Transcriptomic analysis of chicken *Myozenin 3* regulation reveals its potential role in cell proliferation

Maosen Ye*, Fei Ye*, Liutao He‡, Bin Luo‡, Fuling Yang‡, Can Cui‡, Xiaoling Zhao‡, Huadong Yin‡, Diyan Li‡, Hengyong Xu‡, Yan Wang*, Qing Zhu*

Farm Animal Genetic Resources Exploration and Innovation Key Laboratory of Sichuan Province, Sichuan Agricultural University, Chengdu Campus, Chengdu, China

* These authors contributed equally to this work.
‡ These authors also contributed equally to this work.
* as519723614@163.com (YW); zhuqing@sicau.edu.cn (QZ)

Abstract

Embryonic muscle development and fibre type differentiation has always been a topic of great importance due to its impact on both human health and farm animal financial values. *Myozenin3 (Myoz3)* is an important candidate gene that may regulate these processes. In the current study, we knocked down and overexpressed *Myoz3* in chicken embryonic fibroblasts (CEF*s) and chicken myoblasts, then utilized RNA-seq technology to screen genes, pathways and biological processes associated with *Myoz3*. Multiple differentially expressed genes were identified, including *MYH10, MYLK2, NFAM1, MYL4, MYL9, PDZLIM1*; those can in turn regulate each other and influence the development of muscle fibres. Gene ontology (GO) terms including some involved in positive regulation of cell proliferation were enriched. We further validated our results by testing the activity of cells by cell counting kit-8 (CCK-8) and confirmed that under the condition of *Myoz3* overexpression, the proliferation rate of CEFs and myoblasts was significantly upregulated, in addition, expression level of fast muscle specific gene was also significantly upregulated in myoblasts. Pathway enrichment analysis revealed that the PPAR (Peroxisome Proliferator-Activated Receptor) pathway was enriched, suggesting the possibility that *Myoz3* regulates muscle fibre development and differentiation through the PPAR pathway. Our results provide valuable evidence regarding the regulatory functions of *Myoz3* in embryonic cells by screening multiple candidate genes, biological processes and pathways associated with *Myoz3*.

Introduction

Muscle fibres are highly diverse in colour, contractile properties and metabolic mechanisms. Based on those properties, muscle fibres can be generally divided into 2 categories, namely, white and red muscle fibres. White muscle fibres are characterized by glycolytic metabolism and are specialized for fast and transitory activities. Red muscle fibres are rich in myoglobin and oxidative enzymes and possess the capacity for continuous activity but contract relatively...
slowly [1, 2]. Muscle fibre type can influence multiple important physiological and pathological properties regarding skeleton and cardiac muscle, including muscle hypertrophy [3, 4], exercise endurance, speed and glucose tolerance [5, 6]. In the field of farm animal research, muscle fibre type is also one of the key factors that influence the meat quality, including meat colour and water holding capacity [7–9]. Different metabolic mechanisms result in different basal metabolic rates, a point of importance for maximizing the feed conversion ratios of farm animals. Myozenin 3 (Myoz3) is one of the candidate genes that may influence muscle fibre type.

There are three members in the Myozenin (Myoz) family, including Myozenin 1 (Myoz1), Myozenin 2 (Myoz2), and Myozenin 3 (Myoz3); they encode the calsarcin-2 (FATZ1), calsarcin-1 (FATZ2) and calsarcin-3 (FATZ3) proteins, respectively. Since its discovery by three independent groups in the period of 2 years, the Myoz family has emerged as one of the most intensively studied gene families controlling muscle fibre type. The Z-disc plays important roles in both sarcomere structure and signal transduction and is formed by dozens of proteins, including the Myoz family [10–12]. Myoz can interact with multiple Z-disc proteins including α-actinin 2, telethonin, γ-filamin/ABP-L [10], myotilin [13], and calcineurin [11]. In addition, calsarcin-3 is able to interact with the PDZ-LIM domain protein ZASP/Cypher/Oracle [14]. The myozenin family shows a muscle fibre type preference in its expression patterns: while Myoz2 is mainly expressed in slow-twitch skeletal muscle and cardiac muscle, Myoz1 and Myoz3 are predominantly expressed in fast-twitch skeletal muscle.

Studies of the Myoz family’s role in muscle fibre type diversity have mainly focused on their negative regulatory effect on calcineurin (CaN) activity by direct binding. Activation of CaN in skeletal myocytes selectively upregulates slow-fibre-specific gene promoters [15] and hence drives a transition from fast to slow muscle fibres. Myoz2 knockout mice showed inappropriate CaN activity, which resulted in an excess of slow skeletal muscle fibres and enhanced the cardiac growth response to pressure overload [16]. Myoz1-deficient mice showed substantially reduced body weight and fast-twitch muscle, and they displayed markedly improved performance and enhanced running distances, also due to aberrant CaN/NFAT overactivation [17]. Inhibition of the CaN signalling pathway by calsarcin-1 can also help prevent cardiomyocyte hypertrophy induced by Ang-II [18]; four residues of calsarcin-1 undergo phosphorylation during pressure overload, resulting in enrichment in cardiac nuclei [19]. In addition, Myoz2 is a candidate gene for hypertrophic cardiomyopathy (HCM), as established by haplotype mapping of 516 HCM probands [20].

Due to key roles in muscle fibre differentiation, the Myoz family is also of great interests among farm animal researchers. Expression profiling of the Myoz family in mammals such as swine [21–23] and goats [24] reveal an expression pattern similar to that in mice. An association study also indicates that SNPs of Myoz are associated with traits that are influenced by muscle fibre types [25]. However, no research regarding avian Myoz3 has been reported[26].

The chicken is one of the most financially important birds and also among the most thoroughly studied in terms of genetics, its genome having been published in 2004. Additionally, the chicken is the classic model for the study of vertebrate embryonic development. In the current study, we separately knocked down and overexpressed Myoz3 in chicken embryonic fibroblasts (CEFs, which are widely used to study chicken embryonic development). Then, we took advantage of the rapid development of high-throughput mRNA sequencing technology to investigate the chicken Myoz3 gene’s regulatory role at the transcriptome level to identify novel pathways and genes that respond to changes in the expression level of chicken Myoz3. We hope to provide evidence to expand our knowledge regarding Myoz3.
Materials and methods

siRNA synthesis and vector construction

To knock down the expression level of Myoz3, three pairs of short interfering RNAs (siRNAs) targeting Myoz3’s CDS (coding sequence) and one NC (non-specific control) were designed and synthesized by Sangon Biotech (Table 1) (Shanghai, China). An overexpression vector was constructed by cloning Myoz3 CDS into the lentivirus vector GM-1013L050 (pLVX-3Flag-MCS-IRE5-ZsGreen1), provided by Genomiditech (Shanghai, China).

Cell cultivation, transfection and proliferation test

All the fertilized SPF (specific pathogen free) chicken eggs used in this study were purchased from Meili Breeding Corporation (Beijing, China).

Primary chicken embryonic fibroblasts were obtained from 9-day-old SPF (specific-pathogen-free) chicken embryos. Chicken embryos were first removed of their head and abdominal organs and bones, the remaining tissues were minced and digested with 0.25% trypsin, the suspension were filtered and plated in 12 wells and 96 wells cell culture plates.

Primary chicken myoblasts were isolated as described by Shumao Lin et al.[27]. In brief, leg muscle was harvested from 11-day-old chicken embryos and minced for further 0.1% collagenase type I (Invitrogen, Carlsbad, CA, USA) digestion. Then the suspension was subjected to a density gradient centrifugation in three discontinuous layers with 20, 30 and 55% Percoll (Solarbio Beijing China), and the cell suspension between the interface of 30 and 55% Percoll was collected and plated in 12 wells and 96 wells cell culture plates.

CEFs and myoblasts were cultured in Dulbecco’s modified Eagle’s medium (Thermo Scientific, USA) with 10% fetal bovine serum (Thermo Scientific, USA) that was filtered with 20 nm filters before use. In addition, 100 μg/mL streptomycin and 100 U/mL penicillin (Thermo Scientific, USA) were added to the cell culture medium. The cells were cultured in 12-wells plates at 37°C in a humified incubator (Thermo Fisher Scientific, USA) with CO2 concentration set to 5%. Then, following the manufacturer’s protocol, 20 pmol of each siRNA and NC and 20 μg of expression vector and control vector were transfected into $5 \times 10^5$ CEFs using Lipofectamine 3000 (Invitrogen, USA) in three biological replicates. After 48 hours, RNA was extracted from the transfected cells using TRIzol reagent (Invitrogen, USA). Cell proliferation was tested using a Cell Counting Kit-8 (CCK-8) in the condition of overexpression, following the manufacturer’s instructions. Cells for the proliferation test were cultured in 96-well plates and maintained at 40% of maximum density before transfection.

RNA-seq and data analysis

Treated CEFs were subjected to RNA-seq, the total RNA was extracted using TRIzol reagent and diluted with RNase-free H2O (Tiangen, China). The concentration and purity of the

| siRNA     | Direction | Sequence                        |
|-----------|-----------|---------------------------------|
| Myoz3-427 | Sense     | GGAUUGAGCGCUUUGCUUUTT           |
|           | Antisense | AAAGCAAAGCGUUGCACCTT            |
| Myoz3-809 | Sense     | GCGCAUGAACCUCCCACAUTT           |
|           | Antisense | AUGUGGAGGUUUGACGGCTT            |
| Myoz3-983 | Sense     | GGUUCUGCCUGAGUGAUTC             |
|           | Antisense | AUCACUCUGGAGGACACCTT            |
| NC (Non-specific control) | Sense | UUUCGCGACGGUGUCGACUTT          |
|           | Antisense | ACGUGACACGUGUGGGAATT            |

[https://doi.org/10.1371/journal.pone.0189476.t001](https://doi.org/10.1371/journal.pone.0189476.t001)
extracted RNA was measured using the NanoDrop 2000 (Thermo Scientific, USA), and RNA degradation and contamination were assessed on 1% agarose gels.

A total of 12 libraries (one library per transfected sample) were constructed, including 3 control samples transfected with control vector, named Control; 3 overexpression samples transfected with overexpression vectors, named Over; 3 knockdown samples transfected with siRNA of Myoz3, named Inter; and 3 NC samples transfected with non-specific control siRNA, named NC. A total of 3 μg of RNA per sample was used as input material for preparations. Sequencing libraries were generated using a NEBNext Ultra™ RNA Library Prep Kit for Illumina (NEB, USA) following the manufacturer’s recommendations, and index codes were added to attribute sequences to each sample. After library preparation, an Illumina HiSeq 4000 was used to generate 120 bp/150 bp paired-end reads. The data files from RNA-seq analysis have been deposited in NCBI’s Gene Expression Omnibus, and are accessible through GEO Series accession number GSE99146 (https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE99146).

The raw data in fastq format were first subjected to the FastQC for quality assessment, then we applied a in house python script to remove reads containing adapter or poly-N sequences as well as low quality reads and obtain clean reads. Reference genome and gene model annotation files were downloaded from the genome website directly. An index of the reference genome was built using Bowtie v2.2.3 [28], and paired-end clean reads were aligned to the reference genome using TopHat v2.0.12 [29, 30]. HTSeq v0.6.1 was used to count the number of reads mapped to each gene and generate a count table for further analysis [31].

Differentially expressed genes between groups were analysed using edgeR [32], applying LRT (likelihood ratio test) methods, and the P values were adjust using the BH method with a cutoff value of \( q < 0.05 \). Function annotations for significantly differentially expressed genes were performed using the DAVID website [33]. The enriched gene ontology (GO) terms on biological processes and the pathways obtained from DAVID functional analysis were filtered for significance by gene counts \( \geq 3 \), \( p-value < 0.05 \). Protein interaction analyses were performed on the STRING website (http://www.string-db.org/).

Quantitative reverse transcription PCR
For qRT-PCR, first-strand cDNA was synthesized from 1 μg of total RNA using the Prime-Script RT Reagent Kit (Perfect Real-Time) (TaKaRa, Biotechnology Co. Ltd., Dalian, China). The reactions were performed under the following conditions: 42˚C for 2 min, 37˚C for 15 min and 85˚C for 5 s.

qRT-PCR was conducted with two pairs of primers designed with Primer Premier 5 software (Table 2). GAPDH was chosen as the housekeeping gene for normalization. An 11 μL reaction containing 6 μL of SYBR premix Ex Taq™ (TaKaRa, Biotechnology Co. Ltd., Dalian, China), 1 μL of cDNA, 0.5 μL of forward primer, 0.5 μL of reverse primer and 3 μL of RNase-free H₂O (Tiangen, Beijing, China) was used for qRT-PCR. The reactions were carried out with the following amplification conditions: 95˚C for 10 s followed by 40 cycles of 95˚C for 5 s and 60˚C (or the appropriate annealing temperature) for 30s. Each sample was run in 3 technical replicates. The \( 2^{-\Delta\Delta Ct} \) method was applied to quantify mRNA expression levels.

Results
Knockdown and overexpression of Myoz3 in CEFs and myoblasts
First, we thought to determine whether Myoz3 is expressed in CEFs and to which level is Myoz3 expressed in CEFs compare to myoblasts and adult breast muscles (tissue from our
Our results suggest that Myoz3 is expressed in CEFs, although it’s significantly lower than in breast muscles and myoblasts (about 4 and 2 times, respectively). Our results suggest that Myoz3 play a role in chicken embryonic fibroblasts.

To analyse the function of Myoz3 in vitro, we knocked down and overexpressed the Myoz3 gene in CEFs and myoblasts by transfecting them with siRNA and overexpression vectors, respectively. First, after we tested the efficiency of various siRNAs, siRNA 427 was identified as the most efficient siRNA for use in further experiments because of its knockdown efficiency of over 20% compared with NS (non-specific control). The overexpression vector of Myoz3 was previously publication [34]).

### Table 2. Primers for qRT-PCR.

| Gene    | Direction | Sequence          | AT (en) |
|---------|-----------|-------------------|---------|
| CaN     | Forward   | GTTTTCCCATACAGCTGTCCC | 61.2    |
|         | Reverse   | TGGCAATTGAAAGCTTCTT  |         |
| Myoz3   | Forward   | TTCTAGGATGTTAGAGCAAGAGC | 60      |
|         | Reverse   | AGCTTTGGTTGGCCGTCA  |         |
| MYH10   | Forward   | GATCTGGATCATCAGGCA   | 58.6    |
|         | Reverse   | GCACGATCTCTCTCTCTCTGC |         |
| MYLK2   | Forward   | CTGCAACAGGAAGGGAGAAGG | 62      |
|         | Reverse   | GTTGAACGCACAAACAAAAGG |         |
| MYL4    | Forward   | GGAGGAGTCATGGAAGAGTT | 61.4    |
|         | Reverse   | CAGACCCATCCAGGACCTCG |         |
| MYL9    | Forward   | AACATGTCACAAAGCTG    | 58.6    |
|         | Reverse   | AGGGAAAGAACGAGCAAGCA |         |
| NFAM1   | Forward   | GGCAAGGAGGAACAGACA  | 58.4    |
|         | Reverse   | ACTCGATAGGGTTGGAGGCT |         |
| ITGA8   | Forward   | TTGGGCTGTTGGAGCTG    | 59.4    |
|         | Reverse   | ACAGCCAGGAAACGGAGG   |         |
| GAPDH   | Forward   | AGGAGGCTGCTGCTGCTG   | 60      |
|         | Reverse   | CCTCAAGGAGACACAGG    |         |
| PPM1J   | Forward   | GAGAGGCTGCTGCTGCTG   | 60.2    |
|         | Reverse   | CAGACGAGGCAGCCCATTCTT |       |
| ECM2    | Forward   | AAGAGGCTGCGGACATTCTT | 60      |
|         | Reverse   | CCTTCCATTGCGGACATTCTT |       |
| OASL    | Forward   | CATGACCTGGAGAGGAGG   | 60      |
|         | Reverse   | AGCAAGAGCCAGCTGAGAAA |         |
| FAP     | Forward   | GTGCGATGTGTTGTTGAGG  | 60      |
|         | Reverse   | CCTGCCCATCCTGTTTACT  |         |
| MPRIP   | Forward   | ATCCCTCGTAACACACTCG  | 60      |
|         | Reverse   | GCCCTCTCCAGGCTCTAGG  |         |
| PDLIM1  | Forward   | TACAGGAGAGAGAGAGGGGT | 60      |
|         | Reverse   | GTGACCCACTGCTGCTCTTC |         |
| PKM     | Forward   | AGCAGGAGGAGACACGAGAC | 60.2    |
|         | Reverse   | ATGCCGGTGTTTCTGCAAT  |         |
| MYHC7B  | Forward   | TCAAGCAGGCTAGCTATT   | 60      |
|         | Reverse   | CATCTTCCAGGATGCAAGG  |         |
| MYH1F   | Forward   | ACCTTGGTACCAAGACCC   | 59.6    |
|         | Reverse   | GCTTGGTCTGCGCTTCTAC  |         |

AT: Annealing temperature

https://doi.org/10.1371/journal.pone.0189476.t002
driven by the CMV promotor, and 24 hours after transfection, the expression level was elevated more than 10-fold compared with cells that were transfected with empty vectors (Fig 1).

RNA sequencing
A total of 12 libraries were prepared. Those libraries were Inter (CEF transfected with siRNA), NC (CEF transfected with non-specific RNA), Over (cell transfected with overexpression vector), and Control (cell transfected with empty vector), each of them prepared in 3 biological
replicates. An Illumina Hi-Seq 4000 was used to generate paired-end reads. An average 46.77 million reads were generated for all the libraries, with Q20 more than 97.3% and Q30 more than 93.4% (S1 Table). More than 80% of percent reads were mapped to the galGal5 chicken genome (S2 Table) by TopHat2. The data files from the RNA-seq analysis have been deposited in the NCBI’s Gene Expression Omnibus.

Differentially expressed genes and qRT-PCR validation

A count table containing each the ID and read counts of each gene was generated from aligned reads by HTSeq v0.6.1[35]. In addition, the R package edgeR was applied for differentially expressed gene analysis. First, we generated a multidimensional scaling (MDS) plot to evaluate the leading biological coefficient of variation (BCV). We found that the 3rd replicate of Inter was abnormally far from other Inter samples and close to the NC samples; we suspected that the interference was not sufficient, and so we tested the Myoz3 expression level by qRT-PCR, revealing that the expression level of Myoz3 in the 3rd Inter group was similar to that of NC (Fig 2). Therefore, we excluded the 3rd NC group from further analysis. The cutoff value for differential expression was an adjusted P<0.05.

When Inter was compared with NC, 302 genes were found to be differentially expressed, of which 84 were significantly downregulated and 226 of them were significantly upregulated, including MYL4 (myosin light chain 4) and MYL9 (myosin light chain 9). When Over was compared with Control, 301 genes were downregulated and 127 genes were upregulated. Some genes involved in muscle development including MYH10 (myosin heavy chain 10) and MYLK2 (myosin light chain kinase 2) were among the differentially expressed genes. Thirteen genes were differentially expressed in both knockdown and overexpression conditions; they are VWA5B2 (von Willebrand factor A domain-containing 5B2), SQSTM1 (sequestosome 1), ASPN (asporin), ITGA8 (integrin subunit alpha 8), PLK3 (Polo-like kinase 3), CRLF1 (cytokine receptor like factor 1), LRP3 (LDL receptor related protein 3), TNFRSF6B (TNF receptor superfamily member 6b), THBS (thrombospondin-2 precursor), ENSGALG00000035656, ENSGALG00000028466, ENSGALG00000037711, and ENSGALG00000011668 (Fig 3).

Fig 2. The 3rd sample of Inter is abnormal due to inefficient knockdown. (A) An MDS plot of NC and Inter illustrates the abnormality of the 3rd Inter sample. (B) The expression level was not sufficiently knocked down in the 3rd Inter group. The genes that were differentially expressed between the two experiment conditions are presented. The error bars indicate SEM for three replicates (qRT-PCR). Significance was not calculated because to data for each bar do not represent biological replicates.

https://doi.org/10.1371/journal.pone.0189476.g002
Fig 3. Experimental design and overall results. The expression levels of the top 30 differentially expressed genes are shown in log CPM (copies per million). NC (non-specific control).

https://doi.org/10.1371/journal.pone.0189476.g003
To validate the results that were obtained by RNA-seq, qRT-PCR was performed for each of the 12 libraries. We confirm that except for the 3rd library of Inter, Myoz3 was expressed as intended in other groups, that is, highly expressed in the Over group and scarcely expressed in the Inter group. Then, we validated genes that were identified as differentially expressed by RNA-seq, including MYH10, MYLK2, MYL4, MYL9, ASPN and KLP3, protein phosphatase, Mg2+/Mn2+ dependent, 1J (PPM1J), extracellular matrix protein 2 (ECM2), 2’-5’-oligoadenylate synthetase-like (OASL), fibroblast activation protein, alpha (FAP), PDZ and LIM domain 1 (PDLIM1), myosin phosphatase Rho interacting protein (MPRIP), pyruvate kinase, muscle (PKM), ATPase H+ transporting V1 subunit H (ATP6V1). Of 12 genes that were selected as genes to verify (CaN was chosen as a negative control because it has not been identified as a differentially expressed gene), ITGA8 and PDLM1 were not statistically correspond to our RNA-seq results, but trends in ITGA8 and PDLM1 expression were along similar lines. In addition, all the other results are able to verify our results of differentially expressed gene analysis (Fig 4).

**Gene ontology (GO), pathway and protein interaction analyses**

To better understand the differentially expressed genes we filtered from background, we conduct gene ontology and pathway enrichment analysis. The differentially expressed gene lists for both comparisons were submitted to DAVID (http://david.abcc.ncifcrf.gov/) for enriched functional terms in biological processes and pathways.

Under knockdown condition, pathways such as cardiac muscle contraction, ECM-receptor interaction, PPAR, vascular smooth muscle contraction, and focal adhesion were enriched (S1 Fig). Key words such as calcium and muscle protein were also enriched. Biological process such as mesenchyme migration, tendon development, long-chain fatty acid import was enriched (Fig 5).

Under the overexpression condition, biological process terms such as positive regulation of vascular smooth muscle cell proliferation, loop of Henle development, regulation of energy homeostasis were enriched. Four pathways were found to be enriched, namely, the ErbB signalling pathway, influenza A, herpes simplex infection, and focal adhesion.

To further illustrate the regulation network among differentially expressed genes, we conducted proteins interaction network analysis. In the overexpression condition, we focused on the interaction network that is centred on MYH10 (Fig 6). MYH10 can directly interact with MYLK2, GATA6, SMC4, and others. The secondary network, including proteins that interact with proteins that interact with MYH10 in differentially expressed genes, contains 47 genes. In the knockdown condition, few interesting networks were identified.

**Cell proliferation rates was elevated under Myoz3 overexpression condition**

To further support our results from RNA-seq, we conducted a cell proliferation test in both CEFs and myoblasts, since pathways and biological process involve in cell proliferation were enriched, such as the PPAR pathway as well as biological process terms such as positive regulation of vascular smooth muscle cell (VSMC) proliferation. The results show that when Myoz3 is overexpressed in CEFs and myoblasts, cell proliferation was significantly upregulated. Knockdown of Myoz3 in both cells results in trend of proliferation inhibition. (Fig 7).
Transcriptome analysis of chMyoz3

A. MYL9

B. MYL4

C. PKM

D. MPRIP

E. PDLIM1

F. ATP6V1H

G. MYH10

H. MYLK2

I. PPM1J

J. ECM2

K. OASL

L. FAP

M. CaN

N. ITGA8
The fast muscle specific gene was upregulated upon Myoz3 overexpression

To test whether muscle fibre type specific gene can be regulated by Myoz3, we detected the expression level of MYH7B (myosin heavy chain 7b), which is slow muscle marker gene; MYH1F (myosin heavy chain 1f), which is a fast muscle specific gene. Our results show that in myoblast, overexpression of Myoz3 gene can lead to MYH1F upregulated but not MYH7B.

![Bar chart showing expression levels of genes](https://doi.org/10.1371/journal.pone.0189476.g004)

**Fig 4. Validation of differentially expressed genes by qRT-PCR.** The relative expression level of data was presented by bar plots with error bars represent SEM. All experiments were replicated three times. (A) MYL9, (B) MYL4, (C) PKM, (D) MPRIP, (E) PDLIM1, (F) MYH10, (G) MYLK2, (H) PPM1J, (I) ECM2, (J) OASL, (K) FAP, (L) CaN, (M) ITGA8. *P<0.05, **P<0.01, ***P<0.001, N.S. means not significant.

![Biological process terms enrichment](https://doi.org/10.1371/journal.pone.0189476.g005)

**Fig 5. Enriched biological process (BP) terms.** (A) BP terms enriched under the knockdown condition (NC vs Inter). (B) BP terms enriched under the overexpression condition (Control vs Over).
down regulated. However, no change was observed in Myoz3 knockdown group (Fig 8) and all CEFs groups (data not shown).

**Discussion**

Muscle fibre type differentiation is a delicate process that is regulated both during and after the embryonic period [36, 37]. For animals that have limited activity space after birth/hatching and whose muscle tissue is of great economical value, such as chickens, embryonic development of muscle fibres is a factor of great importance, especially developmental control of muscle fibre types. In this study, we focus on Myoz3, a candidate gene for muscle fibre type differentiation, hoping to understand its regulatory roles in embryonic muscle fibre development. We apply the fast-growing technology of deep sequencing to CEFs under both Myoz3 knockdown and overexpression conditions, hoping to identify pathways and biological process associated with Myoz3 expression level.

CEF is one of the most broadly used cell types for studying the embryonic development of chickens [38, 39], and a model for study both chicken signalling pathways [40, 41]. Myoz3 expression was detected in primary CEFs, however, little is known about Myoz family’s role in fibroblasts. we suspect that Myoz3 play a role in CEFs differentiation, and chicken embryonic development. In addition, previous evident suggest that activation of muscle specific gene in non-muscle tissue by forced expression of muscle specific regulator [42]. Therefore, both CEFs and myoblasts were utilized in this study.

Even though little evidence of Myoz3’s role in muscle fibre type differentiation has been reported, studies regarding other Myoz family proteins can provide valuable information about Myoz3 function. In human, mice and chickens alike, each member of the Myoz family

---

**Fig 6. Protein interaction networks under the overexpression condition.** (A) Biggest network under the overexpression condition, containing 108 genes. (B) Secondary interaction networks of MYH10, containing 47 genes. (C) Network of genes that directly interact with MYH10, containing 11 genes.

https://doi.org/10.1371/journal.pone.0189476.g006
possesses only one domain, calsarcin. *Myoz2*−/− mice showed an excess of slow-twitch skeletal muscle fibre due to calcineurin activity upregulation, and a foetal gene program was activated in *Myoz2* deficient hearts that caused the Z-discs to become ‘fuzzy’ [16]. *Myoz1*−/− mice showed a reduction in body weight and fast-twitch muscle mass; also, their endurance capacity was increased due to a fibre type shift towards slow-twitch oxidative fibres. Similar to *Myoz2*−/− mice, *Myoz1*−/− mice also showed an increase in Calcineurin/NFAT activity [17], despite the different expression pattern between *Myoz2* and *Myoz1*. In the current study, we identified multiple genes and pathways that are possibly regulated by chicken *Myoz3* in CEFs.

Fig 7. Proliferation by CCK-8 under both *Myoz3* knockdown and overexpression conditions. (A) 24 hours after CEFs transfected with empty vector (Control) or overexpression vector, non-specific siRNA (NC) or *Myoz3* siRNA (Inter). (B) 48 hours after CEFs transfected with empty vector (Control) or overexpression vector, non-specific siRNA (NC) or *Myoz3* siRNA (Inter). (C) 24 hours after Myoblast transfected with empty vector (Control) or overexpression vector, non-specific siRNA (NC) or *Myoz3* siRNA (Inter). (D) 48 hours after myoblasts transfected with empty vector (Control) or overexpression vector, non-specific siRNA (NC) or *Myoz3* siRNA (Inter).

https://doi.org/10.1371/journal.pone.0189476.g007
In the overexpression condition, a total of 428 genes were identified as significantly differentially expressed when filtered by the least ratio test (LRT) method applying edgeR, including MYH10, which is significantly upregulated, and MYLK2, which is significantly downregulated. MYH10 (non-muscle myosin II-B) belong to a non-muscle myosin family that has roles in cell migration, adhesion [43], and polarity as well as cardiac and brain development [44]. It was proposed that the first step of myofibril assembly is the formation of integrin-dependent cell-matrix adhesion, in which MYH10 play a key role, MYH10 transgenic mutant mice show defects in their heart sarcomeres [45]. MYH10 can also respond to the Ca\(^{++}\)-calmodulin pathway, which can activate MYLK (myosin light chain kinase), which then phosphorylates the RLCs (regulatory light chains) of MYH10, which can facilitate myosin interaction with actin filaments, leading to great increases in the Mg\(^{2+}\)-ATPase activity of myosin in the presence of actin [46]. The MYLK2 reduction and MYH10 elevation under the Myoz3 overexpression condition may lead to MYH10 hypophosphorylation when the Ca\(^{++}\)-calmodulin pathway is activated. Hence, there is a possibility that Myoz3 is a negative regulator that inhibit another downstream signal molecular of the Ca\(^{++}\) signalling pathway. However, further studies are required regarding how Myoz3 might regulate MYLK2. In addition to phosphorylating MYH10 protein, MYLK2 protein also modulates a variety of contractile processes, including smooth muscle contraction and proliferation [47], so we suspect that MYLK2 is a key components in Myoz3 regulatory network.

Under the knockdown condition, 302 genes were differentially expressed, including MYL9 (myosin light chain 9) and MYL4 (myosin light chain 4), PDLIM (PDZ and LIM domain 1), PKM (pyruvate kinase, muscle), NFAM1 (NFAT activating protein with ITAM motif 1). Myosins are a superfamly of molecular motors that depend on action and are implicated in contraction, cell shape, migration, adhesion, and intracellular transport [48]. MYL9 is reported to be associated with injury and ageing [49], and MYL4 is an atrial-specific myosin light chain gene. Mutation of MYL4 can lead to atrial fibrillation in humans, and MYL4 mutant zebrafish displayed disruption of sarcomeric structure and atrial enlargement [50]. In the current study, Myoz3 knockdown results in MYL4 upregulation, consistent with the fact that Myoz3 is
predominantly expressed in fast-twitch muscle [14], which is not localized in the heart. We suspect that a fast-twitch muscle-specific gene such as Myoz3 can play a key role in embryonic cell differentiation. NFAM1 (NFAT activating protein with ITAM motif 1), screened with NFAT-GFP reporter cells for its activating effect on transcription factor NFAT [51], was reported a decade ago and found to be upregulated under Myoz3 knockdown conditions. Consider that calsarcin-3 can bind with calcineurin and inhibit the activation of NFAT, we suspect that part of the inhibitory role is to negatively regulate the expression level of NFAM1. Furthermore, PDLIM1, known to interact with Myoz3, was downregulated. Both PDLIM1 and Myoz3 are localized in the Z-disc of skeletal muscle, and both contribute to Z-disc formation [52, 53]; therefore, downregulation of both genes may result in aberrant Z-disc signal transduction [54], hence the impact on the differentiation of muscle fibres in the embryonic period.

To better understand the function of Myoz3, we conducted gene ontology (GO) analysis. Under Myoz3 overexpression conditions, terms including positive regulation of vascular smooth muscle cell proliferation were enriched. Under Myoz3 knockdown conditions, on the other hand, the BP term negative regulation of smooth muscle cell proliferation was enriched, so we test whether a change in the expression of Myoz3 can influence the cell proliferation of CEs and myoblasts. As described above, under overexpression conditions, both myoblasts and CEs had significantly higher activity, confirming our results from RNA-seq data. Pathways were also enriched: under knockdown condition, pathways such as the PPAR signalling pathway and ECM-receptor interaction were enriched. Several prior publications regarding PPAR pathway show that three isoforms of PPAR subfamily play a role in skeletal muscle metabolisms and plasticity, including PPARα, PPARβ/δ, PPARγ [55]. The expression level of PPARα reflects differences in type I muscle fibres associated with pathologically and physiologically induced skeletal muscle fibre type differences [56], and vascular smooth muscle cell proliferation can be inhibited by PPARα through suppression of telomerase activity [57]. PPARα-overexpressing transgenic mice showed upregulated expression of genes involved in oxidation in skeletal muscle [58]. Regarding PPARβ/δ, activation of PPARδ in skeleton muscle leads to leads to muscle fibre type transformation, from type II to type I [59], the same study also show that activation of PPARβ can also prevent obesity as a results of metabolism alteration. Another study supported these results, Luquet et al. show that muscle-specific overexpression of PPARβ/δ in mice enhances muscle metabolism (fatty acid flux and b-oxidation) and altered muscle fibre type to increase oxidative type 2a. These mice also show decreased body fat mass and smaller fat cells [60]. Furthermore, PPARβ/δ activation that was induced by agonist resulted in enhances fatty acid oxidation in skeletal muscle cells [61]. C2C12 myotube was enriched upon PPARβ activation along with enhanced mitochondrial biogenesis [62]. In general, PPAR pathway that was enriched under Myoz3 knockdown condition is highly likely to relate to Myoz3 regulation, but further study regarding the mechanism as how Myoz3 regulation works require further study.

Although no specific pathway that relates to muscle fibre was found under Myoz3 overexpression conditions, several muscle development genes were nonetheless found, probably due to the background noise. The pathways and biological processes we identified included not only skeletal muscle regulation but also smooth muscle regulation, so we suspect that Myoz3 also functions in smooth muscle development in the embryonic period. Furthermore, overexpression of Myoz3 in myoblasts leads to significantly increase of fast-muscle specific gene expression, which make us believe that chicken gene/pathway annotation may need further improvement.

Another way to better understand and evaluate the pathway formed by differentially expressed gene is to construct protein-protein interaction (PPI) network, our results show that differentially expressed genes encode a highly interconnected network. And further confirmed
the results that MYH10 and MYLK2 interact with each other is highly likely molecular process that under Myoz3 regulation. In addition, GATA6 can also bind to MYH10, which is another possible interaction that explain the function of Myoz3. GATA6 is a transcriptional factor that regulate cardiomycyte hypertrophy[63], smooth muscle contraction [64]. However, PPI construction can only provide us indirect evidence, further experiments regarding protein interaction including CoIP (Co-Immunoprecipitation) and GST pull-down.

In general, by using highly advanced deep sequencing technology for cells under both overexpression and knockdown conditions, we were able to reveal the function of chicken Myoz3 in embryonic development. To the best of our knowledge, this is the first transcriptomic study applying RNA sequencing technology to study the function of Myoz3 in any species. Our results provide more than 302 candidate genes regulated by Myoz3 under Myoz3 knockdown conditions and 428 candidate genes under Myoz3’s regulation under overexpression conditions. Our results indicate that Myoz3 has the potential to regulate multiple myosin light chain family members, such as MYL4, MYL9. Non-muscle myosin heavy chain MYH10 was also found to be regulated by Myoz3, as was MYLK2, the kinase that phosphorylates MYH10. Pathways were also identified. The PPAR pathway is a very promising pathway that is likely involved in Myoz3-mediated embryonic development and muscle fibre type differentiation. We also confirmed our results by testing the proliferation of CEFs and myoblasts. Furthermore, muscle fibre type specific gene can also be regulated in myoblast upon Myoz3 expression alteration. However, due to the knockdown mechanism, our results may contain some background noise from the siRNA transfection and the remaining Myoz3 that we were unable to be knock down completely; a CRISPR/Cas9-mediated gene knockout method may help us better understand the role of Myoz3 at a cellular and organism level. Important question such as how chicken Myoz3 is regulated and the precise mechanism of how chicken Myoz3 is involved in cell proliferation still need to be answered by further studies.

Supporting information
S1 Fig. Enriched pathways. (A) Pathway enriched under the knockdown condition (NC vs Inter). (B) Pathway enriched under the overexpression condition (Control vs Over). (TIF)
S1 Table. Quality control Raw data and Quality control information. (XLSX)
S2 Table. Reads mapping. Reads mapping information. (XLSX)
S3 Table. Inter DE. Differentially expressed gene ID under knockdown condition. (XLSX)
S4 Table. Over DE. Differentially expressed gene ID under overexpression condition. (XLSX)
S5 Table. Counts table. Counts table that generated by HTseq. (XLSX)

Acknowledgments
We thank Lin Ye and Yao Zhang for provide us valuable ideas about chicken muscle color and help us housing the chicken for our experiment.
Author Contributions

Conceptualization: Diyan Li, Yan Wang.
Data curation: Maosen Ye, Liutao He, Bin Luo.
Formal analysis: Maosen Ye, Fuling Yang.
Funding acquisition: Yan Wang, Qing Zhu.
Investigation: Fei Ye, Qing Zhu.
Methodology: Fei Ye.
Project administration: Fei Ye.
Resources: Qing Zhu.
Supervision: Xiaoling Zhao, Huadong Yin, Diyan Li, Hengyong Xu.
Validation: Fei Ye, Xiaoling Zhao, Huadong Yin, Diyan Li, Hengyong Xu, Qing Zhu.
Visualization: Maosen Ye, Qing Zhu.
Writing – original draft: Maosen Ye, Can Cui.
Writing – review & editing: Maosen Ye.

References

1. Zierath JR, Hawley JA. Skeletal muscle fiber type: influence on contractile and metabolic properties. PLoS Biol. 2004; 2(10):e348. https://doi.org/10.1371/journal.pbio.0020348 PMID: 15486583; PubMed Central PMCID: PMCPMC521732.
2. Schiaffino S, Reggiani C. Fiber types in mammalian skeletal muscles. Physiol Rev. 2011; 91(4):1447–531. https://doi.org/10.1152/physrev.00031.2010 PMID: 22013216.
3. Staron R. S., Malicky E. S., Leonardi M. J., Falkel J. E., Dudley aGA. Muscle hypertrophy and fast fiber type conversions in heavy resistance-trained women. European Journal of Applied Physiology. 1989; (60):71–9.
4. Verdijk LB, Gleeson BG, Jonkers RA, Savelberg HH, Dendale P, et al. Skeletal muscle hypertrophy following resistance training is accompanied by a fiber type-specific increase in satellite cell content in elderly men. J Gerontol A Biol Sci Med Sci. 2009; 64(3):332–9. https://doi.org/10.1093/gerona/gln050 PMID: 19196907; PubMed Central PMCID: PMCPMC2655000.
5. Seto JT, Quinlan KG, Lek M, Zheng XF, Garton F, MacArthur DG, et al. ACTN3 genotype influences muscle performance through the regulation of calcineurin signaling. J Clin Invest. 2013; 123(10):4255–63. https://doi.org/10.1172/JCI67691 PMID: 24091322; PubMed Central PMCID: PMCPMC3784532.
6. Riedl I, Osier ME, Benziane B, Chipalin AV, Zierath JR. Association of the ACTN3 R577X polymorphism with glucose tolerance and gene expression of sarcomeric proteins in human skeletal muscle. Physiol Rep. 2015; 3(3). https://doi.org/10.14814/phy2.12314 PMID: 25780092; PubMed Central PMCID: PMCPMC4393151.
7. Klont R. E., Brocks L, Eikelenboom G. Muscle Fibre Type and Meat Quality. Meat Science. 1998; 49 (Suppl. I):S219–S29.
8. DRANSFIELD E., SOSNICKI AA. Relationship Between Muscle Growth and Poultry Meat Quality. Poultry Science. 1999; 78:743–6. PMID: 10228972
9. Joo ST, Kim GD, Hwang YH, Ryu YC. Control of fresh meat quality through manipulation of muscle fiber characteristics. Meat Sci. 2013; 95(4):828–36. https://doi.org/10.1016/j.meatsci.2013.04.044 PMID: 23702339.
10. Faulkner G, Pallavicini A, Comelli A, Salamon M, Bortoletto G, Ievolella C, et al. FATZ, a filamin-, actinin-, and telethonin-binding protein of the Z-disc of skeletal muscle. J Biol Chem. 2000; 275(52):41234–42. https://doi.org/10.1074/jbc.M007493200 PMID: 10984498.
11. Frey N, Richardson JA, Olson EN. Calsarcins, a novel family of sarcomeric calcineurin-binding proteins. Proc Natl Acad Sci U S A. 2000; 97(26):14632–7. https://doi.org/10.1073/pnas.260501097 PMID: 11114196; PubMed Central PMCID: PMCPMC18970.
12. Takada F, Vander Woude DL, Tong HQ, Thompson TG, Watkins SC, Kuncel LM, et al. Myozin-2: an alpha-actinin- and gamma-filamin-binding protein of skeletal muscle Z lines. Proc Natl Acad Sci U S A. 2001; 98(4):1595–600. https://doi.org/10.1073/pnas.041609698 PMID: 11171996; PubMed Central PMCID: PMC29302.

13. Gontier Y, Taiavainen A, Fontao L, Sonnenberg A, Flier Avd, Carpen O, et al. The Z-disc proteins myotilin and FATZ-1 interact with each other and are connected to the sarcolemma via muscle-specific filamin. Journal of Cell Science. 2005; 118:3739–49. https://doi.org/10.1242/jcs.02484 PMID: 16076904

14. Frey N, Olson EN. Calsarcin-3, a novel skeletal muscle-specific member of the calsarcin family, interacts with multiple Z-disc proteins. J Biol Chem. 2002; 277(16):13998–4004. https://doi.org/10.1074/jbc.M200712200 PMID: 11842093.

15. Chin ER, Olson EN, Richardson JA, Yang Q, Humphries C, Shetton JM, et al. A calcineurin-dependent transcriptional pathway controls skeletal muscle fiber type. GENES & DEVELOPMENT. 1998; 12:4299–509.

16. Frey N, Barrientos T, Shetton JM, Frank D, Rutten H, Gehring D, et al. Mice lacking calsarcin-1 are sensitized to calcineurin signaling and show accelerated cardiomyopathy in response to pathological biomechanical stress. Nat Med. 2004; 10(12):1336–43. https://doi.org/10.1038/nm132 PMID: 15543153.

17. Frey N, Frank D, Lipp S, Kuhn C, Kogler H, Barrientos T, et al. Calsarcin-2 deficiency increases exercise capacity in mice through calcineurin/NFAT activation. J Clin Invest. 2008; 118(11):3598–608. https://doi.org/10.1172/JCI36277 PMID: 18846255; PubMed Central PMCID: PMC2564612.

18. Frank D, Kuhn C, van Eickels M, Gehring D, Hanselmann C, Lipp S, et al. Calsarcin-1 protects against angiotensin-II induced cardiac hypertrophy. Circulation. 2007; 116(22):2587–96. https://doi.org/10.1161/CIRCULATIONAHA.107.711317 PMID: 18025526.

19. Paulsson AK, Franklin S, Mitchell-Jordan SA, Ren S, Wang Y, Vondriska TM. Post-translational regulation of calsarcin-1 during pressure overload-induced cardiac hypertrophy. J Mol Cell Cardiol. 2010; 48(6):1206–14. https://doi.org/10.1016/j.yjmcc.2010.02.009 PMID: 20706660; PubMed Central PMCID: PMC2666759.

20. Osio A, Tan L, Chen SN, Lombardi R, Nagueh SF, Shetton S, et al. Myozin-2 is a novel gene for human hypertrophic cardiomyopathy. Circ Res. 2007; 100(6):766–8. https://doi.org/10.1161/01.RES.0000263008.66799.ae PMID: 17347475; PubMed Central PMCID: PMC2775141.

21. Wang H, Zhu Z, Wang H, Yang S, Mo D, Li K. Characterization of different expression patterns of calsarcin-1 and calsarcin-2 in porcine muscle. Gene. 2006; 374:104–11. https://doi.org/10.1016/j.gene.2006.01.035 PMID: 16574346.

22. Wang Heng, Yang Shulin, Tang Zhonglin, Mu Yulian, Cui W, Li K. Expression Characterization, Polymorphism and Chromosomal Location of the Porcine Calsarcin-3 Gene. Asian-Aust J Anim Sci. 2007; 20(9):1349–53.

23. Yin Liu, Zou Ke, Li Xin-Yun, Zhao Shu-Hong, Cao H-J. Expression profiling of MYOZ1 Gene in Porcine Tissue and C2C12 cells. Journal of animal and veterinary advances. 2011; 10(15):1917–21.

24. Han L, Ma J, Wang N, Wang D, Xu G. Molecular cloning and characterization of different expression of MYOZ2 and MYOZ3 in Tianfu goat. PLoS One. 2013; 8(12):e82550. https://doi.org/10.1371/journal.pone.0082550 PMID: 24367529; PubMed Central PMCID: PMC3867352.

25. Yang Dapeng, Zan Linsen, Wang Hongbao, Ma Y. Genetic variation of calsarcin-1 gene and association with carcass traits in 3 Chinese indigenous cattle. African Journal of Biotechnology. 2009; 8(12):2713–7.

26. Frank D, Frey N. Cardiac Z-disc signaling network. J Biol Chem. 2011; 286(12):3987–904. https://doi.org/10.1074/jbc.R110.174268 PMID: 21257757; PubMed Central PMCID: PMC3060542.

27. Lin S, Luo W, Ye Y, Bekele EJ, Nie Q, Li Y, et al. Let-7b Regulates Myoblast Proliferation by Inhibiting IGF2B3 Expression in Dwarve and Normal Chicken. Front Physiol. 2017; 8(477). https://doi.org/10.3389/fphys.2017.00477 PMID: 28736533

28. Trapnell C, Roberts A, Goff L, Pertea G, Kim D, Kelley DR, et al. Differential gene and transcript expression analysis of RNA-seq experiments with TopHat and Cufflinks. Nat Protoc. 2012; 7(3):562–78. https://doi.org/10.1038/nprot.2012.016 PMID: 22383036; PubMed Central PMCID: PMC3334321.

29. Kim D, Pertea G, Trapnell C, Pimentel H, Kelley R, Salzberg SL. TopHat2: accurate alignment of transcriptomes in the presence of insertions, deletions and gene fusions. Genome biology. 2013.

30. Trapnell C, Pachter L, Salzberg SL. TopHat: discovering splice junctions with RNA-Seq. Bioinformatics. 2009; 25(9):1105–11. https://doi.org/10.1093/bioinformatics/btp120 PMID: 19289445; PubMed Central PMCID: PMC2672628.

31. Schmieder R, Edwards R. Quality control and preprocessing of metagenomic datasets. Bioinformatics. 2011; 27(6):863–4. https://doi.org/10.1093/bioinformatics/btq026 PMID: 21278183; PubMed Central PMCID: PMC3051327.
32. Robinson MD, McCarthy DJ, Smyth GK. edgeR: a Bioconductor package for differential expression analysis of digital gene expression data. Bioinformatics. 2010; 26(1):139–40. https://doi.org/10.1093/bioinformatics/btp616 PMID: 19910308; PubMed Central PMCID: PMCPMC2796818.

33. Huang DW, Sherman BT, Lempicki RA. Systematic and integrative analysis of large gene lists using DAVID bioinformatics resources. Nature Protocols. 2008; 4(1):44–57. https://doi.org/10.1038/nprot.2008.211 PMID: 19131956

34. Ye M, Ye F, He L, Liu Y, Zhao X, Yin H, et al. Molecular Cloning, Expression Profiling, and Marker Validation of the Chicken Myo23 Gene. BioMed Research International. 2017; 2017:10. https://doi.org/10.1155/2017/5930918 PMID: 28584817

35. Anders S, Pyl PT, Huber W. HTSeq—a Python framework to work with high-throughput sequencing data. Bioinformatics. 2015; 31(2):166–9. https://doi.org/10.1093/bioinformatics/btu638 PMID: 25260700; PubMed Central PMCID: PMCPMC4287950.

36. Pette D, Staron RS. Transitions of muscle fiber phenotypic profiles. Histochem Cell Biol. 2001; 115:359–72. https://doi.org/10.1007/s004180100268 PMID: 11449884

37. RUBINSTEIN NA, KELLY AM. Development of Muscle Fiber Specialization in the Rat Hindlimb. HE JOURNAL OF CELL BIOLOGY. 1981; 90:128–44.

38. Kolodney Michael S., Matthew S., Thimgan HM, Honda, Tsai G, aH F, et al. Ca+-independent myosin II phosphorylation and contraction in chicken embryonic fibroblasts. Journal of Physiology. 1999; 515.1:87–92.

39. Merlino GT, McKeong C, Crombrugge Bd, Pastan I. Regulation of the Expression of Genes Encoding Types I, 11, and I11 Collagen during Chick Embryonic Development. THE JOURNAL OF BIOLOGICAL. 1983; 258:10041–8.

40. Kim TH, Zhou H. Functional Analysis of Chicken IRF7 in Response to dsRNA Analog Poly(I:C) by Integrating Overexpression and Knockdown. Plos One. 2015; 10(7):e33405. https://doi.org/10.1371/journal.pone.0133450 PMID: 26186542

41. Lu Y, West FD, Jordan BJ, Jordan ET, West RC, Yu P, et al. Induced pluripotency in chicken embryonic cells: evidence for transdifferentiation. Stem Cells and Development. 2014; 23(15):1755–64. https://doi.org/10.1089/scd.2014.0080 PMID: 24720794

42. WEINTRAUB H, Tapscott Stephen J., DAVIS RL, THAYER MJ, ADAM MA, LASSAR AB, et al. Activation of muscle-specific genes in pigment, nerve, fat, liver, and fibroblast cell lines by forced expression of Myod. Proc Natl Acad Sci U S A. 1989; 86:5434–8. PMID: 2748593

43. Hong H, Kim J, Kim J. Myosin heavy chain 10 (MYH10) is required for centriole migration during the biogenesis of primary cilia. Biochem Biophys Res Commun. 2015; 461(1):180–5. https://doi.org/10.1016/j.bbrc.2015.04.028 PMID: 25881509.

44. Ma X, Adelstein RS. A point mutation in Myh10 causes major defects in heart development and body wall closure. Circ Cardiovasc Genet. 2014; 7(3):257–65. https://doi.org/10.1161/CIRCGENETICS.113.000455 PMID: 24825879; PubMed Central PMCID: PMCPMC4106703.

45. Sparrow JC, Schock F. The initial steps of myofibril assembly: integrins pave the way. Nat Rev Mol Cell Biol. 2010; 10(4):293–8. https://doi.org/10.1038/nrm2786 PMID: 19851336; PubMed Central PMCID: PMCPMCC2834236.

46. Vicente-Manzanares M, Ma X, Adelstein RS, Horwitz AR. Non-muscle myosin II takes centre stage in cell adhesion and migration. Nat Rev Mol Cell Biol. 2009; 10(11):778–90. https://doi.org/10.1038/nrm2786 PMID: 19851336; PubMed Central PMCID: PMCPMCC2834236.

47. Herring BP, El-Mounayri O, Gallagher PJ, Yin F, Zhou J. Regulation of myosin light chain kinase and telokin expression in smooth muscle tissues. Am J Physiol Cell Physiol. 2006; 291(5):C817–27. https://doi.org/10.1152/ajpcell.00198.2006 PMID: 16774989; PubMed Central PMCID: PMCPMCC2836780.

48. Krendel M, Moeseker MS. Myosins: Tails (and Heads) of Functional Diversity. Physiology. 2005; 20:239–51. https://doi.org/10.1152/physiol00014.2005 PMID: 16024512

49. Shehadeh LA, Webster KA, Hare JM, Vazquez-Padron RI. Dynamic regulation of vascular myosin light chain (MYL9) with injury and aging. PLoS One. 2011; 6(10):e25855. https://doi.org/10.1371/journal.pone.0025855 PMID: 22003410; PubMed Central PMCID: PMCPMC3189218.

50. Orr N, Arnaout R, Gula LJ, Spears DA, Leong-Sit P, Li Q, et al. A mutation in the atrial-specific myosin light chain gene (MYL4) causes familial atrial fibrillation. Nat Commun. 2016; 7:11303. https://doi.org/10.1038/ncomms11303 PMID: 27066836; PubMed Central PMCID: PMCPMCC4832069.

51. Ohtsuka M, Arase H, Takeuchi A, Yamazaki S, Shina R, Suenaga T, et al. NFAM1, an immunoreceptor tyrosine-based activation motif-bearing molecule that regulates B cell development and signaling. Proc Natl Acad Sci U S A. 2004; 101(21):8126–31. https://doi.org/10.1073/pnas.0401119101 PMID: 15143214; PubMed Central PMCID: PMCPMCC419568.

52. Frank D, Kuhn C, Katus HA, Frey N. The sarcomeric Z-disc: a nodal point in signalling and disease. J Mol Med (Berl). 2006; 84(6):446–68. https://doi.org/10.1007/s00109-005-0033-1 PMID: 16416311.
53. Katzemich A, Liao KA, Czerniecki S, Schock F. Alp/Enigma family proteins cooperate in Z-disc formation and myofibril assembly. PLoS Genet. 2013; 9(3):e1003342. https://doi.org/10.1371/journal.pgen.1003342 PMID: 23505387; PubMed Central PMCID: PMCPMC3591300.

54. Sheikh F, Bang M-L, Lange S, Chen J. “Z”eroing in on the Role of Cypher in Striated Muscle Function, Signaling, and Human Disease. Trends Cardiovasc Med. 2007; 17(8):258–62. https://doi.org/10.1016/j.tcm.2007.09.002 PMID: 18021935.

55. Manickam R, Wahl W. Roles of Peroxisome Proliferator-Activated Receptor β/d in skeletal muscle physiology. Biochimie. 2017; 136:42–8. https://doi.org/10.1016/j.biochi.2016.11.010 PMID: 27916646.

56. Kramer DK, Ahlsen M, Norrbom J, Jansson E, Hjeltnes N, Gustafsson T, et al. Human skeletal muscle fibre type variations correlate with PPAR alpha, PPAR delta and PGC-1 alpha mRNA. Acta Physiol (Oxf). 2006; 188(3–4):207–16. https://doi.org/10.1111/j.1748-1716.2006.01620.x PMID: 17054660.

57. Gizard F, Nomiyama T, Zhao Y, Findeisen HM, Heywood EB, Jones KL, et al. The PPARalpha/p16INK4a pathway inhibits vascular smooth muscle cell proliferation by repressing cell cycle-dependent telomerase activation. Circ Res. 2008; 103(10):1155–63. https://doi.org/10.1161/CIRCRESAHA.108.186205 PMID: 18818403; PubMed Central PMCID: PMCPMC2756491.

58. Gan Z, Burkart-Hartman EM, Han DH, Finck B, Leone TC, Smith EY, et al. The nuclear receptor PPAR-beta/delta programs muscle glucose metabolism in cooperation with AMPK and MEF2. Genes Dev. 2011; 25(24):2619–30. https://doi.org/10.1101/gad.178434.111 PMID: 22135324; PubMed Central PMCID: PMCPMC3248683.

59. Wang YX, Zhang CL, Yu RT, Cho HK, Nelson MC, Bayuga-Ocampo CR, et al. Regulation of Muscle Fiber Type and Running Endurance by PPARd. PLoS Biol 2004; 2(10):e294 https://doi.org/10.1371/journal.pbio.0020294 PMID: 15328533.

60. Luquet S, Lopez-Soriano J, Holst D, Fredenrich A, Melki J, Rassoulzadegan M, et al. Peroxisome proliferator-activated receptor δ controls muscle development and oxidative capability The FASEB Journal 2003. https://doi.org/10.1096/fj.03-0269fje PMID: 14525942.

61. Schnuck JK, Sunderland KL, Nicholas b G, Kuennen MR, Vaughan RA. Leucine stimulates PPARβ/d-dependent mitochondrial biogenesis and oxidative metabolism with enhanced GLUT4 content and glucose uptake in myotubes. Biochimie. 2016. https://doi.org/10.1016/j.biochi.2016.06.009 PMID: 27345255.

62. Thach TT, Lee C-K, Park HW, Lee S-J, Lee S-J. Syringaresinol induces mitochondrial biogenesis through activation of PPARβ pathway in skeletal muscle cells. Bioorganic & Medicinal Chemistry Letters. 2016; 2016:3878–83. https://doi.org/10.1016/j.bmcl.2016.07.001 PMID: 27450786.

63. Liang Qiangrong, De Windt Leon J., Witt Sandra A., Kimball Thomas R., Markham Bruce E., Molkentin JD. The Transcription Factors GATA4 and GATA6 Regulate Cardiomyocyte Hypertrophy in Vitro and in Vivo. THE JOURNAL OF BIOLOGICAL CHEMISTRY. 2001; 276(32):30245–53. https://doi.org/10.1074/jbc.M102174200 PMID: 11356841.

64. Boopath E, Hypolite JA, A. S, Zderic bCMG, Malkowicz Bruce, Liou Hsiou-Chi, et al. GATA-6 and NF-B Activate CPI-17 Gene Transcription and Regulate Ca2+ Sensitization of Smooth Muscle Contraction. Molecular and Cellular Biology. 2012; 33(5):1085–102. https://doi.org/10.1128/MCB.00626-12 PMID: 23275439.