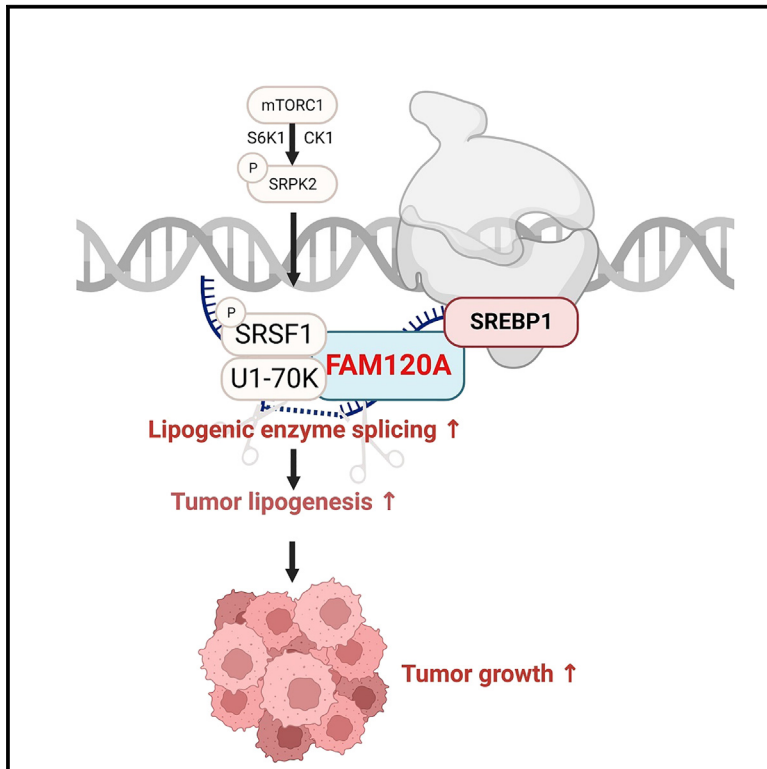


FAM120A couples SREBP-dependent transcription and splicing of lipogenesis enzymes downstream of mTORC1

Graphical abstract



Authors

Sungyun Cho, Yujin Chun, Long He, ..., Cholsoon Jang, John Blenis, Gina Lee

Correspondence

job2064@med.cornell.edu (J.B.),
ginalee@uci.edu (G.L.)

In brief

Cho et al. report FAM120A as a transcription co-activator that connects transcription and splicing of *de novo* lipid synthesis enzymes downstream of mTORC1. Upon mTORC1 activation, SRPK2 phosphorylates splicing factor SRSF1, inducing its binding with FAM120A. Through its interaction with SRSF1, the SREBP1-associated FAM120A promotes efficient splicing of lipogenic genes and facilitates lipid metabolism of cancer cells. These findings reveal FAM120A as a critical downstream of mTORC1 in anabolic gene expression program and cell growth.

Highlights

- FAM120A couples transcription and splicing of lipogenic enzymes downstream of mTORC1
- mTORC1-activated SRPK2 phosphorylates SRSF1 to induce FAM120-SRSF1 interaction
- FAM120A directly interact with SREBP to bring lipogenic genes to RNA-splicing machinery
- mTORC1-driven RNA-binding protein complex promotes lipogenic gene splicing and stability



Article

FAM120A couples SREBP-dependent transcription and splicing of lipogenesis enzymes downstream of mTORC1

Sungyun Cho,¹ Yujin Chun,² Long He,¹ Cuauhtemoc B. Ramirez,^{2,3} Kripa S. Ganesh,⁴ Kyungjo Jeong,⁵ Junho Song,⁵ Jin Gyu Cheong,⁶ Zhongchi Li,¹ Jungmin Choi,^{5,7} Joochwan Kim,² Nikos Koundouros,^{1,8} Fangyuan Ding,^{9,10} Noah Dephoure,⁸ Cholsoon Jang,^{3,10,11} John Blenis,^{1,8,12,*} and Gina Lee^{2,10,11,*}

¹Department of Pharmacology, Weill Cornell Medicine, Cornell University, New York, NY, USA

²Department of Microbiology and Molecular Genetics, School of Medicine, University of California Irvine, Irvine, CA, USA

³Department of Biological Chemistry, School of Medicine, University of California Irvine, Irvine, CA, USA

⁴Department of Biochemistry, Weill Cornell Medicine, Cornell University, New York, NY, USA

⁵Department of Biomedical Sciences, College of Medicine, Korea University, Seoul, South Korea

⁶Department of Pathology and Laboratory Medicine, Weill Cornell Medicine, Cornell University, New York, NY, USA

⁷Department of Genetics, Yale School of Medicine, Yale University, New Haven, CT, USA

⁸Meyer Cancer Center, Weill Cornell Medicine, Cornell University, New York, NY, USA

⁹Department of Biomedical Engineering, Department of Developmental and Cell Biology, Department of Pharmaceutical Sciences, Center for Synthetic Biology, and Center for Neural Circuit Mapping, University of California Irvine, Irvine, CA, USA

¹⁰Center for Complex Biological Systems and Chao Family Comprehensive Cancer Center, University of California Irvine, Irvine, CA, USA

¹¹Center for Epigenetics and Metabolism, University of California Irvine, Irvine, CA, USA

¹²Lead contact

*Correspondence: job2064@med.cornell.edu (J.B.), ginalee@uci.edu (G.L.)

<https://doi.org/10.1016/j.molcel.2023.07.017>

SUMMARY

The mechanistic target of rapamycin complex 1 (mTORC1) is a master regulator of cell growth that stimulates macromolecule synthesis through transcription, RNA processing, and post-translational modification of metabolic enzymes. However, the mechanisms of how mTORC1 orchestrates multiple steps of gene expression programs remain unclear. Here, we identify family with sequence similarity 120A (FAM120A) as a transcription co-activator that couples transcription and splicing of *de novo* lipid synthesis enzymes downstream of mTORC1-serine/arginine-rich protein kinase 2 (SRPK2) signaling. The mTORC1-activated SRPK2 phosphorylates splicing factor serine/arginine-rich splicing factor 1 (SRSF1), enhancing its binding to FAM120A. FAM120A directly interacts with a lipogenic transcription factor SREBP1 at active promoters, thereby bridging the newly transcribed lipogenic genes from RNA polymerase II to the SRSF1 and U1-70K-containing RNA-splicing machinery. This mTORC1-regulated, multi-protein complex promotes efficient splicing and stability of lipogenic transcripts, resulting in fatty acid synthesis and cancer cell proliferation. These results elucidate FAM120A as a critical transcription co-factor that connects mTORC1-dependent gene regulation programs for anabolic cell growth.

INTRODUCTION

Cancer cells rewire metabolic pathways to support increased demands for continuous proliferation.^{1,2} Mechanistic target of rapamycin complex 1 (mTORC1) signaling, which is activated in almost all human cancers, is one of the key mechanisms responsible for such metabolic rewiring. In normal cells, mTORC1 works as a molecular rheostat to promote macromolecule synthesis upon environmental cues, such as growth factors and nutrients. It does so by activating metabolic enzymes directly or through the regulation of metabolic gene expression.^{3,4} For instance, upon insulin stimulation, mTORC1 induces

de novo lipid synthesis by upregulating the expression and activity of sterol regulatory element-binding proteins (SREBPs), a master transcription factor of lipogenic enzymes.^{5–8} In cancer cells, however, overactive mTORC1 signaling constitutively promotes SREBP-dependent lipogenic enzyme expression for tumor cell growth and proliferation.^{9,10}

In addition to regulating transcriptional programs, mTORC1 also activates metabolic enzymes through post-transcriptional RNA processing.^{11–14} Previously, we found that mTORC1-regulated S6 protein kinase 1 (S6K1) phosphorylates serine/arginine-rich protein kinase 2 (SRPK2) to stimulate its nuclear translocation and phosphorylation of downstream serine/arginine-rich



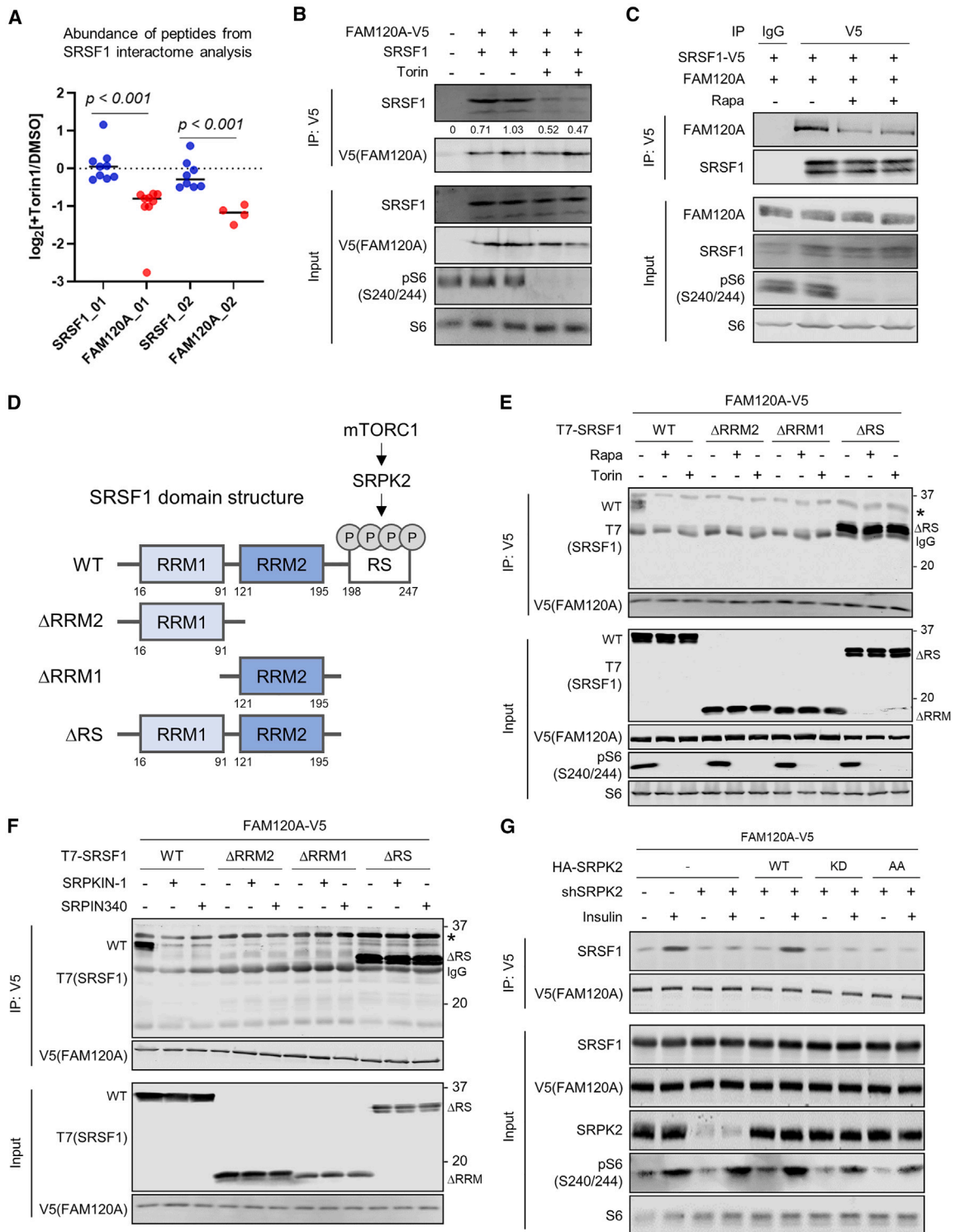


Figure 1. mTORC1-SRPK promotes interaction of SRSF1 with FAM120A

(A) Mass spectrometry analysis of peptides identified in the SRSF1 interactome. HEK293E cells expressing SRSF1-V5 were treated with DMSO or torin1 (250 nM) for 4 h, and SRSF1-binding proteins were co-immunoprecipitated (coIP) by anti-V5 antibody. Each dot represents abundance of individual peptides ($n = 4-9$ for each protein) in $\log_2[\text{Torin1}/\text{DMSO}]$. Data are reanalyzed from a previous publication.¹²
(B and C) CoIP analysis of HEK293E cells expressing FAM120A-V5 and SRSF1 (B) or FAM120A and SRSF1-V5 (C). Cells were treated with torin1 (250 nM) or rapamycin (100 nM) for 4 h. 1% of total cell lysate was loaded as an input.

(legend continued on next page)

splicing factor 1 (SRSF1).¹² Such phosphorylation facilitates SRSF1's interaction with U1 small nuclear ribonucleoprotein (snRNP) 70 kDa (U1-70K), a core spliceosome protein that initiates assembly of the spliceosome complex on target pre-mRNAs.^{15–17} Serine/arginine-rich (SR) proteins are additional components of the spliceosome complex that potentiate co-transcriptional splicing.^{18–21} For example, SRSF2 directly interacts with RNA polymerase II (RNA Pol II) and peroxisome proliferator-activated receptor (PPAR) gamma co-activator 1- α (PGC1 α) for efficient splicing of mitochondrial biogenesis genes.²² Interestingly, the mTORC1-SRPK2-dependent activation of SRSF1 induces lipogenic gene splicing, but the underlying mechanism for this specificity is unclear.

In this study, we identified a family with sequence similarity 120A (FAM120A) as a transcription co-factor and RNA-binding protein (RBP) that determines the specificity of genes spliced downstream of mTORC1-SRPK2 signaling. Upon mTORC1 activation, the SRPK2-phosphorylated SRSF1 binds with FAM120A and U1-70K, forming a multi-protein complex with SREBP1, a lipogenic gene transcription factor, and RNA Pol II to mediate selective splicing of lipogenic genes. Our findings provide mechanistic insights into how oncogenic mTORC1 signaling coordinates the interactions among transcription factors, RBPs, and target gene transcripts to promote cancer cell growth by connecting transcription and pre-mRNA splicing to the translation of metabolic enzymes.

RESULTS

mTORC1-SRPK2 signaling promotes SRSF1 interaction with FAM120A

To investigate how mTORC1-SRPK2 signaling regulates lipogenic mRNA splicing via SRSF1, we previously performed a proteomics-based SRSF1 interactome analysis with or without torin1 (mTOR catalytic inhibitor). Most SRSF1-bound proteins were well-known RNA-processing proteins, including a spliceosome protein, U1-70K, which we characterized as a key mTORC1-dependent SRSF1 interactor in our previous study.¹² Additionally, we found FAM120A, whose peptide abundance in the SRSF1-bound fraction was decreased by torin1 (Figure 1A). FAM120A is an RBP that localizes in the nucleus and associates with RNA Pol II and spliceosome proteins.^{23–26} Co-immunoprecipitation (coIP) analysis confirmed FAM120A binding with SRSF1, which was suppressed by torin1 (Figure 1B). Rapamycin, an mTORC1-specific inhibitor, also blunted their association (Figure 1C).

SRSF1 contains two RNA recognition motifs (RRM1 and RRM2) and a C-terminal arginine/serine (RS)-rich domain (Fig-

ure 1D).²⁷ CoIP experiments revealed that SRSF1 mutants lacking either of the two RRM domains (SRSF1- Δ RRM1 or SRSF1- Δ RRM2) do not interact with FAM120A, suggesting that RNA binding is essential for the association of these two proteins (Figures 1E and S1A). In contrast, deletion of the RS domain of SRSF1 (SRSF1- Δ RS) dramatically enhanced the binding between SRSF1 and FAM120A that was insensitive to rapamycin or torin1, indicating an inhibitory function of RS domain in FAM120A-SRSF1 interaction (Figures 1E and S1A). The RS domain of SRSF1 has been shown to form a folded structure with the RRM1/2 domains,^{15,28} and phosphorylation of the RS domain by SRPK opens this folded structure and promotes the binding of SRSF1 with other proteins.^{12,15,29,30} Indeed, suppression of SRSF1 phosphorylation by inhibitors targeting mTORC1 (rapamycin and torin1) or SRPK (SRPKIN-1 and SRPIN340)^{31,32} decreased interaction of full-length SRSF1 (SRSF1-wild type [WT]) with FAM120A, whereas the interaction of RS-domain-lacking mutant (SRSF1- Δ RS) with FAM120A was not affected by these inhibitors (Figures 1E, 1F, S1A, and S1B). Furthermore, insulin-induced SRSF1 phosphorylation and SRSF1-FAM120A association was suppressed by knockdown of endogenous *SRPK2*, which was restored by re-expression of WT SRPK2 but not by kinase dead (KD, SRPK2-K110M) or mTORC1-dependent phosphorylation-site-lacking (AA, SRPK2-S494A/S497A) mutants¹² (Figures 1G and S1C). These results demonstrate that, when mTORC1 is activated, SRPK2 facilitates the association of FAM120A with SRSF1's RRM domain by phosphorylating its RS domain (Figure 1D).

FAM120A promotes lipogenic enzyme expression

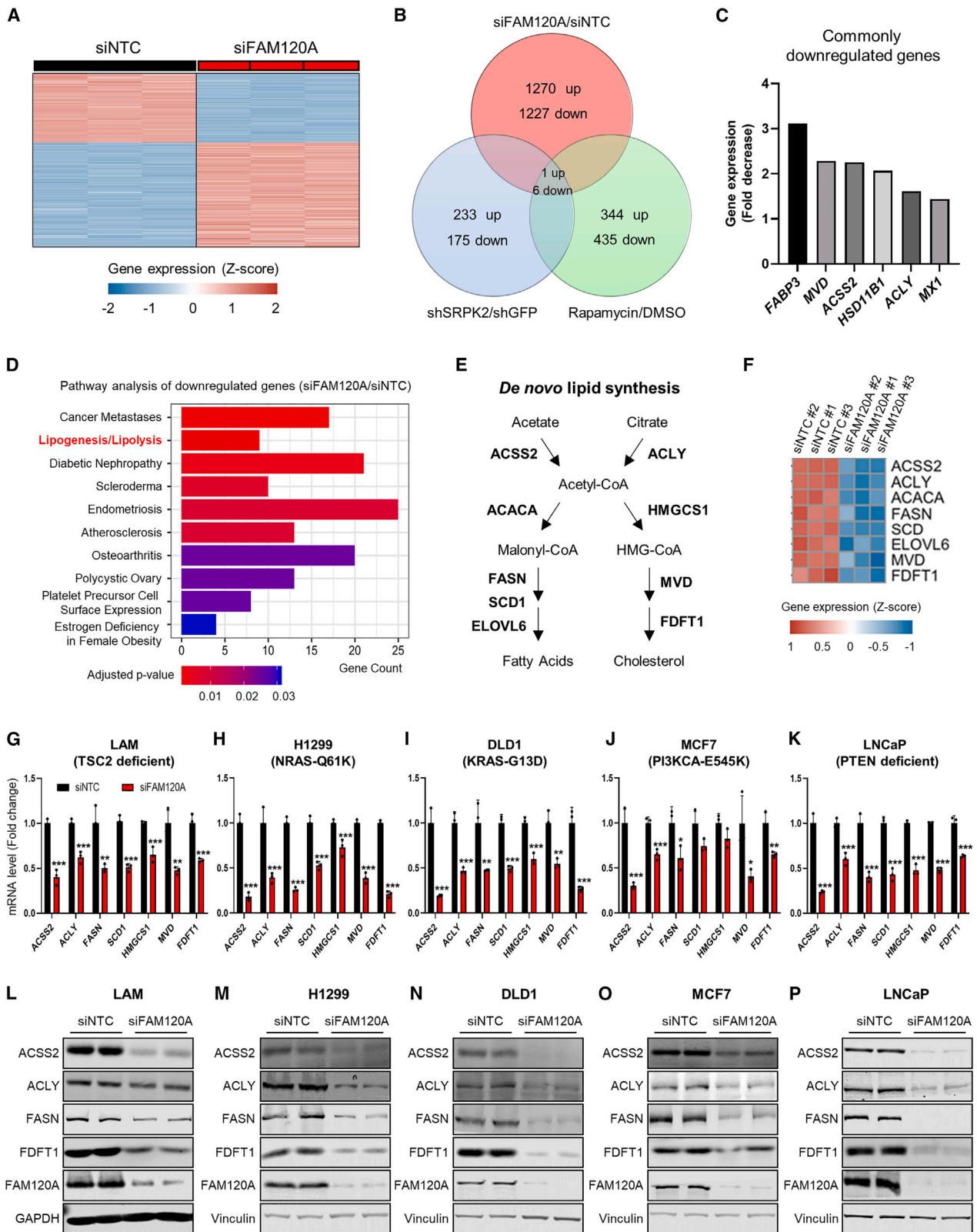
Given the interaction of FAM120A with RNA Pol II²⁴ and mTORC1-SRPK2-dependent interaction of FAM120A with SRSF1 (Figure 1), we reasoned that FAM120A would be involved in the regulation of gene expression downstream of mTORC1-SRPK2. We performed RNA sequencing (RNA-seq) analysis in a kidney-angiomyolipoma-patient-derived cell line (LAM 621-101 or LAM cells where mTORC1 is activated due to *TSC2* deficiency)³³ after transfection with small interfering RNAs (siRNAs) targeting control or *FAM120A* (Figure 2A). To identify the genes downstream of mTORC1-SRPK2-dependent regulation of FAM120A, we compared *FAM120A* knockdown RNA-seq results with our previous whole-transcriptome microarray data from LAM cells treated with rapamycin or *SRPK2* knockdown (GEO: GSE104335).¹² Among the 6 genes commonly down-regulated in these three conditions, 5 were lipid metabolism regulators, including *ATP citrate lyase (ACLY)*, *acyl-coenzyme A (CoA) synthetase short chain family member 2 (ACSS2)*, *fatty-acid-binding protein 3 (FABP3)*, *hydroxysteroid 11- β*

(D) Schematic diagram describing mTORC1-SRPK2-dependent phosphorylation of SRSF1 and the domains of wild-type (WT) and mutant SRSF1 constructs. RRM: RNA recognition motif. RS, arginine/serine-rich domain.

(E and F) CoIP analysis of HEK293E cells expressing FAM120A-V5 and T7-SRSF1 (WT or mutants). Cells were treated with torin1 (250 nM) (E) or SRPK inhibitors (SRPKIN-1, 5 μ M; SRPIN340, 30 μ M) (F) for 4 h. 1% of total cell lysate was loaded as an input. IgG light chain (IgG, 25 kDa) and non-specific bands (asterisks) are indicated.

(G) CoIP analysis of HEK293E cells expressing FAM120A-V5 and HA-SRPK2 (WT [wild type], KD [kinase dead, SRPK2-K110M], or AA [non-phosphorylatable, SRPK2-S394A/S397A]). Endogenous *SRPK2* was knocked down by short hairpin RNA (shRNA) targeting 3' UTR of *SRPK2*. Cells were serum starved overnight followed by insulin (100 nM) stimulation for 4 h. 1% of total cell lysate was loaded as an input.

See also Figure S1.



(legend on next page)

dehydrogenase 1 (HSD11B1), and *mevalonate diphosphate decarboxylase (MVD)* (Figures 2B and 2C). Pathway enrichment analysis of the genes downregulated by siFAM120A also ranked lipogenesis as one of the top FAM120A downstream biological processes (Figure 2D). Because *de novo* fatty acid and cholesterol synthesis enzymes were the major target of mTORC1-SRPK2-SRSF1 signaling,¹² we further looked into the level of individual genes in FAM120A RNA-seq and found that FAM120A depletion significantly decreased the expression of enzymes in *de novo* lipid synthesis pathway (Figures 2E and 2F).

To examine whether FAM120A-dependent lipogenic gene regulation is relevant in other cancers, we knocked down FAM120A in a panel of mTORC1-overactive cancer cell lines derived from diverse tissues of origin and different genetic drivers such as H1299 (NRAS-mutated non-small-cell lung cancer),³⁴ DLD1 (KRAS-mutated colorectal adenocarcinoma),³⁵ MCF7 (phosphatidylinositol 3-kinase [PI3K]-mutated breast cancer),³⁶ and LNCaP (PTEN-deficient prostate cancer).³⁷ Across all these cells, FAM120A depletion decreased lipogenic enzymes at both mRNA and protein levels (Figures 2G–2P and S1D–S1H). In contrast, expression of another metabolic enzyme known to be regulated by mTORC1 for nucleotide synthesis, *ribose 5-phosphate isomerase A (RPIA)*,^{10,38} was not regulated by FAM120A (Figures S1D–S1H), indicating a specific role of FAM120A in regulating lipogenic enzyme expression.

FAM120A promotes RNA splicing and stability of lipogenic enzymes

Next, we sought to elucidate the underlying mechanisms for FAM120A-dependent lipogenic gene regulation. Because FAM120A binds to phosphorylated SRSF1 (Figure 1) and exists in the spliceosome complex,^{23–26} we speculated that FAM120A may regulate lipogenic gene expression via RNA splicing. To assess splicing changes, we performed deep RNA-seq of LAM cells knocked down with FAM120A or SRSF1. Differentially expressed gene analysis identified a total of 2,895 genes whose expression is significantly altered in siFAM120A or siSRSF1 cells compared with control (siNTC) (cutoff: $|\log_2(\text{fold change or FC})| \geq 0.5$ and $q \text{ value} < 0.01$). Among these, 1,585 genes were downregulated by both FAM120A and SRSF1 knockdown (Figures 3A and S2A; Tables S1A and S1B). Pathway

enrichment analysis revealed that lipid-synthesis-related pathways, including cholesterol biosynthesis ($q \text{ value} = 6.45\text{E}-07$), activation of gene expression by SREBP ($q \text{ value} = 1.07\text{E}-06$), and oleate biosynthesis ($q \text{ value} = 0.00822$), were among the top biological processes of the genes commonly regulated by FAM120A and SRSF1 (Figure S2B; Table S1C).

For comprehensive evaluation of splicing events regulated by FAM120A and SRSF1, we analyzed the RNA-seq data using vertebrate alternative splicing and transcription tools (VAST-TOOLS).³⁹ We identified 11,237 differential splicing events by FAM120A knockdown and 5,844 by SRSF1 knockdown, which includes intron retention (IR), alternative 5' splicing site, alternative 3' splicing site, and micro-exon splicing. Among these, IR was the most prevalent splicing events by comprising 85% of the differential splicing events in siFAM120A and 73% in siSRSF1 (Figure S2C). To quantitate IR events more precisely, we calculated IR ratio (IR ratio = intronic abundance/[exonic + intronic abundance]) using IRFinder-S software,⁴⁰ which demonstrated a significant increase of IR by FAM120A or SRSF1 knockdown (Figure 3B). From the cutoff of $q \text{ value} < 0.1$ and $\log_2\text{FC} > 0$, we selected 9,589 (siFAM120A/siNTC) and 2,849 (siSRSF1/siNTC) displaying significant IR events. Finally, by comparing the 1,893 IR events identified in both siFAM120A and siSRSF1 conditions and 1,585 commonly downregulated genes, we found 408 IR events in 186 genes (Figures S2D and S2E; Table S1D). Pathway enrichment analysis of these final candidate genes revealed that IR is enriched in lipid metabolism genes, such as cholesterol metabolism ($q \text{ value} = 0.01145$), omega-9 fatty acid synthesis ($q \text{ value} = 0.01541$), and metabolism of lipids ($q \text{ value} = 0.03637$), suggesting that FAM120A and SRSF1 control expression of lipid metabolism genes by promoting splicing of their introns (Figure S2F; Table S1E).

From in-depth analysis of RNA-seq results on lipogenic genes, we found significantly increased IR by FAM120A or SRSF1 knockdown in FASN, ACSS2, MVD, and HMGCS1 (Figures 3C, 3D, and S2E; Table S1D). We validated these IR events by qPCR using the primers specifically targeting the introns identified from RNA-seq (Figure 3E). Failed splicing and IR lead to degradation of transcripts via nonsense-mediated decay (NMD), which we previously found as the mechanism for mTORC1-SRPK2-SRSF1-dependent regulation of lipogenic

Figure 2. FAM120A is required for the expression of lipogenic enzymes

(A) Heatmap of Z scores for differentially expressed gene levels (cutoff: $|\log_2(\text{fold change or FC})| \geq 1$, $q \text{ value} < 0.005$) in LAM 621-101 cells (hereinafter referred to as LAM cells) transfected with siRNAs targeting FAM120A or non-targeting control (NTC, hereinafter referred to as control) (GEO: GSE207172). $n = 3$. Same RNA-seq results are used in (A)–(F).

(B) Venn diagram analysis of the differentially regulated genes (cutoff: linear fold change ≥ 1.5) identified from the whole-transcriptome microarray and RNA-seq analyses in LAM cells. The gene expression analyses were conducted on cells treated with rapamycin (20 nM) or vehicle for 24 h, on cells stably expressing shRNAs targeting SRPK2 or GFP (GEO: GSE104335),¹² or on cells transfected with siRNAs targeting FAM120A or control (GEO: GSE207172).

(C) Fold decrease of 6 commonly downregulated genes from the Venn diagram analysis in (B). Fold decrease values from siNTC vs. siFAM120A RNA-seq analysis are shown.

(D) Pathway analysis of downregulated genes by FAM120A knockdown.

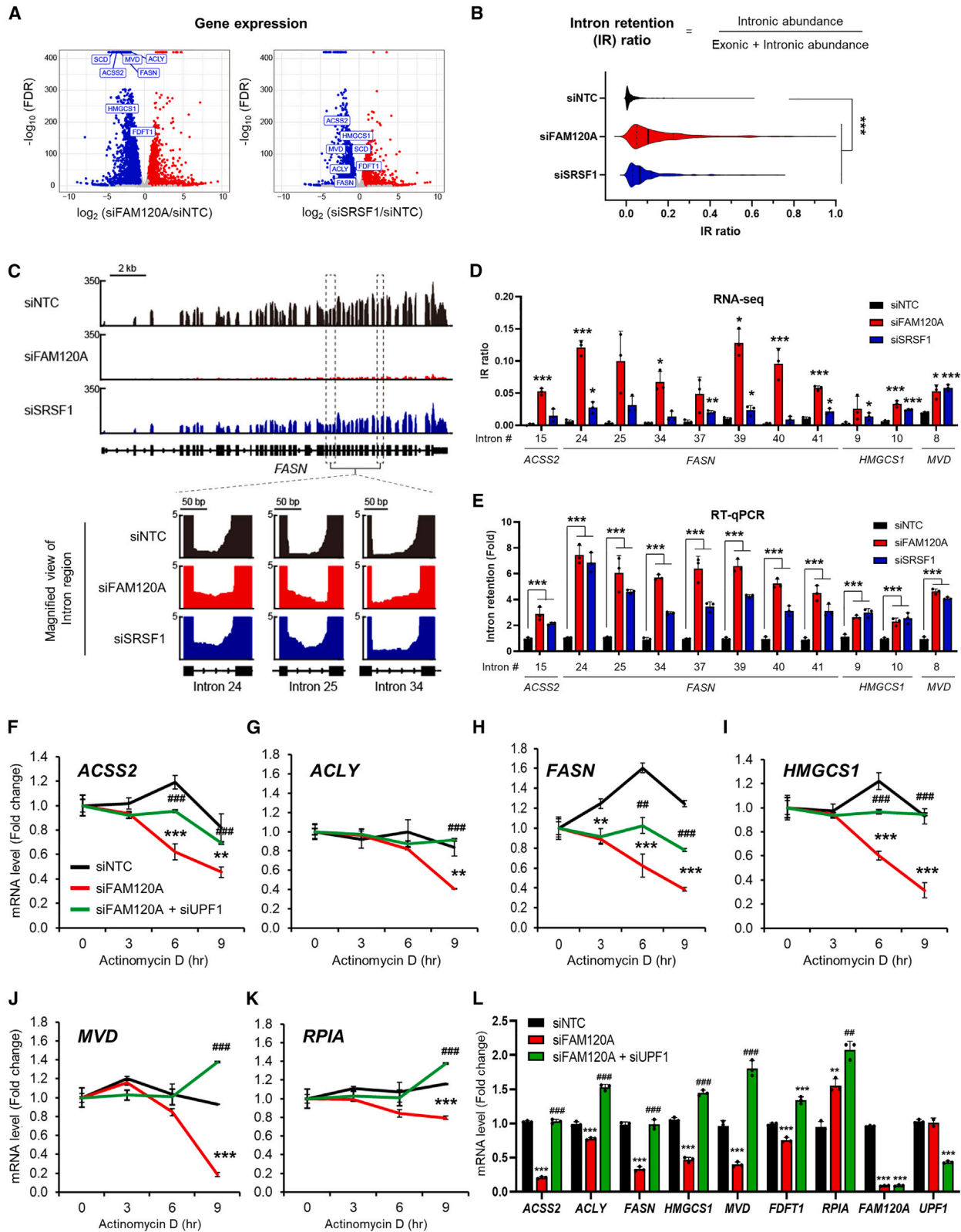
(E) Schematic of *de novo* lipogenesis.

(F) Heatmap of Z scores for the lipogenic gene levels decreased by FAM120A knockdown.

(G–K) qPCR analysis of LAM (G), H1299 (H), DLD1 (I), MCF7 (J), and LNCaP (K) cells transfected with siRNAs targeting FAM120A or control and serum starved overnight. $n = 3$. Data are represented as mean \pm SD. * $p < 0.05$, ** $p < 0.01$, and *** $p < 0.001$.

(L–P) Immunoblot analysis of LAM (L), H1299 (M), DLD1 (N), MCF7 (O), and LNCaP (P) cells transfected with siRNAs targeting FAM120A or control and serum-starved overnight.

See also Figure S1.



(legend on next page)

enzymes.^{12,41,42} To examine whether the same mechanism underlies in FAM120A-dependent regulation of lipogenic genes, we assessed RNA stability by actinomycin D assay.⁴³ FAM120A knockdown accelerated degradation of lipogenic gene transcripts including *ACSS2*, *ACLY*, *FASN*, *HMGCS1*, and *MVD* (Figures 3F–3J). In contrast, the stability of *RPIA*, which is not regulated by mTORC1-SRPK2-FAM120A-SRSF1-dependent splicing, was not dramatically changed (Figure 3K). Suppression of NMD by knocking down a key NMD factor, *up-frameshift 1 (UPF1)*, restored lipogenic gene RNA stability (Figures 3F–3J), as well as their expression levels in FAM120A knockdown cells (Figure 3L). Therefore, FAM120A induces expression of lipogenic enzymes by promoting efficient intron splicing and stability of their RNA.

FAM120A interacts with SRSF1 through lipogenic gene transcripts

Among the FAM120 family proteins, only FAM120A has been identified to be associated with spliceosomes^{23,25,26} and contains an RNA-binding domain (RBD) (Figure 4A).^{44,45} Therefore, we reasoned that RBD would play a critical role in FAM120A-dependent lipogenic gene splicing. Indeed, removal of RBD (FAM120A-ΔRBD) completely abolished its binding with lipogenic gene transcripts (Figures 4B and S3A). FAM120A-ΔRBD failed to bind with SRSF1 (Figure 4C), suggesting that RNA mediates FAM120A-SRSF1 interaction. In line with this notion, treatment of cell lysates with ribonucleases RNase A/T1 disrupted the interaction between FAM120A and SRSF1 (Figure 4D). Removal of SRSF1 RBD (SRSF1-ΔRRM) (Figures 1E and S1A) or ribonuclease treatment to SRSF1-ΔRS also abrogated the binding of SRSF1 with FAM120A (Figure 4D), confirming RNA as an essential component in FAM120A-SRSF1 interaction. In the absence of FAM120A, SRSF1 did not bind with lipogenic transcripts (Figures 4E and S3B), indicating that FAM120A is required for the association of SRSF1 with lipogenic genes.

We then evaluated the functional significance of the binding between FAM120A, SRSF1, and lipogenic transcripts in lipogenic gene expression. The expression of lipogenic enzyme mRNA and protein levels was decreased by FAM120A knockdown in multiple cancer cell lines (Figures 2G–2P), which was rescued by WT FAM120A but not by RNA-binding-deficient FAM120A-ΔRBD mutant (Figures 4F, 4G, and S3C–S3F). Abrogation of FAM120A-SRSF1 interaction by knocking down

SRPK2 and re-expression of mutant SRPK2 proteins led to IR and decreased expression of lipogenic genes (Figures 4H, 4I, and S3G). Collectively, these results suggest that FAM120A facilitates recruitment of lipogenic transcripts to phosphorylated SRSF1, which is essential for efficient splicing and expression of lipogenic enzymes.

FAM120A recruits SRSF1 to lipogenic gene promoter

To regulate transcription and pre-mRNA splicing, SR proteins are recruited to gene promoters and active transcription sites.^{22,46–48} By analyzing previously published chromatin immunoprecipitation sequencing (ChIP-seq) data,⁴⁶ we found that Srsf1 is enriched near the promoter of lipogenic genes including *Acss2*, *Acly*, and *Fasn* (Figures 5A–5C). ChIP-qPCR confirmed that SRSF1 binds lipogenic gene promoters in diverse cancer cells. However, binding of SRSF1 to these promoters was almost completely abolished by FAM120A knockdown (Figures 5D–5F and S4A–S4C), indicating that FAM120A is essential for bringing SRSF1 to these promoters. To understand how FAM120A achieves the specific recruitment of SRSF1 to lipogenic gene promoters, we examined interaction of FAM120A with several lipogenic transcription factors, including SREBP1/2, carbohydrate-response element-binding protein (ChREBP), liver X receptor (LXR) α/β , PPAR α/γ , p300, and X-box-binding protein 1 (XBP1). Among these, only SREBP1 co-immunoprecipitated with FAM120A (Figure 5G). Furthermore, the binding of SREBP1 at lipogenic gene promoters and their promoter activities were substantially decreased by FAM120A depletion (Figures 5H–5J, S4D, and S4E), whereas SREBP1/2 depletion abolished SRSF1 recruitment to lipogenic gene promoters (Figures 5K–5M and S4F–S4H). Therefore, FAM120A is the key mediator that brings SREBP1 and SRSF1 together to the lipogenic genes (Figure S6).

FAM120A connects transcription and splicing machineries to promote efficient splicing of lipogenic genes

Next, we sought to further delve into the nature of the SREBP1-FAM120A-SRSF1 complex. We and others have previously identified U1-70K as the key spliceosome protein recruited to the target genes upon phosphorylation of SRSF1 by mTORC1-SRPK2.^{12,15} Knockdown of *U1-70K* led to the IR of lipogenic genes (Figures 6A and S5A), similar to knockdown of *FAM120A* and *SRSF1* (Figure 3). On the other hand, in the cells

Figure 3. FAM120A is required for the splicing and stability of lipogenic genes

(A) Volcano plot of gene expression from RNA-seq results in LAM cells transfected with siRNAs targeting *FAM120A*, *SRSF1*, or control with overnight serum starvation (GEO: GSE229657). Lipogenic genes are highlighted in rectangles. $n = 3$. Same RNA-seq results are used in (A)–(D).
 (B) Violin plot of 408 intron loci that show significant intron retention (IR) in siFAM120A and siSRSF1 compared with control.
 (C) (Upper) Representative genome browser example of RNA-seq on *FASN1* locus. (Lower) Zoom-in view of intron loci of *FASN* that are retained by *FAM120A* or *SRSF1* knockdown.
 (D) Individual IR ratio calculated from IRFinderS of lipogenic genes from RNA-seq data.
 (E) Validation of intron retention by qPCR analysis of LAM cells transfected with siRNAs targeting *FAM120A*, *SRSF1*, or control. Intron retention = (expression of intron-included region)/(expression of intron-excluded region).
 (F–L) qPCR analysis of LAM cells transfected with siRNAs targeting *FAM120A*, *UPF1*, or control, and serum starved overnight. Actinomycin D (Act D, 5 $\mu\text{g}/\text{mL}$) was treated for the indicated time points for mRNA stability analysis (F–K). $n = 3$. p value (*) was calculated between siNTC and siFAM120A. p value (#) was calculated between siFAM120A and siFAM120A + siUPF1.
 Data are represented as mean \pm SD. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, ## $p < 0.01$, and ### $p < 0.001$.
 See also Figure S2 and Table S1.

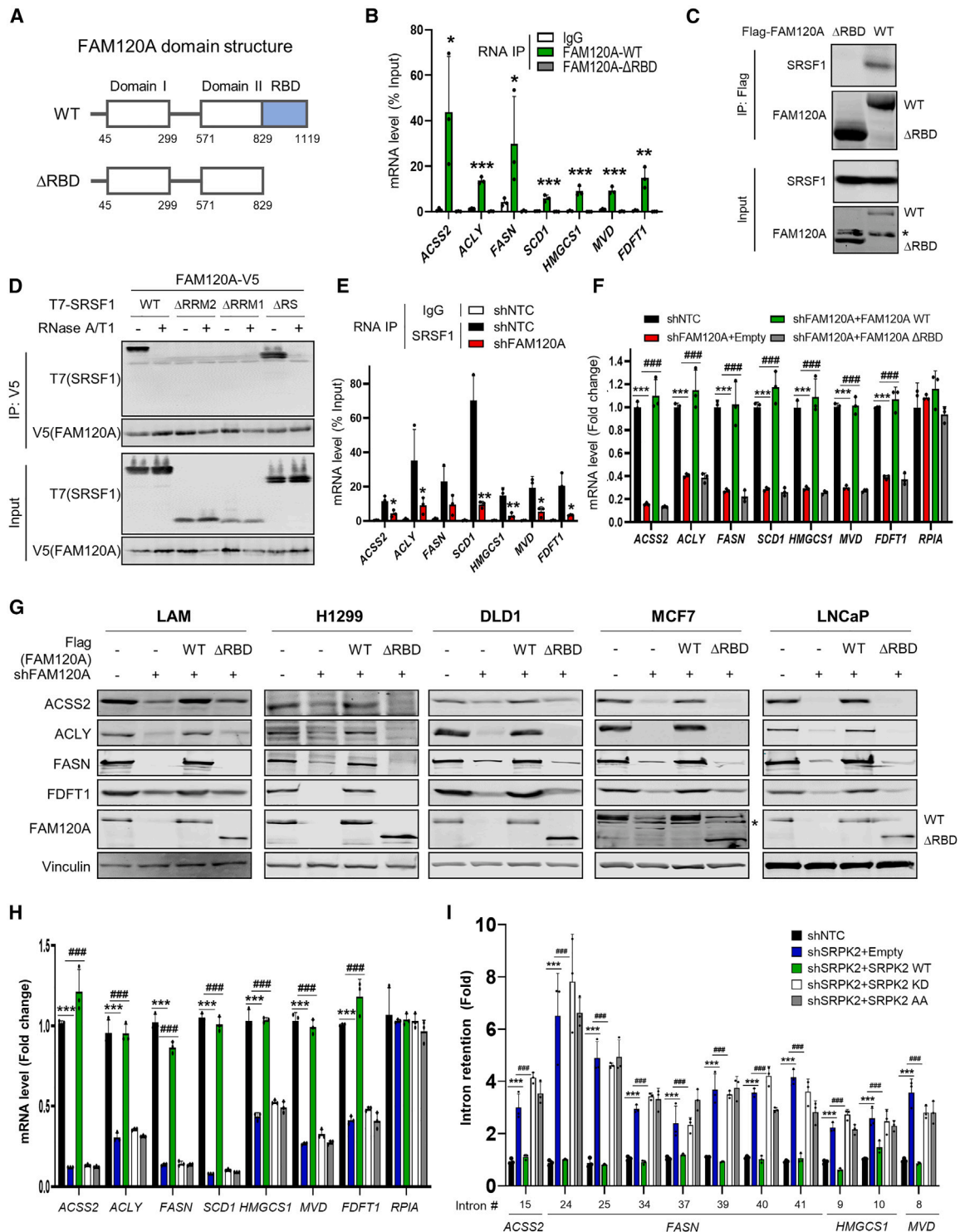


Figure 4. FAM120A interacts with SRSF1 through lipogenic gene RNAs

(A) Schematic diagram of the domains in wild-type (WT) and RNA-binding domain (RBD)-deleted (Δ RBD) FAM120A.

(B) RNA immunoprecipitation qPCR (RIP-qPCR) analysis of LAM cells expressing FAM120A WT or Δ RBD. IP was performed with IgG or anti-FAM120A antibodies. The graph shows qPCR analysis of RIP products. $n = 3$. p value was calculated between IgG and FAM120A-WT.

(C) CoIP analysis of HEK293E cells expressing SRSF1 with FLAG-FAM120A WT or Δ RBD. 1% of total cell lysate was loaded as an input. Asterisk indicates a non-specific band.

(legend continued on next page)

depleted with the transcription factor *SREBP1/2*, *FAM120A* knockdown did not induce IR (Figures 6B and S5B), indicating that transcription is the pre-requisite for the splicing of lipogenic genes. Immunoprecipitation of endogenous *FAM120A* confirmed that RNA Pol II and U1-70K associate with the SREBP1-*FAM120A*-SRSF1 complex (Figure 6C). Ribonuclease treatment abolished the binding among these proteins (Figure 6C), showing that RNA is a key component of this multi-protein co-transcriptional splicing complex. Notably, although deletion of RBD of SRSF1 (SRSF1- Δ RRM) abolished its interaction with *FAM120A*, the binding between *FAM120A* and SREBP1 was intact (Figure 6D). In contrast, deletion of the RBD of *FAM120A* (*FAM120A*- Δ RBD) dissociated binding of all three proteins (Figure 6E); hence, *FAM120A* bridges the binding between SREBP1 and SRSF1. Indeed, SRSF1 did not directly bind with SREBP1 (Figure S5C).

Given that the transcriptionally active nuclear form of SREBP1, SREBP1(M), was identified to interact with *FAM120A* (Figure 5G), we examined whether SREBP1 processing affects its interaction with *FAM120A* or vice versa. Depletion of cholesterol by serum starvation induces cleavage of the precursor SREBP1(P) to the mature SREBP1(M) in cancer cells (Figures 6F and 6G), whereas insulin promotes SREBP1(M) production in non-cancerous HEK293E cells⁴⁹ (Figure 6H). In both cases, inhibition of SREBP1 cleavage by 25-hydroxycholesterol (25-HC) treatment⁵⁰ prevented SREBP1(M) formation and its interaction with *FAM120A* (Figures 6F–6H). However, neither *FAM120A* knockdown nor the RNA-binding-defective *FAM120A* mutant (*FAM120A*- Δ RBD) decreased SREBP1(M) levels (Figures 6E, S5C, and S5D), demonstrating that *FAM120A* does not affect SREBP1 cleavage.

Because SREBP1-*FAM120A* interaction was independent of the SRSF1-*FAM120A* interaction (Figure 6D), we reasoned that the formation of the SREBP1-*FAM120A*-RNA Pol II transcription complex would be independent of splicing complex formation. Unphosphorylated SRSF1 cannot bind with *FAM120A* (Figures 1E–1G and S1A–S1C) or spliceosome protein U1-70K.^{12,15} However, suppression of SRSF1 phosphorylation by SRPK inhibitor (SRPKIN-1) did not disrupt the interaction of *FAM120A* with SREBP1 and RNA Pol II (Figures 6F–6H). Blocking SREBP1 cleavage by 25-HC treatment did not affect the binding of *FAM120A* with SRSF1 and RNA Pol II, either (Figures 6F–6H). Together, these findings suggest that *FAM120A* bridges the two independent transcription and splicing complexes to promote efficient splicing of newly transcribed lipogenic genes (Figure S6).

FAM120A is essential for lipid production and tumor growth

To examine the functional relevance of *FAM120A*-dependent lipogenic gene regulation, we measured levels of the four most abundant fatty acids (C16:0, C16:1, C18:0, and C18:1) produced by *de novo* lipid synthesis using liquid chromatography-mass spectrometry (LC-MS).⁵¹ The knockdown of *FAM120A* markedly depleted these fatty acids in LAM and H1299 cells (Figures 7A–7C, S7A, and S7B). Consistent with the notion that these fatty acids are critical for phospholipid membrane synthesis during cell proliferation,⁵² *FAM120A* knockdown suppressed proliferation of various cancer cell lines (Figures 7D–7H and S7C–S7G). Removal of exogenous lipid supply via lipoprotein-deficient serum (LPDS)⁵³ augmented growth defects in *FAM120A* knockdown cells, which was rescued by supplementation of exogenous lipoproteins or fatty acids (Figures 7I–7M and S7H–S7O). Finally, *FAM120A*-depleted cells showed strongly reduced xenograft tumor growth *in vivo* and re-expression of WT *FAM120A* but not *FAM120A*- Δ RBD mutant partially rescued tumor growth (Figures 7N, 7O, and S7P). Together, these data show that *FAM120A*-dependent lipogenic gene expression is critical for fatty acid synthesis and growth of tumor cells.

DISCUSSION

Here, we discovered *FAM120A* as a unique factor that bridges transcription and splicing of lipogenic genes for tumor cell growth. As a transcription co-factor and RBP, *FAM120A* binds to both transcription factor SREBP and splicing factor SRSF1 to connect efficient splicing of newly transcribed lipogenic gene transcripts. This function of *FAM120A* is stimulated by the mTORC1-activated SRPK2 signaling in various cancers (Figure S6).

Splicing is carried out by a snRNP complex composed of small nuclear RNAs and RNPs.⁵⁴ Although the core spliceosome complex (U1, U2, U4, U5, and U6 snRNPs) recognizes exon-intron borders and splices introns, additional RBPs are involved in determining the target gene specificity and direction of the splicing events. SR proteins are the splicing inducers that bind to exonic splicing enhancer (ESE) sequences for the recruitment of spliceosome complex.⁵⁵ Association of SR proteins with the spliceosome proteins is a highly regulated process, and we previously showed that mTORC1-SRPK2-dependent phosphorylation of SRSF1 promotes its interaction with U1-70K,¹² a U1 snRNP protein involved in spliceosome formation initiation^{15,56}. In this study, we demonstrate that mTORC1-SRPK2 drives the

(D) CoIP analysis of HEK293E cells expressing *FAM120A*-V5 and T7-SRSF1 (WT or mutants). Cell lysates were treated with RNases before IP. 1% of total cell lysate was loaded as an input.

(E) RIP-qPCR analysis of LAM cells stably expressing shRNAs targeting *FAM120A* or control. Immunoprecipitation was performed with IgG or anti-SRSF1 antibodies. The graph shows qPCR analysis of RIP products. $n = 3$. p value was calculated between SRSF1-IP samples from shNTC and sh*FAM120A*.

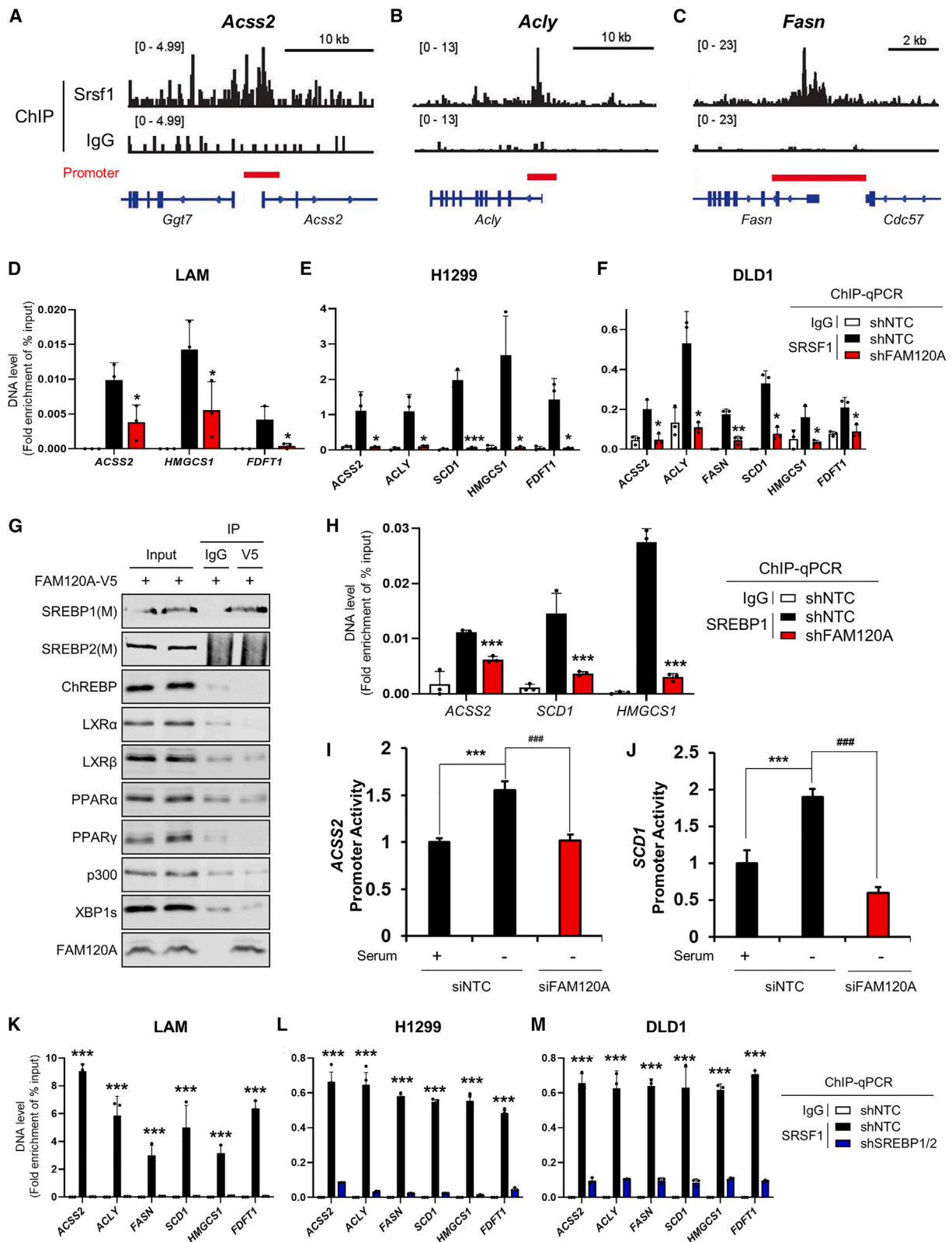
(F) qPCR analysis of LAM cells stably expressing *FAM120A* WT or Δ RBD in *FAM120A* knockdown cells. Cells were serum-starved overnight. $n = 3$.

(G) Immunoblot analysis of LAM, H1299, and DLD1 cells stably expressing *FAM120A* WT or Δ RBD in *FAM120A* knockdown cells. Cells were serum-starved overnight.

(H and I) qPCR analysis of LAM cells stably expressing HA-SRPK2 (WT [wild type], KD [kinase dead, SRPK2-K110M], or AA [non-phosphorylatable, SRPK2-S394A/S397A]). Endogenous *SRPK2* was knocked down by shRNA targeting 3' UTR of *SRPK2*. Cells were serum-starved overnight. Graphs show mRNA levels (H) and intron retention (I). $n = 3$.

Data are represented as mean \pm SD. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$, and #### $p < 0.0001$.

See also Figure S3.



(legend on next page)

selectivity of SRSF1-dependent splicing toward *de novo* lipid synthesis genes through an SREBP1-interacting transcription co-factor, FAM120A. SREBP1 is a transcription factor that recruits transcription preinitiation complex proteins, including Pol II, TATA box-binding protein (TBP), and other general transcription factors, such as transcription initiation factor IIA (TFIIA) and TFIIB. Once Pol II is released from the transcription initiation stage, Pol II enters an active elongation stage and moves along the gene body with transcription elongation associated proteins, including positive-transcription elongation factor b (P-TEFb) and RNA-processing proteins.^{57–59} Therefore, it is unlikely that SREBP1 piggybacks with Pol II and SRSF1 through the transcript during transcription elongation. Instead, our proposed mechanism highlights a key role for FAM120A in coordinating transcription initiation and splicing of target genes by promoting the formation of distinct complexes that augment the efficiency of these processes. In the case of lipogenic gene transcription, one complex is composed of FAM120A-Pol II-SREBP1, whereas the other one responsible for the splicing of newly transcribed genes contains FAM120A-Pol II-U1-70K-SRSF1. Notably, the two complexes form independently of each other, but they cooperate for the FAM120A-mediated co-transcriptional splicing of lipogenic genes (Figures 6 and S6). This provides a *bona fide* mechanism for the efficient coupling of transcription, splicing, and stabilization of SREBP1-regulated lipogenic genes.

As a master regulator of cell growth, mTORC1 is acutely activated by nutrients and growth factors to coordinate macromolecule synthesis for cell growth. However, when mTORC1 is aberrantly and constitutively activated, as in many human cancers, its activation leads to overgrowth and proliferation.^{60,61} Given the vastly broad effect of mTORC1 in biological processes, it may be important to target specific downstream signaling effectors for therapeutic purposes to avoid severe toxicities of mTORC1 targeting drugs.⁶² Our data suggest that suppressing lipogenic gene splicing via FAM120A binding inhibitors or SRPK inhibitors would target a possible metabolic vulnerability. Splicing inhibitors have been in clinical trials based on the findings that splicing is overactivated in tumor cells.⁶³ Because splicing is such a fundamental process that occurs across all mammalian tissues and cell types, splicing inhibitors caused toxic side effects in patients.^{64,65} On the other hand, selectively targeting the SRPK2-SRSF1-FAM120A axis could be an alternative therapeutic strategy to preferentially suppress lipogenic enzymes with

low toxicity. In fact, suppressing fatty acid synthesis by a second generation fatty acid synthase (FASN) inhibitor was well tolerated because *de novo* lipid synthesis is almost inactive in many cells except for the hepatocytes and adipocytes.^{66–68} Looking forward, it will be exciting to investigate if small molecule inhibitors, oligonucleotides, and RNA-PROTACs (proteolysis targeting chimerics)^{69–71} that block FAM120A function and its interaction with lipogenic transcripts (e.g., via its RBD) become important therapies for the treatment of mTORC1-driven cancers.

Limitations of the study

In this study, we found FAM120A as a *bona fide* effector in the mTORC1-SRPK2-SRSF1 signaling cascade that determines the downstream target specificity toward lipid metabolism genes (Figure S6). However, from the pathway enrichment analysis of transcriptome and splicing changes in *FAM120A* and *SRSF1* knockdown cells, we also identified interferon (IFN) signaling and immune response genes, such as IFN stimulated genes (ISGs) and major histocompatibility complex (MHC)-mediated antigen-processing genes, as top biological processes regulated by FAM120A and SRSF1 (Figure S2; Table S1). The potential regulation of the IFN pathway by FAM120A and SRSF1 is particularly interesting because IFN signaling was among the top biological pathways regulated by SRPK2 in our previous study.¹² The ISG, *myxovirus resistance 1 (MX1)*, was, indeed, one of the commonly downregulated genes by rapamycin, SRPK2 knockdown, and *FAM120A* knockdown (Figure 2C). These observations suggest that IFN-mediated immune response could be another important biological function of mTORC1-SRPK2-SRSF1-FAM120A pathway. Similar to *de novo* lipid synthesis enzymes, ISGs are known to be transcribed by master transcription factors such as signal transducer and activator of transcription (STAT) and IFN regulatory factor (IRF).^{72–74} Future studies will be required to identify the immune transcription factor that interacts with FAM120A for the coordination of efficient transcription and splicing of ISGs. Although traditional IFN signaling and immune research have been focused on anti-viral responses, recent studies found that programmed cell-death ligand 1 (PDL1)-associated tumor resistance is linked to the sustained expression of ISGs, including *MX1*.^{75–77} Ultimately, understanding the role of SRPK2 and FAM120A in immune checkpoint control can potentially provide mechanistic insights for improving immunotherapeutic

Figure 5. FAM120A bridges SRSF1 to SREBP1 to induce lipogenic enzyme expression

(A–C) Analysis of Srsf1 chromatin immunoprecipitation (ChIP)-seq results (GEO: GSE45517)⁴⁶ in MEF cells near the transcription start sites of *Acss2* (A), *Acly* (B), and *Fasn* (C).

(D–F) ChIP-qPCR analysis of SRSF1 in target gene promoters. Immunoprecipitation was performed using IgG or anti-SRSF1 antibodies in LAM (D), H1299 (E), and DLD1 (F) cells stably expressing shRNAs targeting *FAM120A* or control. Cells were serum-starved overnight. p value was calculated between shNTC and shFAM120A. n = 3.

(G) CoIP analysis of transcription factors in HEK293E cells expressing FAM120A-V5. 1% of total cell lysate was loaded as an input.

(H) ChIP-qPCR analysis of SREBP1 on target gene promoters. Immunoprecipitation was performed using IgG or anti-SREBP1 antibodies in LAM cells stably expressing shRNAs targeting *FAM120A* or control. Cells were serum starved overnight. p value was calculated between shNTC and shFAM120A. n = 3.

(I and J) Promoter activity analysis of *ACSS2* (I) and *SCD1* (J) in LAM cells transfected with promoter constructs and siRNAs targeting *FAM120A* or control. n = 3.

(K–M) ChIP-qPCR analysis of SRSF1 in target gene promoters. Immunoprecipitation was performed using IgG or anti-SRSF1 antibodies in LAM (K), H1299 (L), and DLD1 (M) cells stably expressing shRNAs targeting *SREBP1/2* or control. Cells were serum-starved overnight. p value was calculated between shNTC and shSREBP1/2. n = 3.

Data are represented as mean ± SD. *p < 0.05, **p < 0.01, ***p < 0.001, and ###p < 0.001.

See also Figure S4.

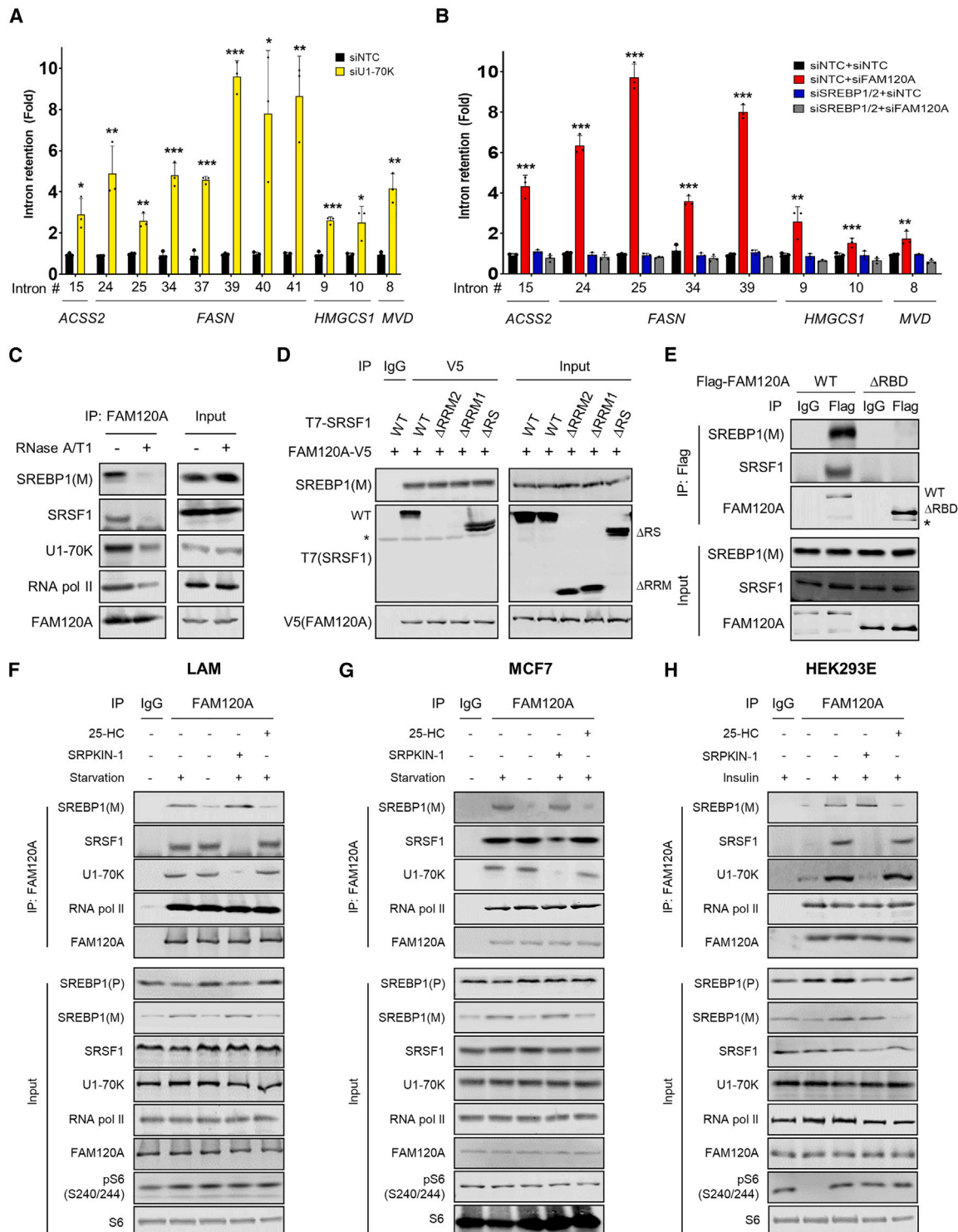


Figure 6. FAM120A promotes efficient splicing of lipogenic genes by associating with transcriptional and splicing machineries

(A and B) qPCR analysis of intron retention in LAM cells transfected with siRNAs targeting *U1-70K* or control (A) and *FAM120A*, *SREBP1/2*, *SREBP1/2* + *FAM120A* or control (B). Cells were serum starved overnight. n = 3. Data are represented as mean ± SD. *p < 0.05, **p < 0.01, and ***p < 0.001. p value was calculated between siNTC and siFAM120A (B). n = 3.

(C) CoIP analysis of HEK293E cells. Immunoprecipitation was performed using anti-FAM120A antibodies. Cell lysates were treated with RNases before IP. 1% of total cell lysate was loaded as an input.

(legend continued on next page)

approaches and patient outcomes in various immune-related diseases and cancers.

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

- **KEY RESOURCES TABLE**
- **RESOURCE AVAILABILITY**
 - Lead contact
 - Materials availability
 - Data and code availability
- **EXPERIMENTAL MODEL AND STUDY PARTICIPANT DETAILS**
 - Cell lines and cell culture
 - Mouse studies
- **METHOD DETAILS**
 - Expression constructs
 - siRNA oligonucleotides
 - Generation of stable cell lines
 - Crystal violet staining
 - Immunoblot analysis
 - Quantitative RT-PCR
 - mRNA stability assay
 - Luciferase promoter activity assay
 - RNA immunoprecipitation qPCR
 - Co-immunoprecipitation of proteins
 - Ribonuclease treatment
 - Chromatin immunoprecipitation qPCR
 - Lipid supplementation
 - Mass spectrometry analysis of SRSF1 interactome
 - Mass spectrometry analysis of saponified fatty acids
 - RNA-seq sample preparation
 - RNA-seq and pathway enrichment analyses
 - Intron retention analysis
 - Analysis of published ChIP-seq and whole transcriptome microarray results
- **QUANTIFICATION AND STATISTICAL ANALYSIS**

SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.molcel.2023.07.017>.

ACKNOWLEDGMENTS

We thank members of the Blenis and Lee laboratories for technical assistance and discussions. We are grateful to Drs. Elizabeth Henske, Adrian Krainer,

and Yongsheng Shi for sharing reagents and scientific discussion. This research was supported by the following grants: NIH R01GM051405 (J.B.), R01CA046595 (J.B.), R01AA029124 (C.J.), K22CA234399 (G.L.), 1DP2GM149554 (F.D.), DOD TS200022 (G.L.), Mary Kay Ash Foundation (G.L.), and STARR Foundation (J.B). This work was also made possible in part through access to CFCCC GRTH core (P30CA062203). J.C. was supported by a Basic Science Research Program (2022R1A4A2000827) through the National Research Foundation of Korea. C.B.R. was supported by predoctoral fellowships from UCI IMSD (R25GM055246) and IDCR (T32CA009054) programs. Schematics were created using BioRender.

AUTHOR CONTRIBUTIONS

G.L., S.C., and J.B. designed the study. S.C. conducted all experiments unless otherwise indicated. Y.C., K.J., J.S., J.G.C., and J.C. performed bioinformatics analysis. C.B.R. and C.J. performed the LC-MS analysis. L.H., K.S.G., Z.L., J.K., and N.K. performed some of the cellular and molecular biological experiments. N.D. analyzed the interactome data. F.D. advised on RNA-seq analysis. G.L., S.C., and J.B. wrote the manuscript. All authors discussed the results and commented on the manuscript.

DECLARATION OF INTERESTS

J.B. is an advisory board member for *Molecular Cell*.

Received: July 28, 2022

Revised: May 23, 2023

Accepted: July 15, 2023

Published: August 17, 2023

REFERENCES

1. Saxton, R.A., and Sabatini, D.M. (2017). mTOR signaling in growth, metabolism, and disease. *Cell* *169*, 361–371. <https://doi.org/10.1016/j.cell.2017.03.035>.
2. Shimobayashi, M., and Hall, M.N. (2014). Making new contacts: the mTOR network in metabolism and signalling crosstalk. *Nat. Rev. Mol. Cell Biol.* *15*, 155–162. <https://doi.org/10.1038/nrm3757>.
3. Ben-Sahra, I., Hoxhaj, G., Ricoult, S.J.H., Asara, J.M., and Manning, B.D. (2016). mTORC1 induces purine synthesis through control of the mitochondrial tetrahydrofolate cycle. *Science* *351*, 728–733. <https://doi.org/10.1126/science.aad0489>.
4. Dibble, C.C., and Manning, B.D. (2013). Signal integration by mTORC1 coordinates nutrient input with biosynthetic output. *Nat. Cell Biol.* *15*, 555–564. <https://doi.org/10.1038/ncb2763>.
5. Han, J., Li, E., Chen, L., Zhang, Y., Wei, F., Liu, J., Deng, H., and Wang, Y. (2015). The CREB coactivator CRTC2 controls hepatic lipid metabolism by regulating SREBP1. *Nature* *524*, 243–246. <https://doi.org/10.1038/nature14557>.
6. Li, S., Brown, M.S., and Goldstein, J.L. (2010). Bifurcation of insulin signaling pathway in rat liver: mTORC1 required for stimulation of lipogenesis, but not inhibition of gluconeogenesis. *Proc. Natl. Acad. Sci. USA* *107*, 3441–3446. <https://doi.org/10.1073/pnas.0914798107>.

(D) CoIP analysis of HEK293E cells expressing FAM120A-V5 and T7-SRSF1 (WT or mutants). 1% of total cell lysate was loaded as an input. Asterisk indicates a non-specific band.

(E) CoIP analysis of HEK293E cells expressing FLAG-FAM120A WT or Δ RBD. 1% of total cell lysate was loaded as an input. Asterisk indicates a non-specific band.

(F–H) CoIP analysis of LAM (F) and MCF7 (G) cells serum starved overnight (to induce SREBP cleavage) and treated with SREBP cleavage inhibitor (25-HC, 10 μ M) or SRPK inhibitor (SRPKIN-1, 5 μ M) for 4 h (F and G); HEK293E cells serum starved overnight followed by insulin stimulation (100 nM) (to induce SREBP cleavage) with or without 25-HC (10 μ M) or SRPKIN-1 (5 μ M) for 4 h (H). Note that serum starvation decreases mTORC1 activity in HEK293E cells (H) but does not affect constitutively active mTORC1 activity in LAM and MCF7 cancer cells (F and G). Immunoprecipitation was performed using IgG or anti-FAM120A antibodies. 1% of total cell lysate was loaded as an input. SREBP1(P), precursor SREBP1; SREBP1(M), cleaved and matured SREBP1.

See also [Figure S5](#).

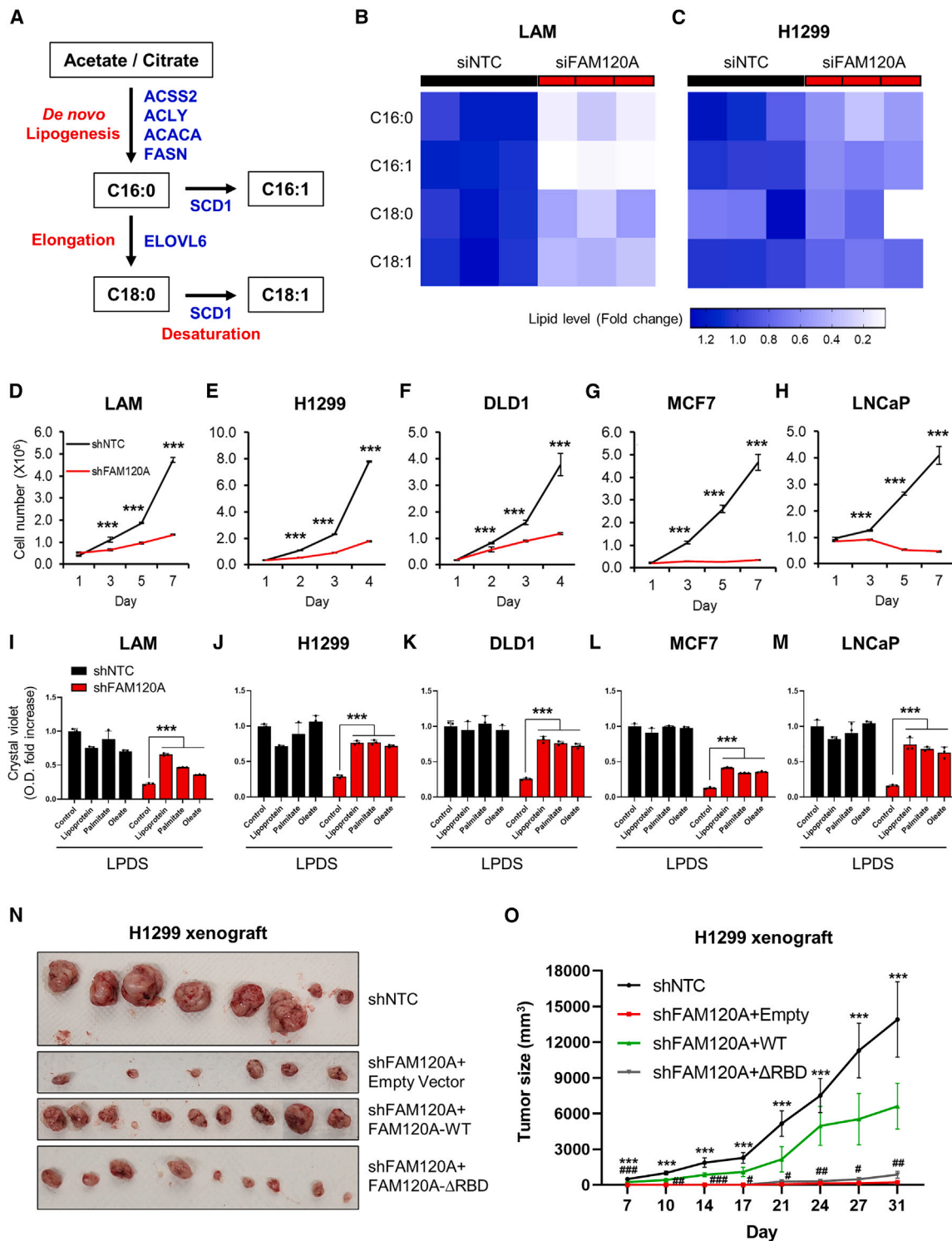


Figure 7. FAM120A RNA binding is essential for lipogenesis and tumor growth

(A) Schematic of *de novo* fatty acid synthesis pathway.

(B and C) Heatmap of fatty acid levels. LC-MS analysis was performed on LAM cells transfected with siRNAs targeting *FAM120A* or control (B) and H1299 cells stably expressing shRNAs targeting *FAM120A* or control (C) with overnight serum starvation. $n = 3$.

(D–H) Cell proliferation analysis of LAM (D), H1299 (E), DLD1 (F), MCF7 (G), and LNCaP (H) cells stably expressing shRNAs targeting *FAM120A* or control. $n = 3$. Data are represented as mean \pm SD.

(legend continued on next page)

7. Owen, J.L., Zhang, Y., Bae, S.H., Farooqi, M.S., Liang, G., Hammer, R.E., Goldstein, J.L., and Brown, M.S. (2012). Insulin stimulation of SREBP-1c processing in transgenic rat hepatocytes requires p70 S6-kinase. *Proc. Natl. Acad. Sci. USA* *109*, 16184–16189. <https://doi.org/10.1073/pnas.1213343109>.
8. Peterson, T.R., SenGupta, S.S., Harris, T.E., Carmack, A.E., Kang, S.A., Balderas, E., Guertin, D.A., Madden, K.L., Carpenter, A.E., Finck, B.N., et al. (2011). mTOR complex 1 regulates lipin 1 localization to control the SREBP pathway. *Cell* *146*, 408–420. <https://doi.org/10.1016/j.cell.2011.06.034>.
9. Zhang, Y., Nicholatos, J., Dreier, J.R., Ricoult, S.J.H., Widenmaier, S.B., Hotamisligil, G.S., Kwiatkowski, D.J., and Manning, B.D. (2014). Coordinated regulation of protein synthesis and degradation by mTORC1. *Nature* *513*, 440–443. <https://doi.org/10.1038/nature13492>.
10. Düvel, K., Yecies, J.L., Menon, S., Raman, P., Lipovsky, A.I., Souza, A.L., Triantafellow, E., Ma, Q., Gorski, R., Cleaver, S., et al. (2010). Activation of a metabolic gene regulatory network downstream of mTOR complex 1. *Mol. Cell* *39*, 171–183. <https://doi.org/10.1016/j.molcel.2010.06.022>.
11. Heintz, C., Doktor, T.K., Lanjuin, A., Escoubas, C., Zhang, Y., Weir, H.J., Dutta, S., Silva-Garcia, C.G., Bruun, G.H., Morantte, I., et al. (2017). Splicing factor 1 modulates dietary restriction and TORC1 pathway longevity in *C. elegans*. *Nature* *541*, 102–106. <https://doi.org/10.1038/nature20789>.
12. Lee, G., Zheng, Y., Cho, S., Jang, C., England, C., Dempsey, J.M., Yu, Y., Liu, X., He, L., Cavaliere, P.M., et al. (2017). Post-transcriptional regulation of de novo lipogenesis by mTORC1-S6K1-SRPK2 signaling. *Cell* *171*, 1545–1558.e18. <https://doi.org/10.1016/j.cell.2017.10.037>.
13. Sinvani, H., Haimov, O., Svitkin, Y., Sonenberg, N., Tamarkin-Ben-Harush, A., Viollet, B., and Dikstein, R. (2015). Translational tolerance of mitochondrial genes to metabolic energy stress involves TISU and eIF1-eIF4G cooperation in start codon selection. *Cell Metab.* *21*, 479–492. <https://doi.org/10.1016/j.cmet.2015.02.010>.
14. Tang, H.W., Hu, Y., Chen, C.L., Xia, B., Zirin, J., Yuan, M., Asara, J.M., Rabinow, L., and Perrimon, N. (2018). The TORC1-regulated CPA complex rewires an RNA processing network to drive autophagy and metabolic reprogramming. *Cell Metab.* *27*, 1040–1054.e8. <https://doi.org/10.1016/j.cmet.2018.02.023>.
15. Cho, S., Hoang, A., Sinha, R., Zhong, X.Y., Fu, X.D., Krainer, A.R., and Ghosh, G. (2011). Interaction between the RNA binding domains of Ser-Arg splicing factor 1 and U1-70K snRNP protein determines early spliceosome assembly. *Proc. Natl. Acad. Sci. USA* *108*, 8233–8238. <https://doi.org/10.1073/pnas.1017700108>.
16. Kohtz, J.D., Jamison, S.F., Will, C.L., Zuo, P., Lührmann, R., Garcia-Blanco, M.A., and Manley, J.L. (1994). Protein-protein interactions and 5'-splice-site recognition in mammalian mRNA precursors. *Nature* *368*, 119–124. <https://doi.org/10.1038/368119a0>.
17. Wang, H.Y., Lin, W., Dyck, J.A., Yeakley, J.M., Songyang, Z., Cantley, L.C., and Fu, X.D. (1998). SRPK2: A differentially expressed SR protein-specific kinase involved in mediating the interaction and localization of pre-mRNA splicing factors in mammalian cells. *J. Cell Biol.* *140*, 737–750.
18. Das, R., Dufu, K., Romney, B., Feldt, M., Elenko, M., and Reed, R. (2006). Functional coupling of RNAP II transcription to spliceosome assembly. *Genes Dev.* *20*, 1100–1109. <https://doi.org/10.1101/gad.1397406>.
19. Das, R., Yu, J., Zhang, Z., Gygi, M.P., Krainer, A.R., Gygi, S.P., and Reed, R. (2007). SR proteins function in coupling RNAP II transcription to pre-mRNA splicing. *Mol. Cell* *26*, 867–881. <https://doi.org/10.1016/j.molcel.2007.05.036>.
20. de la Mata, M., and Kornblihtt, A.R. (2006). RNA polymerase II C-terminal domain mediates regulation of alternative splicing by SRp20. *Nat. Struct. Mol. Biol.* *13*, 973–980. <https://doi.org/10.1038/nsmb1155>.
21. Sapra, A.K., Ankó, M.L., Grishina, I., Lorenz, M., Pabis, M., Poser, I., Rollins, J., Weiland, E.M., and Neugebauer, K.M. (2009). SR protein family members display diverse activities in the formation of nascent and mature mRNPs in vivo. *Mol. Cell* *34*, 179–190. <https://doi.org/10.1016/j.molcel.2009.02.031>.
22. Monsalve, M., Wu, Z., Adelmant, G., Puigserver, P., Fan, M., and Spiegelman, B.M. (2000). Direct coupling of transcription and mRNA processing through the thermogenic coactivator PGC-1. *Mol. Cell* *6*, 307–316. [https://doi.org/10.1016/s1097-2765\(00\)00031-9](https://doi.org/10.1016/s1097-2765(00)00031-9).
23. Chen, Y.I., Moore, R.E., Ge, H.Y., Young, M.K., Lee, T.D., and Stevens, S.W. (2007). Proteomic analysis of in vivo-assembled pre-mRNA splicing complexes expands the catalog of participating factors. *Nucleic Acids Res.* *35*, 3928–3944. <https://doi.org/10.1093/nar/gkm347>.
24. Möller, A., Xie, S.Q., Hosp, F., Lang, B., Phatnani, H.P., James, S., Ramirez, F., Collin, G.B., Naggert, J.K., Babu, M.M., et al. (2012). Proteomic analysis of mitotic RNA polymerase II reveals novel interactors and association with proteins dysfunctional in disease. *Mol. Cell. Proteomics* *11*, M111.011767. <https://doi.org/10.1074/mcp.M111.011767>.
25. Rappsilber, J., Ryder, U., Lamond, A.I., and Mann, M. (2002). Large-scale proteomic analysis of the human spliceosome. *Genome Res.* *12*, 1231–1245. <https://doi.org/10.1101/gr.473902>.
26. Sharma, S., Kohlstaedt, L.A., Damianov, A., Rio, D.C., and Black, D.L. (2008). Polypyrimidine tract binding protein controls the transition from exon definition to an intron defined spliceosome. *Nat. Struct. Mol. Biol.* *15*, 183–191. <https://doi.org/10.1038/nsmb.1375>.
27. Das, S., and Krainer, A.R. (2014). Emerging functions of SRSF1, splicing factor and oncoprotein, in RNA metabolism and cancer. *Mol. Cancer Res.* *12*, 1195–1204. <https://doi.org/10.1158/1541-7786.MCR-14-0131>.
28. Zhou, Z., and Fu, X.D. (2013). Regulation of splicing by SR proteins and SR protein-specific kinases. *Chromosoma* *122*, 191–207. <https://doi.org/10.1007/s00412-013-0407-z>.
29. Xiao, S.H., and Manley, J.L. (1997). Phosphorylation of the ASF/SF2 RS domain affects both protein-protein and protein-RNA interactions and is necessary for splicing. *Genes Dev.* *11*, 334–344. <https://doi.org/10.1101/gad.11.3.334>.
30. Yeakley, J.M., Tronchère, H., Olesen, J., Dyck, J.A., Wang, H.Y., and Fu, X.D. (1999). Phosphorylation regulates in vivo interaction and molecular targeting of serine/arginine-rich Pre-mRNA splicing factors. *J. Cell Biol.* *145*, 447–455. <https://doi.org/10.1083/jcb.145.3.447>.
31. Fukuhara, T., Hosoya, T., Shimizu, S., Sumi, K., Oshiro, T., Yoshinaka, Y., Suzuki, M., Yamamoto, N., Herzenberg, L.A., Herzenberg, L.A., et al. (2006). Utilization of host SR protein kinases and RNA-splicing machinery during viral replication. *Proc. Natl. Acad. Sci. USA* *103*, 11329–11333. <https://doi.org/10.1073/pnas.0604616103>.
32. Hatcher, J.M., Wu, G., Zeng, C., Zhu, J., Meng, F., Patel, S., Wang, W., Ficarro, S.B., Leggett, A.L., Powell, C.E., et al. (2018). SRPKIN-1: a covalent SRPK1/2 inhibitor that potently converts VEGF from pro-angiogenic to

(I–M) Crystal violet (CV) analysis of LAM (I), H1299 (J), DLD1 (K), MCF7 (L), and LNCaP (M) cells stably expressing shRNAs targeting *FAM120A* or control. Cells were grown in lipoprotein-deficient serum (LPDS)-containing media supplemented with lipoprotein (25 $\mu\text{g}/\text{mL}$), palmitate-albumin (10 μM , palmitate), oleate-albumin (50 μM , oleate), or fatty acid-free albumin (25 μM , control). The graph shows the quantified absorbance of solubilized CV stain. $n = 3$. Data are represented as mean \pm SD.

(N and O) Xenograft tumor growth assay of H1299 cells stably expressing shNTC or shFAM120A with *FAM120A* WT or ΔRBD . $n \geq 4$. Data are represented as mean \pm SEM. p value (*) was calculated between shNTC and shFAM120A + Empty. p value (#) was calculated between shFAM120A + Empty and shFAM120A + WT. Data are represented as mean \pm SD. *** $p < 0.001$, # $p < 0.05$, ## $p < 0.01$, and ### $p < 0.001$.

See also [Figure S7](#).

- anti-angiogenic isoform. *Cell Chem. Biol.* 25, 460–470.e6. <https://doi.org/10.1016/j.chembiol.2018.01.013>.
33. Yu, J., Astrinidis, A., Howard, S., and Henske, E.P. (2004). Estradiol and tamoxifen stimulate LAM-associated angiomyolipoma cell growth and activate both genomic and nongenomic signaling pathways. *Am. J. Physiol. Lung Cell. Mol. Physiol.* 286, L694–L700. <https://doi.org/10.1152/ajplung.00204.2003>.
 34. Krall, E.B., Wang, B., Munoz, D.M., Ilic, N., Raghavan, S., Niederst, M.J., Yu, K., Ruddy, D.A., Aguirre, A.J., Kim, J.W., et al. (2017). KEAP1 loss modulates sensitivity to kinase targeted therapy in lung cancer. *eLife* 6, e18970. <https://doi.org/10.7554/eLife.18970>.
 35. Lee, J., Lee, I., Han, B., Park, J.O., Jang, J., Park, C., and Kang, W.K. (2011). Effect of simvastatin on cetuximab resistance in human colorectal cancer with KRAS mutations. *J. Natl. Cancer Inst.* 103, 674–688. <https://doi.org/10.1093/jnci/djr707>.
 36. Wu, G., Xing, M., Mambo, E., Huang, X., Liu, J., Guo, Z., Chatterjee, A., Goldenberg, D., Gollin, S.M., Sukumar, S., et al. (2005). Somatic mutation and gain of copy number of PIK3CA in human breast cancer. *Breast Cancer Res.* 7, R609–R616. <https://doi.org/10.1186/bcr1262>.
 37. Carver, B.S., Chapinski, C., Wongvipat, J., Hieronymus, H., Chen, Y., Chandralapaty, S., Arora, V.K., Le, C., Koutcher, J., Scher, H., et al. (2011). Reciprocal feedback regulation of PI3K and androgen receptor signaling in PTEN-deficient prostate cancer. *Cancer Cell* 19, 575–586. <https://doi.org/10.1016/j.ccr.2011.04.008>.
 38. Buj, R., Chen, C.W., Dahl, E.S., Leon, K.E., Kuskovsky, R., Maglakelidze, N., Navaratnarajah, M., Zhang, G., Doan, M.T., Jiang, H., et al. (2019). Suppression of p16 induces mTORC1-mediated nucleotide metabolic reprogramming. *Cell Rep.* 28, 1971–1980.e8. <https://doi.org/10.1016/j.celrep.2019.07.084>.
 39. Irimia, M., Weatheritt, R.J., Ellis, J.D., Parikshak, N.N., Gonatopoulos-Pournatzis, T., Babor, M., Quesnel-Vallières, M., Tapial, J., Raj, B., O'Hanlon, D., et al. (2014). A highly conserved program of neuronal micro-exons is misregulated in autistic brains. *Cell* 159, 1511–1523. <https://doi.org/10.1016/j.cell.2014.11.035>.
 40. Lorenzi, C., Barriere, S., Arnold, K., Luco, R.F., Oldfield, A.J., and Ritchie, W. (2021). IRFinder-S: a comprehensive suite to discover and explore intron retention. *Genome Biol.* 22, 307. <https://doi.org/10.1186/s13059-021-02515-8>.
 41. Chang, Y.F., Imam, J.S., and Wilkinson, M.F. (2007). The nonsense-mediated decay RNA surveillance pathway. *Annu. Rev. Biochem.* 76, 51–74. <https://doi.org/10.1146/annurev.biochem.76.050106.093909>.
 42. Isken, O., and Maquat, L.E. (2008). The multiple lives of NMD factors: balancing roles in gene and genome regulation. *Nat. Rev. Genet.* 9, 699–712. <https://doi.org/10.1038/nrg2402>.
 43. Tani, H., and Akimitsu, N. (2012). Genome-wide technology for determining RNA stability in mammalian cells: historical perspective and recent advantages based on modified nucleotide labeling. *RNA Biol.* 9, 1233–1238. <https://doi.org/10.4161/ma.22036>.
 44. Holden, S., and Raymond, F.L. (2003). The human gene CXorf17 encodes a member of a novel family of putative transmembrane proteins: cDNA cloning and characterization of CXorf17 and its mouse ortholog orf34. *Gene* 318, 149–161. [https://doi.org/10.1016/s0378-1119\(03\)00770-4](https://doi.org/10.1016/s0378-1119(03)00770-4).
 45. Tanaka, M., Sasaki, K., Kamata, R., Hoshino, Y., Yanagihara, K., and Sakai, R. (2009). A novel RNA-binding protein, Ossa/C9orf10, regulates activity of Src kinases to protect cells from oxidative stress-induced apoptosis. *Mol. Cell. Biol.* 29, 402–413. <https://doi.org/10.1128/MCB.01035-08>.
 46. Ji, X., Zhou, Y., Pandit, S., Huang, J., Li, H., Lin, C.Y., Xiao, R., Burge, C.B., and Fu, X.D. (2013). SR proteins collaborate with 7SK and promoter-associated nascent RNA to release paused polymerase. *Cell* 153, 855–868. <https://doi.org/10.1016/j.cell.2013.04.028>.
 47. Lai, M.C., Teh, B.H., and Tarn, W.Y. (1999). A human papillomavirus E2 transcriptional activator. The interactions with cellular splicing factors and potential function in pre-mRNA processing. *J. Biol. Chem.* 274, 11832–11841. <https://doi.org/10.1074/jbc.274.17.11832>.
 48. Pandit, S., Wang, D., and Fu, X.D. (2008). Functional integration of transcriptional and RNA processing machineries. *Curr. Opin. Cell Biol.* 20, 260–265. <https://doi.org/10.1016/j.ceb.2008.03.001>.
 49. Horton, J.D., Goldstein, J.L., and Brown, M.S. (2002). SREBPs: activators of the complete program of cholesterol and fatty acid synthesis in the liver. *J. Clin. Invest.* 109, 1125–1131. <https://doi.org/10.1172/JCI15593>.
 50. Adams, C.M., Reitz, J., De Brabander, J.K., Feramisco, J.D., Li, L., Brown, M.S., and Goldstein, J.L. (2004). Cholesterol and 25-hydroxycholesterol inhibit activation of SREBPs by different mechanisms, both involving SCAP and Insigs. *J. Biol. Chem.* 279, 52772–52780. <https://doi.org/10.1074/jbc.M410302200>.
 51. Kamphorst, J.J., Cross, J.R., Fan, J., de Stanchina, E., Mathew, R., White, E.P., Thompson, C.B., and Rabinowitz, J.D. (2013). Hypoxic and Ras-transformed cells support growth by scavenging unsaturated fatty acids from lysophospholipids. *Proc. Natl. Acad. Sci. USA* 110, 8882–8887. <https://doi.org/10.1073/pnas.1307237110>.
 52. Menendez, J.A., and Lupu, R. (2007). Fatty acid synthase and the lipogenic phenotype in cancer pathogenesis. *Nat. Rev. Cancer* 7, 763–777. <https://doi.org/10.1038/nrc2222>.
 53. Griffiths, B., Lewis, C.A., Bensaad, K., Ros, S., Zhang, Q., Ferber, E.C., Konisti, S., Peck, B., Miess, H., East, P., et al. (2013). Sterol regulatory element binding protein-dependent regulation of lipid synthesis supports cell survival and tumor growth. *Cancer Metab.* 1, 3. <https://doi.org/10.1186/2049-3002-1-3>.
 54. Lee, Y., and Rio, D.C. (2015). Mechanisms and regulation of alternative pre-mRNA splicing. *Annu. Rev. Biochem.* 84, 291–323. <https://doi.org/10.1146/annurev-biochem-060614-034316>.
 55. Fu, X.D., and Ares, M. (2014). Context-dependent control of alternative splicing by RNA-binding proteins. *Nat. Rev. Genet.* 15, 689–701. <https://doi.org/10.1038/nrg3778>.
 56. Xiao, S.H., and Manley, J.L. (1998). Phosphorylation-dephosphorylation differentially affects activities of splicing factor ASF/SF2. *EMBO J.* 17, 6359–6367. <https://doi.org/10.1093/emboj/17.21.6359>.
 57. Kwak, H., and Lis, J.T. (2013). Control of transcriptional elongation. *Annu. Rev. Genet.* 47, 483–508. <https://doi.org/10.1146/annurev-genet-110711-155440>.
 58. Zhou, Q., Li, T., and Price, D.H. (2012). RNA polymerase II elongation control. *Annu. Rev. Biochem.* 81, 119–143. <https://www.annualreviews.org/doi/10.1146/annurev-biochem-052610-095910>.
 59. Saunders, A., Core, L.J., and Lis, J.T. (2006). Breaking barriers to transcription elongation. *Nat. Rev. Mol. Cell Biol.* 7, 557–567. <https://doi.org/10.1038/nrm1981>.
 60. Lam, H.C., Siroky, B.J., and Henske, E.P. (2018). Renal disease in tuberous sclerosis complex: pathogenesis and therapy. *Nat. Rev. Nephrol.* 14, 704–716. <https://doi.org/10.1038/s41581-018-0059-6>.
 61. Mossmann, D., Park, S., and Hall, M.N. (2018). mTOR signalling and cellular metabolism are mutual determinants in cancer. *Nat. Rev. Cancer* 18, 744–757. <https://doi.org/10.1038/s41568-018-0074-8>.
 62. Soefje, S.A., Karnad, A., and Brenner, A.J. (2011). Common toxicities of mammalian target of rapamycin inhibitors. *Target. Oncol.* 6, 125–129. <https://doi.org/10.1007/s11523-011-0174-9>.
 63. Lee, S.C.-W., and Abdel-Wahab, O. (2016). Therapeutic targeting of splicing in cancer. *Nat. Med.* 22, 976–986. <https://doi.org/10.1038/nm.4165>.
 64. Fujita, K.I., Ishizuka, T., Mitsukawa, M., Kurata, M., and Masuda, S. (2020). Regulating divergent transcriptomes through mRNA splicing and its modulation using various small compounds. *Int. J. Mol. Sci.* 21, E2026. <https://doi.org/10.3390/ijms21062026>.
 65. Hong, D.S., Kurzrock, R., Naing, A., Wheler, J.J., Falchook, G.S., Schiffman, J.S., Faulkner, N., Pilat, M.J., O'Brien, J., and LoRusso, P. (2014). A phase I, open-label, single-arm, dose-escalation study of

- E7107, a precursor messenger ribonucleic acid (pre-mRNA) spliceosome inhibitor administered intravenously on days 1 and 8 every 21 days to patients with solid tumors. *Invest. New Drugs* 32, 436–444. <https://doi.org/10.1007/s10637-013-0046-5>.
66. Falchook, G., Infante, J., Arkenau, H.T., Patel, M.R., Dean, E., Borazanci, E., Brenner, A., Cook, N., Lopez, J., Pant, S., et al. (2021). First-in-human study of the safety, pharmacokinetics, and pharmacodynamics of first-in-class fatty acid synthase inhibitor TVB-2640 alone and with a taxane in advanced tumors. *eClinicalMedicine* 34, 100797. <https://doi.org/10.1016/j.eclinm.2021.100797>.
67. Koundouros, N., and Poulgiannis, G. (2020). Reprogramming of fatty acid metabolism in cancer. *Br. J. Cancer* 122, 4–22. <https://doi.org/10.1038/s41416-019-0650-z>.
68. Fhu, C.W., and Ali, A. (2020). Fatty acid synthase: an emerging target in cancer. *Molecules* 25, 3935. <https://doi.org/10.3390/molecules25173935>.
69. Roberts, T.C., Langer, R., and Wood, M.J.A. (2020). Advances in oligonucleotide drug delivery. *Nat. Rev. Drug Discov.* 19, 673–694. <https://doi.org/10.1038/s41573-020-0075-7>.
70. Wang, L., Rowe, R.G., Jaimes, A., Yu, C., Nam, Y., Pearson, D.S., Zhang, J., Xie, X., Marion, W., Heffron, G.J., et al. (2018). Small-molecule inhibitors disrupt let-7 oligouridylation and release the selective blockade of let-7 processing by LIN28. *Cell Rep.* 23, 3091–3101. <https://doi.org/10.1016/j.celrep.2018.04.116>.
71. Ghidini, A., Cléry, A., Halloy, F., Allain, F.H.T., and Hall, J. (2021). RNA-PROTACs: degraders of RNA-binding proteins. *Angew. Chem. Int. Ed. Engl.* 60, 3163–3169. <https://doi.org/10.1002/anie.202012330>.
72. Schoggins, J.W. (2019). Interferon-stimulated genes: What do they all do? *Annu. Rev. Virol.* 6, 567–584. <https://www.annualreviews.org/doi/10.1146/annurev-virology-092818-015756>.
73. Wang, W., Xu, L., Su, J., Peppelenbosch, M.P., and Pan, Q. (2017). Transcriptional regulation of antiviral interferon-stimulated genes. *Trends Microbiol.* 25, 573–584. <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7127685/>.
74. Kobayashi, K.S., and van den Elsen, P.J. (2012). NLRC5: a key regulator of MHC class I-dependent immune responses. *Nat. Rev. Immunol.* 12, 813–820. <https://www.nature.com/articles/nri3339>.
75. Benci, J.L., Xu, B., Qiu, Y., Wu, T.J., Dada, H., Twyman-Saint Victor, C., Cucolo, L., Lee, D.S.M., Pauken, K.E., Huang, A.C., et al. (2016). Tumor interferon signaling regulates a multigenic resistance program to immune checkpoint blockade. *Cell* 167, 1540–1554.e12. <https://doi.org/10.1016/j.cell.2016.11.022>.
76. Minn, A.J., and Wherry, E.J. (2016). Combination cancer therapies with immune checkpoint blockade: convergence on interferon signaling. *Cell* 165, 272–275. <https://doi.org/10.1016/j.cell.2016.03.031>.
77. López-Soto, A., Gonzalez, S., and Folgueras, A.R. (2017). IFN signaling and ICB resistance: time is on tumor's side. *Trends Cancer* 3, 161–163. <https://doi.org/10.1016/j.trecan.2017.01.004>.
78. Fregoso, O.I., Das, S., Akerman, M., and Krainer, A.R. (2013). Splicing-factor oncoprotein SRSF1 stabilizes p53 via RPL5 and induces cellular senescence. *Mol. Cell* 50, 56–66. <https://doi.org/10.1016/j.molcel.2013.02.001>.
79. Kim, D., Paggi, J.M., Park, C., Bennett, C., and Salzberg, S.L. (2019). Graph-based genome alignment and genotyping with HISAT2 and HISAT-genotype. *Nat. Biotechnol.* 37, 907–915. <https://doi.org/10.1038/s41587-019-0201-4>.
80. Kovaka, S., Zimin, A.V., Pertea, G.M., Razaghi, R., Salzberg, S.L., and Pertea, M. (2019). Transcriptome assembly from long-read RNA-seq alignments with StringTie2. *Genome Biol.* 20, 278. <https://doi.org/10.1186/s13059-019-1910-1>.
81. Love, M.I., Huber, W., and Anders, S. (2014). Moderated estimation of fold change and dispersion for RNA-seq data with DESeq2. *Genome Biol.* 15, 550. <https://doi.org/10.1186/s13059-014-0550-8>.
82. Blighe, K. (2022). EnhancedVolcano: publication-ready volcano plots with enhanced colouring and labeling. (GitHub). <https://bioc.ism.ac.jp/packages/3.14/bioc/manuals/EnhancedVolcano/man/EnhancedVolcano.pdf>.
83. Kuleshov, M.V., Jones, M.R., Rouillard, A.D., Fernandez, N.F., Duan, Q., Wang, Z., Koplev, S., Jenkins, S.L., Jagodnik, K.M., Lachmann, A., et al. (2016). Enrichr: a comprehensive gene set enrichment analysis web server 2016 update. *Nucleic Acids Res.* 44, W90–W97. <https://doi.org/10.1093/nar/gkw377>.
84. Agrawal, S., Kumar, S., Sehgal, R., George, S., Gupta, R., Poddar, S., Jha, A., and Pathak, S. (2019). El-MAVEN: a fast, robust, and user-friendly mass spectrometry data processing engine for metabolomics. In *High-Throughput Metabolomics: Methods and Protocols* (Methods in Molecular Biology, A. D'Alessandro, ed. (Springer), pp. 301–321. https://doi.org/10.1007/978-1-4939-9236-2_19.
85. Mauger, O., Lemoine, F., and Scheiffele, P. (2016). Targeted intron retention and excision for rapid gene regulation in response to neuronal activity. *Neuron* 92, 1266–1278. <https://doi.org/10.1016/j.neuron.2016.11.032>.
86. Kamburov, A., Wierling, C., Lehrach, H., and Herwig, R. (2009). ConsensusPathDB—a database for integrating human functional interaction networks. *Nucleic Acids Res.* 37, D623–D628. <https://doi.org/10.1093/nar/gkn698>.

STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Mouse monoclonal anti-SRPK2	BD Biosciences	Cat#BD611118; RRID: AB_398429
Rabbit polyclonal anti-ACLY	Cell Signaling Technology	Cat#4332; RRID: AB_2223744
Rabbit monoclonal anti-FASN	Cell Signaling Technology	Cat#3180; RRID: AB_2100796
Mouse monoclonal anti-S6	Cell Signaling Technology	Cat#2317; RRID: AB_2238583
Rabbit monoclonal anti-pS6(S240/S244)	Cell Signaling Technology	Cat#5364; RRID: AB_10694233
Rabbit polyclonal anti-SCD1	Cell Signaling Technology	Cat#2438; RRID: AB_823634
Rabbit polyclonal anti-CHREBP	Novus	Cat# NB400-135;RRID:AB_10002435
Rabbit polyclonal anti-FAM120A	GeneTex	Cat#GTX120824; RRID:AB_10723014
Rabbit polyclonal anti-FAM120A	Thermo Fisher Scientific	Cat# A303-889A; RRID:AB_2620239
Rabbit polyclonal anti-LXR alpha	Thermo Fisher Scientific	Cat#PA1-330;RRID:AB_325600
Mouse monoclonal anti-SREBP2	BD Biosciences	Cat#557037; RRID:AB_396560
Rabbit polyclonal anti-LXR beta	Thermo Fisher Scientific	Cat#PA1-333;RRID:AB_2154770
Rabbit polyclonal anti-PPAR alpha	Thermo Fisher Scientific	Cat#PA1-822A; RRID:AB_2165595
Rabbit polyclonal anti-PPAR gamma	Abcam	Cat#ab209350; RRID:AB_2890099
Rabbit monoclonal anti-p300	Cell Signaling Technology	Cat#54062; RRID:AB_2799450
Rabbit monoclonal anti-XBP-1s	Cell Signaling Technology	Cat#12782;RRID:AB_2687943
Mouse monoclonal anti-GAPDH	Sigma-Aldrich	Cat#G8795; RRID: AB_1078991
Rabbit polyclonal anti-ACSS2	Sigma-Aldrich	Cat#HPA004141; RRID: AB_1078094
Mouse monoclonal anti-V5	Sigma-Aldrich	Cat#V8012; RRID:AB_261888
Mouse monoclonal anti-Vinculin	Sigma-Aldrich	Cat#V9264; RRID: AB_10603627
Rabbit polyclonal anti-RNA polymerase II	Abcam	Cat#ab26721; RRID:AB_777726
Rabbit monoclonal anti-FDFT1	Abcam	Cat#ab109723; RRID: AB_10859772
Mouse monoclonal anti-U1-70k	Santa Cruz Biotechnology	Cat#sc-390899; RRID:AB_2801569
Rabbit polyclonal anti-SREBP1	Santa Cruz Biotechnology	Cat#sc-8984; RRID: AB_2194223
Mouse monoclonal anti-SRSF1	Santa Cruz Biotechnology	Cat#sc-33652; RRID: AB_628248
Mouse monoclonal anti-T7	EMD Millipore	Cat#69522; RRID:AB_11211744
Mouse monoclonal anti-Phosphoepitope SR proteins	EMD Millipore	Cat#MABE50;RRID:AB_10807429
Rabbit Normal IgG	Cell Signaling Technology	Cat#2729, RRID:AB_1031062
Mouse Normal IgG	Thermo Scientific	Cat#10-400-C; RRID:AB_2532980
IRDye 800CW Donkey anti-Rabbit IgG	LI-COR Biosciences	Cat# 926-32213; RRID:AB_621848
IRDye 680RD Donkey anti-Mouse IgG	LI-COR Biosciences	Cat# 926-68072; RRID:AB_10953628
IRDye 680RD Detection Reagent	LI-COR Biosciences	Cat#:926-69100
Bacterial and virus strains		
<i>E. coli</i> BL21	New England Biolabs	Cat#C2527
Chemicals, peptides, and recombinant proteins		
Rapamycin	Calbiochem	Cat#553210
Torin1	Tocris Bioscience	Cat#4247
Actinomycin D	Sigma-Aldrich	Cat#A1410
PhosSTOP	Roche	Cat#04906837001
RNase inhibitor	Invitrogen	Cat#10777019
RNase T1	Thermo Fisher Scientific	Cat#AM2283
RNase A	Thermo Fisher Scientific	Cat#12091-021
Anti-V5 agarose affinity gel	Sigma-Aldrich	Cat#A7345

(Continued on next page)

Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Anti-Flag agarose affinity gel	Sigma-Aldrich	Cat# A2220
Lipofectamine™ RNAiMAX Transfection Reagent	Invitrogen	Cat#13778075
DNase I	Sigma-Aldrich	Cat#AMPD1
TRIzol	Invitrogen	Cat#15596018
Protein A/G magnetic beads	Thermo Fisher Scientific	Cat#88802
Protein A/G PLUS-Agarose	Santa Cruz Biotechnology	Cat#sc-2003
Proteinase K	New England Biolabs	Cat#P8107
Lipofectamine 2000 reagent	Invitrogen	Cat#11668500
Phenol:Chloroform:IAA, 25:24:1, pH 6.6	Thermo Fisher Scientific	Cat#AM9730
Insulin solution human	Sigma-Aldrich	Cat#I9278
Crystal Violet	Sigma-Aldrich	Cat#C3886
SRPIN340	Selleck Chemicals	Cat#S7270
SRPKIN-1	MedChem Express	Cat#HY-116856
Lipoprotein, low density from human plasma	Sigma-Aldrich	Cat#L7914
Oleic acid-albumin	Sigma-Aldrich	Cat#O3008
Fatty acid-free albumin	Sigma-Aldrich	Cat#A8806
Lipoprotein-deficient serum	Alfa Aesar	Cat#BT-907
Sodium palmitate	Sigma-Aldrich	Cat#P9767
Methanol-free 10% formaldehyde	Polysciences	Cat#04018-1
25-Hydroxycholesterol	MedChem Express	Cat#HY-113134
Critical commercial assays		
PureLink RNA Mini kit	Ambion	Cat#12183018A
iScript cDNA synthesis kit	Bio-rad	Cat#170-8891BUN
Illumina® Stranded mRNA Prep Kit	Illumina	Cat#20040534
Gateway LR clonase II enzyme mix	Invitrogen	Cat#11791
QuickChange site-directed mutagenesis kit	Stratagene	Cat#200521
RNA Clean & Concentrator	Zymo Research	Cat#R1016
Illumina® Stranded Total RNA Prep, Ligation with Ribo-Zero Plus	Illumina	Cat#20040529
LightSwitch dual assay kit (promoter assay)	Active Motif	Cat#32035
Chromatin IP DNA purification kit	Active Motif	Cat#58002
SYBR Green PCR Master Mix	Life Technologies	Cat#4312704
QuikChange II XL Site-Directed Mutagenesis Kit	Stratagene	Cat#200521
ChIP-IT Express Kit	Active Motif	Cat#53009
Deposited data		
Raw RNAseq data (siNTC and siFAM120A)	This paper	GSE207172
Raw RNAseq data (siNTC, siFAM120, and siSRSF1)	This paper	GSE229657
Raw data of Images and WB	This paper, Mendeley Data	https://doi.org/10.17632/rfgfx9xb8w.1
Experimental models: Cell lines		
Human: Renal angiomyolipoma-derived LAM 621-101 (TSC2 ^{-/-})	Drs. Jane Yu and Elizabeth Henske	Yu et al. ³³ ; RRID: CVCL_S879
Human: HEK293E	ATCC	Cat#293 c18; RRID: CVCL_6974
Human: HEK293T	GenHunter	Cat#Q401; RRID: CVCL_0063
Human: MCF7	ATCC	Cat#HTB-22; RRID: CVCL_0031
Human: DLD1	ATCC	Cat#CCL-221; RRID: CVCL_0248
Human: NCI-H1299	ATCC	Cat#CRL-5803; RRID: CVCL_0060
Human: LNCaP	ATCC	Cat#CRL-1740; RRID:CVCL_1379
Experimental models: Organisms/strains		
nu/nu athymic mice	Envigo	Cat#069

(Continued on next page)

Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Oligonucleotides		
siRNA negative control-#1(siNTC)	Sigma-Aldrich	SIC001
siRNA negative control-#2(siNTC)	Sigma-Aldrich	SIC002
siRNA (Human SRSF1)	Sigma-Aldrich	SASI_Hs01_00115059
siRNA (Human SRSF1)	Sigma-Aldrich	SASI_Hs02_00342609
siRNA (Human UPF1)	Sigma-Aldrich	SASI_Hs01_00101017
siRNA (Human UPF1)	Sigma-Aldrich	SASI_Hs01_00101018
siRNA (Human U1-70k/SNRNP70)	Sigma-Aldrich	SASI_Hs02_00335028
siRNA (Human U1-70k/SNRNP70)	Sigma-Aldrich	SASI_Hs01_00165101
siRNA (Human FAM120A)	Sigma-Aldrich	SASI_Hs01_00149752
siRNA (Human FAM120A)	Sigma-Aldrich	SASI_Hs02_00345898
siRNA (Human FAM120A)	Sigma-Aldrich	SASI_Hs02_00345899
siRNA (Human SREBP1)	Sigma-Aldrich	SASI_Hs01_00051828
siRNA (Human SREBP1)	Sigma-Aldrich	SASI_Hs01_00051829
siRNA (Human SREBP1)	Sigma-Aldrich	SASI_Hs01_00051830
siRNA (Human SREBP2)	Sigma-Aldrich	SASI_Hs01_00075424
siRNA (Human SREBP2)	Sigma-Aldrich	SASI_Hs01_00075425
siRNA (Human SREBP2)	Sigma-Aldrich	SASI_Hs01_00075426
Primers for qPCR analysis, see Table S2	This paper	N/A
Recombinant DNA		
ACSS2 promoter construct	SwitchGear Genomics	Cat#S715263
SCD1 promoter construct	SwitchGear Genomics	Cat#S714484
pLX304-SRSF1-V5	Harvard plasmid	Cat#HsCD00420441
Lentiviral packaging and envelope plasmids	Dr. David Baltimore	N/A
pcDNA3-EGFP	Addgene	Cat#13031
pKH3-HA-SRPK2-WT, K110M	Dr. John Blenis	Lee et al. ¹²
pLNCX-HA-SRPK2-S494A/S497A	Dr. John Blenis	Lee et al. ¹²
pLX304-FAM120A-V5	Harvard plasmid	HsCD00440255
pENTR1A-Flag-FAM120A(WT)	This paper	N/A
pENTR1A-Flag-FAM120A(Δ RBBD)	This paper	N/A
pLenti-CMV-Puro-DEST	Addgene	Cat#17452
pENTR223-SRSF1	Harvard plasmid	Cat#HsCD00367645
pLenti-CMV-Puro-SRSF1	This paper	N/A
pLenti-CMV-Puro-FAM120A	This paper	N/A
pLenti-CMV-Puro-Flag-FAM120A(WT)	This paper	N/A
pLenti-CMV-Puro-Flag-FAM120A(Δ RBBD)	This paper	N/A
pENTR223-FAM120A	Harvard plasmid	Cat#HsCD00438604
T7-SRSF1(WT)	Dr. Adrian Krainer	Fregoso et al. ⁷⁸
T7-SRSF1(Δ RRM1)	Dr. Adrian Krainer	Fregoso et al. ⁷⁸
T7-SRSF1(Δ RRM2)	Dr. Adrian Krainer	Fregoso et al. ⁷⁸
T7-SRSF1(Δ RS)	Dr. Adrian Krainer	Fregoso et al. ⁷⁸
pLKO.1-puro-Non-Mammalian shRNA Control	Sigma-Aldrich	SHC002
pLKO.1-puro-shSRPK2	Sigma-Aldrich	TRCN0000006274
pLKO.1-puro-shFAM120A	Sigma-Aldrich	TRCN0000074918
pLKO.1-puro-shSREBP1	Sigma-Aldrich	TRCN0000055325
pLKO.1-puro-shSREBP2	Sigma-Aldrich	TRCN0000020667
Software and algorithms		
IRFinder-S	Lorenzi et al. ⁴⁰	https://europepmc.org/article/med/34749764

(Continued on next page)

Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
HISAT2 v2.2.1	Kim et al. ⁷⁹	https://www.nature.com/articles/s41587-019-0201-4
StringTie2 v2.1.4	Kovaka et al. ⁸⁰	https://genomebiology.biomedcentral.com/articles/10.1186/s13059-019-1910-1
DESeq2 (v1.30.1)	Love et al. ⁸¹	https://genomebiology.biomedcentral.com/articles/10.1186/s13059-014-0550-8
Enhanced Volcano v1.8.0 R package	Blighe ⁸²	https://bioconductor.org/packages/devel/bioc/vignettes/EnhancedVolcano/inst/doc/EnhancedVolcano.html
Adobe Photoshop	Adobe	Adobe
Odyssey imaging system	LI-COR Biosciences	LI-COR Biosciences
enrichR v3.0 package	Kuleshov et al. ⁸³	https://git.ecdf.ed.ac.uk/sgrannem/pycrac
EI-MAVEN, Version 0.12.0	Agrawal et al. ⁸⁴	Elucidata
VAST-tools	Irimia et al. ³⁹	https://www.sciencedirect.com/science/article/pii/S0092867414015128?via%3Dihub
BioRender	BioRender	https://www.biorender.com/

RESOURCE AVAILABILITY**Lead contact**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, John Blenis (job2064@med.cornell.edu).

Materials availability

All reagents generated in this study are available from the [lead contact](#) without restriction.

Data and code availability

- Sequencing data have been deposited at GEO with identifier GSE207172 and GSE229657. The original images have been deposited at Mendeley Data (<https://doi.org/10.17632/rgfgx9xb8w.1>). All data are publicly available as of the date of publication. Accession numbers and DOI are listed in the [key resources table](#).
- This paper does not report any original code.
- Any additional information required to reanalyze the data reported in this paper is available from the [lead contact](#) upon request.

EXPERIMENTAL MODEL AND STUDY PARTICIPANT DETAILS

Details of cell lines used have been provided under section “[cell lines and cell culture](#)” and details of mice used have been provided under “[mouse studies](#).”

Cell lines and cell culture

HEK293E and HEK293T cells were obtained from ATCC and GenHunter, respectively. Human renal angiomyolipoma-derived LAM 621-101 cell line (referred to as LAM cells in the manuscript) was provided by Dr. Henske (Harvard Medical School).³³ H1299, DLD1, MCF7, and LNCaP cells were obtained from ATCC. Cells were grown in DMEM with 10% FBS at 37°C with 5% CO₂ unless otherwise indicated.

Mouse studies

All animal procedures were approved by the Institutional Animal Care and Use Committee (IACUC) at Weill Cornell Medicine. 6-week-old female athymic nu/nu mice were purchased from Envigo. 1x10⁶ cells resuspended in 100 μl of PBS (phosphate-buffered saline): Growth Factor Reduced Matrigel (Corning) mixture (1:1) were subcutaneously inoculated into the posterior back regions of the mouse. Once the tumor is formed two to three weeks post-inoculation, tumor length and width were measured 3 times a week using a digital caliper. Tumor volume was calculated using the formula: Tumor volume = (Length × Width²)/2.

METHOD DETAILS

Expression constructs

Expression constructs were transfected overnight using Lipofectamine 2000 (Invitrogen). pENTR223-FAM120A and pENTR223-SRSF1 were obtained from Harvard PlasmID. pENTR223-FAM120A and pENTR223-SRSF1 recombined with plx304-V5-DEST or pLenti-puromycin-DEST vector by LR reaction (Gateway LR clonase II enzyme mix, Invitrogen) to generate expression constructs. To generate FAM120A RNA deletion mutant construct, the coding sequence of human FAM120A with or without RNA binding domain (NM_014612.5) was cloned into pENTR1A vector (NotI and Sall). The SRSF1 WT and mutant constructs were a gift from Dr. Krainer (Cold Spring Harbor Laboratory).⁷⁸

siRNA oligonucleotides

30 nM siRNA (Sigma-Aldrich; see list in [key resources table](#)) was transfected using Lipofectamine RNAiMAX reagent (Invitrogen).

Generation of stable cell lines

Lipofectamine 2000 reagent (Invitrogen) was used to co-transfect with the viral plasmid of interest with lentiviral packaging and envelope plasmids in HEK293T cells to generate lentivirus. After 60 hr of transfection, virus-containing supernatant was collected and treated to the cells with 8 $\mu\text{g}/\text{mL}$ polybrene overnight. Cells infected with pLKO.1-puromycin shRNA constructs (Sigma-Aldrich; see list in [key resources table](#)) or pLenti-puromycin infected cells were treated with 2 $\mu\text{g}/\text{mL}$ (H1299, DLD1, MCF7, and LNCaP) or 10 $\mu\text{g}/\text{mL}$ (LAM) puromycin for 2-3 days to select the infected cells. pLenti-plx304-V5 infected cells were selected with 10 $\mu\text{g}/\text{mL}$ (H1299, DLD1, MCF7, and LNCaP) or 30 $\mu\text{g}/\text{mL}$ (LAM) blasticidin. See construct list in [key resources table](#).

Crystal violet staining

4% methanol-free formaldehyde (Polysci) was used to fix the cells for 10 min and incubated with 0.1% crystal violet solution (Sigma-Aldrich) for 20 min at room temperature. The plates were washed with PBS 5 times, and the plates were scanned for image analysis after overnight drying. To quantify the staining intensity, crystal violet was eluted by incubating the cells with methanol for 20 min. Absorbance was measured at 570 nm (Envision plate reader, PerkinElmer).

Immunoblot analysis

After 2 times of cold PBS wash, cells were incubated on ice for 30 min with a RIPA (40 mM HEPES [pH7.4], 1 mM EDTA, 120 mM NaCl, 0.5 mM DTT, 0.1% Brij-35, 0.1% deoxycholate, and 0.5% NP-40) lysis buffer supplemented with protease inhibitors (250 μM PMSF, 5 $\mu\text{g}/\text{mL}$ pepstatin A, 10 $\mu\text{g}/\text{mL}$ leupeptin, and 5 $\mu\text{g}/\text{mL}$ aprotinin) and phosphatase inhibitors (10 mM β -glycerophosphate, 1 mM NaF, and 1 mM Na_3VO_4) to lysis cells. Soluble cell lysates were obtained after centrifugation at 21,130 $\times g$ at 4°C for 30 min to clear the lysates. Protein concentration was measured with detergent compatible (DC) protein assay (Bio-rad) and the same amounts of proteins were boiled for 10 min with Laemmli sample buffer to denature the protein structure. Equal amounts of proteins (15-50 μg) were separated by SDS-PAGE gels and the proteins were transferred to nitrocellulose membranes (Amersham Biosciences). The transferred nitrocellulose membrane was blocked with Odyssey blocking solution (LI-COR Biosciences) for an hour at room temperature, and further incubated with primary and IRDye secondary antibodies (see antibody list on [key resources table](#)). Odyssey imaging system (LI-COR Biosciences) was used to detect and quantify immunoblot signals. Immunoblot images show the representative images of at least two independent experiments.

Quantitative RT-PCR

Total RNA was harvested using PureLink RNA isolation kit (Life Technologies) according to the manufacturer's protocol. DNase I (Sigma-Aldrich) was treated to eliminate genomic DNA. cDNA was synthesized from total RNA by reverse-transcription reaction using iScript (Bio-rad). QuantStudio6 Real-Time PCR system (Life Technologies) was used to analyze the resulting cDNA using SYBR green master mix (Life Technologies). Delta-delta CT method was used to calculate relative mRNA levels by normalization of house-keeping genes ACTIN, GAPDH, PPIB, and TBP as controls. The primer list is in [Table S2](#).

mRNA stability assay

Actinomycin D (5 $\mu\text{g}/\text{mL}$) was treated to cells for the indicated time points. Total RNA was extracted using Purelink RNA miniprep kit and mRNA levels were analyzed by qPCR.

Luciferase promoter activity assay

LightSwitch dual assay system was used according to the manufacturer's protocol (SwitchGear Genomics). Cells plated on a 6-well plate were transfected with siRNAs targeting FAM120A or non-targeting control through Lipofectamine RNAiMax transfection reagent. After 2 days, the cells were co-transfected with 1 μg of renilla construct containing the promoter of the interested gene and 0.2 μg of control cypridina construct using FuGENE HD transfection reagent (Promega). 24 hr after transfection, the activity of renilla and cypridina luciferases was measured (Envision plate reader, PerkinElmer). For calculation of the promoter activities, the renilla luciferase activity was normalized by the cypridina luciferase activity.

RNA immunoprecipitation qPCR

Cells on a 15-cm plate were washed twice with ice-cold PBS and harvested into 15 mL tube. Cells were collected by centrifugation at 1,000 x g at 4 °C for 5 min. Cell pellets were resuspended with 1 ml lysis buffer (1 mM EDTA [pH8.0], 50 mM HEPES-KOH [pH7.5], 140 mM NaCl, 0.1% Sodium deoxycholate, 1% Triton X-100) supplemented with protease inhibitors (250 μM PMSF, 5 μg/mL pepstatin A, 10 μg/mL leupeptin, 5 μg/mL aprotinin) and phosphatase inhibitors (10 mM β-glycerophosphate, 1 mM NaF, 1 mM Na₃VO₄), and RNase inhibitor (160 unit/mL, Invitrogen), and incubated on ice for 30 min. After centrifugation at 21,130 x g at 4°C for 30 min, the supernatant was pre-cleared by incubating with 50 μL protein A/G beads. The pre-cleared lysate was incubated with 5 μg of IgG (Mouse: Thermo Scientific, Rabbit: Cell Signaling Technology), anti-SRSF1, or anti-FAM120A antibody (Bethyl Laboratories) overnight at 4°C. Then, 50 μL protein A/G magnetic beads were added to the lysate and incubated at 4°C for 1.5 hr to pull down antibody-associated complexes. The beads were washed 3 times with lysis buffer and RNA was eluted by incubating with 150 μL lysis buffer containing 1% SDS and 1.2 mg/mL proteinase K (New England Biolabs) at 55°C for 30 min. The RNA elute was purified by phenol/chloroform extraction and precipitated with ammonium acetate and ethanol. Input and immunoprecipitated RNAs were treated with DNase I (Sigma-Aldrich) and reverse-transcribed (iScript cDNA synthesis kit, Bio-rad), and the resulting cDNA was analyzed by qPCR. The amount of transcripts (%) bound to the antibody was calculated: $100 \times 2^{\Delta[Ct(input) - Ct(IP)]}$.

Co-immunoprecipitation of proteins

Cells on a 10-cm plate were harvested through centrifugation at 1,000 x g at 4°C for 5 min after three times washing with ice-cold PBS. One mL of lysis buffer (20 mM Tris HCl [pH8.0], 137 mM NaCl, 1% Nonidet P-40, 2 mM EDTA) supplemented with protease inhibitors (250 μM PMSF, 5 μg/mL pepstatin A, 10 μg/mL leupeptin, 5 μg/mL aprotinin) and phosphatase inhibitors (10 mM β-glycerophosphate, 1 mM NaF, 1 mM Na₃VO₄) was added, and incubated on ice for 30 min to lyse the cells with mechanical disruption by syringe homogenization. After centrifugation at 21,130 x g at 4°C for 30 min, protein concentration was measured, and 1 mg of lysates were incubated with anti-V5 (Sigma-Aldrich) or anti-Flag agarose affinity gels (Sigma-Aldrich) at 4°C for 1.5 h with gentle agitation to pull down immune complexes. For endogenous IP, 1mg of lysates were incubated with 4 μg of anti-FAM120A (Thermo Fisher Scientific) or rabbit IgG (Cell Signaling Technology) for overnight at 4°C h with gentle agitation to pull down immune complexes. 50 μL protein A/G beads were added to the lysate and incubated at 4°C with gentle agitation for 1.5 hr to pull down antibody-associated complexes. The immunoprecipitated beads were then washed 3 times with lysis buffer to remove non-specific binding and the immune complexes were eluted by boiling with 2X Laemmli sample buffer for 5 min. The boiled beads were centrifuged at 845 x g for 5 min and supernatants were subjected to immunoblot analysis.

Ribonuclease treatment

Cells on a 10-cm plate were washed twice with ice-cold PBS and harvested by centrifugation at 1,000 x g at 4°C for 5 min. Cells were lysed and physically disrupted by syringe homogenization in one mL of lysis buffer (20 mM Tris HCl [pH8.0], 137 mM NaCl, 1% Nonidet P-40, 2 mM EDTA) supplemented with protease inhibitors (250 μM PMSF, 5 μg/mL pepstatin A, 10 μg/mL leupeptin, 5 μg/mL aprotinin) and phosphatase inhibitors (10 mM β-glycerophosphate, 1 mM NaF, 1 mM Na₃VO₄), and incubated on ice for 30 minutes. After centrifugation at 21,130 x g at 4°C for 30 min, the clarified lysate was treated with a mixture of RNase A (100 μg/ml) (Invitrogen) and RNase T1 (1 unit/μL) (Invitrogen) for 1 hr at room temperature prior to co-IP analysis.

Chromatin immunoprecipitation qPCR

Cells were incubated with 1% formaldehyde to fix the cells at room temperature for 10 min followed by quenching with 0.125 M glycine for 5 min at room temperature. ChIP assays were performed using a commercially available kit (ChIP-IT Express, Active Motif). First, lysis buffer was used to lyse the cells using a Dounce homogenizer to disrupt cells for harvesting chromatin. Isolated chromatin was sheared by an enzyme for 15 min at 37°C (ChIP-IT Express Enzymatic Shearing Kit, Active Motif). Immunoprecipitation was performed using anti-SRSF1 (32-4500, Invitrogen) or anti-SREBP1 (sc-13551, Santa Cruz) antibodies. Elution buffer containing SDS buffer was used to elute complex from the beads and incubated with RNase and proteinase K after several washes. Reverse crosslinking was performed by incubation overnight at 65°C, and a Chromatin IP DNA purification kit (Active Motif) was used to purify ChIP DNA. QuantStudio6 Real-Time PCR system (Life Technologies) was used to analyze the resulting DNA samples by qPCR. The amount of DNA (%) bound to the antibody was calculated: $100 \times 2^{\Delta[Ct(input) - Ct(IP)]}$.

Lipid supplementation

To generate albumin conjugated palmitate, 10 mL of sodium palmitate (10 mM) was mixed with 10 mL of fatty acid-free BSA (1.66 mM BSA solution) and incubated for 1 hr at 70°C. Cells stably expressing shRNAs targeting FAM120A or control were cultured in DMEM with 10% lipoprotein-deficient serum (LPDS) supplemented with lipoprotein (25 μg/mL), oleate-albumin (50 μM; oleate), palmitate-albumin (10 μM; palmitate), or fatty acid-free albumin (25 μM; control) to the media.

Mass spectrometry analysis of SRSF1 interactome

We previously reported the SRSF1 interactome using reductive dimethylation isotopic labeling to provide quantitative comparisons between treated (torin1) and untreated cells.¹² Those data were reanalyzed at the individual peptide level for the bait protein, SRSF1-V5, and FAM120A. Data were manually extracted from each of two replicate analyses for all quantified peptides.

A stringent signal-to-noise (SN) filter was applied, retaining only peptide quantifications where the sum SN of treated and untreated samples was >100. Figure 1A shows the $\log_2[+TORIN/DMSO]$ for all passing peptides for each protein. Statistical analysis was performed in Prism using unpaired t-tests.

Mass spectrometry analysis of saponified fatty acids

90% confluent cells on 60 mm cell culture dish were incubated with 0.5 ml of 0.3 M KOH in 90% methanol at 80°C for 1 hr in a 2 mL glass vial. Then, formic acid (100 μ L) was added for neutralization. The saponified fatty acids were extracted by adding 1 mL of hexane. After vortexing, the top hexane layer was transferred to a new glass vial. Samples were then dried under a nitrogen gas stream and redissolved in 100 μ L of 1:1 isopropanol:methanol for LC-MS analysis. Fatty acids were detected with a quadrupole orbitrap mass spectrometer (Thermo Fisher Q-Exactive) operating in a negative ion mode with electrospray ionization and used to scan from m/z 200 to 600 at 1 Hz, with a 140,000 resolution. LC separation was achieved on a Poroshell 120 EC-C18 column (2.1 x 150 mm², 2.7 μ m particle size; Agilent) using a gradient of solvent A (90:10 = water: methanol with 1 mM of ammonium acetate and 0.2% acetic acid) and solvent B (90:10 = methanol:isopropanol with 1 mM of ammonium acetate and 0.2% acetic acid). The LC gradient was 0 min, 25% B; 2 min, 25% B; 4 min, 65% B; 16 min, 100% B; 20 min, 100% B; 21 min, 25% B; 22 min, 25% B; and 25 min, 25% B. Flow rate was 150 μ L/min. The autosampler temperature was 5°C and the injection volume was 3 μ L. Mass spectrometry data was analyzed using the EI-MAVEN open source software⁸⁴ (Version 0.12.0, Elucidata). Briefly, LC-MS data was converted from raw files to.mzXML and then loaded into the EI-MAVEN software. Using our annotated MS library, containing retention times and m/z values, we selected metabolite peaks of interest on EI-MAVEN. Peak shape and quality were evaluated and Area Top of the metabolites of interest were recorded.

RNA-seq sample preparation

For GSE207172, we used 100 ng of high-quality total RNA and the Illumina® TruSeq Stranded Total RNA with Ribo-Zero Globin kit (Cat# 20040529) from Illumina. For GSE229657, the Illumina® Stranded mRNA Prep Kit (Cat# 20040534) and 100 ng of high-quality total RNA were used. RNA-seq library was prepared by Weill Cornell Medicine Core facility for the Illumina NovaSeq 6000 system sequencer at paired-end 2x100 cycles.

RNA-seq and pathway enrichment analyses

RNA-seq in Figure 2 was performed by using the Illumina NovaSeq 6000 system with 40M 2x101 bp paired-end (PE) read. Using the HISAT2 v2.2.1 software,⁷⁹ the sequencing reads were mapped onto the human GRCh38/hg38 reference genome, from which the counts matrix was generated with the default parameters using StringTie2 v2.1.4.⁸⁰ Differentially expressed genes (DEGs) were identified using the DESeq2 (v1.30.1)⁸¹ based on a negative binomial generalized linear model. The data visualization was performed by using the EnhancedVolcano v1.8.0 R package.⁸² DEGs that satisfy $|(Log_2FC)| > 1$ and adjusted p -value (q -value) < 0.005 were considered as statistically significant. Pathway enrichment was measured by Fisher's exact test against Elsevier pathway collection using the enrichR v3.0 package.⁸³ RNA-seq data files are accessible through Gene Expression Omnibus (GEO) accession number GSE207172.

In order to detect intron retention events more precisely, we did deep RNA sequencing in Figure 3 using the Illumina NovaSeq 6000 system with 100M 2x101 bp PE read. Analysis of deep RNA-seq was performed as previously described.⁸⁵ In brief, the quality of RNA-seq reads were checked using FastQC and the reads were aligned to the human GRCh38/hg38 reference genome using STAR (2.7.10b) with following parameters: `-outFilterMismatchNmax 2--outFilterMultimapNmax 10 -outSJfilterOverhangMin 30 888 -outSAMunmapped Within KeepPairs`. Genome coverage files (bigwigs) were generated using `bam2wig.py` from RSeQC (v5.0.1) with following parameter: `"1+-,1-+,2++,2--" -t 1000000000`. Differential gene expression analysis was done by extracting counts using `featureCounts` (v2.0.3) and running DESeq2 (v1.34.0) to find differentially expressed genes with the cutoff of q -value < 0.01 and $|(Log_2FC)| \geq 0.5$. Pathway analyses were performed using ConsensusPathDB⁸⁶ (KEGG, Reactome, Wikipathways, Biocarata, Ehm, Humancyc and Inoh) and only pathways with q -value < 0.05 were considered. RNA-seq data files dataset generated in this study is accessible through Gene Expression Omnibus (GEO) accession number GSE229657.

Intron retention analysis

To examine the global splicing changes, raw fastq files from deep RNA-sequencing data were aligned to VAST-tools library (human vastdb.hs2.23.06.20) using VAST-tools (2.5.1) and splicing events were detected using VAST-tools tidy with following parameters: `-min N 2--p_IR 0.1 -min_SD 1`.³⁹ To precisely determine the intron retention events, we performed IRFinder-S (2.0.1) using BAM files from the STAR alignment step and calculated intron retention events with following parameters: `-wl 1 -ir 0.01`. Significantly increased intron retention events were selected with the cutoff of q -value < 0.1 and $Log_2FC > 0$.⁴⁰

Analysis of published CHIP-seq and whole transcriptome microarray results

Previously published CHIP-seq dataset GSE45517⁴⁶ and whole transcriptome microarray dataset GSE104335¹² were obtained from GEO dataset.

QUANTIFICATION AND STATISTICAL ANALYSIS

LC-MS, qPCR, interactome, immunoblot, and crystal violet staining assays data are statistically analyzed using Student's t-test and the graphs show mean \pm SD. Tumor burdens from the xenograft experiment are statistically analyzed using Student's t-test and the error bars show mean \pm SEM. Detailed methods and p-value are described in the Figure Legends and Methods sections.