Adenylate kinase 1 gene deletion disrupts muscle energetic economy despite metabolic rearrangement


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Efficient cellular energy homeostasis is a critical determinant of muscle performance, providing evolutionary advantages responsible for species survival. Phosphotransfer reactions, which couple ATP production and utilization, are thought to play a central role in this process. Here, we provide evidence that genetic disruption of AK1-catalyzed β-phosphoryl transfer in mice decreases the potential of myofibers to sustain nucleotide ratios despite up-regulation of high-energy phosphoryl flux through glycolytic, guanylate and creatine kinase phosphotransfer pathways. A maintained contractile performance of AK1-deficient muscles was associated with higher ATP turnover rate and larger amounts of ATP consumed per contraction. Metabolic stress further aggravated the energetic cost in AK1−/− muscles. Thus, AK1-catalyzed phosphotransfer is essential in the maintenance of cellular energetic economy, enabling skeletal muscle to perform at the lowest metabolic cost.

Keywords: adenylate kinase/creatine kinase/energy homeostasis/knockout mice/phosphoryl transfer

Introduction

Adenine nucleotides are the primary high-energy phosphoryl-carrying molecules in a living cell. Along with their energetic role, ATP, ADP and AMP are major regulators of essential cellular processes, including gene expression, ion channel gating, and receptor- or protein kinase-mediated signal transduction (Tullson and Terjung, 1991; Hardie and Carling, 1997; Dzeja and Terzic, 1998). Although the energetic and regulatory functions of adenine nucleotides are dependent on nucleotide ratios and rates of nucleotide exchange between cellular compartments (van Deursen et al., 1993; Steeghs et al., 1997; Saks et al., 1998; Saupe et al., 1998; Dzeja et al., 1999), the molecular mechanisms governing the dynamics of adenine nucleotide metabolism remain poorly understood.

Adenylate kinase (AK; EC 2.7.4.3), by virtue of its ability to catalyze the reaction ATP + AMP ↔ 2ADP, has been considered a key enzyme in the synthesis, equilibration and regulation of adenine nucleotides (Noda, 1973; Schulz et al., 1986). Through its unique property of transferring and providing for utilization both β- and γ-phosphoryls of ATP, AK doubles the energetic potential of the ATP molecule (Zeleznikar et al., 1990; Dzeja et al., 1998). By regulating ADP and AMP levels, AK renders these nucleotides active components in metabolic signaling determining the cellular response to stress (Elvir Mairena et al., 1996; Olson et al., 1996; Hardie and Carling, 1997; Dzeja and Terzic, 1998; Dzeja et al., 1999). Several AK isoforms have been cloned, and exhibit different levels of tissue expression with distinct intracellular distribution (Tanabe et al., 1993). Tissues with high energy demand, such as skeletal muscle, are particularly rich in AK1, the major enzyme isoform. AK1 is localized in the cytosol, clustered within myofibrils or bound to membranes (Wegmann et al., 1992; Tanabe et al., 1993; Elvir Mairena et al., 1996; Collavin et al., 1999). Other isoforms, such as AK2 and AK3, are less abundant in muscle and are generally confined to the mitochondrial intermembrane space or mitochondrial matrix (Noda, 1973; Nobumoto et al., 1998). The presence of AK in different compartments implicates this enzyme in promoting intracellular energy transfer and in communication between ATP-consuming and ATP-utilizing cellular sites (Bessman and Carpenter, 1985; Zeleznikar et al., 1990; Dzeja et al., 1998). In fact, the AK system has been found to facilitate high-energy phosphoryl transfer from mitochondria to myofibrils (Dzeja et al., 1999). Moreover, evidence is accumulating in support of an integrated intracellular high-energy phosphoryl transfer network comprised of AK, creatine kinase (CK) and glycolytic phosphotransfer enzymes (Wallimann et al., 1992; Savabi, 1994; Dzeja et al., 1998). Although AK appears to be essential in cellular energetics, the significance of this enzyme in maintaining intracellualr energy flow has not been determined.

To obtain direct information on mechanisms regulating the economy of cellular energetics, we generated knockout mice lacking the major AK isoform, AK1. Lack of AK1 disrupted muscle energetic economy and induced remodeling in pathways for ATP production, high-energy phosphotransfer and ATP utilization, underscoring the significance of AK in sustaining efficient cellular energetics. The observed metabolic rearrangement in glycolytic, CK and guanine nucleotide phosphotransfer systems in the AK1-deficient mice demonstrates a high plasticity of muscle energetics, and may provide a mechanism for the maintenance of energetic homeostasis under metabolic stress.
Results

Cloning of the genomic AK1 gene and generation of AK1<sup>-/-</sup> mutant mice

By screening a genomic mouse 129/Sv phage library, several phages were isolated that contained the entire AK1 gene, including upstream and downstream flanking segments. The structural organization of the AK1 gene in these phages (Figure 1A), which spans a total of 12 kbp and comprises seven exons, has an exon–intron distribution that appears highly conserved when compared with other species, including man (Matsuura et al., 1989). For disruption of the AK1 gene, a targeting vector (Figure 1A) was constructed in which the left and right arms of homology spanned 2.6 and 4.1 kbp, respectively. Exons 3–5 (encoding the ATP binding site) plus adjacent intron segments were deleted from the vector and replaced by a hygromycin B resistance cassette, inserted in antiparallel orientation. By gene targeting applied to mouse E14 embryonic stem (ES) cells (van Deursen and Wierenga, 1992; van Deursen et al., 1993; Steeghs et al., 1997), five cell clones with the desired genotypic modification were obtained. All ES clones had undergone correct homologous recombination events, downstream as well as upstream of the replacement mutation, as confirmed by Southern blot analysis (Figure 1B and C). Successful disruption of the AK1 gene was confirmed at the RNA level by northern blot analysis (Figure 1D), and at the protein level by zymogram analysis of protein extracts (Figure 1E–G). No residual mRNA was detected upon hybridization with an AK1 cDNA probe spanning exons 5–7, suggesting that a true null mutation was generated with respect to RNA and protein production. It is of note here that lack of AK1 in animals kept under normal housing conditions did not induce overt developmental or physiological abnormalities (not illustrated).

Absence of AK-catalyzed phosphotransfer in skeletal muscle from AK1<sup>-/-</sup> mice

Genetic deletion of AK1 produced loss of 18O incorporation into β-phosphoryls of ATP, indicating a lack of AK-catalyzed phosphotransfer in AK1-knockout skeletal muscle (Figure 2). The percentage of β-ATP 18O labeling was 4.43 ± 0.33 in wild type (n = 6) and 0.04 ± 0.02 in AK1-knockout (n = 6, p <0.001; Figure 2). AK phosphotransfer flux in wild type was 4.25 ± 0.31 nmol ADP/min/mg protein or 20% of total ATP turnover in contracting (2 Hz) muscle. In AK1-knockout, AK phosphotransfer flux was 0.05 ± 0.01 nmol ADP/min/mg protein or <0.2% of total ATP turnover. Adenine nucleotide-dependent AK activity was 3621 ± 218 nmol ATP/min/mg protein in the wild type, but only 11 ± 3 nmol ATP/min/mg protein in the AK1-knockout (p <0.001). Total AK activity in the AK1-knockout muscle was diminished by 99.7% when compared with wild type. The remaining 0.3% of AK activity could be attributed to minor AK isoforms still present in AK1-knockout muscle. Thus, the AK1-knockout gastrocnemius–plantaris–soleus (GPS) muscle is characterized by the absence of AK-catalyzed β-phosphoryl transfer, which remains uncompensated by remaining AK isoforms.

Adenine nucleotide levels and metabolism in AK1-knockout skeletal muscle

AK1 has been implicated in adenine nucleotide synthesis and metabolism (Noda, 1973; Schulz et al., 1986). AK1<sup>-/-</sup> knockout skeletal muscle demonstrated maintained ATP levels: 28.3 ± 1.5 nmol/mg protein compared with 26.4 ± 0.8 nmol/mg protein in the wild type (n = 6). The total sum of adenine nucleotides (ATP, ADP and AMP) was also preserved in AK1-deficient muscle: 33.9 ± 1.8 nmol/mg protein compared with 31.2 ± 0.9 nmol/mg protein for the wild type (n = 6). De novo adenine nucleotide synthesis is a rather slow process (Tullson et al., 1998) and thus could be supported by minor AK isoforms present in AK1-knockout muscle. However, in non CONTRACTING AK1<sup>-/-</sup> gastrocnemius muscle, the ATP/ADP ratio was higher than in the wild type: 8.8 ± 0.2 versus 8.0 ± 0.2, respectively (p <0.03, n = 5). Moreover, in GPS muscle contracting at 2 Hz, the ATP/ADP and AMP/ADP ratios were 5.6 ± 0.2 versus 6.7 ± 0.4 (p <0.05, n = 6), and 9.3 ± 0.8 versus 5.4 ± 0.2 (p <0.001, n = 6), in AK1<sup>-/-</sup> deficient and wild type, respectively (Figure 3). This suggests that AK1-deficient muscle has a lower potential to sustain nucleotide ratios during functional load.

AK-produced AMP is metabolized to IMP by AMP deaminase (AMPD), a critical enzyme in the adenine nucleotide cycle (Tullson et al., 1998). The maximal activity of AMPD was lower in AK1<sup>-/-</sup> gastrocnemius compared with wild type (214 ± 23 versus 330 ± 30 μmol/min/g muscle, respectively; p <0.02, n = 5). Western blot analysis of the AK1<sup>-/-</sup> gastrocnemius showed a 50% reduction in the expression of the 80 kDa AMPD protein (p <0.01) compared with wild type (not illustrated). Thus, although adenine nucleotide levels are maintained in AK1-deficient muscle, the dynamics of adenine nucleotide metabolism catalyzed by nucleotide metabolizing and/or phosphotransfer enzymes may be altered upon deletion of AK1.

Fig. 1. (A) Targeted mutagenesis of AK1 with wild-type (top) and mutant (bottom) AK1 genes shown after homologous recombination. Exons: shaded or black; introns or extragenic regions: white. In the targeting vector (middle), a selectable HygROB cassette replaces the 5.5 kb BamHI–HindIII gene fragment (exons 3–5) that encodes the ATP-binding domain of AK1. A HSV tk cassette was fused to the right arm of homology to facilitate selection of properly targeted ES clones. Lengths of restriction fragments (arrowed lines) from wild-type and mutant alleles are indicated. (B and C) Southern blot of genomic DNAs from wild-type and mutated E14 ES clones. DNAs were cleaved with KpnI–Xhol or BamHI restriction enzymes, resolved by agar gel electrophoresis, and blot-analyzed by hybridization with probes (indicated in A) that discriminate between recombination events at the 3'-(B) and 5'-(C) segments of homology. Diagnostic KpnI–Xhol digestion yields 11.5 and 5.6 kbp fragments, whereas BamHI digestion results in 4 and 7 kbp fragments for wild-type and mutant alleles. (D) Northern blot analysis of RNAs from wild-type and AK1<sup>-/-</sup> tissues. Blots with equivalent amounts of total RNA (10 μg) from skeletal muscle (SK), heart (H) and brain (B) probed with AK1 cDNA. Note the absence of AK1 mRNA in homoygous mutants. (E, F and G) Zymogram analysis of homogenates from brain (B), heart (H) and gastrocnemius (G)–plantaris (P)–soleus (S) muscle of wild-type, AK1<sup>-/-</sup> (homoygous) and AK1<sup>-/-</sup> (heterozygous) mice. AK1 and CK isoenzymes were separated by agarose gel electrophoresis under native conditions and their migration positions revealed by activity staining. Note the complete absence of staining at the AK1 position for AK1-deficient mice.
Creatine kinase in AK1-knockout skeletal muscle

Along with AK, CK provides a major pathway for regulation of nucleotide ratios and energy transfer (Wallimann et al., 1992; van Deursen et al., 1993; Veklerov et al., 1995; Dzeja et al., 2000). Wild-type and AK1-deficient GPS muscle extracts contained essentially equal levels of creatine phosphate (CrP): 50.4 ± 3.0 (n = 6) and 47.8 ± 3.2 (n = 6) nmol CrP/mg protein, respectively. Moreover, in vivo changes in CrP levels, measured by 31P-NMR spectroscopy, during metabolic transitions exhibited similar profiles in wild-type and AK1−/− hind leg muscles (Figure 4A). However, in AK1-knockout GPS, the percentage of CrP 18O labeling was 21.9 ± 1.7 (n = 6) compared with 15.8 ± 1.3 (n = 6) in the wild type (p < 0.05; Figure 4B), indicating increased turnover of substrates in the CK-catalyzed reaction. This translated into a moderate increase in net CK-catalyzed phosphotransfer flux (13.9 ± 1.4 and 16.5 ± 1.1 nmol CrP/min/mg protein in wild type and AK1-knockout, respectively). Thus, while CrP is maintained at wild-type level in the AK1-knockout skeletal muscle, the relative CK-catalyzed CrP turnover appears increased upon deletion of AK1.

Glycolytic phosphotransfer and glycolytic intermediates in AK1-knockout muscle

AK phosphotransfer is coupled to glycolysis and regulates the activity of ATP- and AMP-sensitive glycolytic enzymes (Otway and Mowbray, 1977; Zeleznikar et al., 1995). In turn, glycolysis also serves a high-energy phosphotransfer function in parallel with AK and CK (Dzeja et al., 1998, 2000). Here, deletion of AK1 affected glycolytic phosphotransfer, as well as levels of glycolytic intermediates. Glycolytic phosphotransfer was assessed by hexokinase-catalyzed incorporation of 18O into glucose-6-phosphate (G-6-P), which increased from 8.4 ± 0.3 (n = 6) in wild type to 11.2 ± 0.6 (n = 6) in AK1-knockout muscle (p < 0.05), at a labeling rate of 0.66 ± 0.03 and 0.151 ± 0.05 nmol/min/mg protein × 10−3 (p < 0.01; Figure 5A), respectively. The estimated net glycolytic flux was increased from 1.31 ± 0.06 to 3.00 ± 0.09 nmol G-6-P/min/mg protein in wild-type and AK1-knockout muscle, respectively (p < 0.01). Increased glycolytic flux was accompanied by higher muscle levels of G-6-P (7.8 ± 1.0 and 13.6 ± 1.6 nmol/ mg protein in wild type and AK1-knockout; n = 6, p < 0.001) and lactate (8.3 ± 0.7 and 12.6 ± 2.1 nmol/ mg protein in wild type and AK1-knockout; n = 6, p < 0.001). Alterations in glycolytic metabolism were
Adenylate kinase 1 and muscle energetic economy

Figure 5. (A) Increased hexokinase-catalyzed G-6-P turnover in AK1-knockout muscle. Glucose-6-phosphate phosphoryl oxygen replaced with $^{18}$O, as an indicator of glycolytic phosphotransfer, in wild-type ($n = 6$) and AK1-knockout ($n = 6$) GPS muscle. (B and C) Intracellular pH (B) and PME levels (C) in wild-type (WT) and AK1−/− mice, analyzed by $^{31}$P-NMR and expressed in pH units or as resonance peak heights normalized to the mean value of the $\beta$-ATP intensity of the first four spectra. Series were recorded before, during, and after ischemia of the lower hind limb muscles in wild-type (dashed line, open squares) and AK1-deficient (solid line, closed triangles) mice. After four reference spectra, an ischemic period of 25 min was introduced succeeded by a recovery of 16 min. Note a faster pH decrease and increased PME levels during the ischemic period in AK1−/− compared with wild-type mice.

Further assessed in vivo by monitoring phosphomonoester (PME)-containing glycolytic intermediates and intracellular pH by $^{31}$P-NMR spectroscopy. Ischemia induced a faster muscle acidification in AK1-mutant mice (Figure 5B), suggesting that the block in $\beta$-phosphoryl exchange between ADP and ATP is associated with accelerated anaerobic glycolysis and production of lactate. Moreover, the increase in PME levels was significantly more pronounced in AK1-deficient mutant mice (Figure 5C). This increase in the intracellular concentration of glycolytic intermediates, along with increased G-6-P turnover, indicates elevated glycolytic metabolism in AK1-knockout muscle.

Guanine nucleotide metabolism in AK1-knockout skeletal muscle

Adenine nucleotide metabolism is coupled to guanine nucleotide metabolism through the nucleoside diphosphate kinase (NDPK)-catalyzed phosphotransfer (Ray and Mathews, 1992). There is an important link between GTP production by substrate level phosphorylation in the Krebs cycle and ATP production in mitochondria. AK1-deficient muscle had a 50% higher intracellular GTP concentration compared with wild type, 0.75 ± 0.06 versus 0.43 ± 0.02 nmol/mg protein, respectively ($p < 0.001$, $n = 6$). Also, the GDP concentration was 17% higher in AK1−/− muscle compared with wild type: 0.20 ± 0.02 and 0.16 ± 0.04 nmol/mg protein, respectively ($p < 0.03$, $n = 6$). This resulted in higher muscle levels of guanine nucleotides (GTP plus GDP) in AK1−/− compared with wild type: 0.95 ± 0.07 versus 0.59 ± 0.02 nmol/mg protein, respectively. Also, the GTP/GDP ratio was elevated in AK1-knockout muscle compared with wild type: 3.7 ± 0.2 versus 2.7 ± 0.1, respectively ($p < 0.003$, $n = 6$; Figure 6A). Concomitantly, the ATP/GTP ratio was lower in AK1−/− muscle compared with wild type (38 ± 1 and 61 ± 2; $p < 0.001$, $n = 6$), indicating increased guanine nucleotide contribution to muscle energetics (Figure 6A). The NDPK activity, however, was unaltered in AK1−/− gastrocnemius [19.9 ± 1.5 × 10$^3$ relative light units (RLU)/min/mg protein; $n = 3$] compared with wild type [21.4 ± 1.7 × 10$^3$ RLU/min/mg protein; $n = 3$]. To assess the dynamics of guanine nucleotide metabolism, the rates of $[^{18}$O$]$/phosphoryl labeling of $\gamma$- and $\beta$-GTP/GDP were determined (Figure 6B). In AK1−/− GPS, the percentage of $\gamma$-GTP $^{18}$O labeling was 26.4 ± 1.6 ($n = 6$), significantly higher than the 18.5 ± 0.5 ($n = 6$) measured in the wild type ($p < 0.05$; Figure 6B). Such an increase in $\gamma$-GTP turnover indicates that phosphorl flux through enzymes catalyzing GTP production, such as succinyl-CoA synthase of the Krebs cycle and/or NDPK, is elevated following deletion of the AK1 gene. NAD$^+$ levels were significantly higher in freshly isolated AK1−/− muscle ($n = 5$) compared with wild type ($n = 5$), 3.4 ± 0.2 versus 2.8 ± 0.6 nmol/mg protein, respectively ($p < 0.02$). Moreover, the NAD$^+$/NADH ratio was higher in in vitro incubated AK1-deficient muscle, 1.1 ± 0.1 versus 0.7 ± 0.1 for the wild type ($p < 0.01$; $n = 6$), suggesting elevated aerobic Krebs cycle metabolism. Guanylate kinase, the AK analog in guanine nucleotide metabolism, catalyzes $\beta$-GTP/GDP turnover, providing energy from $\beta$-phosphoryls. The percentage of combined $\beta$-GTP/GDP $^{18}$O labeling was
has been considered an important enzyme in maintaining cellular energetic economy (Noda, 1973; Dzeja et al., 1998). To assess whether AK1 gene deletion affected muscle energetic efficiency, muscle performance and ATP turnover rate were determined. AK1-deficient gastrocnemius had an isometric (twitch and tetanic) force production equivalent to that of the wild type (Table I). Also, muscle speed was not different between groups, as indicated by a similar maximal rate of force development, half-relaxation time and maximal shortening velocity (Table I). Repeated isometric contractions resulted in a similar decrease in the rate and amplitude of force generation in wild-type and mutant muscle (Table I). Thus, the overall contractile performance of an AK1-deficient muscle appears rather normal. However, γ-ATP 18O labeling, reflecting muscle energy transduction rate, was significantly increased from 14.1 ± 0.9% (n = 6) in the wild-type to 18.4 ± 0.5% in AK1-mutant muscle (n = 6, p <0.002; Figure 7A). Furthermore, the total cellular ATP turnover, determined from 18O incorporation into major high-energy phosphoryls including γβ-ATP, CrP, inorganic phosphate, G-6-P and γβ-GTP/GDP, was significantly increased from 21.6 ± 0.9 nmol/min/mg protein in wild-type (n = 6) to 26.3 ± 1.2 nmol/min/mg protein in AK1-knockout (n = 6) GPS, respectively (p <0.02; Figure 7B). Thus, the overall ATP production and utilization was elevated in AK1-knockout muscle. This translates into an increased ATP consumption per contraction from 180 ± 2 pmol ATP/mg protein in the wild-type to 219 ± 1 pmol ATP/mg protein in AK1-knockout muscles contracting at 2 Hz (p <0.02, n = 6). Such a rise in the energetic cost of contraction, by 22%, indicates reduced efficiency of cellular energetics in the absence of AK1-catalyzed phosphotransfer.

**Aberrant redistribution of phosphotransfer flux and increased energy expenditure in AK1-knockout muscle during hypoxic stress**

Metabolic stress induces redistribution of phosphotransfer in skeletal muscle (Zeleznikar et al., 1995). AK function is believed to increase under stress to regenerate ATP and maintain muscle energetic efficiency (Dzeja et al., 1996, 1999). Here, the contribution of AK-mediated phosphotransfer to total cellular ATP turnover increased from 20 ± 2 to 52 ± 5% in response to hypoxia (2 mM KCN treatment) in wild-type GPS (Figure 8A). In AK1-knock-
Therefore, under hypoxia, the AK1-deficient muscle had a higher energy expenditure, by 31%, compared with normal muscle.

**Discussion**

Efficient cellular energetics is a critical determinant of muscle performance and thus an evolutionary necessity for species survival (Mommaerts, 1969; Gibbs and Barclay, 1998). Here, we provide first evidence that disruption of AK-catalyzed phosphotransfer by gene deletion leads to overt reduction in energetic efficiency of contractile performance associated with metabolic remodeling in glycolytic, CK and guanine nucleotide phosphotransfer systems. Thus, phosphotransfer reactions are an integral component of cellular energetics required for nucleotide homeostasis, optimal high-energy phosphoryl transfer and efficient ATP utilization.

Although the molecular structure and kinetic properties of the AK protein have been established (Schulz et al., 1986; Yan and Tsai, 1999), the biological significance of AK catalysis in a living cell has remained poorly understood. Here, deletion of AK1 was associated with a dramatic reduction in total muscle AK activity and lack of β-phosphoryl turnover, providing definite evidence that β-phosphoryl transfer is an exclusive property of AK catalysis that cannot be supported by other phosphotransfer reactions.

Throughout evolution, AK has been highly conserved and implicated in de novo adenylic nucleotide synthesis and equilibration (Noda, 1973). In yeast, absence of mitochondrial AK activity compromises export of ATP from mitochondria (Bandlow et al., 1988), and generalized AK mutation can be lethal due to uncompensated disruptions in β-phosphoryl energetics and adenylic nucleotide synthesis (Konrad, 1993). Here, in AK1-knockout muscle, ATP and total adenylic nucleotide levels were rather preserved. Thus, the marginal AK-like activity that remains in AK1-knockout muscle is probably sufficient to support adenylic nucleotide synthesis (Tullson et al., 1998). However, AK1-deficient muscle was unable to sustain ATP/ADP and ADP/AMP ratios, underscoring the importance of AK in regulating adenylic nucleotide metabolism.

Through NDPK and guanylate kinase, adenylic nucleotide metabolism is coupled to guanylic nucleotide metabolism, which in turn serves a critical role in signal transduction (Ray and Mathews, 1992). In fact, metabolic consequences induced by NDPK deficiency can be compensated in part through AK (Lu and Inouye, 1996). Although NDPK activity was not notably changed, guanylic levels as well as guanylic nucleotide β- and γ-phosphoryl turnover were significantly increased in AK1−/− muscle. This increase in guanylic nucleotide turnover could compensate for ~10% of lost adenylic nucleotide β-turnover, indicating that metabolic flux through guanylic nucleotide-dependent Krebs cycle reactions and guanylate kinase are altered following deletion of the AK1 gene. In fact, increased [γ-32P]GTP labeling in AK1-knockout muscle was associated with increased activity of succinyl CoA synthetase, a Krebs cycle enzyme that couples cleavage of succinyl CoA to phosphorylation of GDP and contributes to labeling at the γ-phosphoryl
Fig. 8. Aberrant redistribution of phosphotransfer flux in AK1-knockout skeletal muscle in response to hypoxia. (A) Hypoxia markedly elevates AK-catalyzed flux in wild-type, but not in AK1-knockout GPS. AK-catalyzed phosphotransfer, expressed as a percentage of total cellular ATP turnover, in wild-type (n = 6) and AK1-knockout (n = 6) GPS muscle. (B) Increase in AK-catalyzed phosphotransfer in hypoxic AK1-deficient compared with hypoxic wild-type GPS. Creatine kinase-catalyzed phosphotransfer, expressed as a percentage of total ATP turnover, in wild-type (n = 6) and AK1-knockout (n = 6) GPS muscle. (C) Contribution of combined AK- and CK-catalyzed phosphotransfer to cellular ATP turnover is reduced in normoxic and hypoxic AK1-knockout GPS. The sum of AK- and CK-catalyzed phosphotransfer is expressed as a percentage of total ATP turnover measured by the [18O]phosphoryl labeling technique. (D) ATP turnover rates in wild-type and AK1-knockout GPS under hypoxia. Hypoxia was induced by KCN (2 mM), an inhibitor of mitochondrial respiration.

position in GTP (Zeleznikar and Goldberg, 1991). Also, altered mitochondrial NAD+/NADH ratios suggest elevated Krebs cycle activity and increased mitochondrial involvement in AK1−/− muscle. Electron microscopy-ultrastructural analysis of fast-twitch fibers confirmed that these metabolic changes are paralleled by an increase in mitochondrial volume (not shown). Taken together, the observed increase in steady-state levels of guanine nucleotides and their metabolic dynamics upon disruption of adenine nucleotide homeostasis suggest an increased contribution of guanine nucleotides in AK1-knockout muscle.

In many cell types, AK-catalyzed phosphotransfer is interrelated with CK and glycolytic phosphotransfer systems (Savabi, 1994; Dzeja et al., 1998). In insulin-secreting pancreatic β-cells, increased glycolytic and CK flux, which occurs in response to elevated extracellular glucose, results in suppression of AK-catalyzed phosphotransfer (Olson et al., 1996; Dzeja and Terzic, 1998). Conversely, in skeletal muscle, inhibition of CK and/or glycglysis results in a marked increase in AK-catalyzed phosphotransfer (Dzeja et al., 1996). Here, deletion of AK1 induced adaptive changes in CK-catalyzed phosphotransfer, and up-regulation of high-energy phosphoryl flux through glycolytic metabolism. Specifically, the glycolytic flux rate more than doubled and was accompanied by a similar increase in the concentrations of glycolytic intermediates, such as G-6-P and lactate. Similarly, levels of PME compounds (glycerol phosphate, G-1-P and G-6-P), measured in vivo in ischemic AK1−/− muscle, were twice as high as in the wild type, and were accompanied by faster accumulation of lactate and a faster decrease in intracellular pH. Although CrP levels were maintained, CK-catalyzed phosphotransfer was moderately increased in AK1−/− muscle compared with the wild type. Thus, the AK1-knockout muscle adapts to the lack of AK1-catalyzed phosphotransfer through up-regulation of glycolytic, along with CK and guanine nucleotide phosphotransfer systems.

This is in accord with previous studies with CK-knockout mice that also indicate a high plasticity of muscle energetic systems in adapting to genetic disruption of an energy supply pathway (van Deursen et al., 1993; Veksler et al., 1995; Dzeja et al., 1996; Steeghs et al., 1998). However, despite metabolic and cyto-architectural adaptation, deletion of the M-CK and/or ScCKmit CK isoforms resulted in an abnormal muscle contractile response, intracellular Ca2+ handling and mitochondrial respiration (van Deursen et al., 1993; Kay et al., 2000). Moreover, in heart muscle, deletion of CK genes reduced the ability to sustain adenine nucleotide ratios, and rendered cardiac work more ‘energetically costly’ in terms of high-energy phosphate use and decreases in free energy released from ATP hydrolysis (Saupe et al., 1998). Here, we demonstrate that AK1-deficient muscles exhibit a higher rate of [18O-γ]phosphoryl labeling and total ATP
turnover. As the overall level of phosphoryl-containing metabolites was not significantly changed upon deletion of AK1, this indicates that the energetic efficiency of AK1-knockout muscle is lower than that of wild type.

AK1-deficient muscles did not exhibit any apparent deviation in contractile and fatigue performance compared with control muscle, suggesting that the same work output can be generated in both muscles. However, the AK1-knockout muscle did consume a larger amount of ATP per contraction, indicating lower energetic efficiency of muscle performance following AK1 deletion. Metabolic stress, which is associated with aberrant redistribution of phosphotransfer flux, further aggravated the energetic cost in an AK-deficient muscle, providing evidence for the crucial role of AK in optimal cellular energy allocation.

In conclusion, loss of AK-supported β-phosphoryl energetics disrupts adenine and guanine nucleotide homeostasis and associated metabolic circuits (i.e., the AMPD reaction), and results in redistribution of energy flow through alternative glycolytic and CK phosphotransfer systems. Such metabolic plasticity of myocytes provides an apparent compensatory potential that limits cellular energetic failure and preserves muscle contractile performance. However, intracellular metabolic rearrangement is not sufficient to sustain energetic economy in AK1-deficient muscle, in particular under metabolic stress, suggesting a specific energetic function for AK. Thus, this study provides new evidence that phosphotransfer reactions in general, and adenylate kinase in particular, are essential for cellular energetic economy, enabling skeletal muscle to perform with high thermodynamic efficiency under physiological and pathophysiological conditions.

**Materials and methods**

**AK1 targeting vector**

Genomic DNA fragments of the AK1 gene were obtained by screening an EMBl3 phage library of partially digested genomic DNA of the mouse strain 129/Sv using rat AK1 cDNA spanning exons 5–7 as a probe (provided by Dr. N.D. Goldberg and A.Nakazawa). Plagues from positive plaques were purified to homogeneity and characterized using standard procedures (Steeghs et al., 1997). Exons were identified using Southern blot hybridization with exon-specific oligonucleotides, and positioned by restriction mapping followed by partial sequence analysis of genomic DNA fragments. Appropriate segments that covered the entire exon 1–7 region of the AK1 gene were subcloned into plasmid pGEM3 (Promega). The 2.6 and 4.1 kb SalI–BamHI and HindIII–BamHI fragments were used as left and right arms of homology to generate the targeting vector (Figure 1). In the construct, the exon 3–5 area, encoding the ATP binding site, was replaced by a BamHI–HindIII hygromycin B resistance (hygroB) cassette (van Deursen and Wieringa, 1992). To enable selection for homologous targeting, a negative selection cassette containing the 2.0 kb herpes simplex virus thymidine kinase gene (HSV-tk; van Deursen and Wieringa, 1992) was inserted at the 3’ BamHI site. The AK1 targeting vector was linearized at the XhoI site located at the vector’s multiple cloning site upstream of the 5’ AK1 insert.

**AK1−/− embryonic stem cells**

Wild-type E14 ES cells were grown on a layer of irradiated SNL19 feeder cells, cultured and electroporated (van Deursen and Wieringa, 1992; Steeghs et al., 1997). Twenty-four hours after electroporation, cells were grown in the presence of 300 µg/ml hygromycin B (Life Technologies) and 0.2 µM Fiu (Bristol-Meyer), and allowed to grow for another 8–10 days before resistant colonies were isolated. For detection of the desired AK1 mutation, genomic DNAs from individual ES clones were prepared according to established procedures (Steeghs et al., 1997). DNA was digested with either BamHI or KpnI–XhoI restriction enzymes for the identification of targeted homologous recombination events at both sides of the replacement cassette (Steeghs et al., 1997). The homologous recombination event at the 5’ side of the AK1 locus was determined using a unique 800 bp SalI–Kpn fragment as probe. For identification of the 3’ event, a 1700 bp BamHI–XhoI fragment (Figure 1) was used.

**AK1−/− knockout mice**

Targeted ES cells (AK1−/−) were injected into blastocysts derived from C57BL/6 mice to obtain chimeric, heterozygous and homozygous offspring (van Deursen et al., 1993; Steeghs et al., 1997). AK1−/− mice were kept on a mixed 50%–50% C57BL/6 × 129/Ola inbred background during breeding. Southern analysis was performed on DNA from tail biopsies used for genotyping. A PCR assay was designed where three primers were used to discriminate between wild-type, heterozygous and homozygous mutant mice. Forward primer was AK1M1 located in intron 1 (5’-GGAGCCTGACATTACTTGC-3’). Reverse primers were AK1M2 (5’-GAATTCTGTCGTTCCGTATAGC-3’) and H1 (5’-GGCTGG-CACCTCTGCGATAACC-3’) located in intron 2 and the hygromycin box region, respectively. PCR reactions were carried out in 50 µl containing 10 mM Tris–HCl pH 8.3, 50 mM KCl, 1.5 mM MgCl2, 0.01% gelatin, 100 ng of each primer, 100 ng of DNA template and 1 µl of Taq polymerase. Amplified products generated from the wild-type and AK1 mutant locus were 384 and 626 bp in length, respectively. GPS muscles from age-matched homozygous AK1-deficient and wild-type control animals, also having a 50%–50% C57BL/6 × 129/Ola mixed inbred background, were used throughout experiments. Under standard housing conditions, we saw no abnormalities in birth weight, weight gain during growth into adulthood, or daily food intake in first- and second-generation AK1−/− animals. Also, fiber type distribution was normal in AK1 mice. Mice were anesthetized with 2.2% tribromoethanol (350 mg/kg i.p.) or pentobarbital (100 mg/kg i.p.). All procedures conformed to the Guidelines for the Care and Use of Laboratory Animals of the Dutch Council and the National Institutes of Health, and were approved by the Institutional Animal Care and Use Committee at the University of Nijmegen, the Mayo Clinic and the Vrije University, Amsterdam.

**Northern blot analysis**

Total RNAs from freshly isolated skeletal GPS muscle, as well as heart and brain, were extracted using the lithium chloride–urea method (Steeghs et al., 1997) and analyzed by northern blotting (Steeghs et al., 1997). RNA blots were hybridized with a fragment of rat AK1 cDNA, spanning exons 5–7, and a 1.3 kb rat glyceraldehyde-3-phosphate dehydrogenase (GAPDH) cDNA probe. Hybridization was carried out overnight at 65°C in 0.5 M NaPO4 buffer pH 7.2, 7% (v/v) SDS, 1 mM EDTA, and blots were washed to a final stringency of 1× SSC/0.1% (v/v) SDS at 65°C and exposed on Kodak X-Omat films.

**Zymogram analysis**

Homogenates from freshly excised GPS muscles (10% w/v) were prepared in SET buffer (250 mM sucrose, 2 mM EDTA, 10 mM Tris–HCl pH 7.4) at 4°C. GPS extracts were diluted 1:1 in 30 mM NaPO4 buffer pH 7.4, containing 0.05% v/v Triton X-100, 0.3 M diethanol and a protease inhibitor cocktail (1 mM phenylmethylsulfonyl fluoride, 10 µg/ml leupeptin, 1 µg/ml aprotinin; Boehringer Mannheim #1697498). Muscle extracts were incubated for 30 min at room temperature, centrifuged for 20 min at 11,000 g and an aliquot (1–5 µl) applied to agarose gels (Alameda, CA). AK1 and CK isoenzymes were separated electrophoretically, and stained for enzyme activity (Steeghs et al., 1997).

**Adenylate kinase and NDPK activity**

AK activity was measured spectrophotometrically in GPS muscle extracts (Dzeja et al., 1999). In addition, AK activity was measured by a luciferase-based ATP bioluminescence assay (Sigma) in the presence of 2 mM ADP using a BioOrbit 1253 luminometer. The NDPK activity, expressed in RELU/min/mg protein, was determined with the same bioluminescence assay in the presence of the AK inhibitor diadenosine pentaphosphate (Ap5A; 200 µM), and the reaction started with GTP (2 mM).

**AMPD**

AMPD activity was measured using GPS muscle extracts (Rush et al., 1999). AMPD activity was measured in a reaction buffer containing 15 mM AMP, and formation of the reaction product, IMP, monitored by HPLC. Western blot analysis of AMPD was performed on muscle extracts probed with a polyclonal antiseraum raised against purified rat
skeletal muscle AMPD. Developed nitrocellulose membranes were digitized and analyzed using NIH Image (Tullson et al., 1998).

High-energy phosphoryl transfer
ATP turnover and phosphoryl flux through AK, CK and glycolytic systems were measured in intact GFS muscle complex using the [14C]phosphoryl labeling technique (Zelevinski and Goldberg, 1991; Zelevinski et al., 1995; Dzeja et al., 1996, 1999). Isolated mouse GFS muscle was pre-incubated for 15–20 min at room temperature in media containing 137 mM NaCl, 5 mM KCl, 2 mM CaCl2, 1 mM MgCl2, 1 mM NaH2PO4, 20 mM HEPES, 0.05 mM EDTA, 5 mM glucose, 24 mM NaHCO3 pH 7.4. The muscle complex was then rapidly transferred into media enriched with 20% [14C]water (Isotec Inc.), and paced at 2 Hz. In separate experiments, hypoxia was simulated by pre-incubating muscles in 2 mM KCl for 3 min, and then transferring muscles into 1O2-enriched media with 2 mM KCN for an additional 3 min (at continuous 2 Hz pacing). Cellular ATP, ADP, GTP, GDP, inorganic phosphosphate, CrP and G-6-P were purified and quantified using high-performance liquid chromatography (HPLC) (Zelevinski and Goldberg, 1991; Dzeja et al., 1996). Samples containing phosphoryls of γ-ATP, β-ATP, β-ADP, γ-GTP, β-GTP, β-GDP, inorganic phosphate and CrP, as glycerol-3-phosphate, were converted to trimethylsilyl derivatives. 1O2 enrichment of phosphoryls in glycerol-3-phosphates was determined with a Hewlett-Packard 5980B gas chromatograph mass spectrometer. Total cellular ATP turnover, along with net AK- and CK-catalyzed phosphoryltransfer, were determined from the rate of appearance of 18O-containing phosphoryls (Zelevinski et al., 1990; Zelevinski and Goldberg, 1991; Dzeja et al., 1996). The rate of net phosphoryl flux through the glycolytic system was measured by the appearance of 18O-labeled phosphoryls in G-6-P, which was purified by HPLC (Dzeja et al., 1996).

Metabolite levels
ATP, ADP, AMP, GTP and GDP levels were quantified in perfused muscle extracts by HPLC (Zelevinski and Goldberg, 1991; Dzeja et al., 1999). CrP, muscle lactate and G-6-P levels were determined using coupled enzyme assays (van Deursen et al., 1993; Dzeja et al., 1996). Muscle inorganic phosphate level was determined using the EnzChek Phosphate Assay kit (Molecular Probes).

In vivo 31P-NMR spectroscopy
In mice anesthetized with 1.5% enflurane, 31P-NMR spectra were determined in vivo in hind limb muscles using a 7T S.M.I.S. spectrometer. Recordings were performed prior to, during 25 min of ischemia and following 16 min of reperfusion. Ischemia was induced by occluding hind limb circulation. Spectra were analyzed using time domain fitting, and normalized to the mean value of the β-ATP intensity of the first four spectra.

Contractile muscle properties
Peak force, maximal rate of force rise and half-relaxation time were obtained under isometric conditions in median gastrocnemius muscles (at 35°C) from wild-type and AK1-deficient mice (n = 20) as described (de Haan et al., 1989). Maximal shortening velocity was obtained in a subgroup by extrapolation of the force–velocity relationship. Fatigability was investigated by applying a series of 20 isometric contractions (150 ms duration, 150 Hz) within 10 s.

Statistics
Data are presented as mean ± SE, unless indicated otherwise. Student’s t-test for unpaired samples was used for statistical analysis, and p < 0.05 was considered significant.

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References

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