Adrenalectomy enhances the insulin sensitivity of muscle protein synthesis

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Adrenalectomy enhances the insulin sensitivity of muscle protein synthesis. Am J Physiol Endocrinol Metab 284: E102–E109, 2003. First published October 1, 2002; 10.1152/ajpendo.00028.2002.—After confirming that adrenalectomy per se does not affect skeletal muscle protein synthesis rates, we examined whether endogenously produced glucocorticoids modulate the effect of physiological insulin concentrations on protein synthesis in overnight-fasted rats 4 days after either a bilateral adrenalectomy (ADX); ADX with dexamethasone treatment (ADX + DEX), or a sham operation (Sham; n = 6 each). Rats received a 3-h euglycemic insulin clamp (3 mU·min⁻¹·kg⁻¹). Rectus muscle protein synthesis was measured at the end of the clamp, and the phosphorylation states of protein kinase B (Akt), eukaryotic initiation factor 4E-binding protein 1 (4E-BP1), and ribosomal protein S6 kinase (p70S6K) were quantitated before and after the insulin clamp. The basal phosphorylation states of Akt, 4E-BP1, and p70S6K were similar between ADX and Sham rats. Insulin significantly enhanced the phosphorylation of Akt (P < 0.03), 4E-BP1 (P = 0.003), and p70S6K (P < 0.002) in ADX but not in Sham rats. Protein synthesis was significantly greater after insulin infusion in ADX than in Sham rats (P = 0.01). Glucocorticoid replacement blunted the effect of insulin on Akt, 4E-BP1, and p70S6K phosphorylation and protein synthesis. In conclusion, glucocorticoid deficiency enhances the insulin sensitivity of muscle protein synthesis, which is mediated by increased phosphorylation of translation initiation-regulatory proteins. Dexamethasone; translation initiation; protein kinase B; eukaryotic initiation factor 4E-binding protein-1; p70 S6 kinase (PI 3-kinase)-mammalian target of rapamycin (mTOR) pathway involved in the phosphorylation and activation of 4E-BP1 and p70S6K. It is downstream of PI 3-kinase and itself promotes the phosphorylation and activation of mTOR (4, 7, 48, 57). 4E-BP1, in its unphosphorylated form, functions as an mRNA translation initiation repressor by binding to eIF4E. Phosphorylation of 4E-BP1 frees eIF4E, which can then associate with eIF4G to initiate mRNA translation. Phosphorylation of p70S6K increases the phosphorylation of ribosomal protein S6 (44), which facilitates 5′-terminal oligopyrimidine mRNA translation and increases the synthesis of some ribosomal proteins, translation initiation factors, and elongation factors that play important roles in protein synthesis (28, 43).

Glucocorticoid excess induces negative nitrogen balance in experimental animals and humans by enhancing proteolysis, decreasing transport of amino acids into muscle, and inhibiting protein synthesis (10, 45, 54, 56). Glucocorticoid administration acutely inhibits muscle protein synthesis in rats as early as 4 h (46, 49, 55). The mechanism(s) underlying this effect on protein synthesis remains incompletely understood. Recent studies suggest that the major site for glucocorticoid’s negative action on protein synthesis is the inhibition of mRNA translation initiation (45, 49). In rats, high-dose dexamethasone treatment (100 μg/100 g body wt) acutely inhibits protein synthesis, decreases the phosphorylation of both 4E-BP1 and p70S6K, and lowers the amounts of eIF4E bound to eIF4G (49). Interestingly, more prolonged, but lower (though still pharmacological), doses of dexamethasone (2 mg every 6 h for 3 days) did not affect postabsorptive protein synthesis or the phosphorylation status of either 4E-BP1 or p70S6K in humans (31, 34). However, insulin and amino acids induced activation of 4E-BP1, and/or p70S6K was significantly impaired, both in rats (33) and in humans (31). These observations have shed some light on the potential mechanisms for the interactions between the stimulatory effect of insulin and the inhibitory effect of glucocorticoids on protein synthesis. However, most studies have used pharmacological doses of insulin

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IT IS WELL RECOGNIZED that insulin exerts anabolic actions and glucocorticoids exert catabolic effects on skeletal muscle protein metabolism. Both protein synthesis and degradation are affected by these two hormones. However, the cellular mechanisms underlying the interactions between these factors remain poorly defined.

Insulin activates several key signal intermediates that regulate protein synthesis (47), including protein kinase B (PKB or Akt) (48), eukaryotic initiation factor 4E (eIF4E)-binding protein-1 (4E-BP1, or PHAS-I) (26, 27), and ribosomal protein S6 kinase (p70S6K) (42). Akt is a key element of the phosphatidylinositol 3-kinase

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and/or glucocorticoids, making the physiological significance of these findings less certain. Considering insulin, for example, many investigators have failed to demonstrate a stimulatory effect of physiological insulin concentrations on either whole body or skeletal muscle protein synthesis in adult rats or humans in vivo (1, 9, 16, 20, 36, 37, 39, 41, 59, 62). We have previously reported (22, 32) that physiological hyperinsulinemia stimulates the phosphorylation of p70S6K, but not 4E-BP1, in rat and human skeletal muscle. Whether physiological concentrations of glucocorticoids modulate the activation of the protein synthetic pathway by physiological concentrations of insulin has not been defined.

The major purpose of this study was to examine whether endogenous glucocorticoids modulate insulin-stimulated muscle protein synthesis at physiological stress concentrations in vivo and whether the signaling pathway that regulates protein synthesis via phosphorylation of Akt, 4E-BP1, and p70S6K is affected by adrenalectomy. We used an adrenalectomized rat model with or without glucocorticoid replacement to study the effect of insulin on protein synthesis and the insulin signal transduction pathway. The results showed that insulin at physiological concentrations significantly stimulated skeletal muscle protein synthesis and increased the phosphorylation of Akt, 4E-BP1, and p70S6K in adrenalectomized rats compared with sham-operated animals. Dexamethasone, at a dose selected to replace endogenous glucocorticoids in adrenalectomized rats, blunted the muscle’s response to insulin. These findings suggest that endogenous glucocorticoids play a significant role in blunting insulin’s stimulatory effect on protein synthesis at physiologic concentration in vivo.

MATERIALS AND METHODS

Animal preparation and experimental protocols. Male Sprague-Dawley rats, weighing 225–250 g, were studied 4 days after either a bilateral adrenalectomy or a sham operation under generalized anesthesia using pentobarbital sodium (50 mg/kg ip; Abbott Laboratories, North Chicago, IL). Two separate studies were conducted.

In study 1, as a preliminary study, we examined whether adrenalectomy per se affects the rates of skeletal muscle protein synthesis in two groups of rats (n = 5 each). One group underwent adrenalectomy and the other group had sham operation. All surgeries were done through a midabdominal incision. In sham-operated rats, both adrenal glands were isolated and then left in place. After surgery, rats were maintained on a 12:12-h light-dark cycle with food and water ad libitum. Normal saline was provided instead of water to all adrenalectomized rats to prevent dehydration. Four days after surgery, rats were fasted overnight and then anesthetized with intraperitoneal pentobarbital sodium (50 mg/kg). The external jugular vein, internal carotid artery, and trachea were exposed and cannulated through a midline neck incision. The arterial catheter was connected through a three-way stopcock to a pressure probe. Heart rate and mean arterial pressure were monitored throughout the study (Transonic Systems, Ithaca, NY). Pentobarbital sodium was infused at a variable rate to maintain a steady level of anesthesia throughout the study. After a 45-min baseline period to assure hemodynamic stability and a stable level of anesthesia, all rats received a 3-h saline infusion, and a phenylalanine bolus (150 μmol/100 g body wt) containing [3-¹⁴C]phenylalanine (375 μCi) was injected through the jugular vein 10 min before the completion of saline infusion. Gastrocnemius muscle was removed at the end of the infusion period and was freeze-clamped in liquid nitrogen. All muscle samples were stored at −70°C until analysis.

In study 2, three groups of rats were studied to examine the effect of adrenalectomy on insulin-stimulated translation initiation and protein synthesis. One group of rats (n = 6) had a bilateral adrenalectomy (ADX), another group of animals (n = 6) had a bilateral adrenalectomy and was placed on 5 μg/100 g body wt dexamethasone subcutaneously twice daily (ADX + DEX), and a third group of rats (n = 6) had sham operation (Sham). In ADX + DEX rats, the dose of dexamethasone was considered a physiological stress dose that approximates the glucocorticoid level found in stressed rats (8, 35). Four days after surgery, all rats were prepared as described for study 1. After a 45-min baseline period to assure hemodynamic stability and a stable level of anesthesia, a bolus of rectus muscle phosphorylated 4E-BP1 was injected through the jugular vein, and the infusion period and was freeze-clamped in liquid nitrogen. After another 30-min stabilization period, rats received a 3-mU·kg⁻¹·min⁻¹ euglycemic clamp for 3 h. Whole blood glucose was monitored every 10 min (Accu-Chek; Roche Diagnostics, Indianapolis, IN) throughout the insulin infusion, and 30% dextrose was infused at a variable rate to maintain blood glucose within 10% of basal (13). A phenylalanine bolus (150 μmol/100 g body wt) containing [3-¹⁴C]phenylalanine (375 μCi) was injected through the jugular vein 10 min before the completion of insulin infusion. Rectus muscle was again biopsied at the end of the infusion period and was freeze-clamped in liquid nitrogen. Both muscle biopsies were taken from lower abdominal rectus muscle that was not wounded by prior surgery.

All muscle samples were stored at −70°C until analysis. Plasma insulin concentrations were measured by radiomunoassay in plasma obtained at the beginning and the completion of the study. Plasma corticosterone concentrations were measured by radioimmunoassay in plasma obtained at the beginning of the study to confirm the success of adrenalectomy.

Measurement of skeletal muscle protein synthesis. Protein synthesis was estimated using a [³¹⁴C]phenylalanine flooding technique as described previously (19). In brief, the phenylalanine concentration and the specific radioactivity in plasma and the muscle protein were measured by HPLC and β-counting. The protein synthesis rate is estimated from the radioactivity incorporated into the muscle protein over the 10-min period divided by the tracer specific activity and is presented as a fractional turnover rate (%/day). This method uses the tissue free amino acid pool as the precursor pool for the calculations. The phenylalanine bolus results in a rapid rise in specific radioactivity of phenylalanine in tissues, and two recent studies have demonstrated that the phenylalanine specific activity of this pool is similar to that of the true precursor pool in skeletal muscle in neonatal pigs (12) and dogs (5).

Western immunoblotting technique. Pieces (~30 mg) of frozen rectus muscle were weighed and powdered in frozen 25 mM Tri-HCl buffer (26 mM KF and 5 mM EDTA, pH 7.5). Tissue was then disrupted by sonication with the use of a microtip probe (0.5 s on and 0.5 s off for 45 s total) at a 3.0 power setting on the Fisher XL2020 sonicator (Fisher Scientific, Pittsburgh, PA). The homogenate was centrifuged at 2,000 rpm for 2 min, and the protein content of the supernatant...
tant was determined using the Bradford method (3). For Akt, one aliquot of the supernatant, containing ~60 µg of protein, was diluted with an equal volume of SDS sample buffer and electrophoresed on an 8% polyacrylamide gel. For 4E-BP1, one aliquot of the supernatant, containing ~60 µg of protein, was diluted with an equal volume of SDS sample buffer and electrophoresed on a 15% polyacrylamide gel. For p70^{S6K}, one aliquot of supernatant, containing ~50 µg of protein, was diluted with an equal volume of SDS sample buffer and electrophoresed on an 8% polyacrylamide gel. Proteins were then electrophoretically transferred to nitrocellulose membranes. After being blocked with 5% low-fat milk in Tris-buffered saline plus Tween 20, membranes were incubated with either rabbit polyclonal Akt antibody or phospho-Akt (Ser^{473}) antibody (New England BioLabs, Beverly, MA) overnight at 4–8°C, or rabbit anti-4E-BP1 or p70^{S6K} (Santa Cruz Biotechnology, Santa Cruz, CA) for 1 h at room temperature. This was followed by a donkey anti-rabbit IgG coupled to horseradish peroxidase, and the blot was developed using an enhanced chemiluminescence Western blotting kit (Amer sham Life Sciences, Piscataway, NJ).

**Quantitation of Akt, 4E-BP1, and p70^{S6K} phosphorylation status.** Autoradiographic films were scanned densitometrically (Molecular Dynamics, Piscataway, NJ) and quantitated using ImageQuant 3.3. Figure 1 illustrates the Akt, 4E-BP1, and p70^{S6K} phosphorylation status based on Western blots analysis. For Akt, both the total and phospho-Akt (Ser^{473}) densities were quantitated, and the ratios of phosphospecific density to total density were calculated. To quantify the extent of phosphorylation of 4E-BP1, we measured the ratio of the intensity of the most slowly migrating species (γ) to that of the total intensity (α + β + γ). For p70^{S6K}, we determined the ratio of the more slowly migrating forms (β + γ) to the total (α + β + γ). Exploiting the different electrophoretic behaviors of proteins with various amounts of phosphorylation on SDS-PAGE allows the simultaneous quantification of multiple forms of both proteins, as well as internal normalization both for the recovery of target proteins from tissue and for loading of gels. The available data support a good correlation between activity and electrophoretic mobility for both 4E-BP1 (15, 40, 42) and p70^{S6K} (61). We have observed good reproducibility of the fractional phosphorylation ratios for 4E-BP1 and p70^{S6K} after loading gels with different amounts of protein (33).

![Fig. 1. Rat skeletal muscle phospho-Akt (Ser^{473}) (A), total Akt (B), eukaryotic initiation factor 4E-binding protein-1 (4E-BP1) (C), and ribosomal protein S6 kinase (p70^{S6K}) (D) gel patterns on SDS-PAGE. For 4E-BP1 and p70^{S6K}, the α-band is the least phosphorylated portion and has the most rapid electrophoretic mobility. The γ-band is the most phosphorylated form and moves the slowest. The β-band is phosphorylated more than the α-band and less than the γ-band, and shows up in between on SDS-PAGE. The Western blot shows the pattern from sham-operated (Sham; lanes 1 and 2), adrenalectomized (ADX; lanes 3 and 4) and ADX + treatment with dexamethasone (ADX + DEX; lanes 5 and 6) rats. Lanes 1, 3, and 5, basal period; lanes 2, 4, and 6, after insulin infusion.](image)
Effect of adrenalectomy and insulin infusion on phosphorylation of Akt. The basal phosphorylation of Akt was comparable between Sham and ADX rats (0.212 ± 0.04 vs. 0.162 ± 0.03, respectively, P = 0.25). Glucocorticoid replacement slightly decreased basal phosphorylation state of Akt to 0.11 ± 0.02, though this was not statistically significantly different from Sham or ADX rats (P = 0.11 and 0.47 respectively). Insulin treatment significantly increased the phosphorylation of Akt in ADX rats by 112% (0.162 ± 0.03 vs. 0.345 ± 0.08, P < 0.03), but not in sham rats. Dexamethasone treatment abolished this insulin-stimulated Akt phosphorylation in ADX + DEX rats (Fig. 2).

Effect of adrenalectomy and insulin infusion on phosphorylation of 4E-BP1. To quantitate the phosphorylation status of 4E-BP1, we calculated the ratio of the intensity of the most slowly migrating γ-band to that of the total integrated intensity (α + β + γ; Fig. 3). The γ-band is the highly phosphorylated form of 4E-BP1, and an increase in the quantity of this form would correspond to an increase in the phosphorylation of 4E-BP1 and a greater amount of eIF4E available to initiate translation. Basal phosphorylation status of 4E-BP1 was comparable among all three groups of rats, suggesting that adrenalectomy per se does not change the phosphorylation status of this protein. Physiological hyperinsulinemia did not stimulate 4E-BP1 phosphorylation in Sham rats (0.26 ± 0.02 vs. 0.29 ± 0.04, basal vs. insulin respectively, P = 0.24) but did significantly increase the phosphorylation status of 4E-BP1 in ADX rats (0.23 ± 0.02 vs. 0.39 ± 0.03, P = 0.003). Dexamethasone treatment blunted this increase in 4E-BP1 phosphorylation in ADX + DEX rats (0.26 ± 0.02 vs. 0.34 ± 0.06, P = 0.24).

Effect of adrenalectomy and insulin infusion on phosphorylation of p70S6K. The extent of phosphorylation of p70S6K was quantitated by measuring the ratio of the intensity of the more slowly migrating species (β + γ) to that of the total integrated intensity (α + β + γ; Figs. 2 and 4). The overall p70S6K activity is dependent on the phosphorylation of at least seven Ser/Thr residues at three separate domains (44, 61). The more slowly migrating β- and γ-bands represent the more highly phosphorylated forms of p70S6K and generally correspond to species with greater kinase activity. Similar to the effect of adrenalectomy on 4E-BP1 phosphorylation, adrenalectomy appears to have no effect on the baseline phosphorylation of p70S6K, which was comparable among all three groups of rats. Insulin treatment did not stimulate p70S6K phosphorylation in Sham rats (0.50 ± 0.06 vs. 0.46 ± 0.07, basal vs. insulin respectively, P = 0.42) but did significantly increase the phosphorylation status of p70S6K in ADX rats (0.42 ± 0.05 vs. 0.63 ± 0.04, Fig. 4). Dexamethasone treatment again blunted this increase in p70S6K phosphorylation in ADX + DEX rats (0.40 ± 0.09 vs. 0.49 ± 0.07, P = 0.24).

Effect of adrenalectomy on insulin-stimulated protein synthesis. Figure 5 illustrates the rates of protein synthesis observed in Sham, ADX, and ADX + DEX rats infused with insulin. After 3 h of euglycemic hyperinsulinemia, the skeletal muscle fractional protein synthesis rate in ADX rats was 30% higher than that in Sham rats. *P < 0.03 vs. basal.
Fig. 5. Effect of adrenalectomy on insulin-stimulated skeletal muscle protein synthesis. Protein synthesis was estimated using the flooding bolus of \(^{3}H\)phenylalanine technique and expressed as fractional \%/day. Insulin treatment significantly increased the protein synthesis rate in ADX rats. *P = 0.01 vs. Sham.

Sham rats (8.8 ± 0.9 vs. 6.1 ± 0.4%/day, P = 0.01). Glucocorticoid replacement with dexamethasone blunted this increase in insulin-stimulated protein synthesis to 7.8 ± 1.4%/day.

DISCUSSION

The adrenal glands produce glucocorticoids, mineralocorticoids, catecholamines, and small amounts of sex steroids. Both glucocorticoids and catecholamines antagonize insulin's hypoglycemic effect. Mineralocorticoids induce synthesis of specific proteins but have no known effect on bulk muscle protein metabolism. Catecholamines acutely promote protein anabolism by decreasing proteolysis (17, 29). It appears that the major adrenal hormones with negative impact on protein synthesis are glucocorticoid hormones. Glucocorticoids at pharmacological concentrations attenuate translation initiation and protein synthesis (50–52) and blunt the anabolic action of insulin, mixed amino acids, and branch-chain amino acids by inhibiting the phosphorylation of 4E-BP1 and p70\(^{S6K}\) (31, 33). Whether physiological stress concentrations of glucocorticoids modulate the anabolic action of insulin is not clear. In the present study, 3 h of physiological euglycemic hyperinsulinemia in adrenalectomized rats significantly increased the skeletal muscle protein synthesis and the phosphorylation of Akt, 4E-BP1, and p70\(^{S6K}\), three key regulatory proteins in the insulin-stimulated protein synthesis signal pathway, but had no effect in sham-operated rats. Dexamethasone treatment to replace physiological stress levels of glucocorticoids blunted insulin's action on all parameters observed. These findings suggest that glucocorticoid insufficiency produced by adrenalectomy enhances the insulin sensitivity of muscle protein synthesis and that endogenously present glucocorticoids can modulate insulin's anabolic action on muscle protein synthesis in vivo.

At pharmacological doses, insulin has consistently been shown to stimulate protein synthesis both in vivo and in vitro (23–25, 38). A similar effect of physiological hyperinsulinemia is consistently seen in immature animals (11). Insulin activates the PI 3-kinase-mTOR pathway and increases the phosphorylations of 4E-BP1 and p70\(^{S6K}\) (26, 33, 53). Within this pathway, Akt, 4E-BP1, and p70\(^{S6K}\) are three proteins with pivotal roles in insulin-stimulated protein synthesis. Akt is activated in response to activation of PI 3-kinase by phosphorylation at Thr\(^{308}\) and Ser\(^{473}\), and its activation is necessary for insulin-stimulated mTOR protein kinase activity (48). Both 4E-BP1 and p70\(^{S6K}\) are downstream of Akt-mTOR in the insulin signal transduction pathway, and phosphorylation of these two proteins is critical in the initiation and maintenance of protein synthesis. Phosphorylation of 4E-BP1 increases the amount of eIF4E available for the formation of the translation initiation complex (30), a necessary step for the translation of mRNAs with \(m^{7}GTP\) at the 5′ cap, and phosphorylation of p70\(^{S6K}\) correlates strongly with increases in ribosomal S6 kinase activity (61), which appears to be necessary for maintaining the apparatus required for ongoing protein synthesis. In sham-operated rats, insulin did not stimulate protein synthesis, nor did it activate the phosphorylation of Akt, 4E-BP1, and p70\(^{S6K}\), confirming the absence of a stimulatory effect of physiological hyperinsulinemia on bulk protein synthesis in skeletal muscle in vivo. However, our results clearly demonstrated a significant increase in protein synthesis after insulin administration in adrenalectomized rats. In accord with this increased protein synthesis, insulin also increased the phosphorylation status of Akt, 4E-BP1, and p70\(^{S6K}\) in adrenalectomized rats. These findings suggest that adrenalectomized rats have heightened sensitivity to the effect of physiological concentrations of insulin on protein metabolism. Our observation of 4-day adrenalectomy per se having no significant impact on the rates of protein synthesis in the gastrocnemius muscle in *study 1* is entirely consistent with our findings of no significant difference in the basal phosphorylation of Akt, 4E-BP1, and p70\(^{S6K}\) in the rectus among all treatment groups in *study 2*. A previous report (35) has also shown that the whole body protein synthesis rates are essentially the same in adrenalectomized rats and adrenalectomized rats receiving stress dose dexamethasone (same dose as ours).

It is interesting that the glucose infusion rates were not different among the three groups of rats studied. Akt activation is thought to play a critical role in insulin-stimulated GLUT4 translocation and glucose uptake (57, 60). With significantly increased phosphorylation of Akt in the ADX rats following insulin treatment, one would expect that the glucose infusion rates would be higher in this group of rats than in the Sham rats. However, multiple confounding factors may have contributed to the lack of higher glucose infusion rates in the ADX rats in the present study. First, the basal blood glucose concentrations were significantly lower in ADX rats than in Sham rats, and the glucose was infused to maintain the blood glucose concentrations at their respective basal levels. Second, ADX rats have lower body weight and less muscle mass, the site for insulin-stimulated glucose disposal. Also, ADX rats,
being more stressed than Sham rats, may have higher levels of growth hormone, which is a counterregulatory hormone and antagonistic toward insulin's hypoglycemic effect. The elevated basal insulin concentrations and slightly lower, though not statistically significant, glucose infusion rate observed in ADX + DEX rats than in Sham and ADX rats are consistent with relative insulin resistance associated with physiological stress concentrations of glucocorticoids, a well-known clinical observation in patients with persistent and severe physical stress.

That dexamethasone treatment alone, administered at doses selected to replace glucocorticoids to physiological stress concentrations without mineralocorticoid or catecholamine replacement, blunted insulin-induced protein synthesis and phosphorylation of Akt, 4E-BP1, and p70S6K in adrenalectomized rats suggests that the increment in insulin sensitivity observed in adrenalectomized rats is secondary to endogenous glucocorticoid deficiency, not mineralocorticoid or catecholamine deficiency. Inasmuch as it has been previously observed that glucocorticoids given in excess blunt the action of amino acids (31, 33) and pharmacological doses of insulin (33) to stimulate protein synthesis, the present findings suggest that glucocorticoids modulate protein synthesis over the entire range of their availability. In addition, the present findings suggest that the presence of endogenous glucocorticoid hormones may be largely responsible for the lack of stimulatory effect of insulin on protein synthesis observed in most in vivo studies using physiological concentrations of insulin (1, 2, 6, 14, 18, 36, 37, 41).

We have previously observed (22, 32) that pharmacological doses of insulin enhanced the phosphorylation of both 4E-BP1 and p70S6K, whereas physiological hyperinsulinemia differentially phosphorylated p70S6K, but not 4E-BP1, in both human and rat skeletal muscle. In the present study, the physiological hyperinsulinemic clamp did not increase the phosphorylation of p70S6K in the sham-operated rats. The lack of effect of insulin on downstream signaling pathways in the sham-operated rats could be due, in part, to experimental conditions, including the involvement of increased concentrations of glucocorticoids. Abdominal surgery and prolonged anesthesia are associated with significant stress and increased endogenous production of glucocorticoid hormones sufficient to cause insulin resistance and may have diminished the response to insulin in sham-operated rats. Excessive glucocorticoids decrease total tyrosine phosphorylation of the insulin receptor and insulin receptor substrate 1 (IRS-1) as well as the protein content of IRS-1 (21), decrease the phosphorylation of both 4E-BP1 and p70S6K (50), and impair insulin-induced phosphorylation of 4E-BP1 and p70S6K in rats (33). Shah et al. (51) have demonstrated that p70S6K is more sensitive to inhibition by glucocorticoids under growth-promoting conditions than is 4E-BP1.

In the present study, we used the rectus muscle to assay the phosphorylation state of Akt, 4E-BP1, and p70S6K. Rectus muscle was selected as it allows repeated measurements from snip biopsies with minimal trauma to the other portions of the muscle. In the present study, there would be an additional concern, inasmuch as an abdominal incision was used for the adrenalectomy four days before the infusion study. However, we believe that this surgery per se had little impact on the results observed for the following reasons. 1) Each animal served as its own control to examine the phosphorylation status of signaling intermediates. 2) All animals, including the sham-operated animals, received the same abdominal surgery, and there were no differences in the basal phosphorylation status of these proteins among all groups. 3) Aseptic inflammation, as might be expected at the site of abdominal incision, does not appear to affect the phosphorylation of 4E-BP1 (58). Finally, 4) the muscle biopsies from the rectus were made at sites in the lower abdominal rectus muscle, making every effort to avoid previously incised areas of muscle.

In conclusion, the present results suggest an important action of physiological stress concentrations of glucocorticoids in the regulation of protein synthesis by physiological hyperinsulinemia. In particular, our results suggest that low doses of glucocorticoids are able to specifically block the activation by insulin of three proteins that regulate mRNA translation: Akt, 4E-BP1, and p70S6K. This action of endogenous glucocorticoids to counterpoise the effect of insulin on the phosphorylation of translation-regulatory proteins may modulate the effect of physiological doses of insulin to stimulate protein synthesis in vivo.

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