

## Benzimidazole derivative small-molecule 991 enhances AMPK activity and glucose uptake induced by AICAR or contraction in skeletal muscle

Bultot, Laurent; Lai, Yu Chiang; Jensen, Thomas E.; Madsen, Agnete L.B.; Collodet, Caterina; Kviklyte, Samanta; Deak, Maria; Yavari, Arash; Foretz, Marc; Ghaffari, Sahar; Bellahcene, Mohamed; Ashrafian, Houman; Rider, Mark H.; Richter, Erik A.; Sakamoto, Kei

DOI:

[10.1152/ajpendo.00237.2016](https://doi.org/10.1152/ajpendo.00237.2016)

License:

Creative Commons: Attribution (CC BY)

*Document Version*

Publisher's PDF, also known as Version of record

*Citation for published version (Harvard):*

Bultot, L, Lai, YC, Jensen, TE, Madsen, ALB, Collodet, C, Kviklyte, S, Deak, M, Yavari, A, Foretz, M, Ghaffari, S, Bellahcene, M, Ashrafian, H, Rider, MH, Richter, EA & Sakamoto, K 2016, 'Benzimidazole derivative small-molecule 991 enhances AMPK activity and glucose uptake induced by AICAR or contraction in skeletal muscle', *American Journal of Physiology - Endocrinology and Metabolism*, vol. 311, no. 4, pp. E706-E719.  
<https://doi.org/10.1152/ajpendo.00237.2016>

[Link to publication on Research at Birmingham portal](#)

### General rights

Unless a licence is specified above, all rights (including copyright and moral rights) in this document are retained by the authors and/or the copyright holders. The express permission of the copyright holder must be obtained for any use of this material other than for purposes permitted by law.

- Users may freely distribute the URL that is used to identify this publication.
- Users may download and/or print one copy of the publication from the University of Birmingham research portal for the purpose of private study or non-commercial research.
- User may use extracts from the document in line with the concept of 'fair dealing' under the Copyright, Designs and Patents Act 1988 (?)
- Users may not further distribute the material nor use it for the purposes of commercial gain.

Where a licence is displayed above, please note the terms and conditions of the licence govern your use of this document.

When citing, please reference the published version.

### Take down policy

While the University of Birmingham exercises care and attention in making items available there are rare occasions when an item has been uploaded in error or has been deemed to be commercially or otherwise sensitive.

If you believe that this is the case for this document, please contact [UBIRA@lists.bham.ac.uk](mailto:UBIRA@lists.bham.ac.uk) providing details and we will remove access to the work immediately and investigate.

Download date: 23. Apr. 2024

## Benzimidazole derivative small-molecule 991 enhances AMPK activity and glucose uptake induced by AICAR or contraction in skeletal muscle

Laurent Bultot,<sup>1</sup> Thomas E. Jensen,<sup>3</sup> Yu-Chiang Lai,<sup>2</sup> Agnete L. B. Madsen,<sup>3</sup> Caterina Collodet,<sup>1,4</sup> Samanta Kviklyte,<sup>2</sup> Maria Deak,<sup>1</sup> Arash Yavari,<sup>5</sup> Marc Foretz,<sup>6,7,8</sup> Sahar Ghaffari,<sup>5</sup> Mohamed Bellahcene,<sup>5</sup> Houman Ashrafi,<sup>5</sup> Mark H. Rider,<sup>2</sup> Erik A. Richter,<sup>3</sup> and Kei Sakamoto<sup>1,4</sup>

<sup>1</sup>Nestlé Institute of Health Sciences SA, EPFL Innovation Park, Lausanne, Switzerland; <sup>2</sup>Université catholique de Louvain and de Duve Institute, Brussels, Belgium; <sup>3</sup>Department of Nutrition, Exercise and Sports, Faculty of Science, University of Copenhagen, Copenhagen, Denmark; <sup>4</sup>School of Life Sciences, Ecole Polytechnique Fédérale de Lausanne, Lausanne, Switzerland; <sup>5</sup>Experimental Therapeutics and Division of Cardiovascular Medicine, Radcliffe Department of Medicine, University of Oxford, Oxford, United Kingdom; <sup>6</sup>U1016, Institut National de la Santé et de la Recherche Médicale, Institut Cochin, Paris, France; <sup>7</sup>UMR8104, Centre National de la Recherche Scientifique, Paris, France; and <sup>8</sup>Université Paris Descartes, Paris, France

Submitted 20 June 2016; accepted in final form 19 August 2016

**Bultot L, Jensen TE, Lai Y, Madsen AL, Collodet C, Kviklyte S, Deak M, Yavari A, Foretz M, Ghaffari S, Bellahcene M, Ashrafi H, Rider MH, Richter EA, Sakamoto K.** Benzimidazole derivative small-molecule 991 enhances AMPK activity and glucose uptake induced by AICAR or contraction in skeletal muscle. *Am J Physiol Endocrinol Metab* 311: E706–E719, 2016. First published August 30, 2016; doi:10.1152/ajpendo.00237.2016.—AMP-activated protein kinase (AMPK) plays diverse roles and coordinates complex metabolic pathways for maintenance of energy homeostasis. This could be explained by the fact that AMPK exists as multiple heterotrimer complexes comprising a catalytic  $\alpha$ -subunit ( $\alpha 1$  and  $\alpha 2$ ) and regulatory  $\beta$  ( $\beta 1$  and  $\beta 2$ )- and  $\gamma$  ( $\gamma 1$ ,  $\gamma 2$ ,  $\gamma 3$ )-subunits, which are uniquely distributed across different cell types. There has been keen interest in developing specific and isoform-selective AMPK-activating drugs for therapeutic use and also as research tools. Moreover, establishing ways of enhancing cellular AMPK activity would be beneficial for both purposes. Here, we investigated if a recently described potent AMPK activator called 991, in combination with the commonly used activator 5-aminoimidazole-4-carboxamide riboside or contraction, further enhances AMPK activity and glucose transport in mouse skeletal muscle *ex vivo*. Given that the  $\gamma 3$ -subunit is exclusively expressed in skeletal muscle and has been implicated in contraction-induced glucose transport, we measured the activity of AMPK $\gamma 3$  as well as ubiquitously expressed  $\gamma 1$ -containing complexes. We initially validated the specificity of the antibodies for the assessment of isoform-specific AMPK activity using AMPK-deficient mouse models. We observed that a low dose of 991 (5  $\mu$ M) stimulated a modest or negligible activity of both  $\gamma 1$ - and  $\gamma 3$ -containing AMPK complexes. Strikingly, dual treatment with 991 and 5-aminoimidazole-4-carboxamide riboside or 991 and contraction profoundly enhanced AMPK $\gamma 1/\gamma 3$  complex activation and glucose transport compared with any of the single treatments. The study demonstrates the utility of a dual activator approach to achieve a greater activation of AMPK and downstream physiological responses in various cell types, including skeletal muscle.

AMP-activated protein kinase; LKB1; compound 13; A769662; 991; ex229; 5-aminoimidazole-4-carboxamide riboside

AMP-ACTIVATED PROTEIN KINASE (AMPK) is an important energy sensor that maintains energy homeostasis by coordinating

metabolic pathways/processes in response to energy supply and demand (38). During metabolic stress, the cellular AMP-to-ATP ratio increases, leading to the activation of AMPK, which, in turn, switches off energy-consuming anabolic pathways and switches on catabolic pathways to restore ATP levels. One of the most physiological metabolic stresses that activates AMPK is skeletal muscle contraction, which contributes at least partly to an increase in glucose uptake by promoting glucose transporter (GLUT)4 translocation to the plasma membrane by phosphorylating candidate mediators such as TBC1 domain family member 1 and FYVE domain-containing phosphatidylinositol 3-phosphate 5-kinase (15, 21, 25, 30, 32).

AMPK is a heterotrimer comprising a catalytic  $\alpha$ -subunit and regulatory  $\beta$ - and  $\gamma$ -subunits. Multiple isoforms encoded by distinct genes exist ( $\alpha 1$  and  $\alpha 2$ ,  $\beta 1$  and  $\beta 2$ , and  $\gamma 1$ ,  $\gamma 2$ , and  $\gamma 3$ ), which can theoretically form several distinct heterotrimeric combinations (14, 38). Isoform expression varies among cells/tissues and also species (40, 45), with  $\alpha 1$ ,  $\beta 1$ , and  $\gamma 1$  appearing in general to be ubiquitously expressed isoforms. Some isoforms are expressed in a cell/tissue-specific/restricted manner. For example,  $\alpha 2$  is more predominantly expressed in skeletal muscle (26) and  $\gamma 3$  is exclusively detected in skeletal muscle, especially in glycolytic muscle fibers (5, 49). The  $\gamma$ -subunits contain four tandem cystathionine  $\beta$ -synthase (CBS) repeats that bind adenine nucleotides. AMPK activity increases upon phosphorylation of a conserved threonine residue in the T-loop (T172) (14). Binding of ADP and/or AMP to CBS repeats induces conformational changes that promote net T172 phosphorylation by the enhancement of T172 phosphorylation and suppression of T172 dephosphorylation (13, 27, 47). Moreover, the binding of AMP (but not ADP) further increases AMPK activity by direct allosteric stimulation (13). The major upstream kinase phosphorylating T172 in metabolic tissues (e.g., muscle and liver) is a complex containing LKB1 (33, 37), which appears to be constitutively active (1, 31).

AMPK activation has favorable effects on carbohydrate and lipid metabolism in the skeletal muscle and liver, and there has been keen interest in developing AMPK-activating drugs for therapeutic use in the treatment of metabolic diseases, such as type 2 diabetes. Cool et al. (6) described the identification of the thienopyridone A769662, the first small-molecule direct

Address for reprint requests and other correspondence: K. Sakamoto, Nestlé Institute of Health Sciences SA, EPFL Innovation Park, Bâtiment G, Lausanne 1015, Switzerland (e-mail: Kei.Sakamoto@rd.nestle.com).

activator of AMPK. A769662, like AMP, inhibits T172 dephosphorylation (12, 34), although it does not bind to adenine nucleotide-binding sites on the  $\gamma$ -subunit. The oral absorption of this compound is poor. However, when injected intraperitoneally, it displayed antidiabetic effects, including reduced weight gain, decreased plasma glucose and triglycerides, and decreased liver triglycerides in obese mice (6). In addition, A769662 is a valuable research tool, although its utility is limited by the marked selectivity toward complexes containing  $\beta 1$  rather than  $\beta 2$  isoforms (36). Consequently, it robustly stimulates AMPK in mouse hepatocytes (where  $\beta 1$  is predominantly expressed) and modulates lipid metabolism via an AMPK-dependent mechanism (9, 18). In mouse skeletal muscle,  $\beta 2$  is predominantly expressed, whereas only a modest amount of  $\beta 1$  is detectable (39, 43). It has been shown that incubation of isolated mouse skeletal muscle *ex vivo* with A769662 failed to stimulate glucose uptake at 100–200  $\mu$ M. At a much higher concentration (2 mM), A769662 stimulated glucose uptake; however, it was mediated through an AMPK-independent/phosphoinositide 3-kinase-dependent mechanism due to unknown off-target effect(s) (43). Another small-molecule AMPK activator, 991 [also known as ex229 (22)], has recently been described. Structural studies have shown that 991 binds to a site formed between the small lobe of the  $\alpha$ -subunit kinase domain and the  $\beta$ -subunit carbohydrate-binding module (4, 46), which is also the binding site for A769662, termed the allosteric drug and metabolite-binding site (ADaM) (23). 991 is 5- to 10-fold more potent than A769662 in cell-free assays (46) and has been shown to stimulate AMPK activity of both  $\beta 1$ - and  $\beta 2$ -containing complexes and to increase glucose transport in isolated mouse skeletal muscle *ex vivo* (22).

Establishing a means to augment cellular AMPK activation would potentially enhance the therapeutic effects of future AMPK activators and in fundamental research allow more robust biochemical and functional analyses in intact cells/tissues. We (8, 11, 41) and others (35) have shown that dual AMPK-activating compound treatment [A769662 and 5-aminoimidazole-4-carboxamide riboside (AICAR)] resulted in additive/synergistic increases in cellular AMPK activity and downstream physiological effects, such as inhibition of lipogenesis in hepatocytes (8). One limitation of the A769662 and AICAR dual treatment is that the additive or synergistic effects can only be produced in cells where  $\beta 1$  is predominantly expressed (e.g., mouse hepatocytes), as the effects were absent in  $\beta 1$ -deficient cells (8). In the present study, we explored whether treatment with the potent ADaM site-binding agent 991 would enhance the effect of the canonical adenine nucleotide binding-mediated activation of AMPK by AICAR and contraction in mouse skeletal muscle. Given that activation of  $\gamma 3$ -containing complexes (especially  $\alpha 2\beta 2\gamma 3$ ) is associated with AICAR- and contraction-mediated glucose uptake (2, 3), we assessed the effect of 991 alone and in combination with AICAR or contraction on activating AMPK $\gamma 1$  and AMPK $\gamma 3$  complexes. We demonstrated that a low dose of 991 results in only a modest or negligible activation of  $\gamma 1$ - and  $\gamma 3$ -containing AMPK complexes in mouse skeletal muscle. However, strikingly, dual treatment with 991 and AICAR or 991 and contraction profoundly enhanced AMPK activation and glucose transport compared with single agonist treatments.

## MATERIALS AND METHODS

**Materials.** AICAR was from Toronto Research Chemicals. A769662 was from Selleck Chemicals. Compound 13 was obtained as previously described (18). 991 (5-[[6-chloro-5-(1-methylindol-5-yl)-1H-benzimidazol-2-yl]oxy]-2-methyl-benzoic acid; CAS no. 129739-36-2) as obtained as previously described (7). Protein G Sepharose was from GE Healthcare, and FLAG-M2 resin from Sigma-Aldrich. ECL reagent and P81 filter papers were obtained from GE Healthcare. [ $\gamma$ - $^{32}$ P]-ATP was from Perkin-Elmer. AMARA, LKBtide, and Sakamoto substrate peptides were synthesised by GL Biochem. The COS1 cell line was obtained from American Type Culture Collection, and the C<sub>2</sub>C<sub>12</sub> cell line was obtained from Sigma-Aldrich. Frozen tissues or extracts from AMPK $\alpha 1$ -/ $\alpha 2$ -, AMPK $\beta 1$ -/ $\beta 2$ -, and AMPK $\gamma 3$ -deficient mice were obtained from Benoit Viollet and Marc Foretz (Institut National de la Santé et de la Recherche Médicale, Institut Cochin, Paris, France), Gregory Steinberg (McMaster University, Hamilton, ON, Canada), and Alexander Chibalin and Juleen Zierath (Karolinska Institutet, Stockholm, Sweden), respectively. All cell culture reagents were purchased from Thermo Fisher Scientific, and all other chemicals were from Sigma-Aldrich unless otherwise stated.

**Antibodies.** AMPK $\alpha 1/\alpha 2$  (no. 2532), phosphorylated (p)T172 AMPK $\alpha$  (no. 2535), AMPK $\beta 1$  (no. 4182), AMPK $\beta 2$  (no. 4148), AMPK $\beta 1/\beta 2$  (no. 4150), pS79 acetyl-CoA carboxylase (ACC; no. 3661), ACC (no. 3676), pS792 regulatory associated protein of mechanistic target of rapamycin (RAPTOR; no. 2083), and RAPTOR (no. 2280) were from Cell Signaling Technology. FLAG (F7425) and  $\alpha$ -tubulin (T6074) antibodies were from Sigma-Aldrich. AMPK $\gamma 2$  (sc-19141) and creatine kinase (sc-15161) antibodies were from Santa Cruz Biotechnology. AMPK $\alpha 1$  (no. 07-350) and AMPK $\alpha 2$  (no. 07-363) antibodies were obtained from Millipore. For immunoprecipitation, the antibody for AMPK $\beta 1$  (A4856) was from Sigma-Aldrich and the antibody for AMPK $\beta 2$  (MAB3808) was from R&D Systems. AMPK $\alpha 1$  and AMPK $\alpha 2$  antibodies used for immunoprecipitation were as previously described (33). LKB1 and salt-inducible kinase (SIK)3 antibodies have also been previously described (31). The following polyclonal antibodies were generated by YenZym Antibodies by immunizing rabbits with the indicated peptides: a combination of human AMPK $\gamma 3$  peptide [residues S45–S65: \*CS-SERIRGKRRKALRWTRQKS; a terminal cysteine (\*C) was added to the peptide sequence to allow peptide conjugation to carrier proteins], mouse AMPK $\gamma 3$  peptide (residues S44–S64: \*CSSERT-CAIRGVKASRWTRQEA), and AMPK $\gamma 1$  peptide (residues E7–S26 of mouse AMPK $\gamma 1$ : \*CESSPALENEHFQETPESNNS). The produced antibodies were affinity purified using respective antigen-specific peptide columns by YenZym Antibodies. Horseradish peroxidase-conjugated secondary antibodies were from Jackson ImmunoResearch.

**Kinase inhibition screening.** Kinase inhibition screening was performed by the International Centre for Kinase Profiling (University of Dundee) using [ $\gamma$ - $^{32}$ P]phosphotransferase assays (MRC Protein Phosphorylation and Ubiquitylation Unit, <http://www.kinase-screen.mrc.ac.uk/>).

**Molecular cloning.** The coding region of human AMPK $\gamma 1$  (NM\_002733), AMPK $\gamma 2$  (NM\_016203.3), and AMPK $\gamma 3$  (NM\_017431) was amplified from muscle RNA (Agilent) using a Superscript III 1 step RT PCR kit (Invitrogen). The resulting PCR products were either ligated into intermediate vector (pSC Agilent) or digested with the relevant restriction enzymes and ligated directly into the mammalian expression vector (pCMV5 acc. AF239249), which was modified to incorporate a NH<sub>2</sub>-terminal FLAG tag. Sequences of all clones were verified in house at the Nestlé Institute of Health Sciences using the BigDye Terminator 3.1 kit and 3500XL Genetic analyzer (Applied Biosystems).

**Recombinant muscle-specific AMPK trimeric complex.** Human AMPK  $\alpha 2\beta 2\gamma 3$  was expressed in *Escherichia coli* using a tricistronic



construct activated with  $\text{Ca}^{2+}$ /calmodulin-dependent protein kinase kinase- $\beta$  in vitro and then purified and assayed as previously described (18).

**Animals.** Animal experiments were approved by the local ethics committee and conducted in accordance with the European Convention for the Protection of Vertebrate Animals used for Experimental and Other Scientific Purposes. Protocols used were approved by the Service Vétérinaire Cantonal (Lausanne, Switzerland) under license VD2841 or approved by the University of Oxford Animal Care and Ethical Review Committee and conformed with the United Kingdom Animals Scientific Procedures Act 1986, incorporating European Directive 2010/63/EU (no. 30/2977), or approved by the Danish Animal Experiments Inspectorate or approved by the University of Paris-Descartes ethics committee (no. CEEA34.BV.157.12) and performed under French authorization to experiment on vertebrates (no. 75-886) in accordance with European guidelines. The generation of AMPK $\gamma 1^{-/-}$  mice has been previously described (10). Global AMPK $\gamma 2^{-/-}$  mice were generated by deleting the entire exon 7 of the gene encoding AMPK $\gamma 2$  in R299Q knockin mice using Sox2cre-driven excision (48). Mice were maintained on a standard chow diet and 12:12-h light-dark cycle.

**Cell and tissue extract preparation.** After treatment, cells were washed once with PBS and scraped into ice-cold lysis buffer [containing 50 mM Tris-HCl (pH 7.5), 1 mM EDTA, 1 mM EGTA, 270 mM sucrose, 1% (wt/vol) Triton X-100, 20 mM glycerol-2-phosphate, 50 mM NaF, 5 mM  $\text{Na}_4\text{P}_2\text{O}_7$ , 1 mM DTT, 0.1 mM PMSF, 1 mM benzamidine Cl, 1  $\mu\text{g}/\text{ml}$  microcystin-LR, 2  $\mu\text{g}/\text{ml}$  leupeptin, and 2  $\mu\text{g}/\text{ml}$  pepstatin A]. Frozen tissues were homogenised using a polytron PT 2500 E (Kinematica) in ice-cold lysis buffer. Cell/tissue lysates were clarified by centrifugation at 3,500 g for 15 min at 4°C, and protein concentration was measured using Bradford reagent and BSA as a standard.

**Cell culture.** COS1, mouse embryonic fibroblast, and C<sub>2</sub>C<sub>12</sub> cells were maintained in DMEM GlutaMAX (Thermo Fisher Scientific) supplemented with 10% (vol/vol) FBS and antibiotics. C<sub>2</sub>C<sub>12</sub> myoblasts were differentiated into myotubes by 7 days of culture in DMEM GlutaMAX supplemented with 2% (vol/vol) horse serum, 100 U/ml penicillin G, and 100  $\mu\text{g}/\text{ml}$  streptomycin. COS1 cells were grown in 6-cm dishes and transfected at 60–70% confluency with 3.7  $\mu\text{g}$  plasmid prebound to 10.5  $\mu\text{g}$  polyethylenimine in 50 mM HEPES (pH 7.4) and 150 mM NaCl. Cell culture medium was changed once at 24 h after transfection, and cells were left for an additional 24 h before compound treatment. Mouse embryonic fibroblast cells were grown in 10-cm dishes and treated as indicated at 80–90% confluency. Primary mouse hepatocytes were isolated from C57BL/6NTac male mice (Taconic) by collagenase perfusion, as previously described (18, 29). Hepatocytes were seeded in medium 199 containing 100 U/ml penicillin G, 100  $\mu\text{g}/\text{ml}$  streptomycin, 0.1% (wt/vol) BSA, 10% (vol/vol) FBS, 10 nM insulin, 200 nM triiodothyronine, and 100 nM dexamethasone. Hepatocytes were left for attachment (3–4 h) and cultured overnight in medium 199 supplemented with antibiotics and 100 nM dexamethasone. Cells were used for experiments the following morning.

**Immunoblot analysis.** Cell or tissue lysates were denatured in Laemmli buffer at 95°C for 5 min and separated by Tris-glycine SDS-PAGE and transferred onto polyvinylidene difluoride membranes. Membranes were blocked for 1 h at room temperature in 20 mM Tris-HCl (pH 7.6), 137 mM NaCl, and 0.1% (vol/vol) Tween 20 (Tris-buffered saline with Tween 20) containing 5% (wt/vol) skimmed milk. Membranes were incubated in primary antibody prepared in Tris-buffered saline with Tween 20 containing 5% (wt/vol) BSA or skimmed milk overnight at 4°C. Signal detection was performed using horseradish peroxidase-conjugated secondary antibodies and ECL.

**Immunoprecipitation and kinase activity assay.** Cell or tissue lysates were incubated with the indicated antibodies (shown in Fig. 2F) precoupled to protein G-Sepharose for 1 h at 4°C. Immune complexes

were pelleted at 500 g for 1 min and washed twice with 0.5 ml lysis buffer containing 500 mM NaCl and twice with 0.5 ml *buffer A* [50 mM HEPES (pH 7.4), 150 mM NaCl, 1 mM EGTA, and 1 mM DTT] and eluted with Laemmli buffer for analysis by immunoblotting or assayed directly for kinase activity. The AMPK activity assay was performed as previously described (19). Briefly, beads were incubated 45 min at 30°C under agitation in *buffer A* supplemented with 10 mM magnesium acetate, 100  $\mu\text{M}$  ATP, and 200  $\mu\text{M}$  AMARA peptide ( $\text{NH}_2\text{-AMARAASAAALARRR-COOH}$ ) in the presence of  $[\gamma\text{-}^{32}\text{P}]\text{ATP}$  (1  $\mu\text{Ci}$ ). Reactions were terminated by spotting reaction mixtures onto P81 filters and immersion in 75 mM phosphoric acid. Washed filters were dried, and  $^{32}\text{P}$  incorporation into the substrate AMARA peptide was measured by Cherenkov counting.

LKB1 and SIK3 assays were performed as previously described (31), and activity was measured using LKBtide and Sakamotoide (16) as substrates, respectively.

**Incubation, contraction, and glucose uptake in isolated mouse skeletal muscle.** Mice were euthanized by cervical dislocation, and extensor digitorum longus (EDL) muscles were rapidly removed and mounted on an incubation apparatus. The EDL muscle was incubated as previously described (20) in the presence of 991 or vehicle (0.1% DMSO) for 60 min with or without AICAR or in the presence of 991 or vehicle (0.1% DMSO) for 45 min with or without contraction evoked via electrical stimulation (1 Hz, 0.1-ms square wave 30-V pulses) (42) for 15 min. 2-Deoxy- $[\text{H}^3]\text{glucose}$  uptake was measured during the last 10 min of the incubation or contraction period of each experiment, as described previously (20).

**Statistical analysis.** Results are expressed as means  $\pm$  SE (unless otherwise stated) for the indicated number of individual experiments. Statistical significance was assessed using a paired two-sided Student's *t*-test (Fig. 3) or one-way ANOVA. When ANOVA revealed significant differences, further analysis was performed using Fisher's least-significant-difference post hoc test for multiple comparisons (Figs. 4–7). Differences between groups were considered statistically significant at  $P < 0.05$ .

## RESULTS

**Validation of AMPK $\alpha$ -,  $\beta$ -, and  $\gamma$ -isoform-specific antibodies for immunoblot analysis.** To investigate the effect of dual treatments on AMPK isoform/complex-specific activity using cell or tissue extracts in subsequent experiments, we initially performed an extensive characterization/validation of the AMPK isoform-specific antibodies. We verified the specificity of AMPK $\alpha 1$ , AMPK $\alpha 2$ , AMPK $\beta 1$ , and AMPK $\beta 2$  antibodies for immunoblot analysis using the respective isoform-specific AMPK knockout (KO) liver tissue extracts as controls (Fig. 1, A and B). The specificity of the AMPK $\gamma$  antibodies has not been well documented, and we first assessed cross-reactivity of the antibodies using recombinant proteins. Individual constructs encoding FLAG-tagged human AMPK $\gamma 1$ , AMPK $\gamma 2$ , or AMPK $\gamma 3$  were transfected in COS1 cells, and all three  $\gamma$ -isoforms were isolated from cell lysates using anti-FLAG antibody coupled to protein G-Sepharose. Immunoblot analysis of the FLAG-immunoprecipitates confirmed that similar amounts of each AMPK  $\gamma$ -isoform were isolated from the lysates (Fig. 1C, top left). We next immunoblotted the immunoprecipitates with AMPK  $\gamma 1$ -,  $\gamma 2$ -, or  $\gamma 3$ -isoform-specific antibodies. As shown in Fig. 1C (top right and bottom), each antibody only detected the corresponding AMPK  $\gamma$ -isoform. We then tested the specificity of the  $\gamma$  antibodies against the endogenous subunit protein using tissue extracts from wild-type (WT) versus isoform-specific AMPK $\gamma$  KO mice. As shown in Fig. 1, D–F, each isoform-specific antibody detected a band at the

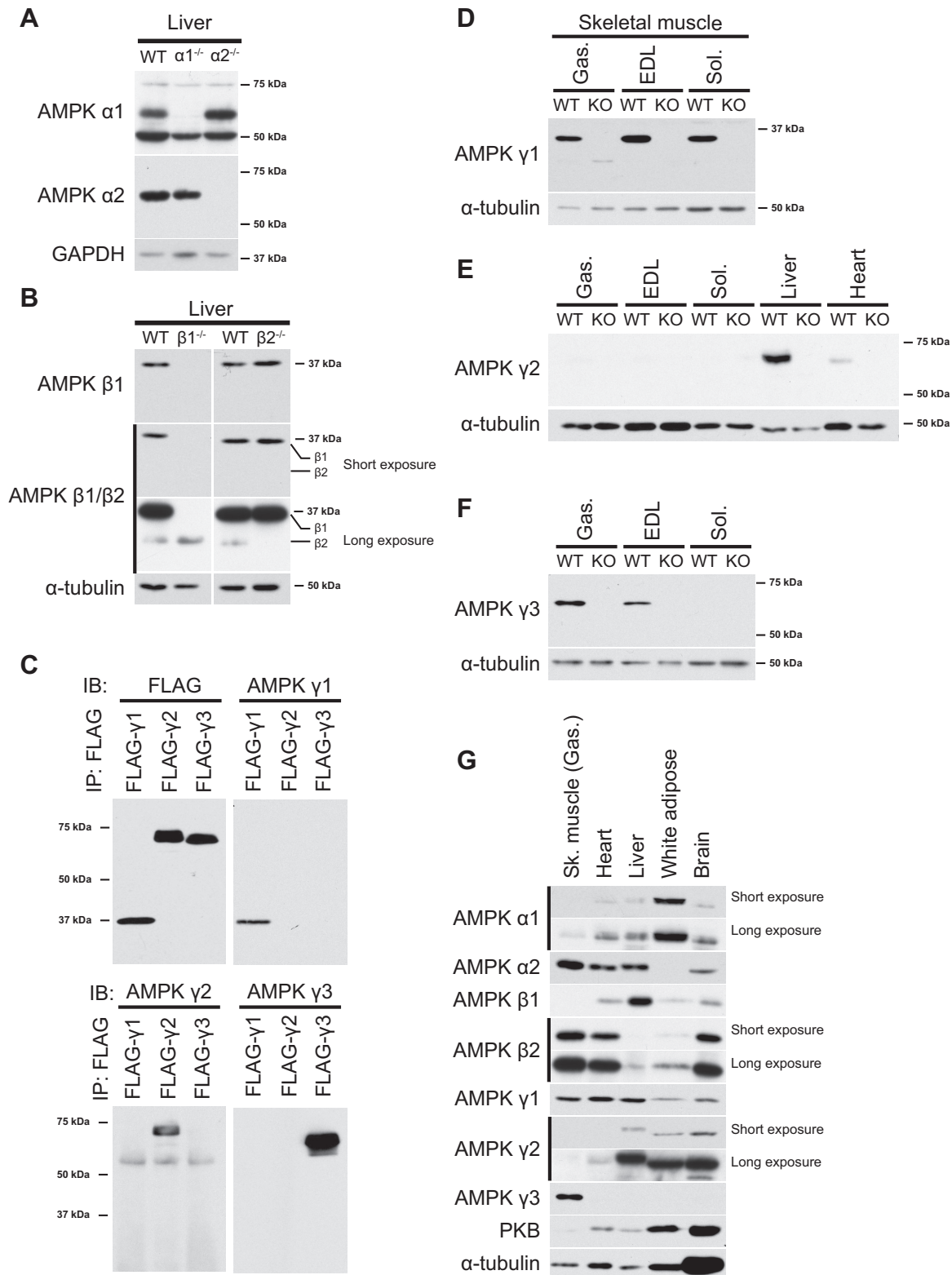


Fig. 1. Validation of AMP-activated protein kinase (AMPK) isoform-specific antibodies for immunoblot (IB) analysis. *A* and *B*: IB analysis was performed with 20  $\mu$ g liver tissue protein extracts from wild-type (WT) and AMPK $\alpha1/2$  knockout (KO) mice (*A*) as well as AMPK $\beta1/\beta2$  KO mice (*B*) using the indicated antibodies. IP, immunoprecipitation. *C*: FLAG-tagged human AMPK $\gamma1$ , AMPK $\gamma2$ , and AMPK $\gamma3$  were ectopically expressed in COS1 cells, and cell extracts were generated. All three AMPK $\gamma$  isoforms were immunoprecipitated with anti-FLAG antibody and immunoblotted with the indicated antibodies. *D–F*: IB analysis was performed with 20  $\mu$ g of the indicated tissue extract proteins from WT and isoform-specific AMPK $\gamma$  KO mice using the indicated antibodies. Gas, gastrocnemius; EDL, extensor digitorum longus; Sol, soleus. *G*: a panel of mouse tissue extracts (male C57BL/6NTac mice) was immunoblotted with the indicated antibodies. Sk, skeletal PKB, protein kinase B. Representative blots are shown from  $n = 1–3$  mice/tissues.

expected relative molecular mass in WT tissue extracts but not in the corresponding KO tissue extracts. AMPK $\gamma$ 1 was expressed similarly in the three muscle types measured: gastrocnemius (glycolytic/oxidative fibers mixed), EDL (predominantly containing glycolytic fibers), and soleus (predominantly containing oxidative fibers). AMPK $\gamma$ 3 was expressed in the gastrocnemius and EDL but not in soleus (Fig. 1F), consistent with previous reports (2, 49). Of note, AMPK $\gamma$ 2 was not detectable in skeletal muscle, although it could be detected in the liver and heart (Fig. 1E). Immunoblot analysis of a selection of mouse tissue extracts using the validated AMPK isoform-specific antibodies (Fig. 1, A–F) verified the exclusive expression of AMPK $\gamma$ 3 in skeletal muscle and predominant expression of AMPK $\alpha$ 1 in white adipose (Fig. 1G). Predominant expression of AMPK $\beta$ 1 in the liver and AMPK $\beta$ 2 in the skeletal muscle, as well as the ubiquitous expression of AMPK $\gamma$ 1, was also confirmed. It has been previously reported that in mouse glycolytic skeletal muscle (EDL),  $\beta$ 1 expression represents only ~5% of total  $\beta$ -subunit expression (43), and  $\beta$ 1 expression was not detectable with immunoblot analysis (20  $\mu$ g lysates; Fig. 1G), even with longer film exposure (data not shown). AMPK $\gamma$ 2 was broadly expressed, but its expression appeared relatively low (considering the levels of loading control:  $\alpha$ -tubulin and protein kinase B) in the heart and was not detectable in skeletal muscle (Fig. 1, E and G).

**Validation of AMPK  $\beta$ - and  $\gamma$ -isoform-specific antibodies for immunoprecipitation.** We next examined the specificity and efficiency of the AMPK  $\beta$ - and  $\gamma$ -isoform-specific antibodies for immunoprecipitation using mouse tissue extracts [specificity of AMPK $\alpha$ 1/ $\alpha$ 2 antibodies for immunoprecipitation has been previously described (33)]. We verified that each isoform-specific antibody immunoprecipitated the corresponding AMPK  $\beta$ - and  $\gamma$ -subunit proteins (using extracts from WT animals) in a specific and nearly complete manner (Fig. 2, A–E). There was no immunoreactive band/signal detected in extracts from the corresponding AMPK KO lysates and only a negligible amount of proteins detected in the supernatant that had been incubated with the corresponding antibody. Moreover, we confirmed that each AMPK  $\beta$ - and  $\gamma$ -isoform-specific antibody could precipitate its known interacting AMPK  $\alpha$ - and  $\beta$ -subunit isoform(s) [i.e., AMPK $\alpha$  and AMPK $\beta$  proteins were identified in  $\gamma$  antibody immunoprecipitates (43); Fig. 2A] from the tissue extracts (Fig. 2, A–E).

In summary, we validated AMPK isoform-specific antibodies for both immunoblot analysis and immunoprecipitation. Given that we used several antibodies for different applications in the present study, a summary of the antibody list is shown in Fig. 2F.

**Effect of a small-molecule AMPK activator, 991, on a panel of protein kinases in vitro.** We screened compound 991, a recently described potent AMPK activator (22, 46), at a concentration of 1  $\mu$ M in cell-free assays against a panel of 139 protein kinases, including AMPK (Fig. 3A). As anticipated, recombinant AMPK complex [human  $\alpha$ 1(11–559) $\beta$ 2(1–272)- $\gamma$ 1(11–331)] was activated ~70% (highlighted in gray in Fig. 3A), whereas none of the other kinases, including upstream kinases for AMPK (LKB1 and  $\text{Ca}^{2+}$ /calmodulin-dependent protein kinase- $\beta$ , highlighted in gray) and members of AMPK-related kinases (1) [BR serine/threonine kinase (BRSK)1, BRSK2, SIK2, SIK3, microtubule affinity regulating kinase (MARK)2, MARK3, MARK4, and maternal embryonic leucine zipper

kinase, highlighted in gray] were significantly affected (Fig. 3A). We have confirmed that 991 robustly stimulates recombinant muscle-specific AMPK $\alpha$ 2 $\beta$ 2 $\gamma$ 3 complex in a separate gold standard kinase activity assay in vitro [vehicle (DMSO):  $496 \pm 123$  (SD) vs. 991,  $3,380 \pm 679$  pmol  $\text{P}_i$  incorporated  $\text{min}^{-1} \cdot \text{mg}^{-1}$ ,  $P < 0.05$ ]. Furthermore, in mouse embryonic fibroblasts 991 increased the phosphorylation of AMPK $\alpha$  and its substrates ACC and RAPTOR (Fig. 3B). Consistent with this observation (Fig. 3B) and in vitro screen data (Fig. 3A), 991 robustly stimulated AMPK $\alpha$ 1 activity (Fig. 3C) but not LKB1 or SIK3 (Fig. 3, D and E). These results suggest that 991 is a specific activator of AMPK at least in vitro and to a limited extent in intact cells.

**Enhanced activation of cellular AMPK activity by dual compound treatment in hepatocytes.** We tested if the effect of 991 [which binds to a site formed between the small lobe of the  $\alpha$ -subunit kinase domain and the  $\beta$ -subunit carbohydrate-binding module (46)] on AMPK activity could be further increased when it was coincubated with another compound, AICAR or C13 [which binds to the CBS domain of the  $\gamma$ -subunit, mimicking the action of AMP (18, 24)] in mouse primary hepatocytes. Hepatocytes were incubated with increasing concentrations of 991 (ranging from 0 to 0.3  $\mu$ M) in the presence or absence of AICAR (0.1 mM) or C13 (10  $\mu$ M) for 1 h, and phosphorylation of AMPK and its bona fide substrates (i.e., ACC and RAPTOR) was assessed by immunoblot analysis as surrogate markers of cellular AMPK activity (Fig. 4A). Treatment of hepatocytes with 991 alone resulted in a slight increase in the phosphorylation of AMPK and RAPTOR only at 0.3  $\mu$ M, whereas a robust increase in ACC phosphorylation was readily observed and saturated at a concentration of 0.03  $\mu$ M 991. AICAR or C13 alone robustly increased T172 phosphorylation of AMPK $\alpha$ , and when 991 was coincubated, there was a modest additional dose-dependent increase in AMPK $\alpha$  phosphorylation (as judged by increased band density and slower-migrating species). Single AICAR or C13 treatment caused ACC phosphorylation to saturating levels. In contrast, RAPTOR phosphorylation was modestly increased by AICAR or C13 alone, and it was dose dependently increased when incubations were carried out with 991 (Fig. 4A). These data are consistent with our previous observation that RAPTOR phosphorylation requires a higher threshold of AMPK activity than ACC phosphorylation in mouse hepatocytes (18).

To assess the effects of single or dual agonist treatment on the activities of specific AMPK isoforms/complexes in hepatocytes, we next performed an in vitro quantitative AMPK activity assay after immunoprecipitation with specific AMPK subunit isoform antibodies (Fig. 4, B–G). As anticipated, the activity of both AMPK $\alpha$ 1- and AMPK $\alpha$ 2-containing complexes (Fig. 4, B and C) resembled the results of AMPK $\alpha$  T172 phosphorylation (Fig. 4A) in response to compound treatments. Of note and consistent with our previous observation (18), incubation with C13 alone more robustly stimulated the activity of AMPK $\alpha$ 1-containing complexes, whereas AICAR treatment increased the activation of AMPK $\alpha$ 1 and AMPK $\alpha$ 2 in a relatively similar manner. However, the absolute activity achieved by C13 was similar between  $\alpha$ 1 and  $\alpha$ 2, whereas AICAR caused much higher absolute activation for  $\alpha$ 2 (Fig. 4, B and C). In the mouse liver, AMPK $\beta$ 1 is the predominant isoform, whereas expression of AMPK $\beta$ 2 is marginal (9). Activity of AMPK $\beta$ 1-containing complexes displayed a simi-



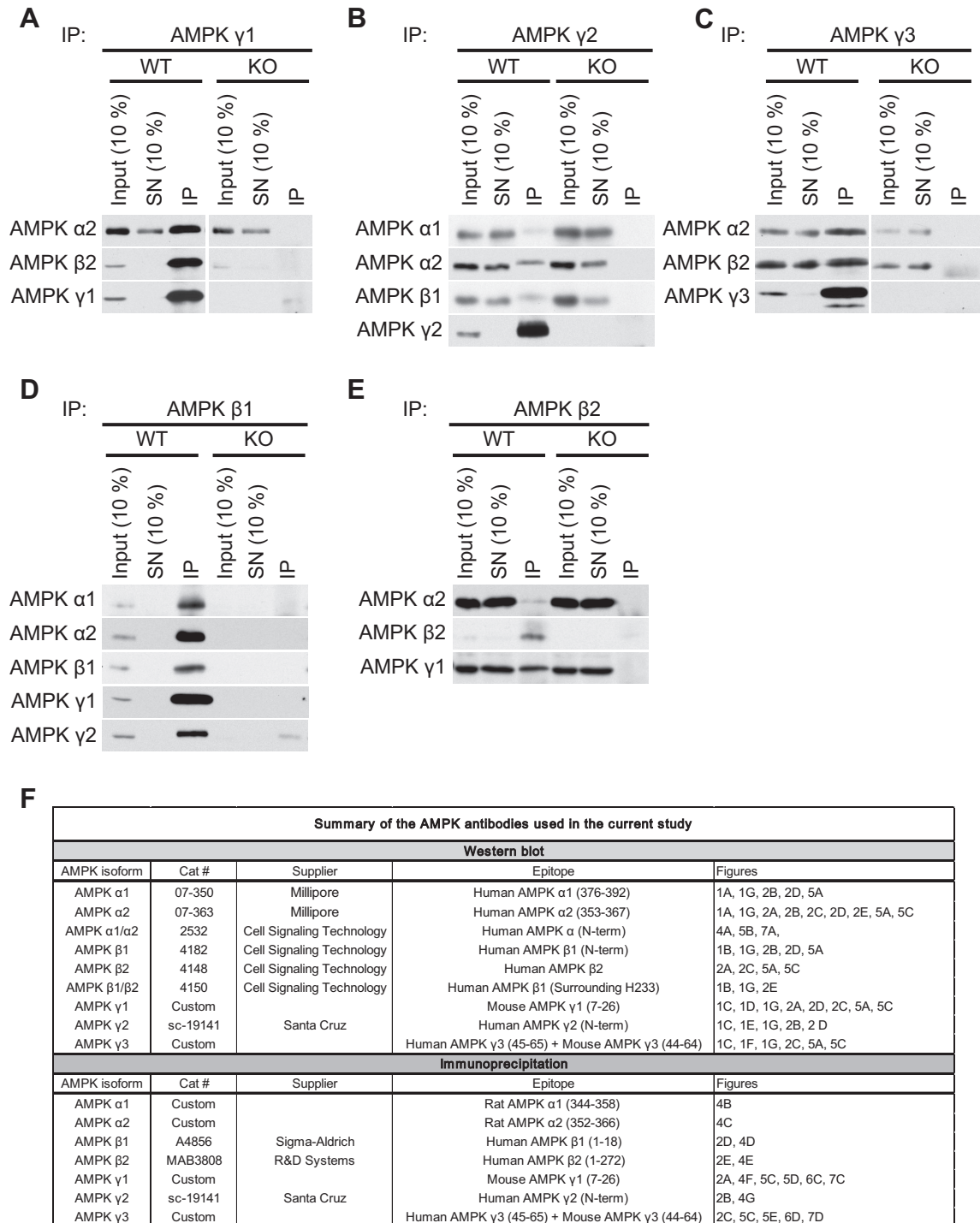


Fig. 2. Validation of AMPK isoform-specific antibodies for IP. A–E: Endogenous AMPK complexes were immunoprecipitated using the indicated antibodies from 200  $\mu$ g extract protein from the gastrocnemius muscle (A and C) or liver (B, D, and E) of the indicated AMPK WT or KO mice. IB analysis was performed on the input (10% = 20  $\mu$ g), the supernatant (SN; 10% = 20  $\mu$ g), and the immunoprecipitates (IP) with the indicated antibodies. F: summary table of the isoform-specific antibodies used in the present study.

lar activation profile (Fig. 4D) to the activity of AMPK $\alpha$ 1- and AMPK $\alpha$ 2-containing complexes (Fig. 4, A and B). As expected, AMPK activity associated with AMPK $\beta$ 2 complexes was much lower than AMPK $\beta$ 1 complexes and was robustly activated by AICAR or C13. However, there was no further activation when 991 was incubated along with AICAR or C13 (Fig. 4E). This might be due to the fact that 991 has a higher

affinity for AMPK $\beta$ 1 (in cell-free assay) (46) and thus the low concentrations of 991 used were not sufficient to stimulate the activity of AMPK $\beta$ 2-containing complexes. We then measured AMPK activity associated with AMPK $\gamma$ 1 or AMPK $\gamma$ 2 and observed that treatment with AICAR alone showed similar and significant increases in AMPK activity of complexes associated with both isoforms, although the extent of activation was

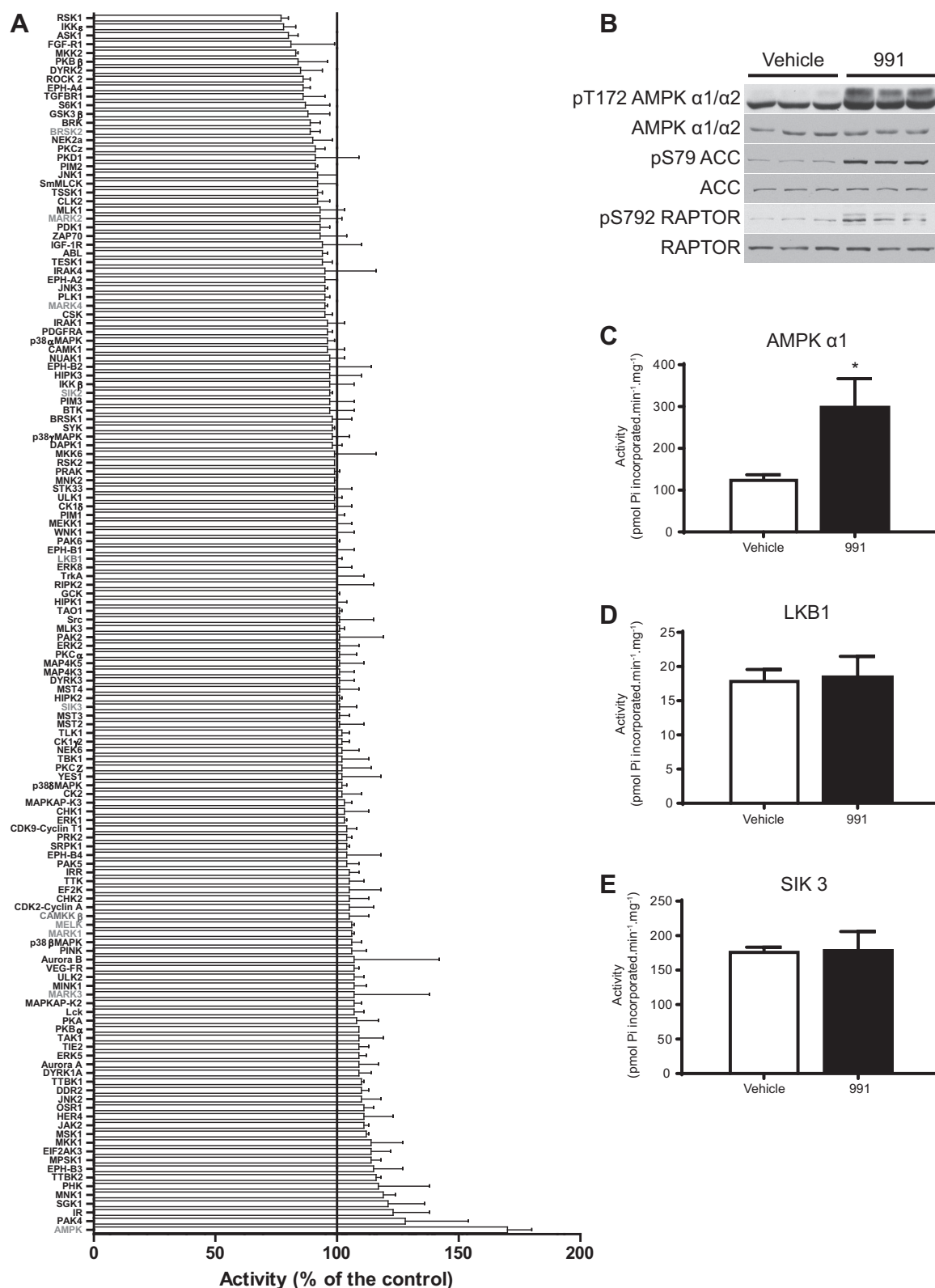


Fig. 3. Effect of the small-molecule AMPK activator 991 on activity of a panel of 139 protein kinases. **A**: a kinase screen was performed by the International Centre for Kinase Profiling (University of Dundee). Assays were performed in the presence of 1  $\mu$ M 991. Results are shown in rank order as mean percent activity compared with controls in the absence of compound  $\pm$  SD. **B–E**: mouse embryonic fibroblasts were treated with vehicle (0.1% DMSO) or 991 (10  $\mu$ M) for 1 h (3 independent plates for each condition). Cell lysates were immunoblotted with the indicated antibodies (**B**). The indicated kinases were immunoprecipitated from cell lysates, and in vitro kinase activity toward substrate peptide was measured (**C–E**) as described in MATERIALS AND METHODS. ACC, acetyl-CoA carboxylase; RAPTOR, regulatory associated protein of mechanistic target of rapamycin; SIK3, salt-inducible kinase. Results are expressed as means  $\pm$  SD. \* $P < 0.05$ .



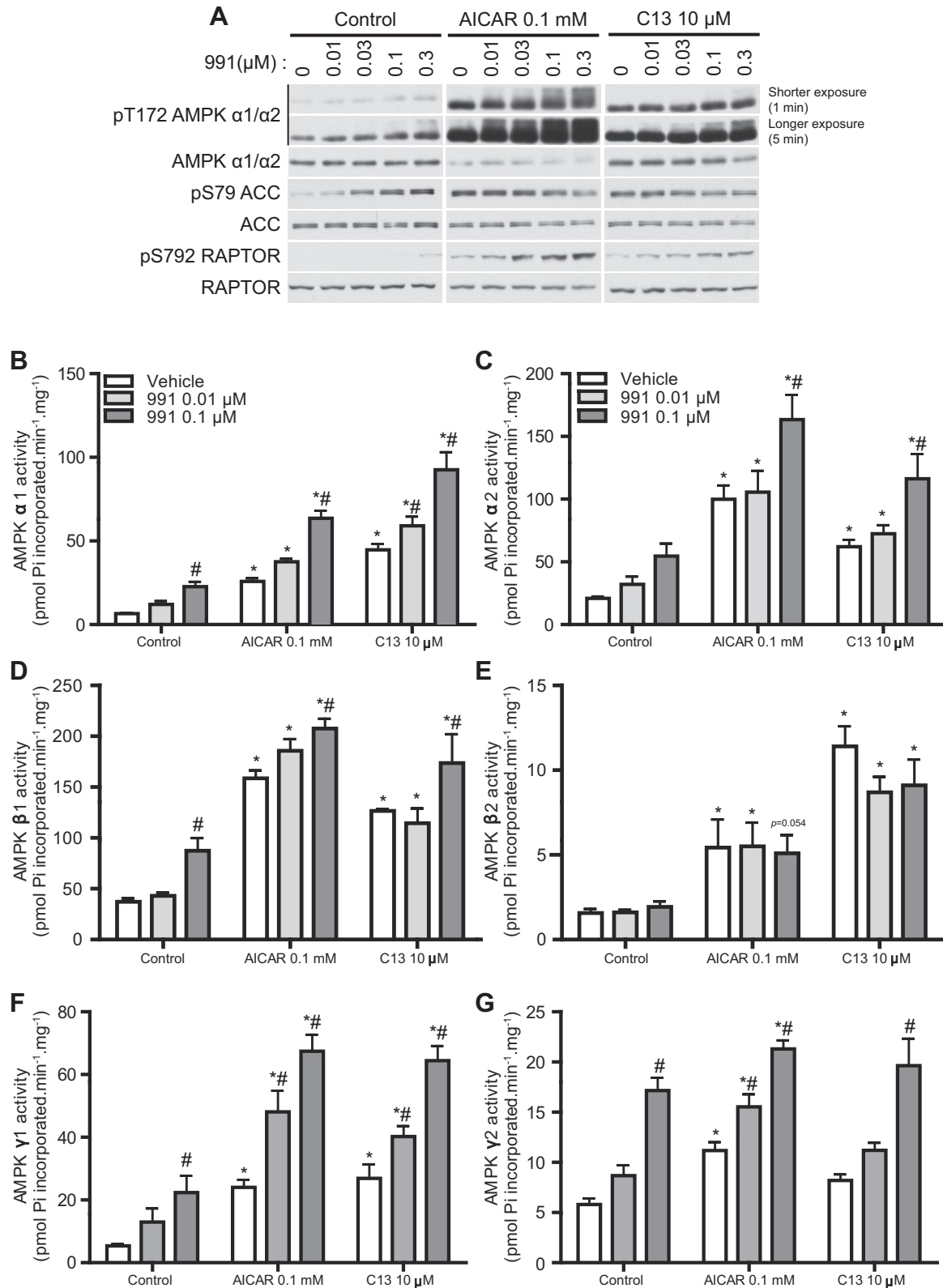


Fig. 4. 991 treatment enhances AMPK activity induced by 5-aminoimidazole-4-carboxamide riboside (AICAR) or C13 in hepatocytes. Hepatocytes were isolated from male C57BL/6NTac mice and cultured overnight as described in MATERIALS AND METHODS. Hepatocytes were untreated (control) or treated with AICAR (0.1 mM) or C13 (10 μM) for 1 h in the presence of the indicated concentrations of 991. **A**: IB analysis was performed with 20 μg lysates using the indicated antibodies. Representative blots are shown;  $n = 3$ . Shorter (1 min) and longer (5 min) exposure indicate the exposure time with X-ray film. **B--G**: AMPK complexes were immunoprecipitated from 50 μg lysates with the indicated antibodies. AMPK activity (in duplicate) was measured in vitro as described in MATERIALS AND METHODS. Results are expressed as means  $\pm$  SE;  $n = 3$ . \*Significance of AICAR or C13 versus the respective control condition (0, 0.01, or 0.1 μM 991); #significance of 991 versus the respective compound (AICAR or C13) or vehicle.  $P < 0.05$ .

higher in AMPK $\gamma$ 1-containing complexes (Fig. 4, *F* and *G*). C13 promoted a robust increase in the activity of the AMPK $\gamma$ 1-containing complex but not the AMPK $\gamma$ 2-containing complex. Moreover, robust increases ( $\sim 2$ - to  $3$ -fold) in AMPK activity of AMPK $\gamma$ 1- or AMPK $\gamma$ 2-containing complexes were observed when 991 was combined in a dose-dependent manner (Fig. 4, *F* and *G*).

We observed that 991 dose dependently ( $0.01$  and  $0.1$   $\mu\text{M}$ ) inhibited lipogenesis ( $34\%$  and  $63\%$ , respectively), which was further reduced when it was coincubated with a low dose of AICAR ( $0.03$  mM) or C13 ( $1$   $\mu\text{M}$ ) (data not shown).

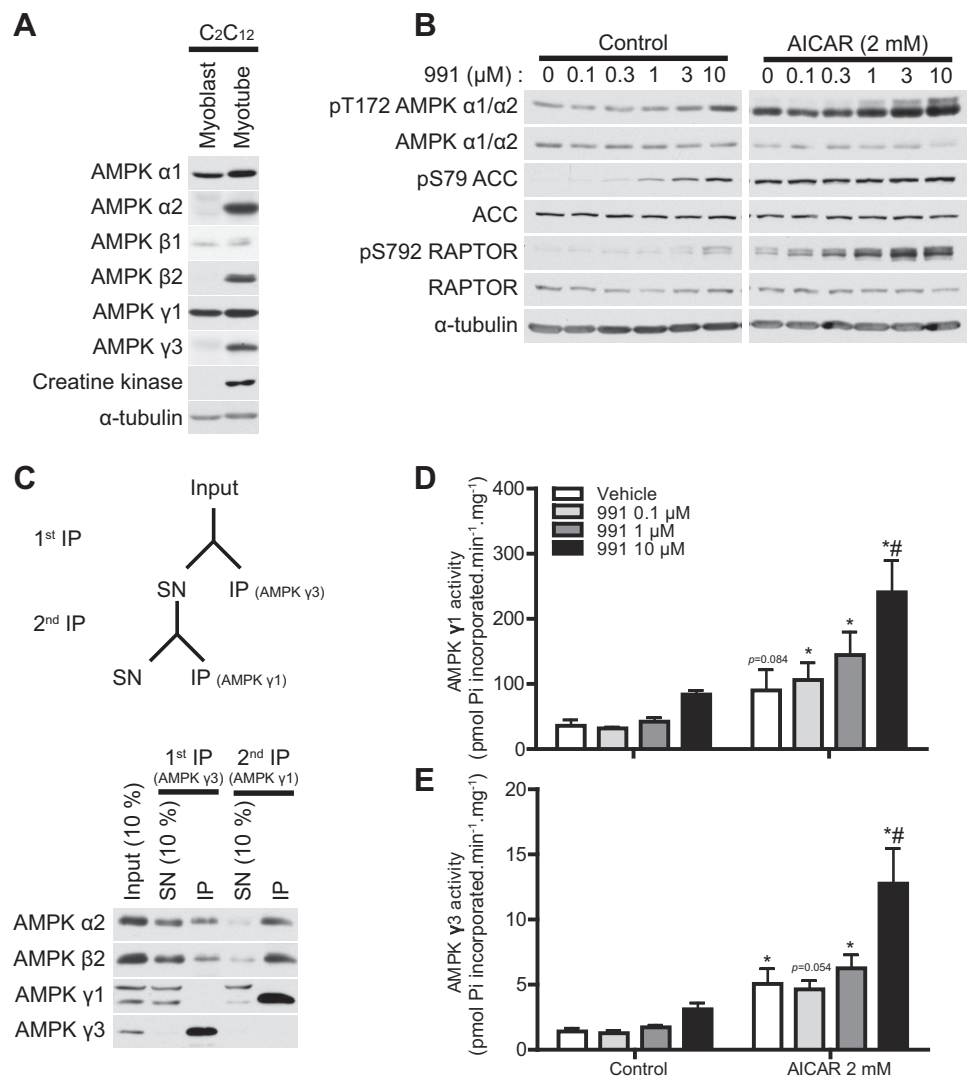
**Enhanced activation of AMPK $\gamma$ 1- and AMPK $\gamma$ 3-containing complexes by dual compound treatment in C<sub>2</sub>C<sub>12</sub> muscle cells.** We next investigated the effects of dual compound treatment on AMPK activity in muscle cells. We first characterized the isoform-specific expression pattern of AMPK in C<sub>2</sub>C<sub>12</sub> mouse myoblasts and myotubes. We observed that in myoblasts, expression of  $\alpha 1$ ,  $\beta 1$ , and  $\gamma 1$  was readily detectable and their levels did not alter when differentiated into myotubes. In contrast, expression of  $\alpha 2$ ,  $\beta 2$ , and  $\gamma 3$  was only detectable in myotubes but not in myoblasts (Fig. 5A). Consistent with the results obtained using mouse skeletal muscle tissue (Fig. 1E),

levels of AMPK $\gamma$ 2 in C<sub>2</sub>C<sub>12</sub> myoblasts and myotubes were undetectable (data not shown).

Incubation of C<sub>2</sub>C<sub>12</sub> myotubes with 991 alone modestly increased AMPK $\alpha$  phosphorylation but only at a concentration of  $10$   $\mu\text{M}$  (Fig. 5B). Treatment with 991 promoted dose-dependent increases in ACC and RAPTOR phosphorylation. Similar to the observations in hepatocytes (Fig. 4A) (18), in muscle cells, phosphorylation of RAPTOR also appeared to require a higher threshold of AMPK activity than ACC phosphorylation (Fig. 5B). Incubation with AICAR robustly promoted AMPK $\alpha$  phosphorylation, which was further increased in a dose-dependent manner in the presence of 991. AICAR ( $2$  mM) alone readily saturated ACC phosphorylation, whereas RAPTOR phosphorylation was further increased in a dose-dependent manner in the presence of 991 (Fig. 5B). We then attempted to establish a sequential immunoprecipitation protocol to measure activities of AMPK $\gamma$ 1- and AMPK $\gamma$ 3-containing complexes from small volumes of muscle cell/tissue extracts. As shown in Fig. 5C and consistent with our earlier observations (Fig. 2, A and C), incubation of lysates with AMPK $\gamma$ 3 antibody (in complex with protein G-Sepharose) selectively immunoprecipitated AMPK $\gamma$ 3-containing com-

Fig. 5. 991 treatment enhances AMPK activity induced by AICAR in C<sub>2</sub>C<sub>12</sub> muscle cells.

**A:** IB analysis was performed with  $20$   $\mu\text{g}$  lysates from C<sub>2</sub>C<sub>12</sub> myoblasts or myotubes with the indicated antibodies. Representative blots are shown;  $n = 2$  for each condition. Creatine kinase was used as a differentiation marker. **B:** IB analysis was performed with  $20$   $\mu\text{g}$  lysates from C<sub>2</sub>C<sub>12</sub> myotubes untreated (control) or treated with AICAR ( $2$  mM) in the presence or absence of the indicated dose of 991 for  $30$  min. Representative blots are shown;  $n = 3$  for each condition. **C:** schematic representation of the sequential immunoprecipitation protocol for the isolation of AMPK $\gamma$ 3 and AMPK $\gamma$ 1 from tissue lysates (*top*). AMPK $\gamma$ 3 and AMPK $\gamma$ 1 immune complexes were sequentially immunoprecipitated from  $200$   $\mu\text{g}$  mouse gastrocnemius muscle lysate protein. IB analysis was then performed on the input ( $10\% = 20$   $\mu\text{g}$ ), the SN ( $10\% = 20$   $\mu\text{g}$ ), and the immunoprecipitates (IP) from the first and second immunoprecipitation using the indicated antibodies (*bottom*). **D:** AMPK $\gamma$ 3 and AMPK $\gamma$ 1 activity (in duplicate) from  $200$   $\mu\text{g}$  C<sub>2</sub>C<sub>12</sub> myotube lysate protein were measured after sequential immunoprecipitations as described in MATERIALS AND METHODS. Results are expressed as means  $\pm$  SE;  $n = 3$ . \*Significance of AICAR versus the respective control condition ( $0$  or  $10$   $\mu\text{M}$  991); #significance of 991 versus the respective control ( $0$  or  $2$  mM AICAR).  $P < 0.05$ .



plexes but not AMPK $\gamma$ 1-containing complexes. Using the supernatant after AMPK $\gamma$ 3 immunodepletion, AMPK $\gamma$ 1-containing complexes were efficiently immunoprecipitated (Fig. 5C). Activity of both AMPK $\gamma$ 1- and AMPK $\gamma$ 3-containing complexes tended to increase after treatment with 991 alone at a concentration of 10  $\mu$ M (Fig. 5, D and E), consistent with the results of AMPK $\alpha$  phosphorylation (Fig. 5B). In contrast, 991 treatment only further enhanced the effect of AICAR on AMPK activity of AMPK $\gamma$ 1- and AMPK $\gamma$ 3-containing complexes at a concentration of 10  $\mu$ M (Fig. 5, D and E).

**Effect of dual compound treatment on AMPK $\gamma$ 1/ $\gamma$ 3 activity and glucose transport in mouse skeletal muscle.** We tested whether dual compound treatment further stimulated AMPK activity and glucose transport in mouse skeletal muscle. Isolated EDL muscle was incubated in the presence or absence of 991 with or without AICAR. 991 alone increased the phosphorylation of AMPK $\alpha$  and RAPTOR in a dose-dependent manner, whereas 5  $\mu$ M 991 already increased phosphorylation of ACC at maximum levels. AICAR (0.3 mM) alone robustly increased the phosphorylation of AMPK $\alpha$ , ACC, and RAPTOR (Fig. 6A). Cotreatment of EDL with 991 and AICAR further promoted the phosphorylation of AMPK $\alpha$  and RAPTOR compared with single treatments. Incubation of EDL with 991 (5 and 30  $\mu$ M) caused an almost twofold increase in 2-deoxyglucose transport, as previously described (22), although there was no dose-response

effect under the condition we tested. AICAR also tended to increase 2-deoxyglucose transport  $\sim$ 1.5-fold (not statistically significant), which was further enhanced ( $\sim$ 1.5-fold) by 991 (5 and 30  $\mu$ M; Fig. 6B). AMPK $\gamma$ 1-containing activity was increased ( $\sim$ 2.5-fold) with 30  $\mu$ M 991 (Fig. 6C). While AICAR (0.3 mM) alone showed no significant increase, when coincubated with a lower concentration of 991 (5  $\mu$ M but not 30  $\mu$ M), it significantly promoted AMPK $\gamma$ 1-containing activity compared with single 5  $\mu$ M 991 treatment (Fig. 6C). 991 (30  $\mu$ M) caused a significant increase in AMPK $\gamma$ 3-containing activity. AMPK $\gamma$ 3-containing activity was robustly increased by incubation with AICAR ( $\sim$ 4-fold) and was further increased in the presence of 991 (5 and 30  $\mu$ M; Fig. 6D).

**Additive effects of 991 treatment and contraction on AMPK activity and glucose transport in mouse skeletal muscle.** We next examined if the effect of 991 on AMPK activity and glucose transport could be further stimulated when combined with contraction in mouse skeletal muscle *ex vivo*. We have previously observed that intense tetanic contraction [100 Hz, which causes a robust ( $\sim$ 4-fold) increase in muscle glucose uptake] failed to elicit an additive effect when combined with various doses of 991 (22). It might be the case that under such a high-intensity contraction protocol, glucose transport through GLUT4 translocation got saturated and thus there was no room for further increase when 991 was added. Therefore, we used

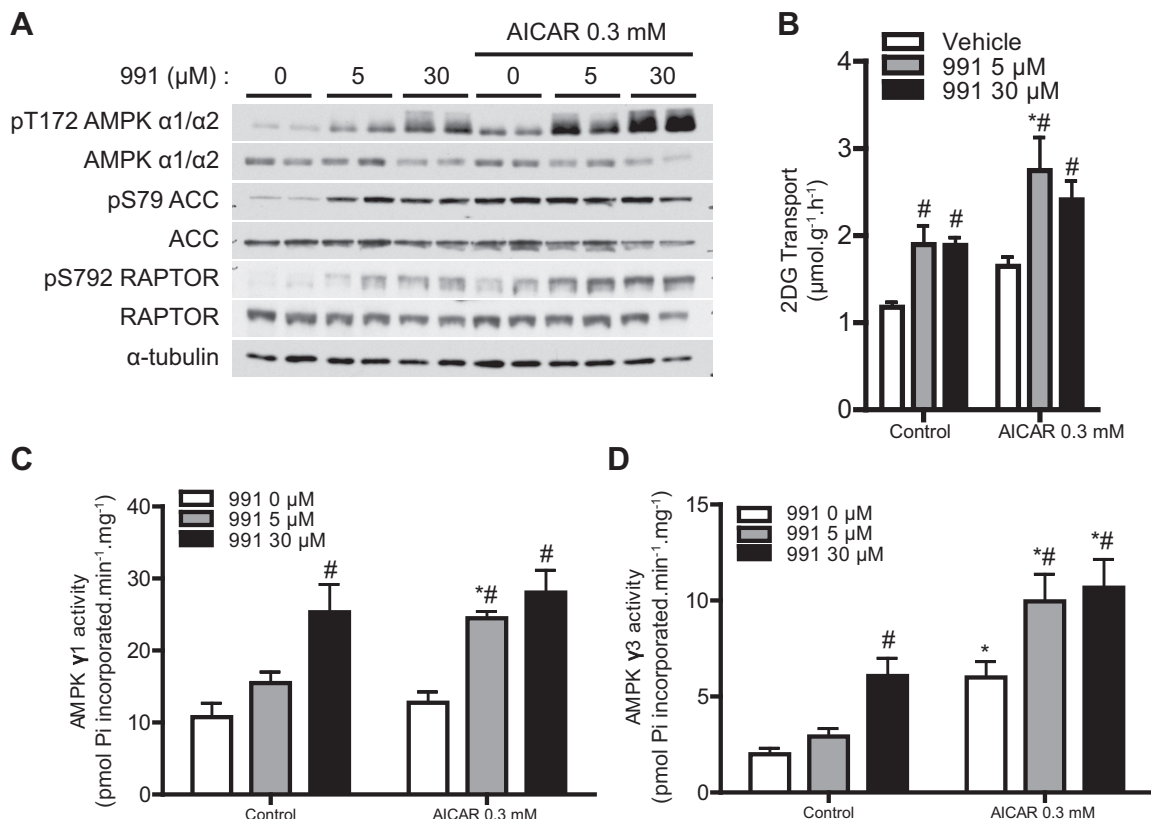


Fig. 6. Dual compound treatment further increases AMPK activity and glucose transport in mouse skeletal muscle compared with single compound treatment. EDL muscles from C57BL/6 mice were isolated and incubated in the presence or absence of the indicated compounds (and concentrations) for 1 h followed by the 2-deoxyglucose (2-DG) transport assay as described in MATERIALS AND METHODS. A: IB analysis was performed with 20  $\mu$ g muscle lysate protein with the indicated antibodies. Representative blots are shown;  $n = 4$ . B: 2-DG transport ( $n = 4$ ). C and D: AMPK complexes were immunoprecipitated from 200  $\mu$ g muscle lysate protein using either AMPK $\gamma$ 1 (C) or AMPK $\gamma$ 3 (D) antibody. AMPK activity was measured as described in MATERIALS AND METHODS. Results are expressed as means  $\pm$  SE;  $n = 4$ . \*Significance of AICAR versus the respective control condition (0, 5, or 30  $\mu$ M 991); #significance between 991 and the respective vehicle condition (0 or 0.3 mM AICAR).  $P < 0.05$ .



a low/moderate-intensity twitch (1 Hz) contraction protocol, which causes submaximal glucose transport (42). Isolated mouse EDL muscle was incubated with 991 (5  $\mu$ M) for 45 min, and the muscle was then either electrically stimulated (contraction) or further incubated (control) with or without 991 for 15 min. As shown in Fig. 7A, 991 treatment alone modestly increased AMPK $\alpha$  phosphorylation (longer exposure), which was associated with increases in the phosphorylation of ACC and RAPTOR. Contraction increased AMPK phosphorylation much more robustly compared with 991, whereas the increases in phosphorylation of ACC and RAPTOR were similar to those induced by 991. Treatment with 991 and electrical stimulation in combination resulted in modest additional increases in the phosphorylation of AMPK $\alpha$  and RAPTOR (Fig. 7A). Treatment with 991 or contraction alone significantly increased 2-deoxyglucose transport to the same extent ( $\sim$ 2-fold; Fig. 7B). When 991 treatment and contraction were combined, an additive increase in 2-deoxyglucose transport was observed.

We next assessed if the activities of AMPK $\gamma$ 1- and AMPK $\gamma$ 3-containing complexes were increased in response to contraction and whether the increase was enhanced when combined with 991 treatment. 991 treatment resulted in a modest but nonsignificant increase in AMPK activity of AMPK $\gamma$ 1- and AMPK $\gamma$ 3-containing complexes (Fig. 7, C and D). Contraction robustly increased activities of both AMPK $\gamma$ 1- and AMPK $\gamma$ 3-containing complexes, which were further in-

creased when 991 treatment and contraction were combined (Fig. 7, C and D).

## DISCUSSION

There has been keen interest in developing AMPK-activating drugs for therapeutic use in treating metabolic diseases such as type 2 diabetes. AMPK is involved in a plethora of biological/physiological events, which could partly be explained by the fact that AMPK exists theoretically as 12 distinct trimeric complexes (excluding splice variants) that are uniquely distributed across different cell types/tissues. Therefore, activators that specifically target different isoforms/complexes of AMPK hold promise in drug development for the treatment of specific symptoms/disorders, possibly with fewer off-target effects. Isoform-specific activators also represent valuable research tools to study AMPK functions in intact cells/tissues. Although several studies have reported expression profiles of AMPK isoforms/complexes in mouse skeletal muscles (43), the specificity of the antibodies used was not perfectly documented and validated. In the present study, we first carried out a comprehensive characterization/validation of AMPK isoform-specific antibodies for immunoblot analysis and immunoprecipitation using tissue extracts from AMPK isoform-specific KO mouse models as negative controls. Taking advantage of the validated antibodies, we also assessed the

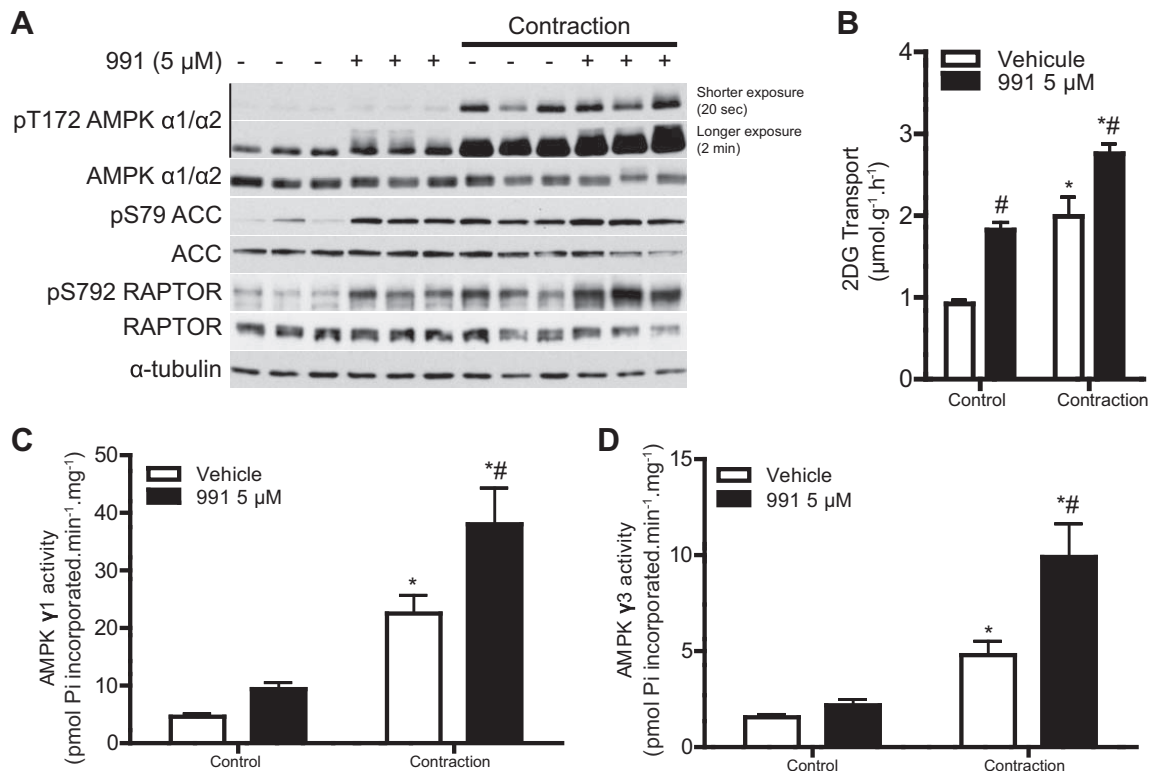


Fig. 7. 991 and contraction additively increase AMPK activity and glucose transport in mouse skeletal muscle. EDL muscles from C57BL/6 mice were isolated and incubated in the presence or absence of 991 (5  $\mu$ M) for 45 min. Muscles were then either electrically stimulated to induce contraction or further incubated (no electrical stimulation) for 15 min. 2-DG transport was measured during the last 10 min of contraction or incubation as described in MATERIALS AND METHODS. A: IB analysis was performed on 20  $\mu$ g muscle lysate protein with the indicated antibodies. Representative blots are shown;  $n = 8$ . Shorter (20 s) and longer (2 min) exposure indicate the exposure time with X-ray film. B: 2-DG transport ( $n = 8$ ). C and D: AMPK complexes were immunoprecipitated from 200  $\mu$ g muscle lysate protein using either AMPK $\gamma$ 1 (C) or AMPK $\gamma$ 3 (D) antibody. AMPK activity was measured as described in MATERIALS AND METHODS. Results are expressed as means  $\pm$  SE. \*Significance of contraction versus the respective control condition (0 or 5  $\mu$ M 991); #significance of 991 versus the respective vehicle condition with or without 1-Hz contraction.  $P < 0.05$ .

expression of AMPK isoforms in different mouse skeletal muscles and other tissues. It has been controversial whether the AMPK  $\gamma 2$ -isoform is expressed in muscle. One study showed the presence/detection of AMPK $\gamma 2$  in mouse skeletal muscle tissues assessed by immunoblot analysis but failed to detect it in AMPK complexes associated with  $\alpha 1/\alpha 2$  or  $\beta 1/\beta 2$  (43). Using a newly generated AMPK $\gamma 2$ -deficient mouse model, we showed that the full-length AMPK $\gamma 2$  variant is not expressed at detectable levels in mouse hindlimb skeletal muscles. We speculate that the previously detected immunoreactivity was nonspecific. Using AMPK $\gamma 3$  KO muscle lysates, we also confirmed that AMPK $\gamma 3$  is only detectable in skeletal muscle and is expressed in glycolytic (e.g., EDL) and glycolytic-oxidative mixed (e.g., gastrocnemius) but not oxidative (e.g., soleus) mouse muscles, as previously described (2, 49). Moreover, we observed that AMPK $\gamma 3$  (as well as AMPK $\alpha 2$  and AMPK $\beta 2$ ) is not detectable in C<sub>2</sub>C<sub>12</sub> myoblasts but is induced upon differentiation into myotubes. Although the molecular mechanism by which AMPK $\gamma 3$  expression is controlled remains unknown, the involvement of coactivator-associated arginine methyltransferase 1/protein arginine methyltransferase 4 has been shown in the regulation of AMPK $\gamma 3$  expression and that of other genes involved in glycogen metabolism in C<sub>2</sub>C<sub>12</sub> muscle cells (44).

We (8, 11, 41) and others (35) have recently shown that dual AMPK-activating compound treatment (A769662 and AICAR) resulted in additive/synergistic increases in cellular AMPK activity and downstream physiological effects such as the inhibition of lipogenesis in hepatocytes (8) and glucose transport in cardiomyocytes (41). One limitation of using A769662 in dual treatment is that it is only effective in cells where  $\beta 1$  is predominantly expressed. Although both A769662 and 991 bind to the ADaM site (4, 23, 46), 991 is much more potent than A769662 (in assays monitoring allosteric activation and protection against dephosphorylation using recombinant  $\alpha 2\beta 1\gamma 1$  complexes) (46). However, it should be noted that even though 991 increases AMPK activity of both  $\beta 1$ - and  $\beta 2$ -containing complexes, it binds to  $\beta 1$ -containing complexes  $\sim 10$  times stronger than to  $\beta 2$ -containing complexes in cell-free assays (46) (which possibly explains its weaker activation of  $\beta 2$ -complexes versus  $\beta 1$ -complexes in vitro and also in intact cells; Fig. 4E). Nonetheless, we demonstrated that 991 increases AMPK activity of both  $\beta 1$ - and  $\beta 2$ -containing complexes and thus, as anticipated, it also activates AMPK $\alpha 1$ -/ $\alpha 2$ - and AMPK $\gamma 1$ -/ $\gamma 2$ -containing complexes in hepatocytes. We also showed that the effects of 991 on cellular AMPK activity in hepatocytes can be further enhanced (as judged by a dose-dependent increase in RAPTOR phosphorylation) in the presence of AICAR or C13. Moreover, we showed here that  $\gamma 1$ - and  $\gamma 3$ -complex activity as well as glucose transport were additively/synergistically enhanced when 991 and AICAR were combined, indicating that the dual treatment approach can work in various different cell systems irrespective of the composition of the  $\beta$ -subunit.

Recently, the effect of another small-molecule AMPK activator, PT-1, on AMPK activity and glucose uptake in mouse skeletal muscle was investigated (20). An unexpected observation reported in that study (20) was that although PT-1 was initially thought to activate AMPK by direct binding between the kinase and auto-inhibitory domains of the  $\alpha$ -subunit (28), Jensen et al. showed that PT-1 failed to activate AMPK

directly but rather indirectly activated it by inhibiting the respiratory chain and increasing cellular AMP-to-ATP and/or ADP-to-ATP ratios (20). Nevertheless, they showed that PT-1-activated AMPK associated only with AMPK $\gamma 1$ - and not AMPK $\gamma 3$ -containing complexes, which resulted in no significant increase in ACC2 and TBC1 domain family member 1 phosphorylation and no increase in glucose transport in mouse skeletal muscle. Although the mechanism by which PT-1 preferentially activates AMPK $\gamma 1$ -containing complexes (particularly in mouse skeletal muscle) is unknown, the study by Jensen et al. (20) indicated that activation of AMPK $\gamma 3$ -containing complexes is required to increase glucose transport in intact mouse skeletal muscle. It has been well documented that AICAR stimulates glucose transport in mouse skeletal muscle in an AMPK-dependent mechanism based on a study using AMPK-deficient genetic mouse models (30). It has recently been shown that 991 treatment stimulates glucose transport in mouse skeletal muscle and that the 991-mediated increase in glucose transport was absent in AMPK $\alpha 1/\alpha 2$  KO cultured myotubes (22). We showed in mouse EDL muscle that both 5  $\mu$ M 991 and contraction significantly increased glucose transport to a similar extent ( $\sim 2$ -fold), although AMPK activity of  $\gamma 1$ - and  $\gamma 3$ -containing complexes was much greater in contracted muscles. However, it should be noted that 991 increases AMPK activity not only through protection against dephosphorylation leading to AMPK $\alpha$  T172 phosphorylation-dependent activation but also via potent allosteric stimulation through the ADaM site (46). Given that measurement of AMPK activity in vitro only assesses phosphorylation (T172)-dependent activation, it is possible that  $\alpha 2\beta 2\gamma 1/\alpha 2\beta 2\gamma 3$  complexes are allosterically activated in intact muscle by 991. This may at least partly explain why phosphorylation of bona fide AMPK substrates (ACC and RAPTOR) and the extent of glucose transport were comparable between 991- and contraction-stimulated muscles. It would be of interest to identify  $\gamma 1$ - and  $\gamma 3$ -complex-specific substrate(s) (if any) and monitor their cellular activity upon 991 treatment through its/their phosphorylation by immunoblot analysis. In addition, whether AMPK $\gamma 3$  is necessary for 991-stimulated glucose uptake, like AICAR, would be interesting to test using an AMPK $\gamma 3$  KO mouse model.

In summary, we demonstrated that 991 treatment increases AMPK activity of both AMPK $\gamma 1$ - and AMPK $\gamma 3$ -containing complexes in mouse skeletal muscle. We also showed that dual treatment with 991 and AICAR or 991 and contraction augments AMPK activation and glucose transport compared with single treatment. The dual treatment approach has proven to be useful to robustly activate cellular AMPK, thereby facilitating the identification of new AMPK substrates in different cell types (7, 17). Next, it would be of interest to test if dual treatment (combinations of various compounds or compound plus exercise/contraction) also promotes AMPK activity and glucose transport in vivo.

#### ACKNOWLEDGMENTS

The authors thank Alexander Chibalin and Juleen Zierath (Karolinska Institutet, Stockholm, Sweden), Benoit Viollet (Institut National de la Santé et de la Recherche Médicale, Institut Cochin, Paris, France), and Gregory Steinberg (McMaster University, Hamilton, ON, Canada) for the kind provision of tissues or lysates from AMPK-deficient mouse models. The authors thank Roger Hunter for hepatocyte isolation.

Present address of Y.-C. Lai: MRC Protein Phosphorylation and Ubiquitylation Unit, College of Life Sciences, University of Dundee, Dundee, UK.

## GRANTS

A. Yavari is supported by the United Kingdom (UK) National Institute for Health Research. A. Yavari (RE/08/004) and H. Ashrafiyan acknowledge support from the British Health Foundation Centre of Research Excellence (Oxford, UK). Y.-C. Lai was supported by the Fonds de la Recherche Scientifique (FNRS; Belgium) and Interuniversity Attraction Poles (IAP) Programme of the Belgian Science Policy (P7/13). S. Kviklyte was also supported by IAP programme P7/13. This work was funded by IAP Belgian Science Policy (P7/13), by the Directorate General Higher Education and Scientific Research French Community of Belgium, and by the FNRS (Belgium) under Grants 3.4518.11 and T.0008.15. T. E. Jensen and A. L. B. Madsen were supported by a Novo Nordisk Foundation Excellence project grant and the Danish Diabetes Academy. E. A. Richter was supported by a Grant 4183-00249B from the Danish Council for Independent Research/Medicine.

## DISCLOSURES

L. Bultot, C. Collodet, M. Deak, and K. Sakamoto are employees of the Nestlé Institute of Health Sciences SA (Switzerland).

## AUTHOR CONTRIBUTIONS

L.B., T.E.J., and K.S. conception and design of research; L.B., T.E.J., Y.-C.L., A.L.M., C.C., S.K., M.D., A.Y., M.F., S.G., and M.B. performed experiments; L.B., T.E.J., Y.-C.L., A.L.M., C.C., E.A.R., and K.S. analyzed data; L.B., T.E.J., Y.-C.L., C.C., A.Y., M.F., H.A., M.R., E.A.R., and K.S. interpreted results of experiments; L.B. and C.C. prepared figures; L.B. and K.S. drafted manuscript; L.B., T.E.J., Y.-C.L., M.D., A.Y., M.F., M.R., E.A.R., and K.S. edited and revised manuscript; L.B., T.E.J., Y.-C.L., A.L.M., C.C., S.K., M.D., A.Y., M.F., S.G., M.B., H.A., M.R., E.A.R., and K.S. approved final version of manuscript.

## REFERENCES

- Alessi DR, Sakamoto K, Bayascas JR. LKB1-dependent signaling pathways. *Annu Rev Biochem* 75: 137–163, 2006.
- Barnes BR, Marklund S, Steiler TL, Walter M, Hjalms G, Amarger V, Mahlapuu M, Leng Y, Johansson C, Galuska D, Lindgren K, Abrink M, Stapleton D, Zierath JR, Andersson L. The 5'-AMP-activated protein kinase gamma3 isoform has a key role in carbohydrate and lipid metabolism in glycolytic skeletal muscle. *J Biol Chem* 279: 38441–38447, 2004.
- Birk JB, Wojtaszewski JF. Predominant alpha2/beta2/gamma3 AMPK activation during exercise in human skeletal muscle. *J Physiol* 577: 1021–1032, 2006.
- Calabrese MF, Rajamohan F, Harris MS, Caspers NL, Magyar R, Withka JM, Wang H, Borzilleri KA, Sahasrabudhe PV, Hoth LR, Geoghegan KF, Han S, Brown J, Subashi TA, Reyes AR, Frisbie RK, Ward J, Miller RA, Landro JA, Londregan AT, Carpino PA, Cabral S, Smith AC, Conn EL, Cameron KO, Qiu X, Kurumbail RG. Structural basis for AMPK activation: natural and synthetic ligands regulate kinase activity from opposite poles by different molecular mechanisms. *Structure* 22: 1161–1172, 2014.
- Cheung PC, Salt IP, Davies SP, Hardie DG, Carling D. Characterization of AMP-activated protein kinase gamma-subunit isoforms and their role in AMP binding. *Biochem J* 346: 659–669, 2000.
- Cool B, Zinker B, Chiou W, Kifle L, Cao N, Perham M, Dickinson R, Adler A, Gagne G, Iyengar R, Zhao G, Marsh K, Kym P, Jung P, Camp HS, Frevert E. Identification and characterization of a small molecule AMPK activator that treats key components of type 2 diabetes and the metabolic syndrome. *Cell Metab* 3: 403–416, 2006.
- Ducommun S, Deak M, Sumpton D, Ford RJ, Nunez Galindo A, Kussmann M, Viollet B, Steinberg GR, Foretz M, Dayon L, Morrice NA, Sakamoto K. Motif affinity and mass spectrometry proteomic approach for the discovery of cellular AMPK targets: identification of mitochondrial fission factor as a new AMPK substrate. *Cell Signal* 27: 978–988, 2015.
- Ducommun S, Ford RJ, Bultot L, Deak M, Bertrand L, Kemp BE, Steinberg GR, Sakamoto K. Enhanced activation of cellular AMPK by dual-small molecule treatment: AICAR and A769662. *Am J Physiol Endocrinol Metab* 306: E688–E696, 2014.
- Dzambo N, van Denderen BJ, Hevener AL, Jorgensen SB, Honeyman J, Galic S, Chen ZP, Watt MJ, Campbell DJ, Steinberg GR, Kemp BE. AMPK beta1 deletion reduces appetite, preventing obesity and hepatic insulin resistance. *J Biol Chem* 285: 115–122, 2010.
- Foretz M, Hebrard S, Guihard S, Leclerc J, Do Cruzeiro M, Hamard G, Niedergang F, Gaudry M, Viollet B. The AMPKgamma1 subunit plays an essential role in erythrocyte membrane elasticity, and its genetic inactivation induces splenomegaly and anemia. *FASEB J* 25: 337–347, 2011.
- Foretz M, Hebrard S, Leclerc J, Zarrinpashneh E, Soty M, Mithieux G, Sakamoto K, Andreelli F, Viollet B. Metformin inhibits hepatic gluconeogenesis in mice independently of the LKB1/AMPK pathway via a decrease in hepatic energy state. *J Clin Invest* 120: 2355–2369, 2010.
- Goossens O, McBride A, Hawley SA, Ross FA, Shpiro N, Foretz M, Viollet B, Hardie DG, Sakamoto K. Mechanism of action of A-769662, a valuable tool for activation of AMP-activated protein kinase. *J Biol Chem* 282: 32549–32560, 2007.
- Gowans GJ, Hardie DG. AMPK: a cellular energy sensor primarily regulated by AMP. *Biochem Soc Trans* 42: 71–75, 2014.
- Hardie DG, Ross FA, Hawley SA. AMPK: a nutrient and energy sensor that maintains energy homeostasis. *Nat Rev Mol Cell Biol* 13: 251–262, 2012.
- Hardie DG, Sakamoto K. AMPK: a key sensor of fuel and energy status in skeletal muscle. *Physiology (Bethesda)* 21: 48–60, 2006.
- Henriksson E, Jones HA, Patel K, Pegg M, Morrice N, Sakamoto K, Goossens O. The AMPK-related kinase SIK2 is regulated by cAMP via phosphorylation at Ser358 in adipocytes. *Biochem J* 444: 503–514, 2012.
- Hoffman NJ, Parker BL, Chaudhuri R, Fisher-Wellman KH, Kleinert M, Humphrey SJ, Yang P, Holliday M, Trefely S, Fazakerley DJ, Stockli J, Burchfield JG, Jensen TE, Jothi R, Kiens B, Wojtaszewski JF, Richter EA, James DE. Global phosphoproteomic analysis of human skeletal muscle reveals a network of exercise-regulated kinases and AMPK substrates. *Cell Metab* 22: 922–935, 2015.
- Hunter RW, Foretz M, Bultot L, Fullerton MD, Deak M, Ross FA, Hawley SA, Shpiro N, Viollet B, Barron D, Kemp BE, Steinberg GR, Hardie DG, Sakamoto K. Mechanism of action of compound-13: an alpha1-selective small molecule activator of AMPK. *Chem Biol* 21: 866–879, 2014.
- Hunter RW, Treebak JT, Wojtaszewski JF, Sakamoto K. Molecular mechanism by which AMP-activated protein kinase activation promotes glycogen accumulation in muscle. *Diabetes* 60: 766–774, 2011.
- Jensen TE, Ross FA, Kleinert M, Sylow L, Knudsen JR, Gowans GJ, Hardie DG, Richter EA. PT-1 selectively activates AMPK-gamma1 complexes in mouse skeletal muscle, but activates all three gamma subunit complexes in cultured human cells by inhibiting the respiratory chain. *Biochem J* 467: 461–472, 2015.
- Jensen TE, Sylow L, Rose AJ, Madsen AB, Angin Y, Maarbjerg SJ, Richter EA. Contraction-stimulated glucose transport in muscle is controlled by AMPK and mechanical stress but not sarcoplasmic reticulum Ca<sup>2+</sup> release. *Mol Metab* 3: 742–753, 2014.
- Lai YC, Kviklyte S, Vertommen D, Lantier L, Foretz M, Viollet B, Hallen S, Rider MH. A small-molecule benzimidazole derivative that potentially activates AMPK to increase glucose transport in skeletal muscle: comparison with effects of contraction and other AMPK activators. *Biochem J* 460: 363–375, 2014.
- Langendorf CG, Kemp BE. Choreography of AMPK activation. *Cell Res* 25: 5–6, 2015.
- Langendorf CG, Ngoei KR, Scott JW, Ling NX, Issa SM, Gorman MA, Parker MW, Sakamoto K, Oakhill JS, Kemp BE. Structural basis of allosteric and synergistic activation of AMPK by furan-2-phosphonic derivative C2 binding. *Nat Commun* 7: 10912, 2016.
- Liu Y, Lai YC, Hill EV, Tyteca D, Carpentier S, Ingvaldsen A, Vertommen D, Lantier L, Foretz M, Dequiedt F, Courtot PJ, Erneux C, Viollet B, Shepherd PR, Tavaré JM, Jensen J, Rider MH. Phosphatidylinositol 3-phosphate 5-kinase (PIKfyve) is an AMPK target participating in contraction-stimulated glucose uptake in skeletal muscle. *Biochem J* 455: 195–206, 2013.
- Musi N, Fujii N, Hirshman MF, Ekberg I, Froberg S, Ljungqvist O, Thorell A, Goodyear LJ. AMP-activated protein kinase (AMPK) is activated in muscle of subjects with type 2 diabetes during exercise. *Diabetes* 50: 921–927, 2001.
- Oakhill JS, Steel R, Chen ZP, Scott JW, Ling N, Tam S, Kemp BE. AMPK is a direct adenylate charge-regulated protein kinase. *Science* 332: 1433–1435, 2011.



28. Pang T, Zhang ZS, Gu M, Qiu BY, Yu LF, Cao PR, Shao W, Su MB, Li JY, Nan FJ, Li J. Small molecule antagonizes autoinhibition and activates AMP-activated protein kinase in cells. *J Biol Chem* 283: 16051–16060, 2008.
29. Patel K, Foretz M, Marion A, Campbell DG, Gourlay R, Boudaba N, Tournier E, Titchenell P, Pegg M, Deak M, Wan M, Kaestner KH, Goransson O, Viollet B, Gray NS, Birnbaum MJ, Sutherland C, Sakamoto K. The LKB1-salt-inducible kinase pathway functions as a key gluconeogenic suppressor in the liver. *Nat Commun* 5: 4535, 2014.
30. Richter EA, Hargreaves M. Exercise, GLUT4, and skeletal muscle glucose uptake. *Physiol Rev* 93: 993–1017, 2013.
31. Sakamoto K, Goransson O, Hardie DG, Alessi DR. Activity of LKB1 and AMPK-related kinases in skeletal muscle: effects of contraction, phenformin, and AICAR. *Am J Physiol Endocrinol Metab* 287: E310–E317, 2004.
32. Sakamoto K, Holman GD. Emerging role for AS160/TBC1D4 and TBC1D1 in the regulation of GLUT4 traffic. *Am J Physiol Endocrinol Metab* 295: E29–E37, 2008.
33. Sakamoto K, Zarrinpashneh E, Budas GR, Pouleur AC, Dutta A, Prescott AR, Vanoverschelde JL, Ashworth A, Jovanovic A, Alessi DR, Bertrand L. Deficiency of LKB1 in heart prevents ischemia-mediated activation of AMPK $\alpha$ 2 but not AMPK $\alpha$ 1. *Am J Physiol Endocrinol Metab* 290: E780–E788, 2006.
34. Sanders MJ, Ali ZS, Hegarty BD, Heath R, Snowden MA, Carling D. Defining the mechanism of activation of AMP-activated protein kinase by the small molecule A-769662, a member of the thienopyridone family. *J Biol Chem* 282: 32539–32548, 2007.
35. Scott JW, Ling N, Issa SM, Dite TA, O'Brien MT, Chen ZP, Galic S, Langendorf CG, Steinberg GR, Kemp BE, Oakhill JS. Small molecule drug A-769662 and AMP synergistically activate naive AMPK independent of upstream kinase signaling. *Chem Biol* 21: 619–627, 2014.
36. Scott JW, van Denderen BJ, Jorgensen SB, Honeyman JE, Steinberg GR, Oakhill JS, Iseli TJ, Koay A, Gooley PR, Stapleton D, Kemp BE. Thienopyridone drugs are selective activators of AMP-activated protein kinase beta1-containing complexes. *Chem Biol* 15: 1220–1230, 2008.
37. Shaw RJ, Lamia KA, Vasquez D, Koo SH, Bardeesy N, Depinho RA, Montminy M, Cantley LC. The kinase LKB1 mediates glucose homeostasis in liver and therapeutic effects of metformin. *Science* 310: 1642–1646, 2005.
38. Steinberg GR, Kemp BE. AMPK in health and disease. *Physiol Rev* 89: 1025–1078, 2009.
39. Steinberg GR, O'Neill HM, Dzamko NL, Galic S, Naim T, Koopman R, Jorgensen SB, Honeyman J, Hewitt K, Chen ZP, Schertzer JD, Scott JW, Koentgen F, Lynch GS, Watt MJ, van Denderen BJ, Campbell DJ, Kemp BE. Whole body deletion of AMP-activated protein kinase  $\beta$ 2 reduces muscle AMPK activity and exercise capacity. *J Biol Chem* 285: 37198–37209, 2010.
40. Stephenne X, Foretz M, Taleux N, van der Zon GC, Sokal E, Hue L, Viollet B, Guigas B. Metformin activates AMP-activated protein kinase in primary human hepatocytes by decreasing cellular energy status. *Diabetologia* 54: 3101–3110, 2011.
41. Timmermans AD, Balteau M, Gelinis R, Renguet E, Ginion A, de Meester C, Sakamoto K, Balligand JL, Bontemps F, Vanoverschelde JL, Horman S, Beauloye C, Bertrand L. A-769662 potentiates the effect of other AMP-activated protein kinase activators on cardiac glucose uptake. *Am J Physiol Heart Circ Physiol* 306: H1619–H1630, 2014.
42. Toyoda T, Nakao K, Ebihara K, Masuzaki H, Hosoda K, Sato K, Fushiki T, Nakao K, Hayashi T. Low-intensity contraction activates the  $\alpha$ 1-isoform of 5'-AMP-activated protein kinase in rat skeletal muscle. *Am J Physiol Endocrinol Metab* 290: E583–E590, 2006.
43. Treebak JT, Birk JB, Hansen BF, Olsen GS, Wojtaszewski JF. A-769662 activates AMPK  $\beta$ 1-containing complexes but induces glucose uptake through a PI3-kinase-dependent pathway in mouse skeletal muscle. *Am J Physiol Cell Physiol* 297: C1041–C1052, 2009.
44. Wang SC, Dowhan DH, Eriksson NA, Muscat GE. CARM1/PRMT4 is necessary for the glycogen gene expression programme in skeletal muscle cells. *Biochem J* 444: 323–331, 2012.
45. Wu J, Puppala D, Feng X, Monetti M, Lapworth AL, Geoghegan KF. Chemoproteomic analysis of inter-tissue and inter-species isoform diversity of AMP-activated protein kinase (AMPK). *J Biol Chem* 288: 35904–35912, 2013.
46. Xiao B, Sanders MJ, Carmena D, Bright NJ, Haire LF, Underwood E, Patel BR, Heath RB, Walker PA, Hallen S, Giordanetto F, Martin SR, Carling D, Gamblin SJ. Structural basis of AMPK regulation by small molecule activators. *Nat Commun* 4: 3017, 2013.
47. Xiao B, Sanders MJ, Underwood E, Heath R, Mayer FV, Carmena D, Jing C, Walker PA, Eccleston JF, Haire LF, Saiu P, Howell SA, Aasland R, Martin SR, Carling D, Gamblin SJ. Structure of mammalian AMPK and its regulation by ADP. *Nature* 472: 230–233, 2011.
48. Yavari A, Stocker CJ, Ghaffari S, Wargent ET, Steeples V, Czibik G, Pinter K, Bellahcene M, Woods A, Martinez de Morentin PB, Cansell C, Lam BY, Chuster A, Petkevicius K, Nguyen-Tu MS, Martinez-Sanchez A, Pullen TJ, Oliver PL, Stockenhuber A, Nguyen C, Lazdam M, O'Dowd JF, Harikumar P, Toth M, Beall C, Kyriakou T, Parnis J, Sarma D, Katritsis G, Wortmann DD, Harper AR, Brown LA, Willows R, Gandra S, Poncio V, de Oliveira Figueiredo MJ, Qi NR, Peirson SN, McCrimmon RJ, Gereben B, Tretter L, Fekete C, Redwood C, Yeo GS, Heisler LK, Rutter GA, Smith MA, Withers DJ, Carling D, Sternick EB, Arch JR, Cawthorne MA, Watkins H, Ashrafian H. Chronic activation of gamma2 AMPK induces obesity and reduces beta cell function. *Cell Metab* 23: 821–836, 2016.
49. Yu H, Fujii N, Hirshman MF, Pomerleau JM, Goodyear LJ. Cloning and characterization of mouse 5'-AMP-activated protein kinase  $\gamma$ 3 subunit. *Am J Physiol Cell Physiol* 286: C283–C292, 2004.