

Relative importance of β_{cyto} - and γ_{cyto} -actin in primary mouse embryonic fibroblasts

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ABSTRACT The highly homologous β (β_{cyto}) and γ (γ_{cyto}) cytoplasmic actins are hypothesized to carry out both redundant and unique essential functions, but studies using targeted gene knockout and siRNA-mediated transcript knockdown to examine β_{cyto} - and γ_{cyto} -isoform-specific functions in various cell types have yielded conflicting data. Here we quantitatively characterized actin transcript and protein levels, as well as cellular phenotypes, in both gene- and transcript-targeted primary mouse embryonic fibroblasts. We found that the smooth muscle α_{sm} -actin isoform was the dominantly expressed actin isoform in WT primary fibroblasts and was also the most dramatically up-regulated in primary β_{cyto} - or $\beta/\gamma_{\text{cyto}}$ -actin double-knockout fibroblasts. Gene targeting of β_{cyto} -actin, but not γ_{cyto} -actin, led to greatly decreased cell proliferation, decreased levels of cellular ATP, and increased serum response factor signaling in primary fibroblasts, whereas immortalization induced by SV40 large T antigen supported fibroblast proliferation in the absence of β_{cyto} -actin. Consistent with *in vivo* gene-targeting studies in mice, both gene- and transcript-targeting approaches demonstrate that the loss of β_{cyto} -actin protein is more disruptive to primary fibroblast function than is the loss of γ_{cyto} -actin.

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INTRODUCTION

Vertebrate actin isoforms, which are encoded by six different genes, share a very high degree of sequence homology and are completely conserved from birds to mammals (Rubenstein, 1990). Four of the actin isoforms are predominately expressed in smooth (α_{sm} - and γ_{sm} -actin), cardiac (α_{ca} -actin), and skeletal (α_{sk} -actin) muscles. The two cytoplasmic actins (β_{cyto} - and γ_{cyto} -actin) are ubiquitously expressed in all cell types and differ at only four of 375 amino acids. Despite the remarkable conservation between the β_{cyto} - and γ_{cyto} -actin sequences, purified recombinant β_{cyto} - and γ_{cyto} -actin proteins display marked differences in dynamics *in vitro* (Bergeron *et al.*, 2010). Furthermore, numerous studies of

transcript/protein localization, protein overexpression, and disease-causing mutations have established a strong argument for unique functions supported by β_{cyto} - or γ_{cyto} -actin (Khaitlina, 2001; Condeelis and Singer, 2005; Rubenstein and Wen, 2014). More recently, gene-targeting studies have been used to test for non-overlapping functions of actin isoforms (Perrin and Ervasti, 2010). With regard to closely related β_{cyto} - and γ_{cyto} -actins, the embryonic lethality of mice homozygous for severely hypomorphic (Shawlot *et al.*, 1998; Shmerling *et al.*, 2005) or null alleles of *Actb* (Bunnell and Ervasti, 2011) demonstrated that β_{cyto} -actin is an essential gene. In contrast, γ_{cyto} -actin-null animals survive to term but display significant perinatal lethality, stunted growth, and decreased survival into adulthood (Belyantseva *et al.*, 2009; Bunnell and Ervasti, 2010). The distinct auditory hair cell and deafness phenotypes in mice in which floxed *Actb* and *Actg1* alleles were selectively ablated also support unique roles for β_{cyto} - and γ_{cyto} -actin in some tissues (Perrin *et al.*, 2010, 2013). On the other hand, mice with skeletal muscle-specific knockouts of either β_{cyto} - or γ_{cyto} -actin each develop a similar myopathy, with progressive myofiber degeneration/regeneration and muscle weakness (Sonnemann *et al.*, 2006; Prins *et al.*, 2011), supporting that some functions of β_{cyto} - and γ_{cyto} -actins are redundant in some tissues.

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Abbreviations used: α_{sm} -actin, α smooth-muscle actin; MEF, mouse embryonic fibroblast; SRF, serum response factor.

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We previously investigated the isoform-specific functions of β_{cyto} - and γ_{cyto} -actin in primary mouse embryonic fibroblasts (MEFs) from *Actg1*^{-/-} animals (Bunnell and Ervasti, 2010; Bunnell et al., 2011) and from mice with floxed *Actb* alleles conditionally ablated by tamoxifen-inducible Cre recombinase (Bunnell et al., 2011). γ_{cyto} -Actin-null MEFs showed mild growth impairment and a small decrease in cell viability but were otherwise comparable to control cells in all other parameters tested (Bunnell and Ervasti, 2010; Bunnell et al., 2011). In contrast, β_{cyto} -actin-knockout MEFs exhibited a wide range of more severe phenotypes (Bunnell et al., 2011). Ablation of β_{cyto} -actin resulted in severe growth impairment and an increase in the percentage of multinucleate cells, suggesting a specific role for β_{cyto} -actin in cell growth and division (Bunnell et al., 2011), as previously supported by localization of β_{cyto} -actin to the cleavage furrow and contractile ring during cell division (Dugina et al., 2009). Furthermore, β_{cyto} -actin-knockout cells exhibited significant migration defects that were not observed in γ_{cyto} -actin-null cells. The altered expression of genes that regulate the cell cycle and cell migration observed in β_{cyto} -actin- but not γ_{cyto} -actin-knockout MEFs provided a potential explanation for the more severe phenotypes in β_{cyto} -actin-knockout cells and early lethality in β_{cyto} -actin-knockout embryos (Bunnell et al., 2011). It is also interesting that β_{cyto} -actin-knockout MEFs exhibited some characteristics reminiscent of myofibroblasts, such as an increase in stress fibers and greatly elevated expression of α_{sm} -actin (Bunnell et al., 2011).

In contrast to our results, four studies reached different conclusions that warrant further investigation. One study of immortalized β_{cyto} -actin-knockout MEF lines transformed by SV40 large T antigen (Tondeleir et al., 2012) reported altered morphology, impaired migration, and up-regulation of α_{sm} -actin similar to what we observed in primary *Actb*-null MEFs (Bunnell et al., 2011) but with no impairment of cell growth. Two studies using small interfering RNA (siRNA) methodologies reported that knockdown of γ_{cyto} -actin more dramatically impaired motility in human subcutaneous fibroblasts, spontaneously immortalized keratinocytes, and neuroblastoma SH-EP cells than did knockdown of β_{cyto} -actin (Dugina et al., 2009; Shum et al., 2011). Most notably, the fourth study demonstrated specific up-regulation of myofibroblast-associated gene products (including α_{sm} -actin) after siRNA knock down of γ_{cyto} -actin, but not β_{cyto} -actin, in several epithelial cell lines (Lechuga et al., 2014). Although differences in methodologies and cell types studied are obvious explanations for the different outcomes, Lechuga et al. (2014) further speculated that the different results could depend on whether the ablated actin isoform was also the most abundant in the cell type under investigation.

To address the bases for differences between our studies (Bunnell and Ervasti, 2010; Bunnell et al., 2011) and those of others (Dugina et al., 2009; Shum et al., 2011; Tondeleir et al., 2012; Lechuga et al., 2014), we quantitatively profiled actin isoforms at both the transcript and the protein level. Because MEFs display extensive developmental and phenotypic heterogeneity (Singhal et al., 2016), we used adenovirus-5-mediated expression of Cre recombinase (Ad5-Cre) to ablate *Actb* and/or *Actg1* in both primary and large T antigen immortalized MEFs controlled against the same MEFs infected with Ad5-GFP. α_{sm} -Actin and β_{cyto} -actin transcripts predominated in primary MEFs, whereas α_{sm} -actin was most abundant at the protein level, with equivalent levels of β_{cyto} - and γ_{cyto} -actin protein expressed despite their marked differences in transcript levels. We also found that immortalization by transformation with SV40 large T antigen appears to negate the essentiality of *Actb* observed in primary MEFs. Finally, siRNA-mediated transcript knockdown of *Actb* and *Actg1* in the same cells used for gene-targeting studies yielded

results similar to those observed after gene targeting. We conclude that the genetic reprogramming associated with immortalization by transformation with SV40 large T antigen can markedly influence the cellular response to β_{cyto} - and γ_{cyto} -actin protein ablation.

RESULTS

Adenoviral Cre-mediated knockout of *Actb* and *Actg1*

Because primary MEFs display extensive developmental and phenotypic heterogeneity (Singhal et al., 2016), each of our biological replicates consisted of E13.5 primary MEFs obtained from a single mouse embryo carrying floxed alleles of *Actb* (*Actb*^{LoxP}; Bunnell et al., 2011), *Actg1* (*Actg1*^{LoxP}; Sonnemann et al., 2006), or both *Actb* and *Actg1* (*Actb*^{LoxP} *Actg1*^{LoxP}). MEFs from each animal were split into two pools, which were treated with Ad5-green fluorescent protein (GFP; Figure 1A) as a negative control or Ad5-Cre (Figure 1B) to conditionally knock out *Actb* (β_{cyto} -actin KO), *Actg1* (γ_{cyto} -actin KO), or both *Actb* and *Actg1* (dKO). Quantitative reverse transcription PCR (qRT-PCR) analysis showed that *Actb* and *Actg1* transcripts were reduced 95% by 3 d postinfection (dpi) in β_{cyto} -actin KO, γ_{cyto} -actin KO, and dKO MEFs compared with controls (Figure 1, C–F). Western blot analysis revealed that β_{cyto} - and γ_{cyto} -actin were each undetectable at 5 dpi in single β_{cyto} -actin KO or γ_{cyto} -actin KO cells, respectively (Figure 1, G and H). In contrast, complete loss of β_{cyto} - and γ_{cyto} -actin in dKO cells was observed at 9 dpi (Figure 1I). Thus Ad5-Cre was an efficient and robust method to eliminate both β_{cyto} - and γ_{cyto} -actin transcripts and proteins in primary MEFs.

β -Actin KO MEFs are growth impaired with reduced ATP levels

We previously showed that tamoxifen-induced, cre-mediated ablation of β_{cyto} -actin caused significant growth impairment in primary MEFs, whereas only a modest decrease in cell growth was observed in *Actg1*^{-/-} MEFs (Bunnell and Ervasti, 2010; Bunnell et al., 2011). However, our previous experiments involved different methods of gene ablation for *Actb* and *Actg1* and compared knockout MEFs with controls derived from different embryos (Bunnell and Ervasti, 2010; Bunnell et al., 2011). Therefore we more directly compared the effects of β_{cyto} - versus γ_{cyto} -actin gene ablation on cell growth, using β_{cyto} -actin KO and γ_{cyto} -actin KO MEFs. Under these more controlled conditions, γ_{cyto} -actin KO MEFs exhibited a normal growth rate, whereas both β_{cyto} -actin KO and dKO MEFs displayed growth impairment by 3 dpi that became significantly different from controls at 7 dpi (Figure 2, A–C). Because growth deficiency was not previously observed in MEFs ablated for β_{cyto} -actin expression after transformation by SV40 large T antigen (Tondeleir et al., 2012), we immortalized *Actb*^{LoxP}, *Actg1*^{LoxP}, and double *Actb*^{LoxP} *Actg1*^{LoxP} MEFs by transformation with SV40 large T antigen and measured cell growth after infection with either Ad5-Cre or Ad5-GFP. Both SV40 large T antigen immortalized β_{cyto} -actin KO and γ_{cyto} -actin KO MEFs exhibited growth rates not different from controls (Supplemental Figure S1, A and B), suggesting that β_{cyto} -actin is important for cell growth in primary MEFs but not in MEFs transformed by SV40 large T antigen. In contrast, SV40 large T antigen immortalized dKO MEFs displayed growth impairment that was significantly different from control at 8–10 dpi (Supplemental Figure S1C) but less severe than in primary MEFs, suggesting that MEFs immortalized by SV40 large T antigen transformation are able to circumvent the essential function supported by β_{cyto} -actin in primary cells and during embryonic development.

To assess whether growth deficiencies seen in β_{cyto} -actin KO and dKO MEFs could potentially be due to decreases in energy availability, we used a luciferase-based assay to measure the relative

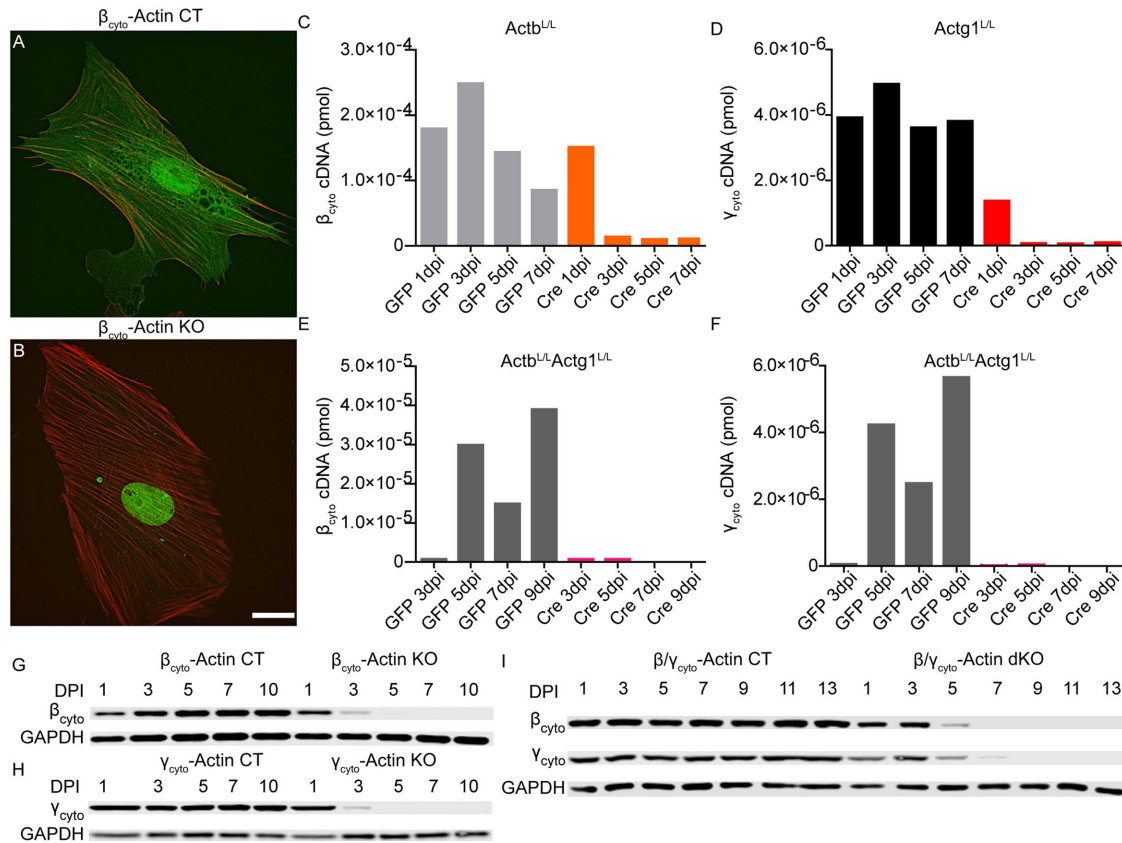


FIGURE 1: Adenoviral Cre efficiently ablated β_{cyto} - and γ_{cyto} -actin in primary MEFs. (A, B) Representative images of β_{cyto} -actin CT and β_{cyto} -actin KO cells at 7 dpi. Scale bar, 20 μm . (C–F) Representative qRT-PCR analysis of β_{cyto} - and γ_{cyto} -actin transcript amount (picomoles) in $\text{Actb}^{\text{L/L}}$, $\text{Actg1}^{\text{L/L}}$, and $\text{Actb}^{\text{L/L}}\text{Actg1}^{\text{L/L}}$ MEFs. (G–I) Representative relative Western blot analysis of MEF lysates probed with β_{cyto} -actin and γ_{cyto} -actin antibodies; GAPDH served as loading control.

levels of cellular ATP in β_{cyto} -actin KO, γ_{cyto} -actin KO and dKO MEFs. Correlating with the deficiencies seen in growth, ATP levels were significantly decreased in β_{cyto} -actin KO and dKO MEFs compared with controls but not in γ_{cyto} -actin KO MEFs (Figure 3A). However, Western blot analysis demonstrated that a number of proteins in the mitochondrial electron transport chain complexes were equivalent across the MEF lines tested (Figure 3, B and C), as were basal and maximal oxygen consumption rates measured in β_{cyto} -actin KO and γ_{cyto} -actin KO MEFs using a Seahorse XF24 extracellular flux analyzer (Figure 3D). Therefore we conclude that the measured decreases in cellular ATP are not due to impaired mitochondrial function in β_{cyto} -actin KO and dKO MEFs.

Quantitation of actin isoform transcript and protein levels in primary MEFs

We used qRT-PCR to measure transcript levels for all six actin isoforms in control, β_{cyto} -actin KO, γ_{cyto} -actin KO, and dKO MEFs at 5 dpi. The amount of transcript for each actin isoform in MEFs was calculated from standard curves that were amplified in parallel (Supplemental Figure S2, A–F). In all control MEFs, α_{sm} - and β_{cyto} -actins were the dominant transcripts (Figure 4, A–C), in contrast to NIH3T3s fibroblasts cultured under the same conditions, where β_{cyto} - and γ_{cyto} -actins were the predominant transcripts (Supplemental Figure S3A). α_{sm} -Actin was the only actin transcript significantly up-regulated in β_{cyto} -actin KO MEFs (Figure 4A), whereas both α_{sm} - and β_{cyto} -actins transcripts were up-regulated in γ_{cyto} -actin KO MEFs (Figure 4B). Similar to β_{cyto} -actin KO MEFs (Figure 4A), α_{sm} -actin was the only transcript significantly up-regulated in dKO MEFs

(Figure 4C). The high levels of α_{sm} -actin transcript suggest a myofibroblast-like phenotype in the primary MEFs that was enhanced to a greater extent by ablation of *Actb* over *Actg1*.

Because α_{sm} -, β_{cyto} -, and γ_{cyto} - were the most abundant actin transcripts in primary MEFs, we measured α_{sm} -, β_{cyto} -, and γ_{cyto} -actin protein levels in both control and KO MEFs via quantitative Western blot analysis at 5 dpi in each single KO and at 9 dpi in the dKO using purified α_{sm} -, β_{cyto} -, and γ_{cyto} -actin proteins as standards. α_{sm} -Actin was the most abundant actin isoform protein expressed, and β_{cyto} - and γ_{cyto} -actin proteins were expressed to similar levels in all control MEFs (Figure 5, B–D). Consistent with transcript analysis (Supplemental Figure S3A), β_{cyto} - and γ_{cyto} -actin were the dominant actin proteins expressed in NIH3T3 fibroblasts cultured under the same conditions (Supplemental Figure S3B). In β_{cyto} -actin KO MEFs, α_{sm} -actin protein was dramatically increased by sixfold over controls, but γ_{cyto} -actin protein was not significantly increased (Figure 5, A and B). In γ_{cyto} -actin KO MEFs, α_{sm} -actin protein was up-regulated by 1.7-fold, whereas β_{cyto} -actin protein was not significantly increased (Figure 5, A and C). α_{sm} -Actin protein was also significantly up-regulated by sevenfold in dKO MEFs (Figure 5, A and D). Thus α_{sm} -actin was prominently expressed in all control primary MEFs and more dramatically up-regulated upon ablation of *Actb* compared with *Actg1*, suggesting that primary MEFs may be predisposed to a myofibroblast-like phenotype that can be most enhanced by selective perturbation of *Actb*. It is also interesting that β_{cyto} -actin transcript was expressed sixfold higher than γ_{cyto} -actin transcript in all control MEFs, and yet the ratio of β_{cyto} -actin protein to γ_{cyto} -actin protein was 1:1 (Figure 6, A–C). The unequal transcript-to-protein

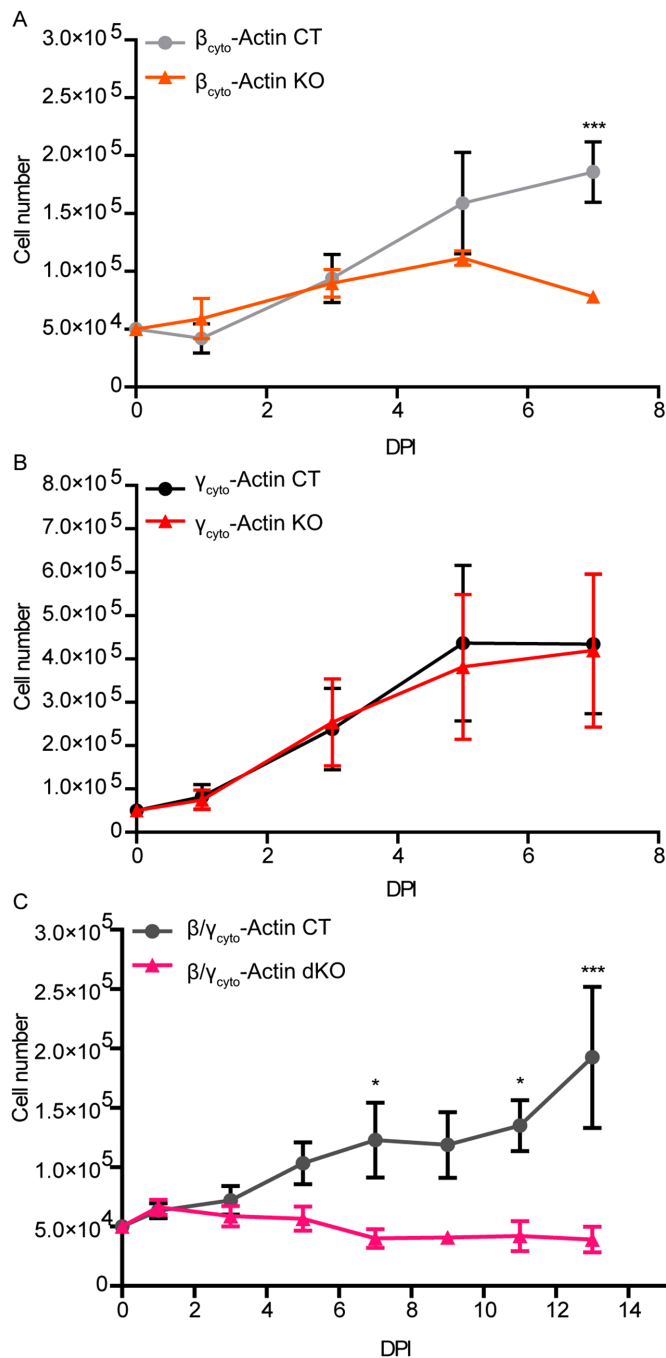


FIGURE 2: β_{cyto} -actin deficient MEFs were growth impaired. (A–C) Growth curve analysis of $\text{Actb}^{\text{L/L}}$, $\text{Actg1}^{\text{L/L}}$, and $\text{Actb}^{\text{L/L}}\text{Actg1}^{\text{L/L}}$ MEFs treated with either Ad5-GFP or Ad5-Cre ($n = 3$, hand counted in duplicate). * $p < 0.05$, *** $p < 0.001$. Two-way ANOVA with Bonferroni posttest; error bars are SEM.

ratio supports other studies concluding that *Actb* and *Actg1* gene expression are differentially regulated posttranscriptionally (Hüttelmaier *et al.*, 2005; Zhang *et al.*, 2010).

siRNA-mediated knockdown of *Actb* and *Actg1*

Because other studies reporting results different from ours used siRNA approaches to knock down *Actb* and *Actg1* transcripts and proteins (Dugina *et al.*, 2009; Shum *et al.*, 2011; Lechuga *et al.*, 2014), we assessed how siRNA-mediated knockdown (KD) of β_{cyto} -

and/or γ_{cyto} -actin in $\text{Actb}^{\text{L/L}}$ (β_{cyto} -actin KD), $\text{Actg1}^{\text{L/L}}$ (γ_{cyto} -actin KD), and $\text{Actb}^{\text{L/L}}\text{Actg1}^{\text{L/L}}$ ($\beta/\gamma_{\text{cyto}}$ -actin double KD [dKD]) primary MEFs affected cell phenotype. On average, we were able to specifically knock down β_{cyto} -actin protein by 70% and γ_{cyto} -actin protein by 55% compared with controls. In dKD MEFs, β_{cyto} -actin protein was reduced by 65% and γ_{cyto} -actin protein by 55% (Figure 7A). Quantitative Western blot analysis showed that α_{sm} -actin protein was dramatically increased ninefold in β_{cyto} -actin KD cells over controls, but γ_{cyto} -actin protein was not significantly increased (Figure 7, A and B). In γ_{cyto} -actin KD cells, neither α_{sm} -actin protein nor β_{cyto} -actin protein was significantly increased (Figure 7, A and C). α_{sm} -Actin protein was also significantly up-regulated fourfold in $\beta/\gamma_{\text{cyto}}$ -actin dKD cells (Figure 7, A and D). Thus siRNA-mediated transcript KD and cre-mediated gene KO gave similar results in primary MEFs.

Actb gene ablation enhances some myofibroblast-like phenotypes of primary MEFs

In response to a variety of perturbations (Small, 2012; Davis and Molkenin, 2013), fibroblasts often differentiate into smooth muscle-like cells called myofibroblasts via a process called the fibroblast-to-myofibroblast transition (FMT). In addition to increased α_{sm} -actin expression, another hallmark of myofibroblasts is increased stress fiber formation (Small, 2012; Davis and Molkenin, 2013), which we observed in β_{cyto} -actin KO, γ_{cyto} -actin KO, and dKO MEFs compared with controls (Figure 8, A–F). To quantify the change in the stress fibers of KO MEFs, we made line scans across the cell width (Figure 8G). Peak-to-valley ratios were taken as an average measure of the fluorescence intensity of the stress fibers (Figure 8H). All KO cell types displayed significant elevation in the average peak-to-valley ratios, indicating increased stress fiber formation (Figure 8, A–F). Because caldesmon, calponin, and transgelin proteins have been shown to be up-regulated during epithelial-to-myofibroblast transition (Lechuga *et al.*, 2014), we measured their protein levels in addition to myosin light chain 2 and phosphorylated myosin light chain 2 protein levels in control and knockout MEFs. Caldesmon expresses a higher-molecular weight isoform in smooth muscle cells (Cald1_{sm}) than in non-smooth muscle cells (Ueki *et al.*, 1987; Frid *et al.*, 1992). In agreement with our previously published data (Bunnell *et al.*, 2011), Cald1_{sm} was up-regulated in β_{cyto} -actin KO, γ_{cyto} -actin KO, and dKO primary MEFs, although more so in β_{cyto} -actin KO MEFs than γ_{cyto} -actin KO MEFs (Figure 9, A–D). In contrast, none of the other myofibroblast marker proteins interrogated were significantly up-regulated (Figure 9, A–D). siRNA-treated primary MEFs showed similar up-regulation of Cald1_{sm} in KD MEFs (Supplemental Figure S4, A–D). Thus increased stress fiber formation and expression of Cald1_{sm} suggests that ablation of *Actb* enhances a partial myofibroblast-like phenotype in primary MEFs.

Finally, actin dynamics is known to play a role in myofibroblast differentiation via the serum response factor (SRF)/myocardin-related transcription factor (MRTF) gene regulatory pathway (Small, 2012). Therefore we investigated whether SRF or MRTF-A protein expression or SRF activity was perturbed when cytoplasmic actins were ablated. In all knockout MEFs, SRF protein levels were not significantly altered compared with controls (Figure 10, A and C). MRTF-A protein level was down-regulated in all gene-knockout MEFs, which was significant in dKO MEFs (Figure 10, A and B). Similar results were observed in siRNA KD MEFs (Supplemental Figure S5, A and B). Although no drastic changes in protein expression were observed, SRF activity was significantly increased by sixfold in both β_{cyto} -actin KO and dKO MEFs compared with controls. SRF activity increased by twofold in γ_{cyto} -actin KO MEFs, although this increase was not significantly different from controls. Our data

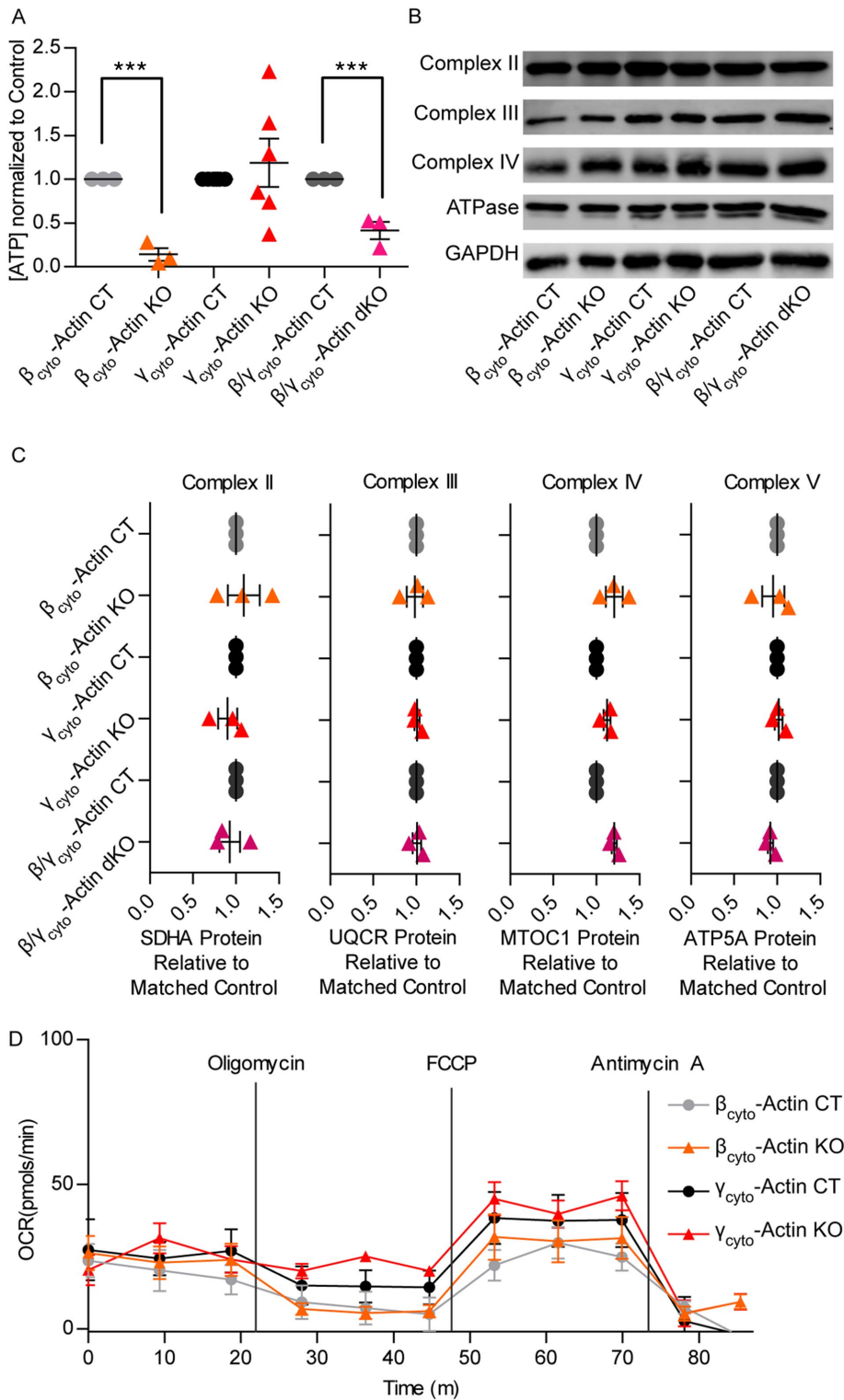


FIGURE 3: β_{cyto} -Actin-deficient MEFs displayed lower ATP levels but maintained ETC protein abundance. (A) ATP levels for CT and KO MEFs determined through a luciferase assay at 7 dpi. Results were normalized to cell number and relative to the paired embryo control ($6 \geq n \geq 3$). (B, C) Western blot and relative quantification of protein levels of succinate dehydrogenase complex, subunit A, flavoprotein (SDHA) for complex II; ubiquinol-cytochrome C reductase core protein I (UQCR) for complex III; mitochondrially encoded cytochrome C oxidase I (MTCO1) for complex IV; and ATP5a ATP synthase, H⁺ transporting, mitochondrial F1 complex, α -subunit (ATP5A) for the F1/F0 ATPase; GAPDH served as loading control. Levels were normalized to GAPDH and relative to the paired embryo control, which was set at 1 ($n = 3$). (D) Oxygen consumption rate of CT and KO MEFs at 7 dpi. Levels normalized to background. *** $p < 0.001$. One sample t test; error bars are SEM.

suggest that Actb knockout more significantly and specifically affects the SRF/MRTF gene regulatory pathway than does the loss of Actg1.

DISCUSSION

β_{cyto} -Actin is believed to be an essential protein, based on the early embryonic lethality observed in Actb gene-targeted mice (Shawlot *et al.*, 1998; Shmerling *et al.*, 2005; Bunnell *et al.*, 2011). In addition, many cell biological studies provide further compelling support to the essentiality of β_{cyto} -actin in a wide array of cellular processes (Zhao *et al.*, 1998; Hofmann *et al.*, 2004; Hu *et al.*, 2004; Karakozova *et al.*, 2006; Leung *et al.*, 2006; Yao *et al.*, 2006). Although γ_{cyto} -actin is clearly important *in vivo*, it is not essential, given that Actg1-null mice can survive up to 2 years (Belyantseva *et al.*, 2009). Here we sought to resolve the different results reported across a group of cell-based studies all seeking to elucidate the respective roles of β_{cyto} - and γ_{cyto} -actin via ablation of protein expression using different targeting approaches in different cell types. Most vexing, β_{cyto} -actin gene knockout in two studies caused essentially the same effects on cell motility and α_{sm} -actin expression (Bunnell *et al.*, 2011; Tondeleir *et al.*, 2012) as those caused by γ_{cyto} -actin knockdown in three other studies (Dugina *et al.*, 2009; Shum *et al.*, 2011; Lechuga *et al.*, 2014). Because gene and transcript targeting cause strikingly different *in vivo* phenotypes for a number of loci (Rossi *et al.*, 2015), we anticipated that actin gene knockout and transcript knockdown experiments carried out in the same cells would explain the discrepant results. Instead, we observed the same experimental outcomes whether we used siRNA-mediated transcript knockdown or Cre-mediated gene knockout. As an alternative explanation for the discrepant results across studies (Dugina *et al.*, 2009; Bunnell *et al.*, 2011; Shum *et al.*, 2011; Tondeleir *et al.*, 2012; Lechuga *et al.*, 2014), Lechuga *et al.* (2014) hypothesized that loss of the most abundantly expressed actin isoform may dictate a given cell type's response to actin gene ablation. However, our quantitative Western blot analysis demonstrated that β_{cyto} - and γ_{cyto} -actin proteins were present in equal abundance in the primary MEFs studied here. We conclude that neither differences in actin isoform-targeting method or endogenous protein levels can explain the dramatically different and often opposite results reported in the six studies from five different groups.

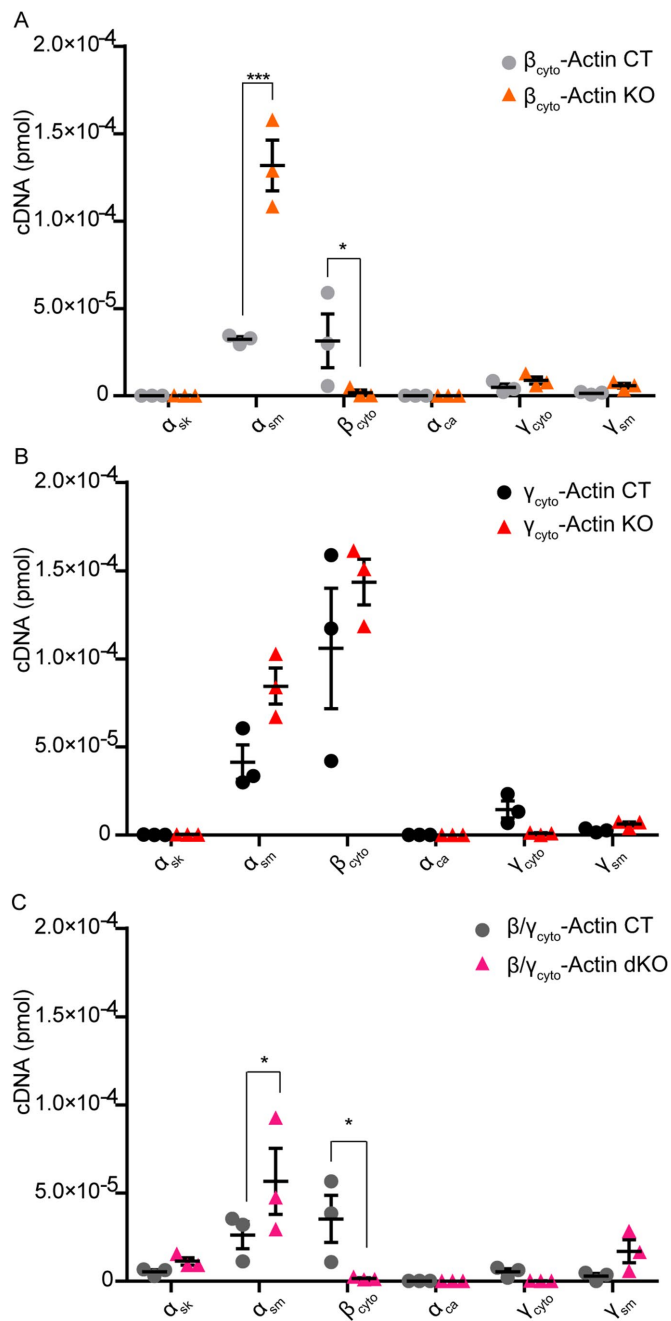


FIGURE 4: α_{sm} -Actin transcript was up-regulated in β_{cyto} -actin-ablated MEFs. (A–C) qRT-PCR analysis of six mouse actin isoforms in CT and KO MEFs at 5 dpi ($n = 3$, in triplicate). Calculated transcript amount (picomoles) were calculated based on the standard curve, amplified in parallel. * $p < 0.05$, *** $p < 0.001$. Two-way ANOVA with Bonferroni posttest; error bars are SEM.

Here (Figure 2) and previously (Bunnell et al., 2011), we measured dramatic cell proliferation defects with ablation of β_{cyto} -actin in primary MEFs, which could account for the embryonic lethality associated with *Actb* gene deletion (Shawlot et al., 1998; Shmerling et al., 2005; Bunnell et al., 2011). We measured significantly decreased levels of cellular ATP associated with knockout of β_{cyto} -actin in the presence of apparently unaffected mitochondrial function (Figure 3). Although the basis for decreased ATP levels in β_{cyto} -actin knockout MEFs is an interesting question for future studies, it will be challenging to answer, given that ablation of functionally disparate

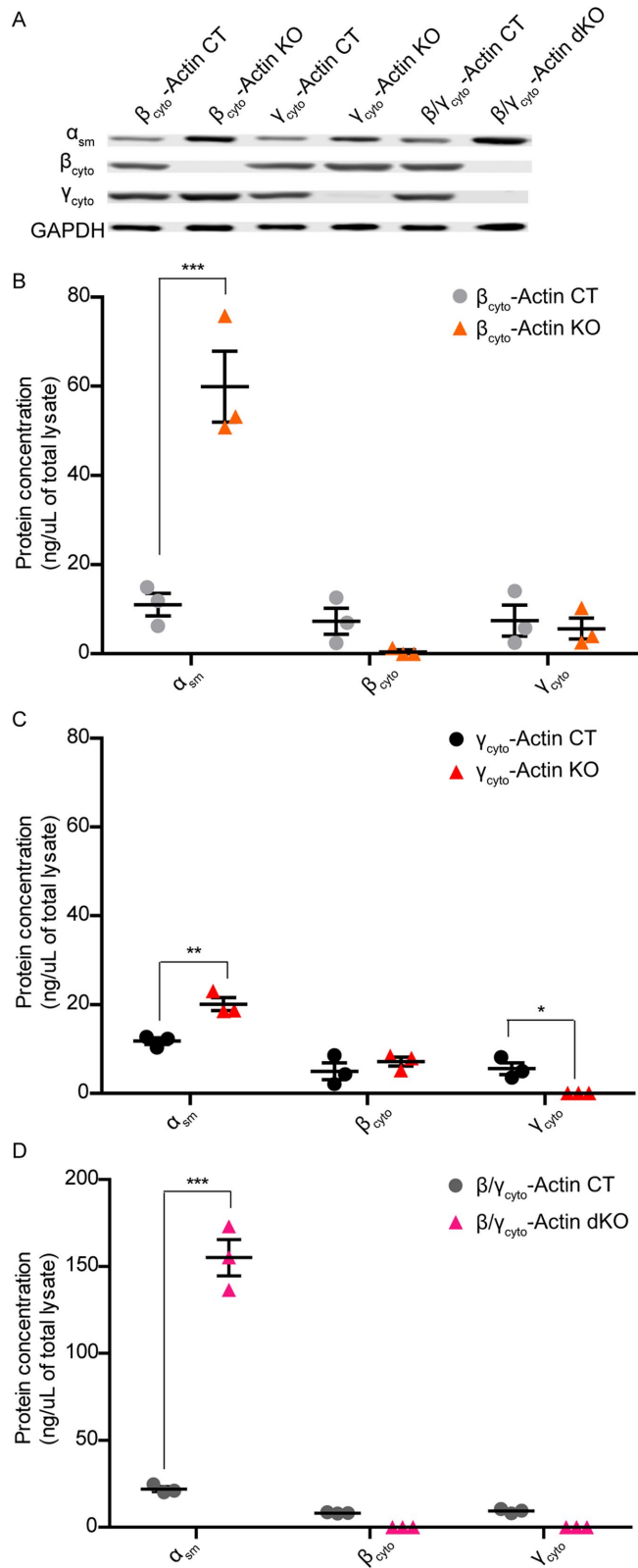


FIGURE 5: α_{sm} -Actin protein was up-regulated in cytoplasmic ablated MEFs. (A) Representative Western blot of CT and KO MEF lysates blotted with α_{sm} -actin, β_{cyto} -actin, and γ_{cyto} -actin antibodies; GAPDH served as loading control. (B–D) Quantitative Western blot analysis of CT and KO MEF lysates ($n = 3$). Calculated protein concentrations (nanograms/microliter lysate) were determined based on the standard curve blotted in parallel. * $p < 0.05$, ** $p < 0.01$, and *** $p < 0.001$. Two-way ANOVA with Bonferroni posttest; error bars are SEM.

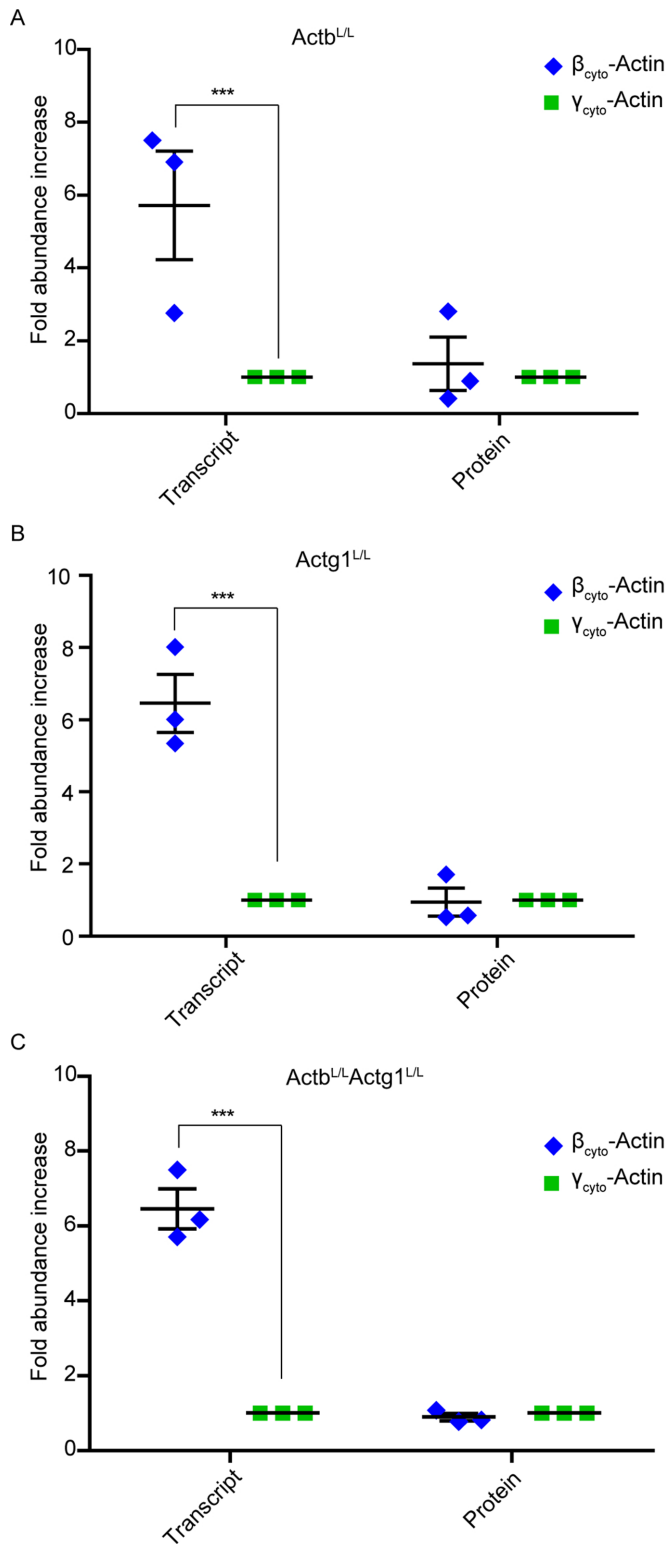


FIGURE 6: Unequal Actb/Actg1 transcript and protein ratios in primary MEFs. (A–C) Calculated transcript and protein ratios between β_{cyto} - and γ_{cyto} -actin in CT MEFs ($n = 3$). Calculations were based on the qRT-PCR and quantitative Western blot data; γ_{cyto} -actin was set at 1. *** $p < 0.001$. One-sample t test; error bars are SEM.

genes in MEFs affects cellular ATP levels independent of corresponding defects in mitochondrial respiration (Gautier *et al.*, 2008; Giaime *et al.*, 2012). We also found that immortalization induced by transformation with SV40 large T antigen rendered Actb nonessential

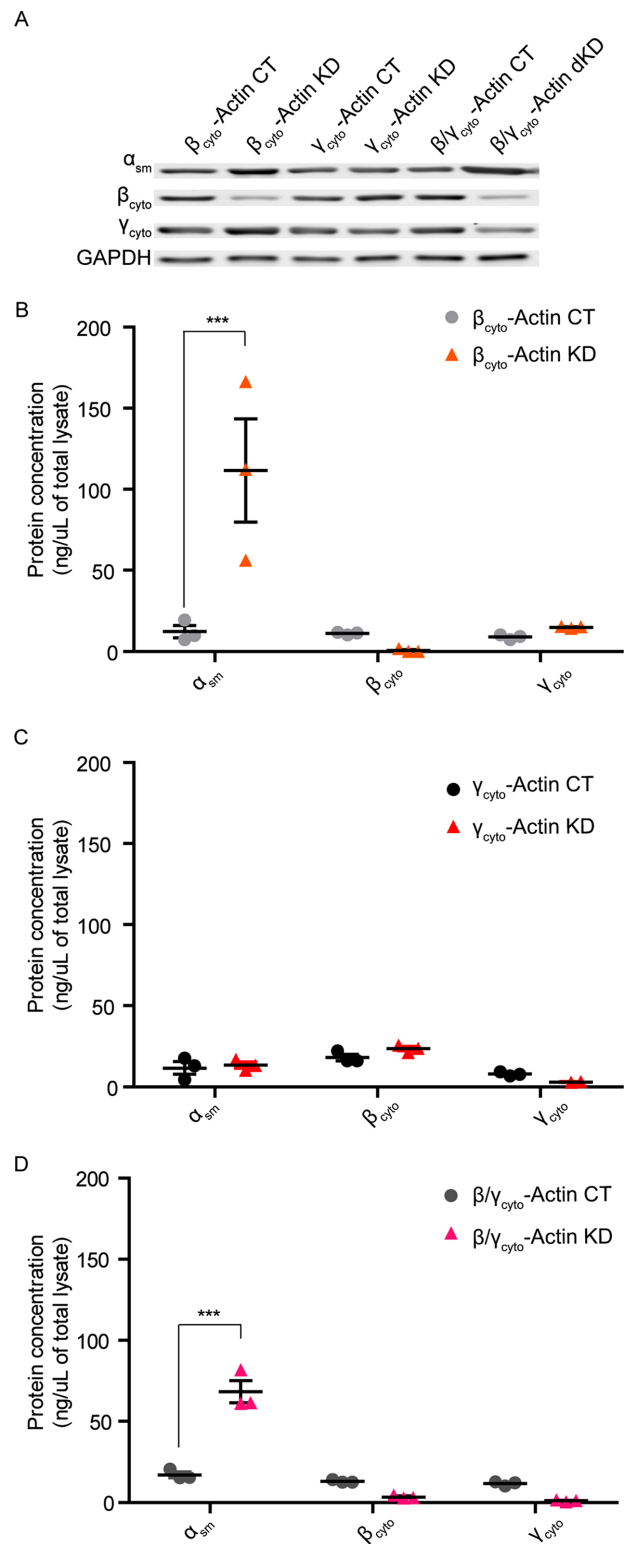


FIGURE 7: α_{sm} -Actin protein was up-regulated in siRNA-mediated β_{cyto} -actin knockdown MEFs. (A) Representative Western blot of CT and β_{cyto} - and/or γ_{cyto} -actin KD MEF lysates at 3 dpi for single KD and 4 dpi for dKD blotted with α_{sm} -actin, β_{cyto} -actin, and γ_{cyto} -actin antibodies; GAPDH served as loading control. (B–D) Quantitative Western blot analysis of control and KD MEF lysates ($n = 3$). Calculated protein concentrations (nanograms/microliter lysate) were determined based on the standard curve, blotted in parallel. *** $p < 0.001$. Two-way ANOVA with Bonferroni posttest; error bars are SEM.

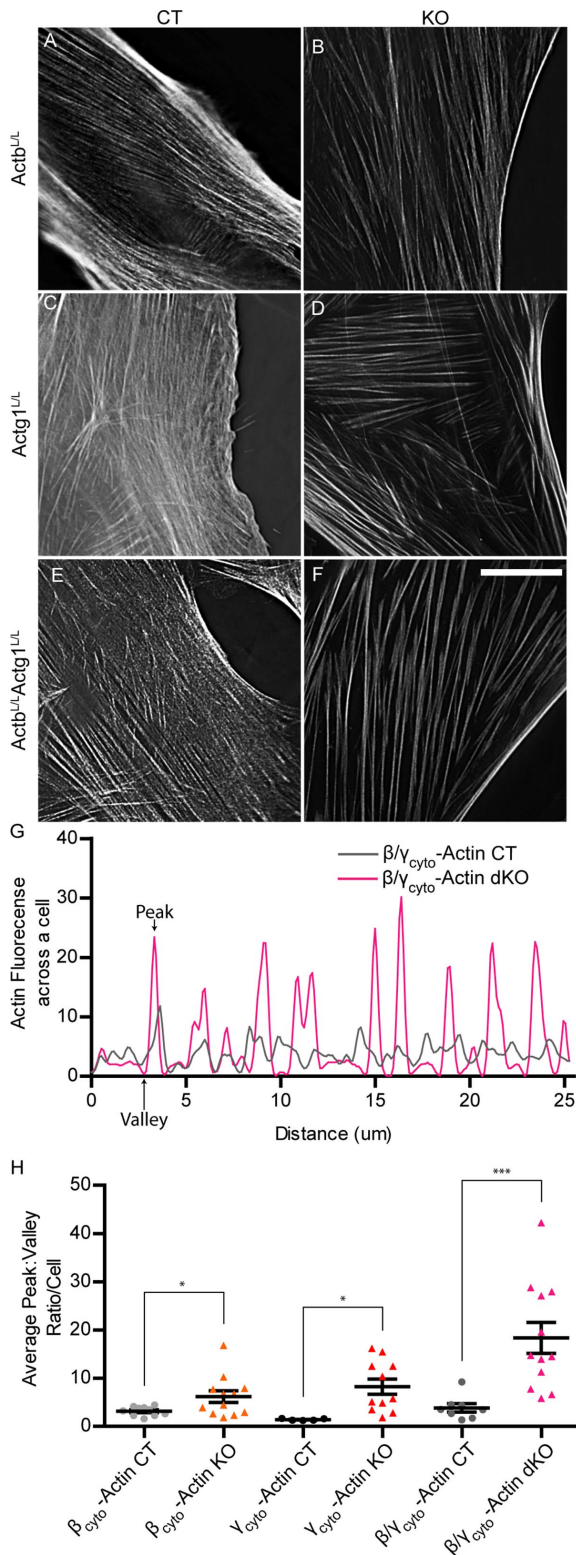


FIGURE 8: The β_{cyto} - and/or γ_{cyto} -actin-ablated MEFs displayed increased stress fiber thickness. (A–F) Representative images of phalloidin-stained actin filaments in Ad5-GFP- or Ad5-Cre-treated primary Actb^{L/L}, Actg1^{L/L}, and Actb^{L/L}Actg1^{L/L} MEFs imaged at 7, 7, and 9 dpi, respectively. Scale bar, 20 μm . (G) Representative line scans of actin fluorescence across the dKO cells in E and F. Arrows denote sample valley and peak. (H) Quantification of differences between valley:peak ratios for all genotypes ($n \geq 8$). * $p < 0.05$, *** $p < 0.001$. Two-way ANOVA; error bars are SEM.

for proliferation (Supplemental Figure S1). Although Tondeleir et al. (2012) noted that cell survival was not impaired in their Actb-null MEFs immortalized by transformation with SV40 large T antigen, a more recent study reported a general delay in cell cycle progression, impaired colony growth, and reduced proliferation rate in the same cell line (Almuzzaini et al., 2016). Although we cannot exclude other effects of transformation with SV40 large T antigen, cell immortalization has been associated with substantial changes in gene expression (Kuo et al., 2012; Gordon et al., 2014). A comparison of primary and spontaneously immortalized MEFs reported almost 2000 transcripts that were up- or down-regulated by threefold (Kuo et al., 2012). Other studies demonstrated that immortalization can significantly affect cell behavior both in the absence (Vasioukhin et al., 2000) and presence of subsequent perturbations (Wade et al., 2002) and even after reversal (Wang et al., 2014). In one highly cited study, primary epithelial cells formed prominent “adhesion zippers” that become challenging to detect after immortalization (Vasioukhin et al., 2000). Moreover, the essential role for p27 in mediating the growth-inhibitory effects of vitamin D in mouse knockout models and primary MEFs was not recapitulated in immortalized MEFs (Wade et al., 2002). Given the unpredictable effects of immortalization and transformation on cell biological function, we suggest that primary MEFs more appropriately model the embryonic lethality associated with Actb gene ablation in mice.

We measured striking differences between the ratios of β_{cyto} - to γ_{cyto} -actin transcript versus protein in primary MEFs: β_{cyto} -actin transcript was sixfold greater than γ_{cyto} -actin transcript, yet β_{cyto} - and γ_{cyto} -actin protein levels were equivalent (Figure 6). β_{cyto} -Actin translation is more rapid than that of γ_{cyto} -actin (Zhang et al., 2010) and also tightly regulated via an RNA regulatory sequence in its 3' untranslated region (UTR) named the “zipcode” (Kislauskis et al., 1993; Ross et al., 1997; Hüttelmaier et al., 2005). Because β_{cyto} -actin protein is the more dynamic of the two cytoplasmic actin isoforms (Bergeron et al., 2010), it is tempting to speculate that the greater abundance of the β_{cyto} -actin transcript is maintained for rapid translation and dynamic cytoskeletal rearrangement. However, the similar levels of β_{cyto} - and γ_{cyto} -actin protein (Figure 6) and similar rates of decay after gene ablation in primary MEFs (Figure 1) suggest that β_{cyto} -actin transcript may play some additional function in cells independent of its role as a template for translation. As one possibility, the large pool of Actb transcript may serve to buffer miRNAs important in regulating the expression of other genes important for cell proliferation, motility, and/or differentiation. Of the >20 different microRNA (miRNA) binding sites collectively predicted by TargetScan, microRNA.Org, and PicTar to be present in the 3' UTR of the Actb transcript but absent from Actg1 (Supplemental Table S1), only mir-205 was identified by all three programs, whereas mir-1 and mir206 were identified by two of three algorithms. Of interest, mir-1, mir-205, and mir-206 were all previously hypothesized to function as tumor suppressors based on experiments demonstrating a correlation between growth inhibition and miRNA overexpression (Hudson et al., 2012; Jalali et al., 2012; Liu et al., 2012; Koshizuka et al., 2017) or increased proliferation after miRNA inhibition (Jalali et al., 2012). Of further relevance to our study, overexpression of mir-1, mir-205, and mir-206 also was associated with significantly decreased cell motility, increased expression of α_{sm} -actin and calponin, and perturbations in actin cytoskeletal dynamics (Hudson et al., 2012; Jalali et al., 2012; Liu et al., 2012). We speculate that ablation of Actb transcript could indirectly drive similar phenotypic changes by freeing up normally sequestered miRNAs. Finally, differential miRNA binding may not be required to explain different phenotypic outcomes for Actb versus

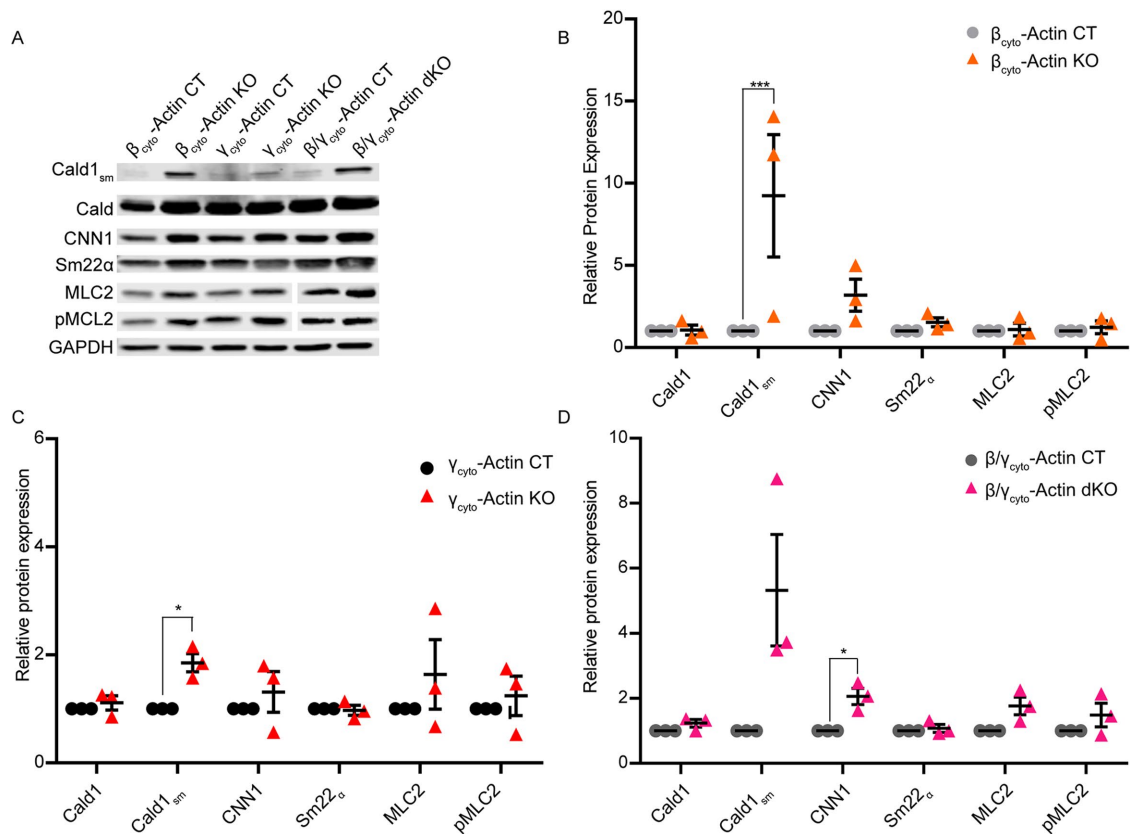


FIGURE 9: Caldesmon smooth muscle isoform protein expression was up-regulated in β_{cyto} -actin-deficient MEFs. (A) Representative Western blot analysis of CT and KO MEFs blotted with caldesmon (Cald1_{sm}), CNN1, Sm22 α , MLC2, and pMLC2; GAPDH served as loading control. (B–D) Relative protein expression normalized to GAPDH and relative to the paired embryo control. * $p < 0.05$, *** $p < 0.001$. One-sample t test, error bars are SEM.

Actg1 transcript ablation. The miRNA mir-145 is predicted to bind both Actb and Actg1 and is important in smooth muscle cell proliferation and differentiation (Cordes *et al.*, 2009), and so it is possible that the more abundant Actb transcript could play the dominant role in sequestering mir-145 during development.

Finally, the differentiation status and/or tissue origin of cells used may also account for some of the differences among studies (Dugina *et al.*, 2009; Bunnell *et al.*, 2011; Shum *et al.*, 2011; Tondeleir *et al.*, 2012; Lechuga *et al.*, 2014). Although Actb and/or β_{cyto} -actin are essential during relatively early stages of embryonic development (Shawlot *et al.*, 1998; Shmerling *et al.*, 2005; Bunnell *et al.*, 2011), neither is specifically required for later-stage development of more differentiated auditory hair cells (Perrin *et al.*, 2010), neurons of the peripheral (Cheever *et al.*, 2011) or central (Cheever *et al.*, 2012) nervous systems, or skeletal muscle (Sonnemann *et al.*, 2006; Prins *et al.*, 2011). These data suggest that cells transition from an Actb-essential state to an Actb-important state as development and differentiation progress *in vivo*. Clearly, the challenge remains to define the *in vivo* biological process modeled by any particular cell in culture.

MATERIALS AND METHODS

Cell culture

Primary MEFs were cultured from E13.5 Actb^{L/L}, Actg1^{L/L}, and double Actb^{L/L}/Actg1^{L/L} mouse embryos as described previously (Bunnell and Ervasti, 2010). Cells were grown to 80% confluency on 10-cm plates and frozen down at passage 1 at 1×10^6 cells/ml in

MEF freezing medium (DMEM supplemented with 10% fetal bovine serum, 1% penicillin/streptomycin, 0.5 $\mu\text{g/ml}$ Fungizone, and 5% dimethyl sulfoxide). Primary MEFs were immortalized by transformation with SV40 large T antigen expression, which was incorporated into the genome via the piggyBac transposase system. MEFs from individual embryos were thawed, cultured in MEF medium (DMEM supplemented with 10% fetal bovine serum, 1% penicillin/streptomycin, 0.5 $\mu\text{g/ml}$ Fungizone), grown to 80% confluency, split into two culture dishes, and treated with Ad5-GFP control virus (Ad5CMV-hrGFP) or Ad5-Cre (Ad5CMV-Cre-eGFP) virus purchased from the University of Iowa Viral Vector Core following their Adenovirus Adfection Protocol (www.medicine.uiowa.edu/vectorcore).

SiRNA transfection

MEFs from individual embryos were thawed, grown to 80% confluency, split into two culture dishes, and treated with control or β_{cyto} - and/or γ_{cyto} -actin ONTARGETplus siRNAs purchased from Dharmacon following their protocol (dharmacon.gelifsciences.com/uploadedFiles/Resources/basic-dharmafect-protocol.pdf). See Supplemental Methods for siRNA sequences.

SRF luciferase activity assay

Actb^{L/L}, Actg1^{L/L}, and double Actb^{L/L}/Actg1^{L/L} MEFs from individual embryos were treated with either Ad5-GFP or Ad5-Cre. Single KO at 3 dpi and dKO at 7 dpi were nucleofected (Amaxa Nucleofector II) with PGL3-Basic-luc (control) or PGL3-FHL2_145-luc (Four and a half LIM domain protein-2) promoter (SRF reporter) constructs

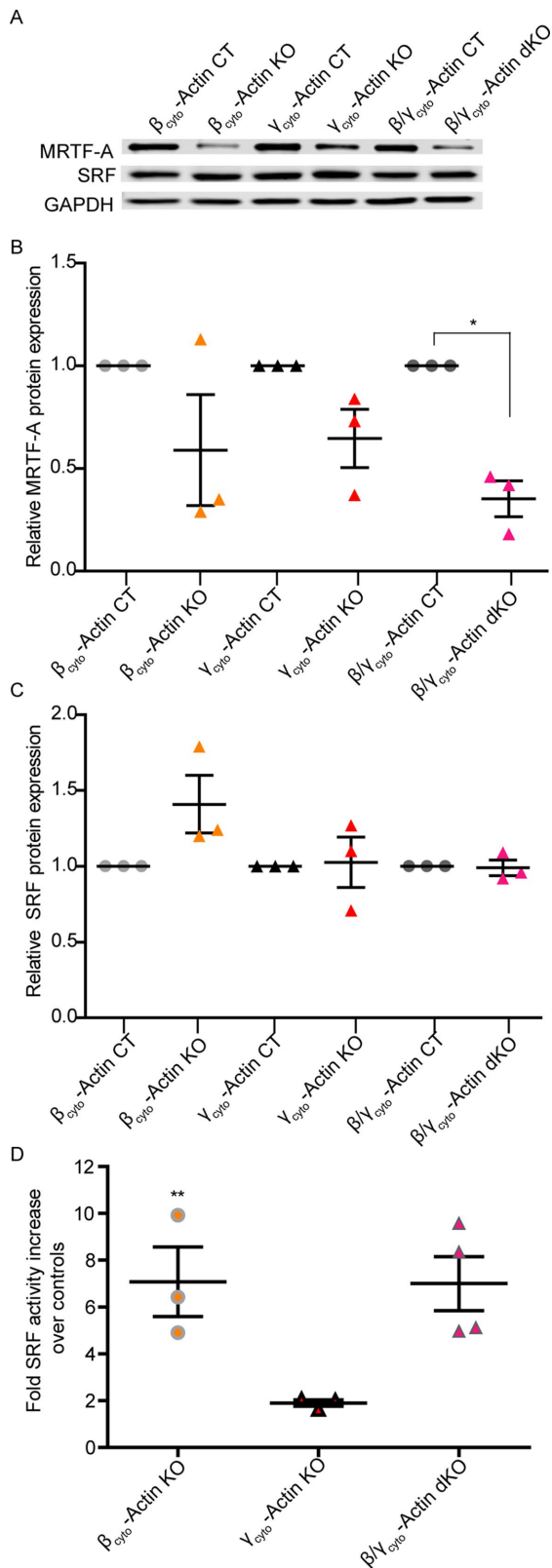


FIGURE 10: SRF activity but not protein was up-regulated in β_{cyto} -actin-ablated MEFs. (A) Representative Western blot analysis of CT and KO MEFs blotted with MRTF-A and SRF; GAPDH served as loading control. (B, C) Relative protein expression normalized to GAPDH and relative to the paired embryo control ($n = 3$). (D) Calculated fold increase in SRF activity, via luciferase assay, in KO over CT MEFs ($n = 3$). * $p < 0.05$, ** $p < 0.01$. One-sample t test; error bars are SEM.

(Schmidt et al., 2012), generously provided by Hannelore V. Heemers (Department of Cancer Biology, Lerner Research Institute). At 2 d postnucleofection, at 5 dpi for single KO and 9 dpi for dKO, cells were lifted and hand counted using a hemacytometer. Equal numbers of cells between Ad5-GFP and Ad5-Cre treated MEFs were washed with phosphate-buffered saline (PBS) and processed based on manufacturer's protocols (Dual-Luciferase Reporter Assay; Promega). Aliquots (20 μl) of cleared processed lysates were used to determine SRF luciferase activity (BioTek Synergy 4 spectrometer with Gen5 2.07 software).

qRT-PCR

Mouse actin isoform cDNAs were amplified from WT MEFs by PCR using the primers listed in the Supplemental Methods and cloned into a pENTR/D-TOPO vector (Life Technologies) to generate a control construct for each actin isoform. Each actin isoform qRT-PCR primer set (Supplemental Methods) was tested for amplification of all control constructs to assess primer specificity (Supplemental Figure S2). Total RNA was extracted from Actb^{L/L}, Actg1^{L/L}, and double Actb^{L/L}/Actg1^{L/L} MEF samples using the Bio-Rad Aurum Total RNA Mini Kit following the manufacturer's instructions. RNA concentration and purity (260/280 ratio) were determined using a NanoDrop spectrophotometer (Wilmington, DE). First-strand cDNA was synthesized with a Bio-Rad iScript Advanced cDNA Synthesis Kit for qRT-PCR using the same initial RNA amount (1 μg) for all samples. Individual control constructs were used in a 10-fold dilution to generate a standard curve, and MEF samples were amplified in parallel with each specific qRT-PCR primer set using Bio-Rad SsoAdvanced Universal SYBR polymerase on the Bio-Rad CFX96 Real Time System C1000 Touch Thermal Cycler to profile each actin isoform transcript amount (picomoles).

Actin isoforms

Platelet non-muscle actin (β_{cyto} - and γ_{cyto} -actin) was purchased from Cytoskeleton (Denver, CO). Recombinant γ_{cyto} - and α_{sm} -actin were expressed in the Bacto-Bac insect cell-expression system and purified as previously described (Perrin et al., 2013). Protein concentrations were determined using a NanoDrop spectrophotometer. Contamination of recombinant γ_{cyto} - and α_{sm} -actins by insect cell actin was measured by electrospray ionization (ESI) mass spectrometry acquired using a QSTAR Pulsar i (Applied Biosystems, Foster City, CA) quadrupole time-of-flight (TOF) mass spectrometer equipped with a turbo-ESI source. Protein samples were loaded directly in 50% acetonitrile and 0.1% formic acid. The ion spray voltage was 1000 V, the TOF region acceleration voltage was 4 kV, and the injection pulse repetition rate was 7.0 kHz. External calibration was performed using renin (monoisotopic mass [MH3⁺] 586.9830 and [MH2⁺] 879.9705; Sigma-Aldrich, St. Louis, MO). Mass spectra were summed and averaged over the injection peak. Deconvolution of the charge envelope was performed with the Bioanalyst Analyst-QS software package (ABI).

Western blotting

For Western blot quantitation of actin isoforms against a standard curve, total MEF protein was extracted with 1% SDS buffer in 1x PBS and a cocktail of protease inhibitors (100 μM aprotinin, 0.79 mg/ml benzamide, 10 nM E-64, 10 μM leupeptin, 0.1 mg/ml pepstatin, 1 mM phenylmethylsulfonyl fluoride), sonicated (Model 150V/T Ultrasonic homogenizer; BioLogics), boiled, and centrifuged to remove the insoluble fraction. An increasing amount (25–300 ng) of purified actin protein and all MEF samples were run on the same blot to ensure consistency (Supplemental Figure S6). LiCor fluorescence

signal versus purified actin protein amount (nanograms) was used to generate a standard curve in order to calculate specific actin isoform concentration (nanograms of protein/microliter of total lysate). All samples were western blotted with β -actin (AC-15; Sigma-Aldrich), γ -actin (mAb 117), and α_{sm} -actin (A14; Sigma-Aldrich). For comparisons of relative immunoreactivities, equal amounts of cleared total MEF lysate protein (25 μ g) were blotted with antibodies to caldesmon 1 (2980; Cell signaling), SM22 (ab14106; Abcam), calponin (ab46794; Abcam), SRF (G-20; Santa Cruz Biotechnology), MRTF-A (H-140; Santa Cruz Biotechnology), MTOC1 (ab14705; Abcam), SDHA (ab14715; Abcam), UQCR (ab110252; Abcam), ATP5A (ab14748; Abcam), MLC2 (ab48003; Abcam), and MLC2 (phosphor S20; ab2480; Abcam), with glyceraldehyde 3-phosphate dehydrogenase (GAPDH; G9545 or G8795; Sigma-Aldrich) as loading control.

Cell fixation and staining

MEFs cultured for 6–8 dpi with either Ad5-GFP or Ad5-CRE were plated on 5 μ g/ml fibronectin-coated coverslips at a density of 1×10^4 cells/coverslip and incubated in MEF culturing media. The next day, cells were fixed with fresh 4% paraformaldehyde in PBS for 15 min at room temperature. To stain actin filaments, fixed cells were treated with 0.2% Triton X-100 and 16.5 nM phalloidin (A22285; Life Technologies) in PBS for 30 min at room temperature. Coverslips were mounted in Prolong (P36931; Life Technologies) mounting medium.

Cell imaging and quantification

Samples were imaged using a DeltaVision PersonalDV microscope under a 60 \times /1.42 numerical aperture objective (GE Technologies). All cells were imaged with the same laser intensity and exposure time. The intensity of the stress fibers was quantified from line scans across the widest part of the cell using ImageJ. The peaks and corresponding valleys were then determined (excluding the first peak, which represents the edge of the cell).

The average peak-to-valley ratio per cell was then plotted using GraphPad Prism.

ATP quantification

At 7 or 9 dpi, cells were lifted and counted. Equal numbers of CT and KO MEFs were lysed using boiling water (Yang *et al.*, 2002). The lysate was spun at 12,000 $\times g$ for 5 min at 4°C, and the supernatant was collected. Following the luciferase-based ATP Determination Kit (A22066) protocol, the reaction solution (reaction buffer, dithiothreitol (DTT), D-luciferin, and firefly luciferase) was added to the cell lysate supernatant. After a 15-min incubation, ATP levels were determined using a BioTek Synergy 4 spectrometer with Gen5 2.07 software and quantified using an ATP standard ranging from 0 nM to 1 μ M.

Analysis of cell growth

A 5×10^5 amount of MEFs/well was treated with Ad5-GFP or Ad5-Cre. Samples were counted every other day beginning at 1 dpi in duplicate, using a hemacytometer to generate growth curves.

Oxygen consumption

Cells were plated in triplicate on 5 μ g/ μ l fibronectin-coated XF24 cell culture microplates (Seahorse Bioscience) at 5×10^4 cells/well 24 h before the experiment. Mitochondrial function was examined using the Seahorse XF24 extracellular flux analyzer and the cell mitochondrial stress assay (Seahorse Bioscience). Oligomycin (2 μ g/ml final), FCCP (carbonyl cyanide 4-(trifluoromethoxy)phenylhydrazone;

0.5 μ M final), and antimycin A (4 μ M final) were injected at time points throughout the experiment according to the company protocol (all compounds from Seahorse Bioscience). Oxygen consumption was measured at multiple points between each injection and normalized to background controls.

Statistics

For quantitative RT-PCR and quantitative Western blots, two-way analysis of variance (ANOVA) with a Bonferroni posttest was used on calculated amounts of transcript (picomoles) and protein (nanograms/microliter of lysate) to determine statistics. For all relative Western blots, immunoreactivity levels for all samples were normalized to GAPDH and relative to the paired embryo control, which was set at 1, and a one-sample *t* test was performed to evaluate statistics.

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