Effect of ACTH and CRH on Plasma Levels of Cortisol and Prostaglandin F\textsubscript{2\alpha} Metabolite in Cycling Gilts and Castrated Boars

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Madej A, Mwanza AM, Kindahl H, Einarsson S: Effect of ACTH and CRH on plasma levels of cortisol and prostaglandin F\textsubscript{2\alpha} metabolite in cycling gilts and castrated boars. Acta vet. scand. 2005, 46, 249-256. – The present study was designed to evaluate the effects of synthetic ACTH (1-24, tetracosactid) and porcine CRH on the plasma levels of cortisol and PGF\textsubscript{2\alpha} metabolite in cycling gilts (n = 3) and castrated boars (n = 3). The experiments were designed as crossover studies for each gender separately. Each animal received, during three consecutive days; 1) ACTH (Synacthen\textsuperscript{®} Depot) at a dose of 10 µg/kg body weight in 5 ml physiological saline, 2) porcine CRH at a dose 0.6 µg/kg body weight in 5 ml physiological saline or 3) physiological saline (5 ml). The test substances were administered via an indwelling jugular cannula in randomized order according to a Latin square. The administration of ACTH to cycling gilts resulted in concomitant elevations of cortisol and PGF\textsubscript{2\alpha} metabolite with peak levels reached at 70.0 ± 10.0 and 33.3 ± 6.7 min, respectively. Similarly, the administration of ACTH to castrated boars resulted in concomitant elevation of cortisol and PGF\textsubscript{2\alpha} metabolite with peak levels reached at 60.0 ± 0.0 and 20.0 ± 0.0 min, respectively. Cortisol peaked at 20 min after administration of CRH in both cycling gilts and castrated boars with maximum levels of 149.3 ± 16.5 nmol/l and 138.3 ± 10.1 nmol/l, respectively. It can be concluded that administration of synthetic ACTH (tetracosactid) to pigs caused a concomitant elevation of cortisol and PGF\textsubscript{2\alpha} metabolite levels in both cycling gilts as well as castrated boars. The administration of CRH to pigs resulted in an elevation of cortisol levels in both cycling gilts and castrated boars. Conversely, PGF\textsubscript{2\alpha} metabolite levels were not influenced by the administration of CRH either in cycling gilts or in castrated boars.

ACTH; CRH; cortisol; PGF\textsubscript{2\alpha} metabolite; gilts; castrated boars

Introduction
Corticotropin releasing hormone (CRH) plays a central role in regulating the release of adrenocorticotropic hormone (ACTH) during a stress response. ACTH acts on the adrenal glands, inducing the secretion of cortisol. In our previous study, we reported that ACTH administration to ovariectomized gilts results in the plasma elevation of cortisol, progesterone and prostaglandin F\textsubscript{2\alpha} metabolite (Mwanza et al. 2000b). How ACTH is capable of stimulating the secretion of PGF\textsubscript{2\alpha} metabolite remains unanswered. However, it was previously suggested by Laychok & Rubin (1975) that ACTH enhances the conversion in vitro of \textsuperscript{3}H-arachidonic acid to prostaglandins in feline adrenocortical cells. The findings of Anthonisen et al. (1997) indi-
cater that prostaglandins in the brain interact in their stimulatory regulation of ACTH secretion. Such an interaction may also be involved in prostaglandins mediation of the ACTH response to immunochallenges. Abraham et al. (1998) reported that stimulation of porcine pituitary cells by relatively low concentrations of prostaglandin E₂ support increased secretion of ACTH but exposure to greater concentrations of this prostaglandin in fact suppresses ACTH secretion. Food deprivation, which is a form of stress, has been shown to result in the plasma elevation of both cortisol and PGF₂α metabolite (Tsuma et al. 1996; Mburu et al. 1998; Mwanza et al. 2000a; Razdan et al. 2001). Intracerebroventricular as well as intravenous injections of CRH resulted in an increased plasma cortisol concentration in pigs (Sakumato et al. 2004; Lang et al. 2004). Previously, it was suggested that CRH may also act directly or indirectly to enhance cortisol secretion beyond the level achieved through adrenal stimulation by ACTH (Minton & Parsons 1993). The objectives of the present study were to evaluate the effects of synthetic ACTH (tetra-cosactid) and porcine CRH on the plasma levels of cortisol and PGF₂α metabolite in cycling gilts and castrated boars.

Materials and Methods

Animals

Six crossbred pigs (Landrace x Yorkshire; three gilts and three castrated boars) aged approximately 6 months weighing between 110 and 125 kg were used for this experiment. The pigs were brought to the Division of Comparative Reproduction, Obstetrics and Udder Health, and were housed in individual pens. The stable had a light period of 12 h starting from 06:30 h and the room temperature varied between 20 and 23°C. The pigs were fed according to the Swedish breeding stock standard (Simonsson 1994). The gilts were stimulated by boars in the vicinity and were expected to come in oestrus within the first week after arrival. After the second oestrus, the experiments were carried out in the early luteal phase (days 5-10). The gilts were checked twice daily at 06:00 h and 18:00 h for signs of oestrus in the presence of a fertile boar by back-pressure test. All the six animals were vein-cannulated (Rodriguez & Kunavongkrit 1983) at about one week before the experiments. The experiments were designed as crossover studies for each gender separately. Each animal received, during three consecutive days; 1) ACTH (1-24) (Synacthen® Depot, Novartis Pharma AG, Basel, Schweiz) at a dose of 10 µg/kg body weight in 5 ml physiological saline, 2) porcine CRH (American Peptide Company, Inc., Sunnyvale, CA, USA) at a dose 0.6 µg/kg body weight in 5 ml physiological saline or 3) physiological saline (5 ml). The test substances were administered via an indwelling jugular cannula in randomized order according to a Latin square. On the day of the experiment, blood samples were taken at -40, -20 min and immediately before injection. Treatment was performed at 10:00 h (time = 0) and blood samples were taken 20, 40, 60, 80, 100, 120, 140, 160, 180, 210, 240, 270, 300, 330 and 360 min after injection. Blood was collected in ten ml heparinised tubes, centrifuged immediately and plasma stored at -20°C until analysed. The care of the animals and the experimental design of this study were approved by the Local Animal Ethics Committee in Uppsala, Sweden.

Hormone assays

Cortisol. Plasma cortisol was determined by radioimmunoassay (Coat-A-Count Cortisol, Diagnostic Products Corporation, Los Angeles, CA, USA) according to the manufacturer’s instructions. Serial dilutions of porcine plasma with high concentrations of cortisol produced displacement curves parallel to the standard curve. The intra-assay coefficients of variation

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calculated from 5 assays were 22% at 14 nmol/l, 14% at 28 nmol/l and decreased below 8% for concentrations between 138 and 552 nmol/l. The inter-assay coefficients of variation for three control samples were 13% (33 nmol/l), 9% (74 nmol/l) and 9% (541 nmol/l). The average detection limit of the assay was 7 nmol/l. Prostaglandin F$_{2\alpha}$ metabolite. The main initial blood plasma metabolite of prostaglandin F$_{2\alpha}$, 15-keto-13,14-dihydro-PGF$_{2\alpha}$ (15-ketodihydro-PGF$_{2\alpha}$), was analysed by radioimmunoassay according to Kunavongkrit et al. (1983). The relative cross-reactions of the antibody were 16% with 15-keto-PGF$_{2\alpha}$ and 4% with 13,14-dihydro-PGF$_{2\alpha}$. The intra-assay coefficients of variation ranged between 3.4 and 7.6% for different ranges of the standard curve and the inter-assay coefficient of variation was around 14%. The practical limit of sensitivity for the assay analyzing 0.2 ml of plasma was 60 pmol/l.

Statistical analyses
Data were examined by analysis of variance using MIXED procedure according to SAS package (Statistical Analysis Systems 1989). Additionally, the area under the curve, peak value, time and duration of the peak were calculated for each animal according to the GraphPad Prism version 3.02 for Windows (GraphPad Software, San Diego, CA, USA). Data are expressed as means ± S.E.M. Probabilities less then 0.05 were considered significant.

Results
No significant (P>0.05) differences were seen in the pretreatment plasma levels of cortisol or PGF$_{2\alpha}$ metabolite between the saline, ACTH and CRH treated cycling gilts (Figures 1 and 2). The administration of ACTH to cycling gilts resulted in concomitant elevations of cortisol (Figure 1) and PGF$_{2\alpha}$ metabolite (Figure 2) with peak levels reached at 70.0 ± 10.0 and 33.3 ± 6.7 min, respectively. The durations of the peaks were 153.3 ± 28.2 and 103.2 ± 11.4 min, respectively and their maximum concentrations were 270.7 ± 16.5 nmol/l and 1517.7 ± 137.2 pmol/l, respectively.

No significant (P>0.05) differences were seen in the pretreatment plasma levels of cortisol or PGF$_{2\alpha}$ metabolite between the saline, ACTH and CRH treated castrated boars (Figures 3 and 4). The administration of ACTH to castrated boars resulted in concomitant elevation of cortisol (Figure 3) and PGF$_{2\alpha}$ metabolite (Figure 4) with peak levels reached at 60.0 ± 0.0 and 20.0 ± 0.0 min, respectively. The durations of these peaks were 199.1 ± 30.0 and 86.3 ± 13.8 min, respectively and their maximum concentrations were 289.0 ± 10.1 nmol/l and 1262.3 ± 53.2 pmol/l, respectively.

The administration of CRH to both cycling gilts and castrated boars resulted in the cortisol peak 20 min later with maximum levels of 149.3 ± 16.5 nmol/l (Figure 1) and 138.3 ± 10.1 nmol/l (Figure 3), respectively. The durations of these peaks were 57.3 ± 18.5 min and 255.2 ± 43.6 min, respectively.

Prostaglandin F$_{2\alpha}$ metabolite levels were not influenced by the injection of CRH either in cycling gilts or castrated boars (Figures 2 and 4). Physiological saline did not alter significantly either cortisol or PGF$_{2\alpha}$ metabolite levels in any animal (Figures 1-4).

No significant (P>0.05) differences were seen in the measured responses between females and males.

Discussion
The present study clearly demonstrates that cortisol reach peak levels much lower and earlier in CRH (approximately after 20 min) than in ACTH (approximately after 70 min) treated cycling gilts or castrated boars. This confirmed earlier results by Beerda et al. (2004) who reported that the cortisol concentration peaked
Figure 1. Plasma cortisol concentrations (LSmeans ± SEM) in cycling gilts given the injection (time = 0) of saline (□, n = 3), ACTH (■, n = 3) and CRH (○, n = 3).

Figure 2. Plasma PGF$_{2\alpha}$ metabolite concentrations (LSmeans ± SEM) in cycling gilts given the injection (time = 0) of saline (□, n = 3), ACTH (■, n = 3) and CRH (○, n = 3).
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Figure 3. Plasma cortisol concentrations (LSmeans ± SEM) in castrated boars given the injection (time = 0) of saline (□, n = 3), ACTH (■, n = 3) and CRH (○, n = 3).

Figure 4. Plasma PGF$_{2\alpha}$ metabolite concentrations (LSmeans ± SEM) in castrated boars given the injection (time = 0) of saline (□, n = 3), ACTH (■, n = 3) and CRH (○, n = 3).
approximately 30 min after administration of CRH and approximately 60-90 min after administration of synthetic ACTH to dairy cows. The present study also demonstrates that the administration of ACTH stimulates a concomitant elevation of both cortisol and PGF$_{2\alpha}$ metabolite levels in both cycling gilts and castrated boars. In addition, peak PGF$_{2\alpha}$ metabolite levels occur earlier than peak cortisol levels. Apparently, it takes approximately twice the time for cortisol than for PGF$_{2\alpha}$ metabolite to reach peak levels following ACTH administration. This is consistent with our previous findings in ovariectomized gilts (Mwanza et al. 2000b) and suggests that ACTH stimulates the secretion of PGF$_{2\alpha}$ earlier than cortisol. The frequency of blood collection may have impact on the occurrence of PGF$_{2\alpha}$ metabolite peak in relation to cortisol peak. When blood samples were taken only at 1-h intervals, both PGF$_{2\alpha}$ metabolite and cortisol peak were seen one hour after ACTH administration in recently ovulated sows (Razdan et al. 2002).

Cooke & Ahmad (1994) have demonstrated that daily administration of ACTH from day 13 to day 16 of the oestrous cycle in multiparous Welsh Mountain ewes suppressed the levels of PGF$_{2\alpha}$ metabolite. They further showed that in ovariectomized multiparous Welsh Mountain ewes, primed first with progesterone and then with oestradiol-17β, ACTH reduced the ability of oxytocin to cause the release of PGF$_{2\alpha}$ into the peripheral circulation. However, there is evidence that feline and rat adrenocortical cells synthesise prostaglandins F$_{2\alpha}$ and E$_{2}$ and that the total prostaglandins synthesis is stimulated by ACTH (Laychock & Rubin 1976; Chanderbhan et al. 1979). Winter et al. (1990) demonstrated that in vitro, the cytokine interleukin-1 enhances the conversion of $^{3}$H-arachidonic acid to prostaglandins by cultured bovine adrenal cells. The secreted prostaglandins i.e. PGD$_{2}$, PGF$_{2\alpha}$ and PGE$_{2}$ were in turn found to stimulate cortisol secretion. Furthermore, Nashuhita et al. (1997) reported that intravenously administered PGE$_{1}$, PGE$_{2}$ or PGF$_{2\alpha}$ had significant ACTH-releasing activity in the rat and suggested that prostaglandins are playing a role in regulating the hypothalmo-pituitary-adrenal axis. In sows, injection of PGF$_{2\alpha}$ after ovulation resulted in a dramatic cortisol elevation, which lasted approximately 1.5 h (Mwanza et al. 2002).

In contrast to ACTH-treated pigs, no peak PGF$_{2\alpha}$ metabolite levels were seen in any CRH treatment. We can speculate that a combination of CRH and lysine vasopressin (LVP) could have been a better option since LVP + CRH was seen to have a better ACTH response than CRF or LVP alone in pigs (Minton & Parsons 1993). It might also simply indicate that CRH does not stimulate the secretion of PGF$_{2\alpha}$.

Interestingly, food deprivation which is a form of stress has been shown to result in the plasma elevation of both cortisol and PGF$_{2\alpha}$ metabolite (Mburu et al. 1998; Mwanza et al. 2000a; Razdan et al. 2001; Tsuma et al. 1996). It is postulated (Silver & Fowden 1982) that in food deprived animals, PGF$_{2\alpha}$ metabolite levels are elevated owing to increased levels of free fatty acids that includes arachidonic acid, the precursor of prostaglandin synthesis. In addition, Madej et al. (2005) reported that during artificial insemination of sows housed in crates, a dramatic elevation of cortisol levels was seen before PGF$_{2\alpha}$ metabolite reached its maximum. It is still unclear what role if any ACTH plays either directly or indirectly in the stimulation of PGF$_{2\alpha}$ production.

It can be concluded from the present study that the administration of synthetic ACTH to pigs at a dose of 10 µg/kg body weight caused a concomitant increase of cortisol and PGF$_{2\alpha}$ metabolite levels in both cycling gilt as well as castrated boars. The administration of CRH to pigs resulted in an elevation of cortisol levels in
both cycling gilts and castrated boars. Conversely, PGF$_{2\alpha}$ metabolite levels were not influenced by the administration of CRH either in cycling gilts or in castrated boars.

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References

Abraham EJ, Morris-Hardeman JN, Sswenson LM, Knoppel EL, Ramanathan B, Wright KJ, Grieger DM, Minton JE: Pituitary function in the acute phase response in domestic farm animals: Cytokines, prostaglandins, and secretion of ACTH. Domest. Anim. Endocrinol. 1998, 15, 389-396.

Anthonisen M, Knigge U, Kjoer A, Warberg J: Histamine and prostaglandin interaction in the central regulation of ACTH secretion. Neuroendocrinology 1997, 66, 68-74.

Beerda B, Kornalijnslijper JE, van der Werf JTN, Nordhuizen-Stassen EN, Hopster H: Effect of milk production capacity and metabolic status on HP A function in early postpartum dairy cows. J. Dairy Sci. 2004, 87, 2094-2102.

Chanderbhan R, Hodges VA, Treadwell CR, Vahouny GV: Prostaglandins synthesis in rat adrenocortical cells. J. Lipid Res. 1979, 20, 116-124.

Cooke RG, Ahmad N: The effect of ACTH on oxytocin-induced release of prostaglandin F$_{2\alpha}$ and on uterine oxytocin receptors in the ewe. Anim. Reprod. Sci. 1994, 35, 201-208.

Kunavongkrit A, Kindahl H, Madej A: Clinical and endocrinological studies in primiparous zero-weaned sows. 2. Hormonal patterns of normal cycling sows after zero-weaning. Zbl. Vet. Med. A 1983, 30, 616-624.

Lang, A, Kaeneke K, Kindahl H, Madej, A, Einarsson S: Influence of CRH and ACTH administration on endocrine profile and ovulation in sows. Reprod. Domest. Anim. 2004, 39, 181-189.

Laychock SG, Rubin RP: ACTH-induced prostaglandin biosynthesis from 3H-arachidonic acid by adrenocortical cells. Prostaglandins 1975, 10, 529-540.

Laychock SG, Rubin RP: Radioimmunoassay measurements of ACTH-facilitated PGE$_2$ and PGF$_{2\alpha}$ release from isolated cat adrenocortical cells. Prostaglandins 1976, 11, 753-765.

Madej A, Lang A, Brandt Y, Kindahl H, Madsen MT, Einarsson S: Factors regulating ovarian function in pigs. Domest. Anim. Endocrinol. 2005, 29, 347-361.

Mburu JN, Einarsson S, Kindahl H, Madej A, Rodriguez-Martinez H: Effects of food post-ovulatory deprivation on oviductal sperm concentration, embryo development and hormonal profiles in the pig. Anim. Reprod. Sci. 1998, 52, 221-234.

Minton JE, Parsons KM: Adrenocorticotrophic hormone and cortisol response to corticotropin-releasing factor and lysine vasopressin in pigs. J. Anim. Sci. 1993, 71, 724-729.

Mwanza AM, Einarsson S, Madej A, Lundeheim N, Rodriguez-Martinez H, Kindahl H: Postovulatory effect of repeated administration of prostaglandin F$_{2\alpha}$ on the endocrine status, ova transport, binding of accessory spermatozoa to the zona pellucida and embryo development of recently ovulated sows. Theriogenology 2002, 58, 1111-1124.

Mwanza AM, Englund P, Kindahl H, Lundeheim N, Einarsson S: Effects of postovulatory food deprivation on the hormonal profiles, activity of the oviduct and ova transport in sows. Anim. Reprod. Sci. 2000a, 59, 185-199.

Mwanza AM, Madej, A, Kindahl H, Lundeheim N, Einarsson S: Plasma levels of cortisol, progesterone, oestradiol-17β and prostaglandin F$_{2\alpha}$ metabolite after ACTH (Synacthen Depot$^{\text{PR}}$) administration in ovariectomized gilts. J. Vet. Med. A 2000b, 47, 193-200.

Nasushita R., Watanobe H, Takebe K: A comparative study of adrenocorticotropic-releasing activity of prostaglandin E1, E2, F2 alpha and D2 in the rat. Prostaglandins Leukot. Essent. Fatty Acids 1997, 56, 165-168.

Razdan P, Mwanza AM, Kindahl H, Hultén F, Einarsson S: Impact of postovulatory food deprivation on the ova transport, hormonal profiles and metabolic changes in sows. Acta vet. scand. 2001, 42, 45-55.

Razdan P, Mwanza AM, Kindahl H, Rodriguez-Martinez H, Hultén F, Einarsson S: Effect of repeated ACTH-stimulation on early embryonic development and hormonal profiles in sows. Anim. Reprod. Sci. 2002, 70, 127-137.

Rodriguez-Martinez H, Kunavongkrit A: Chronic venous catheterisation for frequent blood sampling in unrestrained pigs. Acta. vet. scand. 1983, 42, 45-55.
Sammanfattning

Behandlingseffekt av syntetiskt ACTH (tetracosactid) och CRH på blodplasmakoncentrationerna av kortisol och PGF$_{2\alpha}$ metaboliten hos gyltor med normal brunstcykel och hos kastrerade galtar.

Målsättningen med denna studie var att utvärdera behandlingseffekten av syntetiskt ACTH (1-24, tetracosactid) och CRH från gris på blodplasmakoncentrationerna av kortisol och PGF$_{2\alpha}$ metaboliten hos gyltor med normal brunstcykel och hos kastrerade galtar. Experimenten utfördes enligt crossover modellen separat för varje kön. Varje djur behandlades under tre dagar med 1) ACTH (Synaethen® Depot), 10 µg/kg kroppsvikt i 5 ml fysiologisk koksaltlösning, 2) CRH från gris 0,6 µg/kg kroppsvikt i 5 ml fysiologisk koksaltlösning eller 3) 5 ml fysiologisk koksaltlösning. Testsubstanserna injicierades via en permanent jugularkateter slumpartat enligt Latinkvadrat principen. Behandlingen av gyltor med ACTH resulterade i samtidig stegring av kortisol och PGF$_{2\alpha}$ metaboliten, med högsta koncentrationerna efter 70,0 ± 10,0 respektive 33,3 ± 6,7 minuter. På samma sätt resulterade behandling av kastrerade galtar med ACTH i samtidig stegring av kortisol och PGF$_{2\alpha}$ metaboliten och med högsta koncentrationerna efter 60,0 ± 0,0 respektive 20,0 ± 0,0 minuter. Kortisol nådde sitt högsta värde 20 minuter efter behandling med CRH både hos gyltor (149,3 ± 16,5 nmol/l) och kastrerade galtar (138,3 ± 10,1 nmol/l). Sammanfattningsvis resulterade behandling med syntetiskt ACTH (tetracosactid) i samtidig stegring av kortisol och PGF$_{2\alpha}$ metaboliten hos både gyltor och kastrerade galtar. Behandling med CRH resulterade i stegring av kortisol hos både gyltor och kastrerade galtar. Blodplasmakoncentrationerna av PGF$_{2\alpha}$ metaboliten var oförändrade hos både gyltor och kastrerade galtar efter CRH behandlingen.

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