA of fibroblasts from the patient or the healthy volunteers were extracted using RNAiso Plus (TaKaRa, Cat#9109) following the instructions of the supplier. First-stand cDNA for evaluated mRNA was generated using PrimerScriptTM RT Master Mix (TaKaRa, Cat.No.RR0-36A) and cDNA for miRNA was generated using TanscriptmiRNA First-Strand cDNA Synthesis SuperMix (TransGen Biotech, Lot#N10824). Three primer pairs for qRT-PCR of *TSC2* were applied for quantification of cDNA located in either 5' or 3' terminal and the middle of the mRNA of *TSC2*. The primer pairs for mTOR and VEGF were obtained from Primer Bank (<https://pga.mgh.harvard.edu/primerbank/>).

Western blot

The total protein of fibroblasts was extracted by radio-immunoprecipitation buffer (RIPA, Solarbio, R0020) with 1 mM PMSF, 1 mM dithiothreitol and phosphatase inhibitors followed the manufacturers' instructions. Total protein concentration was measured using a PierceTMBCA protein assay kit (Thermo).

The co-immunoprecipitation was performed as the protocol of IP/Co-IP kit (Absin, Cat. No. abs955). The antibodies involved in this study were as follows: *TSC1* monoclonal antibody (Stanta Cruz, Cat.No.sc-377386), *TSC2* monoclonal mouse antibody (Stanta Cruz, Cat. No. sc-271314), *TSC2* monoclonal rabbit antibody (Cell Signaling, Cat.No.#4308), mouse IgG (Stanta Cruz, Cat.No.sc-2025), mouse anti Myc-tag monoclonal antibody (ABclonal, Cat.No.AE010), DDDDK tag (flag) mouse monoclonal antibody (Proteintech, Cat.No.66008-3-Ig), mTOR (phosphor-S2448) polyclonal antibody (Bioworld, Cat.No.BS4706), Beta actin Polyclonal antibody (Absin, Cat.No.abs132001), 4E-BP1 (53H11) rabbit monoclonal antibody (Cell Signaling, Cat.No.#9644), Phospho-4E-BP1 rabbit monoclonal antibody (Thr37/46) (236B4) (Cell Signaling, Cat.No.#2855), p70S6 Kinase (49D7) rabbit monoclonal antibody (Cell Signaling, Cat. No. #2708), Phospho-p70S6 Kinase rabbit monoclonal antibody (Ser371) (Cell Signaling, Cat. No. #9208), goat anti-rabbit IgG (H + L), AKT mouse monoclonal antibody (Santa cruz, Cat.No.sc-5298), p-AKT mouse monoclonal antibody (Ser 473) (Santa cruz, Cat.No.sc-293125). Secondary antibody, HRP Conjugate (BOSTER, Cat. No. BA1054), HRP-conjugated affinipure goat anti-mouse IgG (Proteintech, Cat.No.SA00001-1).

Running Native gel for the detection of TSC complex was performed using the HEK293T cells. Post-transfection, cells were collected using non-denature lysis buffer on ice. The soluble fraction was then separated by centrifugation at 12,000×g for 10 min at 4 °C. The supernatants were harvested and added with 5×protein non-denaturation loading buffer. All the running buffers and the native gels were without SDS.

Immunofluorescence analysis

Cells grown on coverslips were fixed by 4% paraformaldehyde at 37 °C for 10 min. After permeabilized with 0.5% Triton and X-100 for 15 min, cells were blocked in 5%BSA (Bull Serum Albumin, Solarbio) for 1 h at room temperature. Then the cells were incubated in primary antibody overnight at 4 °C. After washing, the cells were incubated with the secondary antibody for 1 h at room temperature. The cells staining by DAPI (Sigma) after washing, finally mounted on glass slides using gelatin. Images were acquired using a Delta Vision Image Restoration Microscope with a ×63 objective (DeltaVisionElite, GE).

Measurement of apoptosis

Cell apoptosis assays were performed by incubation with Fas-ligand (Peprotech, Cat: #310-03 H-10VG) at 1 µg/ml for 24 h, then extra incubation with 5 µg/ml CHX (Cycloheximide) (MCE, Cat.No. HY-12320) and 50 µg/ml TNFα (MACS, Lot. No. 51809100455) for 24 h. Finally, cells were treated with trypsin (without EDTA) and collected for apoptosis analysis by flow cytometry using AnnexinVPE/7-AAD Apoptosis detection Kit (Biotect, Cat.No.20180426) according to the manufacturer's instructions.

Cell proliferation assay

The fibroblast proliferation activity was measured by Cell Counting Kit-8 (CCK-8, BOSTER, Cat.No. AR1160). Fibroblasts were seeded in 96-well plate at the density of 1,000 cells per well and cultured for 1–7 days depending assay schedules, at every detection time point of cell proliferation, cells were fed with fresh medium with the addition of 10 µl CCK-8 solutions for 1 h at 37 °C, optical densities were measured on a micro plate reader at 450 nm.

Luciferase reporter assay

HEK293T cells were seeded into 24-well plates at a density of 1.5 × 10⁵ per well. The cells were co-transfected psiCheck-2 with insertion of the 3' UTR of mTOR and miR-199-3p inhibitors or negative control. The analysis of luciferase activities was using the TransDetect Double-Luciferase Reporter Assay Kit (Transgen, Beijing, China).

Statistical analysis

All the data are showed as mean ± SD from independent experiments at least three times. The two-tailed Student's *t*-test was applied for the determination of the *p*-values. *p* < 0.05 was considered as significant difference.

Results

Family Pedigree/Patient information

All patients included in this study were recruited from two unrelated Chinese families. The blood samples were obtained for whole-exome sequencing for the detection of the variations in the probands. The information of all the patients was listed in Table 1. The proband in family I was diagnosed as TSC when she was 16 with facial angiofibromas and renal angiomyolipoma. Her daughter was suspected as TSC because of cardiac rhabdomyomas during prenatal ultrasound. The patients in family II were not diagnosed as TSC but the patients in this family were all with epilepsy.

The proband (II-1) of family I (Fig. 1A) showed the typical symptom of TSC-facial angiofibromas (Fig. 1B), however, epilepsy was not present in the proband but in her daughter (III-1). A *de novo* mutation in TSC2 (c.1113delG, p.Gln371Hisfs*18) was detected in this patient as well as her daughter (III-1) (Fig. 1C). The deletion of one base pair caused amino acid change at Gln371, and resulting in frameshift and premature truncation of the protein at the highly conserved region imply that this mutation is very likely to be the major pathogenic cause of TSC (Fig. 1D).

The proband of family II (Fig. 1E) was detected a reported variation (c.2509-2512del, p.Asn837Valfs*11) in TSC1, his mother and his son also detected the same variation in TSC1 by sanger sequencing (Fig. 1F). This variation caused a translational frame shift, resulting in premature truncation at the highly conserved domains of TSC1 (Fig. 1G).

TSC2-Q371fs influence the TSC2 expression and the phenotype of fibroblasts

The heterozygous variation in TSC2 (c.1113delG) resulted in frameshift and a truncated protein (Fig. 2A). The mRNA and protein level of endogenous wild type TSC2 in TSC2-Q371fs fibroblasts were much lower than TSC2-WT fibroblasts (Fig. 2A, B and C). The proliferation assays showed that the proliferation of the fibroblasts with the mutation TSC2-Q371fs from patient was significantly increased compared to TSC2-WT fibroblasts from healthy volunteers (Fig. 2D). The apoptosis analyses revealed that the apoptosis was declined in TSC2-Q371fs fibroblasts (Fig. 2E).

TSC1-N837fs and TSC2-Q371fs impairs the function of TSC complex

The results of qRT-PCR showed that the VEGF and HIF-1α which were regulated by mTOR signaling pathway in fibroblast with mutation TSC2-Q371fs were elevated as reported before²⁰, suggesting that variation (c.1113delG)

Family ID	Patient ID	gender	Gene (nucleotide)	Mutation (nucleotide)	Mutation (protein)	Reference transcript	Mutation reported previously	symptom
Family I	II-1	F	TSC2	c.1113delG	p.Gln371Hisfs*18, p.Q371fs	NM_00548.5	No	facial angiofibromas renal angiomyolipoma
	III-1	F					No	epilepsy cardiac rhabdomyoma
Family II	I-1	F	TSC1	c.2509_2512del	p.Asn837Valfs*11, p.N837fs	NM_000368.5	Yes	epilepsy
	II-1	M					Yes	epilepsy
	III-1	M					Yes	epilepsy

Table 1. The information of the patients involved in this article. The patient symptoms listed in the table are limited to the patient symptoms we have collected and may not be all of the patient's clinical symptoms.

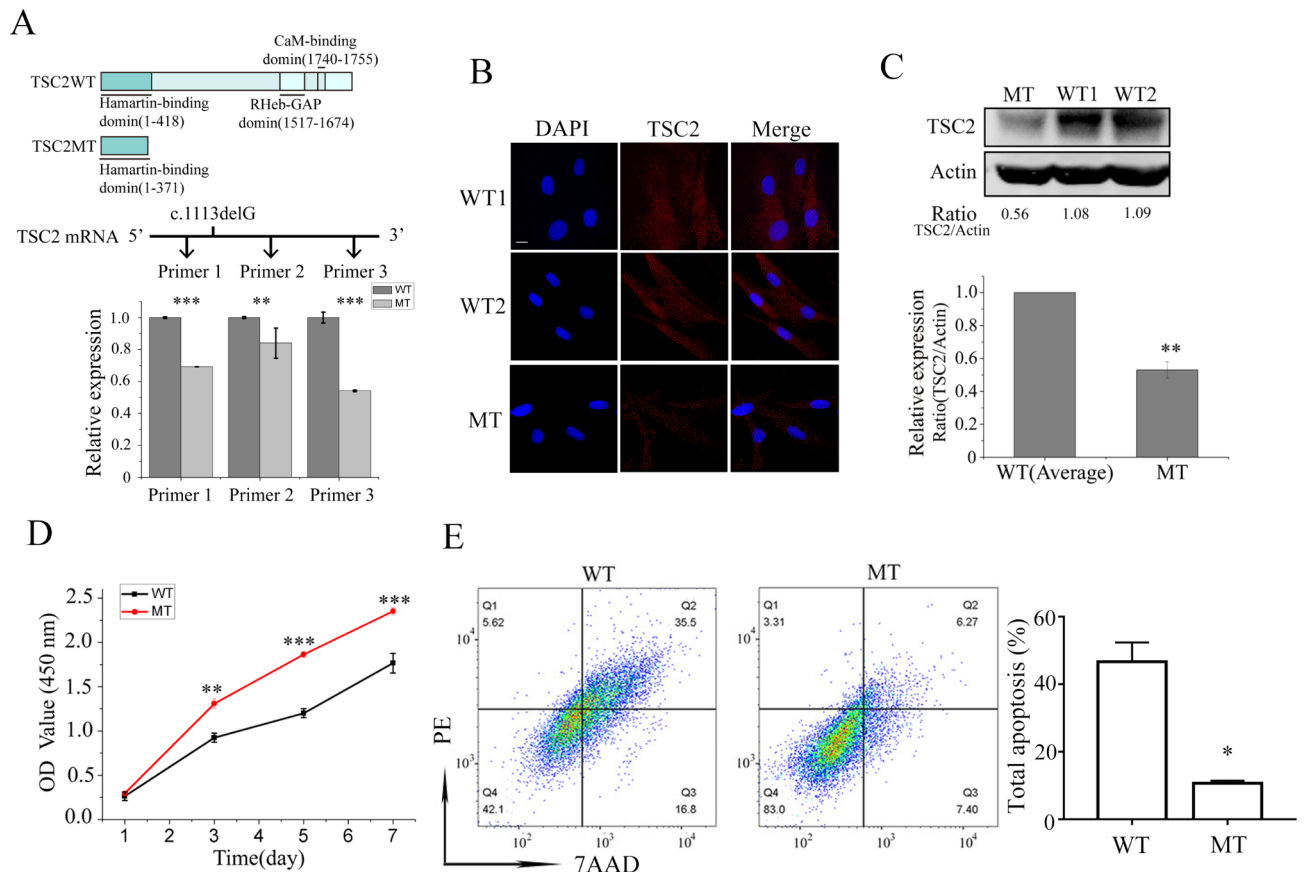


Fig. 1. Family pedigree and sequencing results. **(A, E)** Family pedigree, the families involved in this study. Squares indicate males, circles indicate females, the patients are shown as filled symbols. The proband is indicated by arrows(↗), the individual showed no genotype under the squares or circles indicate the genotype was not tested during this study. **(B)** The facial angiofibromas of the proband in family I. **(C, F)** Sanger sequencing analysis, the gene variation is shown by red arrows. **(D, G)** Evolutionary conservation of amino acid residues altered by the variations across different species.

in TSC2 impairs the function of TSC complex in fibroblast from patient (Fig. 3A). Western blot results revealed that phosphorylation of p70S6K and 4EBP1 were increased, indicating that mTOR was over activated in mutant cells. Supply of TSC2-WT in mutant cells restored the activity of TSC complex, thus less active mTOR subsequently decreased the p70S6K and 4EBP1 phosphorylation (Fig. 3B).

To investigate whether the TSC1-N837fs and TSC2-Q371fs influence the interaction with each other, we constructed plasmids containing with the variations. After the transfection, the co-immunoprecipitation assays suggested that TSC1-N837fs and TSC2-Q371fs can't detected a TSC complex, unlike the overexpressed TSC1-WT and TSC2-WT formed detectable TSC complex respectively, (Fig. 3C and E), and native PAGE showed that the TSC1-N837fs and TSC2-Q371fs failed the formation of TSC complex with the endogenous TSC2 and TSC1 respectively (Fig. 3D and F).

miR-199b-3p directly regulate the expression of mTOR

We found that the expression of mTOR was significantly increased in TSC2-Q371fs fibroblasts compared with TSC2-WT fibroblasts whether in mRNA or protein level (Fig. 4A, B). It has been reported that miR-199b is associated with lower expression of mTOR, we predicted miR-199b-3p at miRbase (<http://www.mirbase.org/>), which may bind to the 3' UTR of mTOR and then promote the mRNA degradation of mTOR (Fig. 4C). We performed qRT-PCR to test miR-199b-3p expressing in fibroblasts from either patient or healthy donors. The results indicated that the expression of miR-199b-3p was dramatically decreased in TSC2-Q371fs fibroblasts compared with that in the TSC2-WT fibroblasts (Fig. 4D). To confirm whether miR-199b-3p directly target the 3'-UTR of mTOR, we co-transfected with miR-199b-3p inhibitor N.C. and mTOR 3' UTR, the luciferase activity had no obvious difference with that transfected with mTOR 3' UTR, but the luciferase activity was significantly increased when co-transfected with miR-199b-3p inhibitor and mTOR 3' UTR (Fig. 4E). These results showed that miR-199b-3p directly target the 3'-UTR of mTOR. Furthermore, the expression of mTOR and phosphorylation proteins level of AKT, and p70S6K was significantly decreased in HEK293T cells after the transfection with miR-199b-3p mimics and was significantly increased upon treatment with the inhibitor

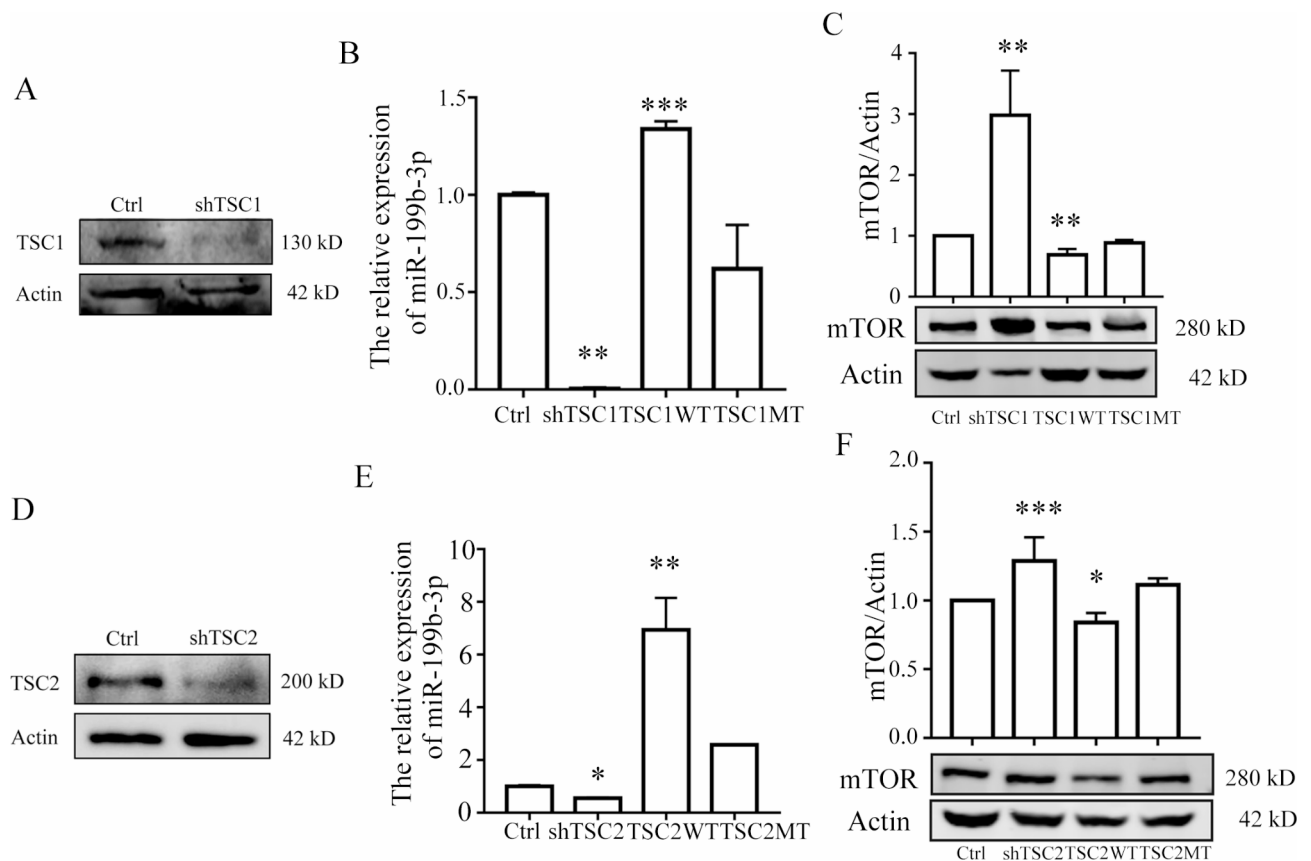


Fig. 2. Expression of TSC2 and phenotypes of the fibroblasts with or without TSC2-Q371fs, the WT was present as the average of all healthy volunteers. (A) Linear schematic of the protein structure of TSC2-WT, TSC2-Q371fs and the location of the primers and the qRT-PCR results, relative expression levels were normalized with *GAPDH*. (B) Immunofluorescence analysis using TSC2 (red) antibody and DAPI (blue) in Bar 15 μ m. (C) The western blot of the endogenous TSC2 and the statistics, the originals gels can be seen in the supplementary figure S1. (D) Proliferation rate of fibroblasts. (E) Apoptosis assay of fibroblasts and the statistics.

specific to miR-199b-3p (Fig. 4E, G, H and I). Taken together, these results suggest that miR-199b-3p may directly regulate the expression of mTOR, and the influence the activation of mTORC1 and mTORC2.

TSC complex regulated the expression of miR-199b-3p

The expression of miR-199b-3p was decreased in TSC2-Q371fs fibroblasts (Fig. 4D). To evaluate whether the TSC complex regulates the expression of miR-199b-3p, we knockdown the expression of either TSC1 or TSC2 for the reduction of TSC complex (Fig. 5A and D), then we performed RT-PCR and found that the expression of miR-199b-3p were down-regulated significantly ($p < 0.01$) in HEK293T-shTSC2 and HEK293T-shTSC1 cell lines (Fig. 5B and E). However, the expression of miR-199b-3p was significantly up-regulated in HEK293T cells with over-expression TSC1-WT or TSC2-WT ($p < 0.001$) whereas the expression of miR-199b-3p was significantly decreased in HEK293T cells with overexpression of TSC1-N837fs or TSC2-Q371fs compared to overexpression of TSC1-WT or TSC2-WT (Fig. 5B and E). These results suggest TSC complex regulates the expression of miR-199b-3p. Opposite with the expression of miR-199b-3p, the expression of mTOR was increased in shTSC1-HEK293T and shTSC2-HEK293T cell lines while decreased in TSC1-WT or TSC2-WT overexpressed cell lines (Fig. 5C and F). Taken together, these results indicated that the content of endogenous miR-199b-3p was regulated by TSC complex.

Discussion

mTORC1 is an important downstream node of TSC complex and the continuous activation of mTORC1 has been reported as the main cause of the disorder Tuberous Sclerosis Complex (TSC). The complex signaling network of mTOR kinase plays a crucial role in cell growth, replication, survival, aging, and metabolism²¹.

In this study, we investigated two variations that involved in TSC. The variation c.2509_2512del in *TSC1* which involved in TSC were reported previously²². But the variation in *TSC2* (c.1113delG) was a *de novo* germline variation which was detected in the proband (II-1) of family 1. Unfortunately, her daughter inherited this variation from her and even showed more serious symptom of TSC. The proband in family I have a severe renal

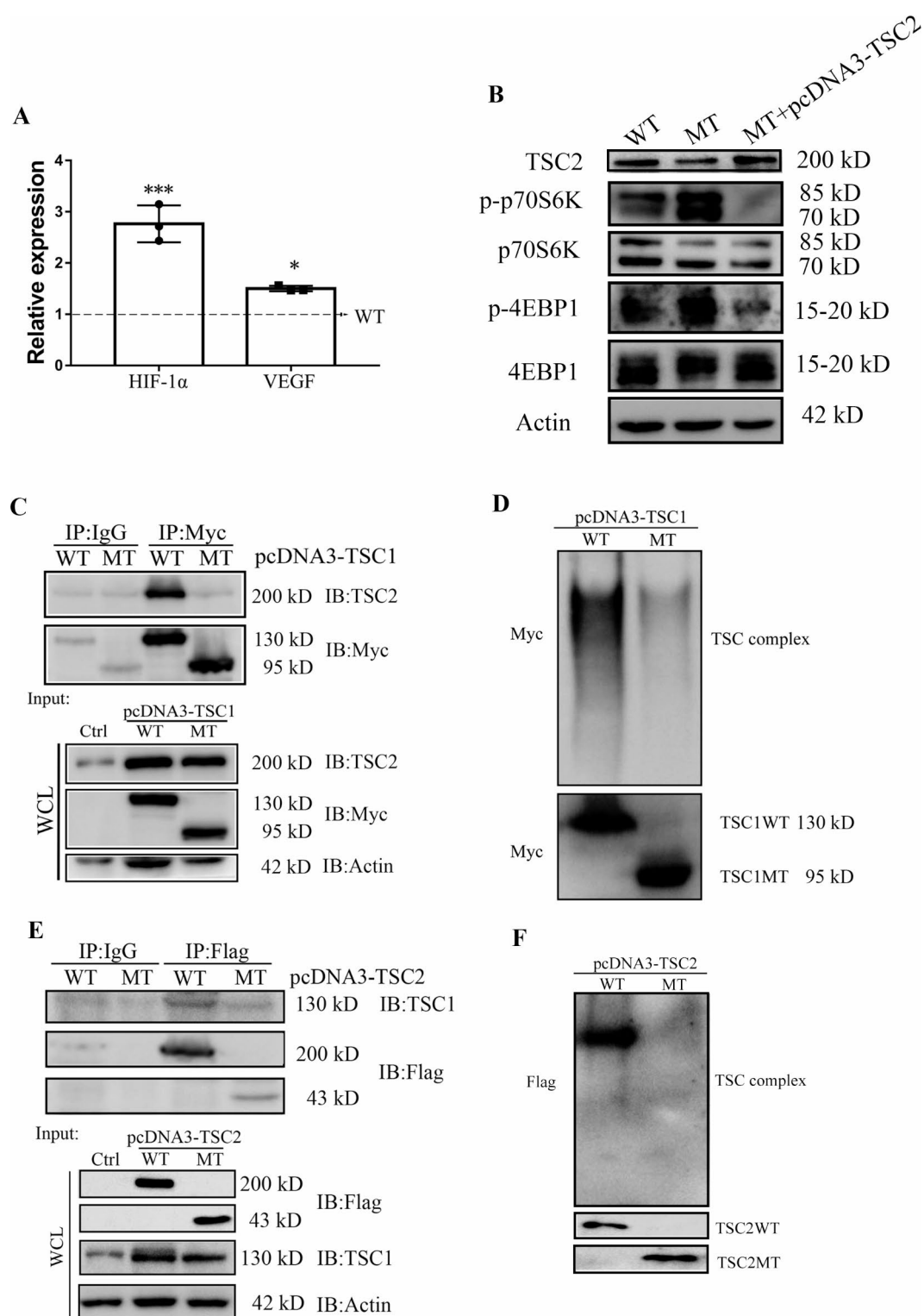
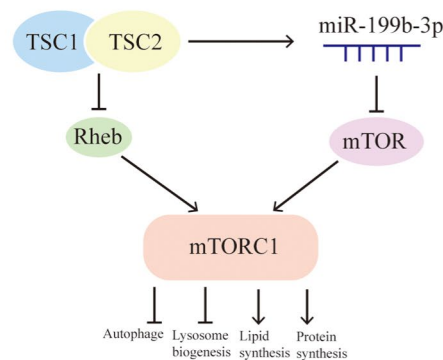


Fig. 3. Functional analysis of the variations (A) Quantification of mRNA level using qRT-PCR, the WT represents the mixture of the equal amount total RNA from the three healthy fibroblasts. Relative expression levels were normalized with *GAPDH*. (B) Evaluation of the rescue experiment by western blot (the WT group was the WT2). The originals gels can be seen in the supplementary figure S2. (C) The co-ip experiment of TSC1-WT and TSC1-N837fs. The originals gels can be seen in the supplementary figure S3. (D) The measurement of TSC complex using non-denaturing PAGE. The originals gels can be seen in the supplementary figure S3 (E) The co-ip experiment of TSC2-WT and TSC2-Q371fs. The originals gels can be seen in the supplementary figure S4 (F) The measurement of TSC complex using non-denaturing PAGE. The originals gels can be seen in the supplementary figure S4.

Physiological model



Pathological model

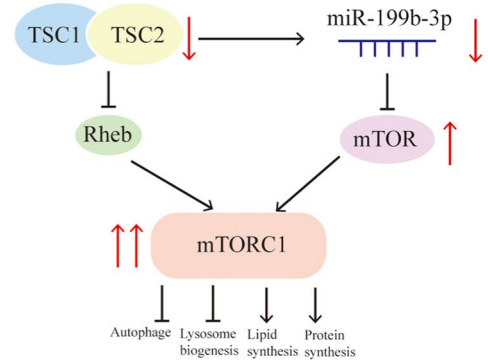


Fig. 4. miR-199b-3p regulate the expression of mTOR (A) Quantitative analysis of mTOR in fibroblasts using qRT-PCR. (B) Quantitative analysis of mTOR in fibroblasts using western blot and the statistics. The originals gels can be seen in the supplementary figure S5. (C) Predicted miR-199b-3p target sequence of the mTOR 3'UTR using miRBase. (D) Quantification of miR-199b-3p and mRNA using qRT-PCR, the WT represents the mixture of the equal amount total RNA from the two healthy fibroblasts. Relative expression levels were normalized with GAPDH. (E). Luciferase reporter assay. (F) Quantitative analysis of mTOR in HEK293T cells at 48 h post-treatment with miR-199b-3p mimics (low panel) and the statistics (up panel). The originals gels can be seen in the supplementary figure S5. (G) Western bolt analysis of p-AKT, AKT, p-p70S6K, p70S6K in HEK293T cells at 48 h post-treatment with miR-199b-3p mimics, β -actin served as a control of total protein. (H) Quantitative analysis of mTOR in HEK293T cells at 48 h post-treatment with the inhibitor specific to miR-199b-3p (low panel) and the statistics (up panel). (I) Western bolt analysis of p-AKT, AKT, p-p70S6K, p70S6K in HEK293T cells at 48 h post-treatment with the inhibitor specific to miR-199b-3p, β -actin served as a control of total protein. The originals gels can be seen in the supplementary figure S5.

angiomyolipoma, this may be because of the abnormal growth of cells. The results of cell proliferation suggested that increased cell growth in TSC2-Q371fs fibroblast was via reduction of cell apoptosis (Fig. 2).

We found that the expression of mTOR in TSC2-WT fibroblasts was less than in TSC2-Q371fs fibroblasts (Fig. 4). It was reported that micro199b might influence the expression of mTOR¹². Online analysis through miRBase (<http://www.mirbase.org/>), we predicted that miR-199b-3p potentially targets 3' UTR of mTOR so that regulates the expression of mTOR post-transcription. To evaluate our hypothesis, we detected the expression of miR-199b-3p in fibroblast, and the results indicated that the expression of miR-199b-3p and mTOR had negative correlation in fibroblasts, at the same time, the activation of the mTORC1 had negative correlation with the expression of miR-199b-3p. The results of dual-luciferase reporter assay suggested that miR-199b-3p directly regulates the expression of mTOR (Fig. 4).

TSC2 regulates mTOR signaling to control cell growth and survival. To test if this process is mediated by miR-199b-3p, we detected the expression of miR-199b-3p and found that TSC1-N837fs and TSC2-Q371fs overexpression could decrease the expression of miR-199b-3p compared with overexpression TSC1-WT or TSC2-WT, respectively. Furthermore, knockdown of either *TSC1* or *TSC2* also decreased the expression of miR-199b-3p (Fig. 5). These results from both overexpression and knockdown of *TSC1* or *TSC2* suggest there is an axis that TSC complex regulates the expression of miR-199b-3p, and the low level of miR-199b-3p introduces the higher level expression of mTOR, causing abnormally active mTOR signal pathways which involves many cellular biological processes, in this axis miR-199b-3p acts a regulator for mTOR expression and then mTOR signaling pathway (Fig. 6).

It seems that the expression of miR-199b-3p was decreased more when knockdown *TSC1* than knockdown *TSC2*. As the same time, the expression of mTOR was increased more when knockdown *TSC1* (Figure 5). These results may suggest that *TSC1* has a larger effect on the expression of miR-199b-3p and mTOR. But in fact, the efficiency of knockdown is the determining factor of this question. The efficiency of knockdown *TSC1* was higher than knockdown *TSC2* in Fig. 5. The difference in effect of regulation the expression of miR-199b-3p and mTOR by *TSC1* and *TSC2* can't get a clean conclusion. But from the difference in Fig. 5, it confirmed further that the negative relationship between miR-199b-3p and mTOR, when miR-199b-3p decreased the expression of mTOR is increase, when miR-199b-3p increased the expression of mTOR is decrease. The results showed that overexpression of *TSC2* has a larger effect on the expression on miR-199b-3p, but not on the expression of mTOR in Fig. 5. It's very funny to see these phenomenons, but we can't get a certain conclusion about the difference between *TSC1* and *TSC2* in the regulation on miR-199b-3p just according to few experiments. We will further explore the difference regulation between *TSC1* and *TSC2* on miR-199b-3p to get a confirmed conclusion in future.

In summary, our studies revealed a novel molecular mechanism of TSC that the failure of the formation of physiological functional TSC complex causes the decrease of miR-199b-3p expression by which introduces the increase of mTOR expression and changes of mTOR signal pathway, followed the effects of changed pathway on all related cellular biological processes (Fig. 6). The significance of our studies on variations in *TSC1* and

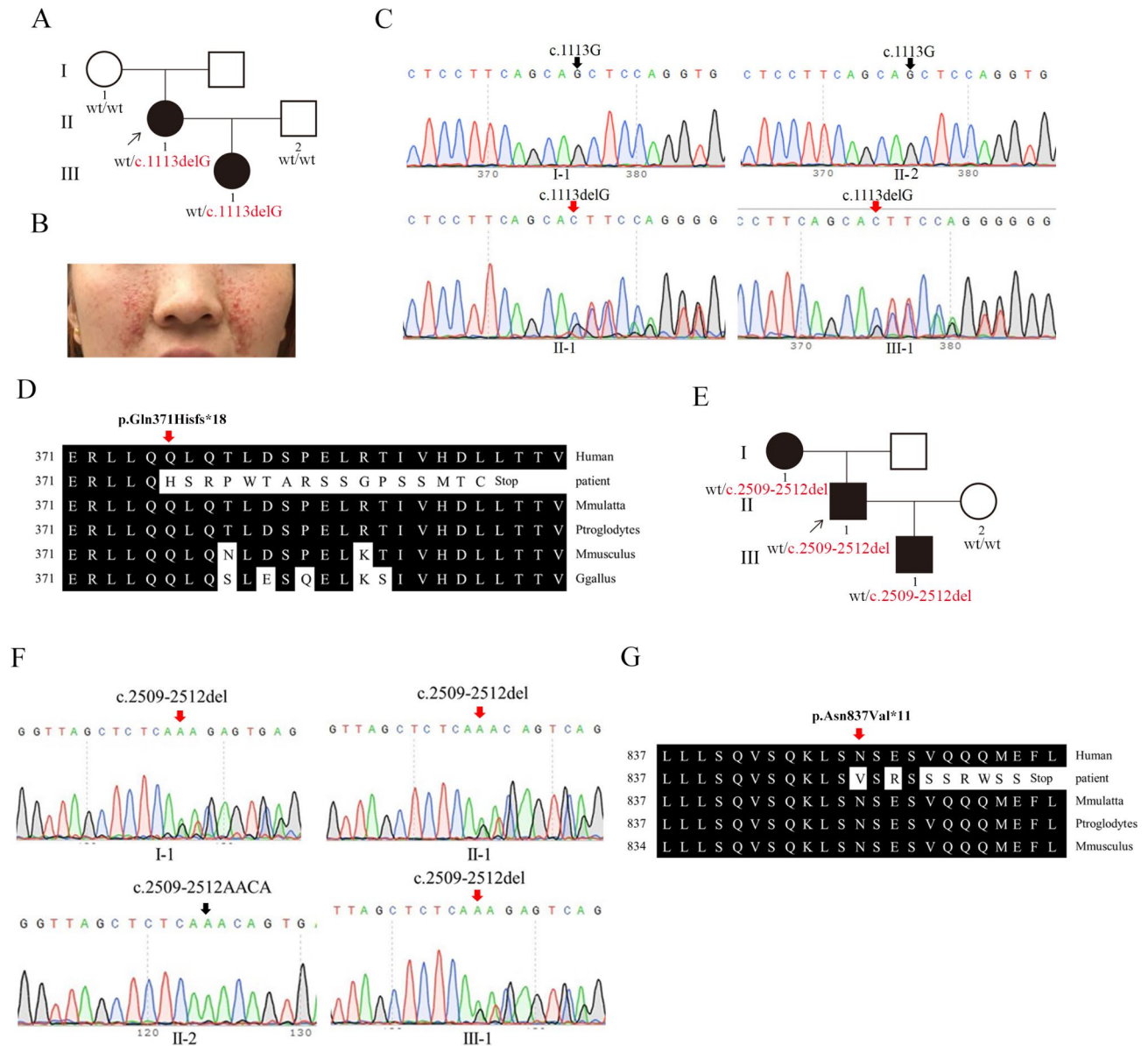


Fig. 5. TSC complex regulates the endogenous content of miR199b-3p, $*p < 0.05$, $**p < 0.01$, $***p < 0.001$ comparison with WT. (A) The efficiency of knockdown TSC1 in shTSC1-HEK293T cell line. The originals gels can be seen in the supplementary figure S6. (B) The expression of miR-199b-3p in shTSC1-HEK293T cells and overexpression TSC1-WT or TSC1-N837fs in HEK293T cell line. (C) Quantitative analysis of mTOR in shTSC1-HEK293T cell line and overexpression TSC1-WT or TSC1 N837fs in HEK293T cell line using western blot (low panel) and the statistics (up panel). The originals gels can be seen in the supplementary figure S6. (D) The efficiency of knockdown TSC2 in shTSC2-HEK293T cell line. The originals gels can be seen in the supplementary figure S6. (E) The expression of miR-199b-3p in shTSC2-HEK293T cell line and overexpression TSC2-WT or TSC2-Q371fs in HEK293T cell line. (F) Quantitative analysis of mTOR in shTSC2-HEK293T cell line (low panel) and overexpression TSC2-WT or TSC2-Q371fs in HEK293T cell line using Western blot (low panel) and the statistics analysis (up panel). The originals gels can be seen in the supplementary figure S6.

one *de novo* variation in TSC2 is that we not only defined pathogenesis of those two variations in either *TSC1* or *TSC2*, we also elucidated a novel molecular mechanism of TSC, and discovered the therapeutic potentials of miR-199b-3p for TSC.

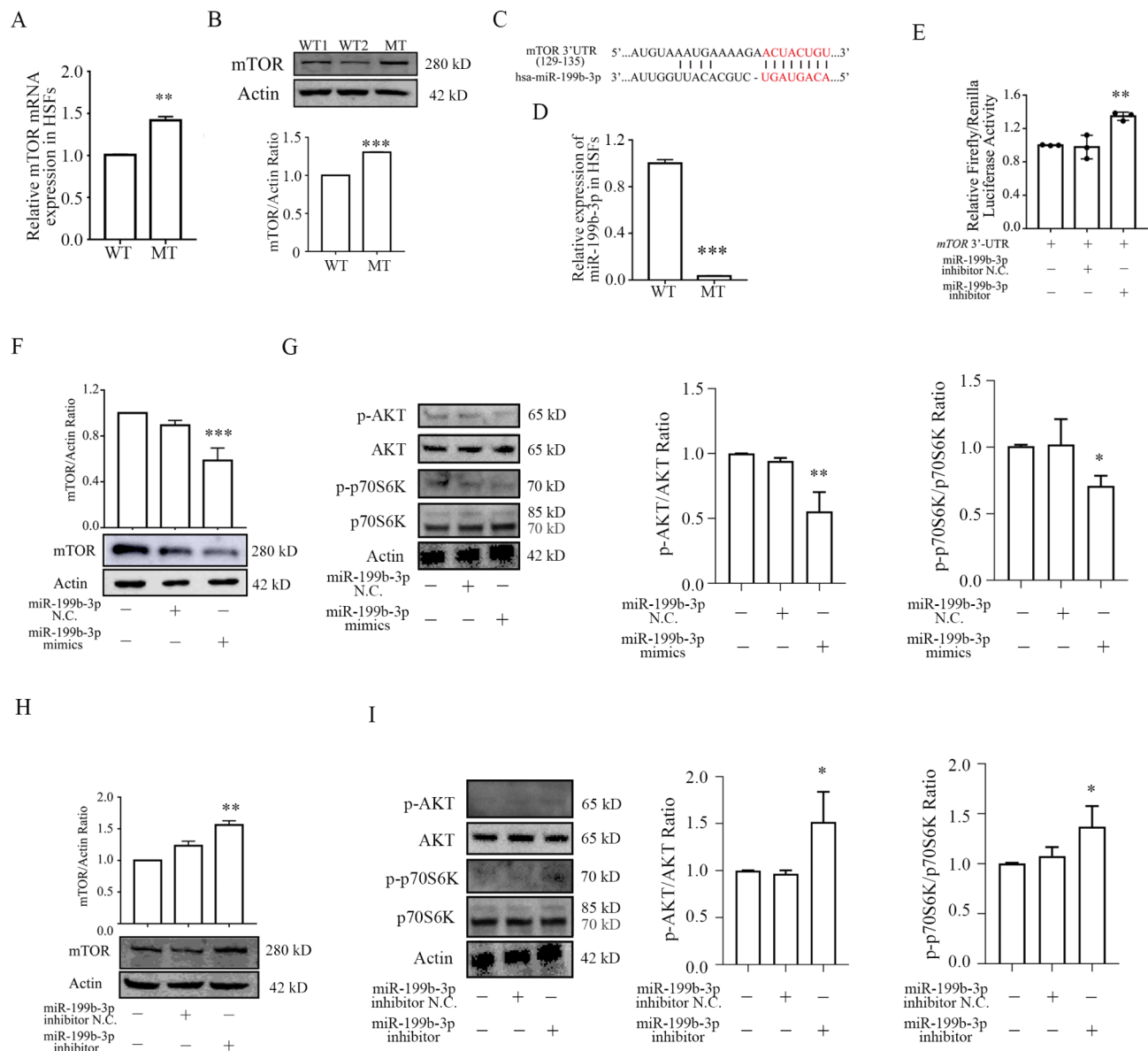


Fig. 6. The molecular mechanism model of TSC complex regulate mTOR signal pathway through miR-199b-3p.

Data availability

All primary data are available upon reasonable request from the corresponding author.

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Author contributions

Conceptualization and designed the experiments: C.W., H.X.; formal analysis and investigation: N.Z.; original draft preparation: N.Z., Q.X.; review and editing, C.W. and P.L.; resources: H.X.; supervision, H.X. and C.W.; funding acquisition: C.W. and P.L. All authors have read and agreed to the published version of the manuscript.

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Declarations

Competing interests

The authors declare no competing interests.

Institutional Review Board Statement

The study was conducted according to the guidelines of the Declaration of Helsinki and approved by the Ethics Committee of Shanxi University.

Informed consent

was obtained from all subjects involved in the study.

Additional information

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