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Frøsig, Christian; Jensen, Thomas Elbenhardt; Jeppesen, Jacob; Pehmøller, Christian; Treebak, Jonas Thue; Maarbjerg, Stine Just; Kristensen, Jonas Møller; Sylow, Lykke; Alsted, Thomas Junker; Schjerling, Peter; Kiens, Bente; Wojtaszewski, Jørgen; Richter, Erik

Published in:
P L o S One

DOI:
10.1371/journal.pone.0062338

Publication date:
2013

Document version
Publisher's PDF, also known as Version of record

Citation for published version (APA):
AMPK and Insulin Action - Responses to Ageing and High Fat Diet

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Abstract

The 5′-AMP-activated protein kinase (AMPK) is considered “a metabolic master-switch” in skeletal muscle reducing ATP-consuming processes whilst stimulating ATP regeneration. Within recent years, AMPK has also been proposed as a potential target to attenuate insulin resistance, although the exact role of AMPK is not well understood. Here we hypothesized that mice lacking α2AMPK activity in muscle would be more susceptible to develop insulin resistance associated with ageing alone or in combination with high fat diet. Young (~4 month) or old (~18 month) wild type and muscle specific α2AMPK kinase-dead mice on chow diet as well as old mice on 17 weeks of high fat diet were studied for whole body glucose homeostasis (OGTT, ITT and HOMA-IR), insulin signaling and insulin-stimulated glucose uptake in muscle. We demonstrate that high fat diet in old mice results in impaired glucose homeostasis and insulin stimulated glucose uptake in both the soleus and extensor digitorum longus muscle, coinciding with reduced insulin signaling at the level of Akt (pSer473 and pThr308), TBC1D1 (pThr590) and TBC1D4 (pThr642). In contrast to our hypothesis, the impact of ageing and high fat diet on insulin action was not worsened in mice lacking functional α2AMPK in muscle. It is concluded that α2AMPK deficiency in mouse skeletal muscle does not cause muscle insulin resistance in young and old mice and does not exacerbate obesity-induced insulin resistance in old mice suggesting that decreased α2AMPK activity does not increase susceptibility for insulin resistance in skeletal muscle.

Introduction

Insulin resistance in peripheral tissues is a hallmark characteristic of obesity-related type 2 diabetes (T2D). In this context, skeletal muscle is a critical organ constituting ~40% of body weight and contributing the majority of whole body insulin-stimulated glucose disposal [1]. Insulin resistance, associated with high fat feeding or obesity, is believed to be the combined result of chronic low-grade inflammation and accumulation of bio-active lipid species within muscle. In turn, this leads to impairment of insulin signaling to GLUT4 translocation and subsequently insulin-stimulated glucose uptake [2,3]. Consistent with a pathogenic role of intracellular lipid accumulation, genetically engineered mouse models with improved capacity for lipid oxidation in muscle are protected against adverse effects of high fat feeding [4,5]. The AMP-activated protein kinase (AMPK) is activated in response to stimuli that increase the intracellular ratio of AMP/ATP, such as exercise, hypoxia, osmotic stress and ischemia [6–8]. AMPK is referred to as “a metabolic master-switch” due to its general ability to reduce ATP consuming anabolic processes while alternative pathways for ATP regeneration are stimulated [9], including stimulation of lipid oxidation in muscle [10,11]. This has led to the speculation that AMPK activation may protect muscle from high fat feeding induced insulin resistance. To support this, Goodyear et al. demonstrated that overexpression of kinase dead α2AMPK in mice on a FVB mouse background (AMPK Ki),

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resulted in massive exacerbation of muscle insulin resistance in response to 30 weeks of high fat feeding, coinciding with an increase in muscle diacylglycerol content [12]. Interestingly, the AMPK Ki mice on chow diet in that study also showed a tendency towards impaired insulin action (~50% reduction vs. wild type (WT); p = 0.07) whereas normal insulin action has previously been reported in AMPK Ki mice after only 15 weeks of chow diet [13].

This suggests that acute lack of 2AMPK activation may not directly trigger impaired insulin action, but rather lack of 2AMPK activity over time leads to a muscle phenotype that is more susceptible for insulin resistance. This would also explain our previous observation that insulin resistance induced by 12 weeks high fat feeding was not exacerbated in young kinase dead 2AMPK mice on a C57BL/6J background (AMPK KD) [14].

In order to test this interpretation and firmly link AMPK with muscle insulin action, we here hypothesized that old AMPK KD mice would develop insulin resistance on a chow diet (as indicated by observations in AMPK Ki mice). Furthermore, we hypothesized that insulin resistance induced by high fat feeding would be exacerbated in old AMPK KD mice (in contrast to observations young AMPK KD mice).

Materials and Methods

Ethics

All animal experiments were approved by the Danish Animal Experimental Inspectorate (No. 2012-15-2934-00310) in compliance with the European Convention for Protection of Vertebrate Animals Used for Scientific Purposes. All efforts were made to minimize suffering during in vivo experiments. Furthermore, surgery was performed under sodium pentobarbital anesthesia and after surgery animals were sacrificed by cervical dislocation.

Animals

Animals used were age 4.2±0.1 month (Young) or 18.0±0.2 month (Old) C57BL/6J male mice overexpressing a muscle-specific, kinase-dead 2AMPK construct (AMPK KD), as described by Mu et al. [15,16] and corresponding WT littermates. Briefly, the animals overexpress a Lys45-to-Arg mutant of the 2AMPK protein, driven by a heart- and skeletal muscle-specific creatine kinase promoter. Average age.

Diet Treatments

Animals were kept on a 10:14-h light-dark cycle with unlimited access to standard rodent diet (60% [of energy] carbohydrates, 27% protein and 15% fat) and water (CHO groups). In the high fat diet group (FA group) standard chow was replaced 17 weeks prior to terminal experiments with a high fat diet (20% [of energy] carbohydrates, 20% protein and 60% fat) specified as 19.7% [of energy] casein, 54.4% lard, 5.5% soybean oil, 12.3% maltodextrin and 6.7% sucrose. Animals were then maintained on this diet to terminal ages.

Whole Body Glucose Homeostasis

Glucose (OGTT) and insulin (ITT) tolerance tests were performed after a 5 hours fast on separate occasions in all mice, 2–3 weeks prior to terminal experiments. Glucose (2 g/kg body weight) was given by oral gavage and insulin (0.5 units/kg body wt; Actrapid, Novo Nordisk, Bagsvaerd, Denmark) for 10 min and next for an additional 10 min in stimulation buffer with insulin and tracers (0.75 µCi/ml [3H]mannitol [PerkinElmer, NJ, USA]). After incubation, muscles were harvested, washed in ice-cold Krebs-Henseleiinger buffer, blotted on filter paper and frozen in liquid N2 for later analyses.

Body Composition and Metabolic Measurements

In order to test this interpretation and firmly link AMPK with muscle insulin action, we here hypothesized that old AMPK KD mice would develop insulin resistance on a chow diet (as indicated by observations in AMPK Ki mice). Furthermore, we hypothesized that insulin resistance induced by high fat feeding would be exacerbated in old AMPK KD mice (in contrast to observations young AMPK KD mice).

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Body Composition and Metabolic Measurements

1–2 weeks prior to terminal experiments, all mice were weighed, and scanned for body composition (EchoMRI-411; EchoMRI, Houston, TX, USA). Following 48 hours of acclimatization to individual metabolic cages with access to food (either CHO or FA diet) and water ad libitum, the metabolic cages were sealed and O2 uptake and CO2 production were measured for 24 hours (Fasted conditions) using a CaloSys apparatus (TSE Systems GmbH, Bad Homburg, Germany). Next, food was removed from the cages (Fasted conditions) and similar measurements were made for an additional 24 hours. Data for VO2 and RER (VCO2/VO2) was calculated as AUC using a weighted average of time points obtained every hour. Following these measurements mice were returned to standard cages and were housed individually for at least one week prior to terminal experiments.

Muscle Incubation and Glucose Uptake

On the day of terminal experiments, animals were anesthetized intraperitoneally by injection of pentobarbital sodium (6 mg/100 g body wt). In all mice, m. Soleus (SOL) and m. Extensor Digitorum Longus (EDL) were quickly excised and suspended in incubation chambers (Multi Myograph system; Danish Myotechnology, Aarhus, Denmark). Muscles were incubated for 40 min in prebuffer (standard Krebs-Henseleiinger-Ringer buffer with addition of 8 mM Mannitol, 2 mM pyruvate, and 0.01% BSA) at 30°C and oxygenated with a gas mixture containing 95% O2 and 5% CO2. Subsequently, muscles were incubated in stimulation buffer (standard Krebs-Henseleiinger-Ringer buffer with addition of 7 mM Mannitol, 1 mM 2-deoxy-D-glucose (2DG) and 0.01% BSA) containing 500 µCi/ml (3 mCi) insulin (Actrapid; Novo Nordisk, Bagsvaerd, Denmark) for 10 min and next for an additional 10 min in stimulation buffer with insulin and tracers (0.75 µCi/ml [2,2-3H]deoxy-D-glucose and 0.32 µCi/ml [1,14C]mannitol (PerkinElmer, MS, USA)). After incubation, muscles were rinsed, and washed in ice-cold Krebs-Henseleiinger buffer, blotted on filter paper and frozen in liquid N2 for later analyses.

Muscle Lysate Preparation

SOL and EDL muscles were dissected free of tendons and homogenized in ice-cold buffer (10% glycerol, 20 mM sodium pyrophosphate, 1% NP-40, 2 mM PMSF, 150 mM sodium chloride, 50 mM Hepes, 20 mM β-glycerophosphate, 10 mM sodium fluoride, 1 mM EDTA, 1 mM EGTA, 10 µg/ml aprotinin, 3 mM benzamidine, 10 µg/ml leupeptin and 2 mM sodium orthovanadate [pH 7.4]). Homogenates were subsequently rotated end-over-end for 1 hour at 4°C before being centrifuged at 16,000 g at 4°C for 25 min. Supernatants were collected and stored at −80°C for later analyses. Total protein concentrations were determined in triplicates with a coefficient of variance maximum of 5% by the bicinchoninic acid method (Fierce Biotechnology, Rockford, IL, USA).

2-deoxy-D-glucose (2DG) Uptake

2DG uptake was measured by mixing 150 µl muscle lysate protein (~600 µg) in 3 ml scintillation fluid (Ultima Gold; PerkinElmer, Waltham, MA, USA). Subsequently radioactivity
Table 1. Body composition and metabolic characterization.

<table>
<thead>
<tr>
<th></th>
<th>Young CHO WT</th>
<th>Old CHO WT</th>
<th>Old FA WT</th>
<th>Young CHO AMPK KD</th>
<th>Old CHO AMPK KD</th>
<th>Old FA AMPK KD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Body weight (g)</td>
<td>33.5±0.8</td>
<td>39.1±1.0</td>
<td>53.0±1.7</td>
<td>32.9±0.5</td>
<td>38.1±0.6</td>
<td>48.6±1.6</td>
</tr>
<tr>
<td>Fat free mass (g)</td>
<td>29.6±0.9</td>
<td>32.8±0.8</td>
<td>31.8±0.7</td>
<td>30.2±0.5</td>
<td>32.0±0.3</td>
<td>29.6±0.6</td>
</tr>
<tr>
<td>VO2 (ml/hr/kg),</td>
<td>2095±160</td>
<td>2061±97</td>
<td>2263±89</td>
<td>2278±143</td>
<td>2289±164</td>
<td>2334±113</td>
</tr>
<tr>
<td>Fed</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>VO2 (ml/hr/kg),</td>
<td>1945±44§</td>
<td>1848±57§</td>
<td>1958±75§</td>
<td>1958±57§</td>
<td>1995±75§</td>
<td>2115±112§</td>
</tr>
<tr>
<td>Fasted</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RER, Fed</td>
<td>0.97±0.05</td>
<td>0.94±0.03</td>
<td>0.80±0.01</td>
<td>0.94±0.03</td>
<td>0.96±0.04</td>
<td>0.81±0.01</td>
</tr>
<tr>
<td>RER, Fasted</td>
<td>0.85±0.04§</td>
<td>0.84±0.03§</td>
<td>0.72±0.00</td>
<td>0.84±0.02§</td>
<td>0.84±0.02§</td>
<td>0.73±0.03§</td>
</tr>
</tbody>
</table>

Body composition, VO2 and RER were determined in young and old AMPK KD mice and WT littermates on chow diet (CHO) or in old mice after 17 weeks of high fat diet (FA).

Results

Body Composition and Metabolic Characterization

As reported in Table 1, body composition (body weight and fat free mass) and metabolic adaptation (VO2 and RER) to feeding and fasting were similar in young WT and AMPK KD mice. When compared to young mice, old mice on CHO diet had increased body weight (~15%, p<0.05) and fat free mass (~10%, p<0.05), whereas metabolic characteristics were similar (NS). In old mice maintained on a FA diet for 17 weeks, a further increase (~30%, p<0.01) in body weight was observed whereas fat free mass slightly decreased (~5%, p<0.01). This is consistent with a direct effect of the FA diet on body fat accumulation. The FA diet also led to an expected decreased RER in both the fed (~20%, p<0.01) and fasted (~15%, p<0.01) state whereas VO2 was unaltered (NS). Interestingly, old AMPK KD mice were slightly smaller (~5%, p<0.05) than old WT mice independent of diet, as indicated by both reduced body weight and fat free mass.

Whole Body Glucose Homeostasis

In order to investigate the effect of genotype, age and diet on whole body glucose homeostasis, mice underwent an OGTT and ITT. Before and after 20 min in the OGTT plasma insulin concentrations were measured and HOMA-IR index was calculated based on basal values.

OGTT Glucose and Insulin Values

When comparing mice on CHO diet, glucose AUC in response to the OGTT was reduced (~10%, p<0.001) with age but was increased (~10%, p<0.005) in both young and old AMPK KD mice compared to WT (Figure 1A). This should be seen in context of increased (~30%, p<0.05) insulin concentrations in old mice as well as decreased (~20%, p<0.05) insulin concentrations in old AMPK KD mice on CHO diet (Figure 1B). In response to the FA diet, glucose AUC in response to the OGTT was unaltered (NS),
however the FA diet resulted in a marked increase (~100%, p<0.001) in insulin concentrations prior to and during the OGTT (Figure 1A and 1B). Furthermore, in old AMPK KD mice, both basal and 20 min insulin concentrations were reduced (~15%, p<0.05) after the FA diet similar to the observations after the CHO diet (Figure 1B).

**ITT Glucose Values**

Glucose AUC in response to the ITT was not influenced by genotype and did not change with ageing in mice on the CHO diet. However, old mice on FA diet were characterized by increased (~20%, p<0.001) glucose AUC in response to the ITT when compared to the CHO diet (Figure 1C).

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**Figure 1. Characterization of whole body glucose homeostasis.** (A) OGTT (2 g/kg body weight) and (C) ITT (0.5 U/kg body weight) were conducted after 5 hours of fasting. Values are expressed as AUC based on weighted means of all glucose measurements (t = 15, 20, 40, 60, 90 and 120 min). Figure 1B shows plasma insulin concentrations before (0 min) and after 20 min in response to the OGTT. Figure 1D illustrates calculated HOMA-IR index values based on basal glucose and insulin concentrations obtained after 5 hour of fasting. Measurements were made in young and old AMPK KD mice and WT littermates on chow diet (CHO) or in old mice after 17 weeks of high fat diet (FA). $: Main effect of age, p<0.001. #: Main effect of diet, p<0.001. †: Main effect of genotype, p<0.005. ‡: Main effect of time, p<0.001. Values are means ± SE. n = 11–17.

doi:10.1371/journal.pone.0062338.g001
Muscle Insulin-stimulated Glucose Uptake

To investigate the role of genotype, age and diet on muscle glucose uptake, insulin-stimulated glucose uptake was measured in SOL and EDL muscles *in vitro* in response to 500 μU/ml insulin. Compared to a subset of muscles stimulated with 10,000 μU/ml; 500 μU/ml was verified as a sub-maximal insulin stimulus leading to ~70% of maximal insulin-stimulated glucose uptake in both muscles (data not shown). In SOL, insulin stimulation resulted in a ~150% increase (p<0.001) in glucose uptake in both young and old mice on CHO diet when compared to basal. In contrast, in old mice on the FA diet, insulin-stimulated glucose uptake was decreased by ~30% (p<0.001) compared with old mice on CHO diet (Figure 2). In EDL, insulin-stimulated glucose uptake was increased by ~40%, (p<0.001) in young and old mice on CHO diet when compared to basal. Although a trend towards generally greater glucose uptake in the old mice was observed, this did not reach statistical significance (p = 0.09). In contrast, glucose uptake in old mice on FA diet was decreased by ~30% (p<0.001) compared with old mice on CHO diet, but the response to insulin was preserved (Figure 2). Notably, in both SOL and EDL muscles, basal and insulin-stimulated glucose uptake in all groups was independent of genotype (NS).

Muscle Insulin Signaling

To elucidate, if adaptations in insulin-stimulated glucose uptake reflected on muscle insulin signaling, phosphorylation of Akt, TBC1D1 and TBC1D4 was evaluated by western blotting. Representative blots can be viewed in Figure 3A and 3B.

**Figure 2. In vitro glucose uptake.** Basal (0 μU/ml) and insulin (500 μU/ml) stimulated glucose uptake measured *in vitro* in m. Soleus (SOL) and m. Extensor Digitorum Longus (EDL). Measurements were made in young and old AMPK KD mice and WT littermates on chow diet (CHO) or in old mice after 17 weeks of high fat diet (FA). *: Main effect of insulin, p<0.001. #: Main effect of diet, p<0.001. †: Interaction between diet and insulin action, p<0.001. Values are means ± SE. n = 11–15. doi:10.1371/journal.pone.0062338.g002
TBC1D1 Phosphorylation

Complimentary to TBC1D4 although less well characterized, Thr590 phosphorylation on TBC1D1 is believed to inhibit TBC1D1 GAP function to allow for vesicle translocation [21]. When expressed per total TBC1D1 protein, we detected a 50% increase (p < 0.001) in phosphorylation of Thr590 in response to insulin stimulation in both young and old mice on the CHO diet (Figure 7). Furthermore, in both SOL and EDL muscle, old mice on FA diet had reduced (240%, p < 0.001) TBC1D1 phosphorylation in basal and insulin-stimulated muscle and a modest reduction (15%, p < 0.05) in TBC1D1 phosphorylation was observed in AMPK KD mice when compared to WT in EDL muscle. At the protein level, muscles of AMPK KD mice were characterized by markedly reduced (~40%, p < 0.001) TBC1D1 protein content whereas TBC1D1 protein content increased (~30%, p < 0.05) in muscle of old mice after the FA diet (Figure 8C).

Muscle Protein Expression

To further evaluate muscle proteins in relation to insulin action, we next measured protein content of GLUT4, HK2 and TRB-3 in basal SOL and EDL muscle (Figure 8A, 8B and 8F). These proteins were more heavily expressed in SOL compared to EDL, but neither changed in response to ageing nor with FA diet.
However, protein content of both HK2 (~20%, p<0.05) and TRB-3 (~50–80%, p<0.001) was suppressed in AMPK KD mouse muscles.

**Discussion**

In response to high fat feeding, mouse skeletal muscle rapidly (within weeks) becomes insulin resistant at the level of glucose uptake. In many, but not all, studies this is associated with impairment of insulin signaling to GLUT4 translocation, detectable at the level of IRS-1 associated PI-3 kinase [22–24], Akt [14,24,25] and aPKC [26]. In the present study we provide novel evidence linking high fat feeding to defects in insulin signaling to Akt (Figure 4; pSer473 and Figure 5; pThr308) and the downstream target TBC1D4 (Figure 6; pThr642) believed to act as a molecular switch for GLUT4 movement in the cell [20,21,27].
In contrast to our hypothesis, muscle-specific overexpression of kinase dead α2AMPK (AMPK KD) did not lead to insulin resistance with ageing. Furthermore, development of insulin resistance in response to high fat feeding was not exacerbated in old AMPK KD mice when compared to WT littermates. Using the same mouse strain, it was previously shown that high fat diet-induced insulin resistance is also not exacerbated in young AMPK KD mice [14], collectively supporting that functional AMPK is not protecting against development of high fat diet-induced insulin resistance in mouse muscle; at least not in AMPK KD mice on a C57BL/6J background.

Interestingly, both young and old AMPK KD mice on chow diet were slightly glucose intolerant, but had normal insulin tolerance, HOMA-IR values and insulin-stimulated glucose uptake in isolated muscle (Figure 1 and 2). In these mice the plasma insulin response during the OGTT was similar or lower (Figure 1B) despite higher blood glucose levels raising the possibility that lack of muscle AMPK activity may influence pancreatic function. Previously, normal glucose tolerance (1 g/kg body wt, 6 hours fasting) has been observed in young (6–9 weeks of age) AMPK KD mice [14] whereas adult (36 weeks of age) mice were slightly glucose intolerant (2 g/kg body weight, 12 hours fasting) [28]. Since glucose intolerance may be more easily masked at low glucose doses [29], the reason for these discrepancies likely relates to procedural differences. Furthermore, the apparent mildness of this metabolic phenotype in AMPK KD mice may further encumber experimental verification.

In the present study ageing per se did not result in development of insulin resistance as indicated by responses to OGTT, ITT and HOMA-IR (Figure 1). Also, both insulin signaling and insulin-stimulated glucose uptake were not impaired in muscle from old vs. young mice. Generally, aging is linked to insulin resistance in both humans and rodents [30,31] and characterized by increased fat accumulation, chronic inflammation and oxidative stress in muscle [32,33]. In the present study, although body weight increased with age, metabolic characteristics (RER, VO2) and importantly body composition were not markedly altered. Thus, our results indirectly support an important role of excessive fat accumulation as a key contributing factor in the etiology of age-induced insulin resistance, as previously speculated [30,34].

When old mice were placed on a high fat diet for 17 weeks they became insulin resistant both at the whole body level and in skeletal muscle, coinciding with a ~30% increase in body weight but unaltered lean body mass (Table 1). Despite excessive fat accumulation, these mice were normally glucose tolerant (Figure 1A), associated with a compensatory increase in circulating insulin concentration (Figure 1C). As described, lack of functional AMPK did not exacerbate the detrimental effect of high fat feeding. If anything AMPK KD mice exhibited slightly improved (although not significant) HOMA-IR values and ITT response compared to WT littermates (Figure 1C). Curiously, these observations are in stark contrast to a study of high fat feeding in mice overexpressing inactive α2AMPK bred on a FVB mouse background (AMPK Ki) [12]. In that study, high fat feeding led to a reduction in insulin-stimulated glucose uptake in WT mice, similar to our observations, whereas insulin-stimulated glucose uptake was abolished in AMPK Ki mice. Considering the implications of placing AMPK in the nexus of diet and insulin action, these contradicting observations deserve further consideration.

![Figure 6. TBC1D4 Thr642 phosphorylation.](https://example.com/figure6.png)
Despite different genetic approaches [15,35] both mouse strains overexpress a non-functional α2AMPK isoform that displaces endogenous α2AMPK and to some extent α1AMPK. As a result, in both mouse strains, basal α2AMPK activity is markedly reduced, and activation of α2AMPK in response to pharmacological activators, hypoxia or muscle contraction is almost completely abolished [13,15,35,36]. Based on the similarities in AMPK dysfunction, in our view, a plausible explanation for the different adaptations to high fat dieting may relate to the differences in mouse strains wherein kinase dead AMPK is induced. As an indication, the FVB strain of mice have previously been characterized as being more resistant to development of high fat diet-induced insulin resistance than mice on a C57BL/6J background [37]. Furthermore, it is noteworthy, that in muscle from high fat fed AMPK Ki mice on a FVB background, reduced protein content of a range of proximal insulin signaling components including the insulin receptor β subunit, IRS-1, and Akt when compared to WT or AMPK Ki mice on a chow diet [12]. Based on similar Akt2 protein content (Figure 8E) and insulin signaling (Figure 4, 5, 6) in the present study, similar adaptations in AMPK KD mice on a C57BL/6J background apparently do not take place.

In contrast, it should be emphasized that in our model both SOL and EDL muscle are characterized by significant reductions of TRB-3 protein content independent of age and diet (Figure 8F). TRB-3, the mammalian homolog of drosophila tribbles, is emerging as an important player in insulin signaling, by its capacity to bind to Akt and prevent phosphorylation in the activation loop (pSer473 and pThr308) [38]. In LKB-1 KO mice, reduced TRB-3 protein content has been suggested as a critical adaptation improving insulin-stimulated glucose uptake [39] and also more recently, improved insulin signaling and insulin-stimulated glucose uptake after exercise in ob/ob mice have been associated with exercise-induced reductions in TRB-3 protein content [40]. The present study indicates a role of AMPK to regulate expression of TRB-3 although by yet undefined mechanisms in mouse muscle. Furthermore, reduced TRB-3 expression in our AMPK KD model may contribute to explaining the normal insulin signaling and insulin-stimulated glucose uptake in our AMPK KD model. Collectively, this study provides evidence that high fat diet-induced insulin resistance in mouse skeletal muscle on a C57BL/6J background is associated with impaired insulin signaling at the level of Akt and importantly also TBC1D4, providing a novel link between insulin signaling defects and impairments in control of GLUT4 translocation. In contrast to our hypothesis, the lack of functional AMPK did not influence insulin-stimulated glucose uptake with ageing or exacerbate insulin resistance after high fat feeding in old mice. Thus, based on elaborate studies of AMPK KD mice, the lack of α2AMPK activity in muscle (both oxidative and glycolytic) does not result in insulin resistance in either lean or obese, young or old mice ([14], present study). This strongly suggests that in contrast to previously indicated, AMPK does not constitute a necessary protective component for normal insulin action in muscle.

**Acknowledgments**

We kindly thank Professor Carol MacKintosh (University of Dundee, UK) for providing invaluable reagents used in this study as well Prof. Morris J. Birnbaum (Howard Hughes Medical Institute and University of Pennsylvania School of Medicine, USA) for providing the AMPK KD mice.

**Author Contributions**

Comments and corrections of manuscript: TEJ JJ CP JTT SJM JMK LS TJA PS BK JFPW. Conceived and designed the experiments: CF TEJ JTT BK JFPW EAR. Performed the experiments: CF TEJ JJ CP JTT SJM JMK LS TJA PS BK JFPW. Analyzed the data: CF JJ TEJ CP JTT EAR PS. Contributed reagents/materials/analysis tools: BK JFPW EAR PS. Wrote the paper: CF EAR.

**Figure 7. TBC1D1 Thr590 phosphorylation.** Basal (0 μU/ml) and insulin (500 μU/ml) stimulated TBC1D1 Thr590 phosphorylation measured by Western blot analyses in m. Soleus (SOL) and m. Extensor Digitorum Longus (EDL). Measurements were made in young and old AMPK KD mice and WT littermates on chow diet (CHO) or in old mice after 17 weeks of high fat diet (FA). *: Main effect of insulin, p<0.001. #: Main effect of diet p<0.001. †: Main effect of genotype, p<0.05. Values are means ± SE. n = 11–15.

doi:10.1371/journal.pone.0062338.g007


