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**Activation of peroxisome proliferator-activated receptors (PPARs)
by their ligands and protein kinase A activators**

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ABBREVIATIONS

AF, activation domain;

Bro, bromopalmitate;

8-Br-cAMP, 8-Bromoadenosine 3':5'-cyclic monophosphate;

CAT, chloramphenicol acetyl-transferase;

CT: cholera toxin;

DBD, DNA binding domain;

FSK, forskolin;

GST, glutathione S-transferase;

H89, N-[2-((p-Bromocinnamyl)amino)ethyl]-5-isoquinolinesulfonamide, 2HCl;

IBMX, isobutyl methylxanthine;

LBD, ligand binding domain;

PAGE, polyacrylamide gel electrophoresis;

PKA, protein kinase A;

PPAR, peroxisome proliferator-activated receptors;

RXR, retinoid X receptor;

TK, thymidine kinase;

ABSTRACT

The nuclear peroxisome proliferator-activated receptors (PPARs) α , β and γ activate the transcription of multiple genes involved in lipid metabolism. Several natural and synthetic ligands have been identified for each PPAR isotype but little is known about the phosphorylation state of these receptors. We show here that activators of protein kinase A (PKA) can enhance mouse PPAR activity in the absence and the presence of exogenous ligands in transient transfection experiments. The activation function 1 (AF-1) of PPARs was dispensable for transcriptional enhancement, whereas the activation function 2 (AF-2) was required for this effect. We also show that several domains of PPAR can be phosphorylated by PKA *in vitro*. Moreover, gel experiments suggest that PKA stabilizes binding of the liganded PPAR to DNA. PKA inhibitors decreased not only the kinase dependent induction of PPARs but also their ligand-dependent induction, suggesting that the ligands may also mobilize the PKA pathway to lead to maximal transcriptional induction by PPARs. Moreover, comparing PPAR α KO with PPAR α wild-type mice, we show that the expression of the ACO gene can be regulated by PKA-activated PPAR α in liver. These data demonstrate that the PKA pathway is an important modulator of PPAR activity and we propose a model associating this pathway in the control of fatty acid β -oxidation under conditions of fasting, stress and exercise.

INTRODUCTION

PPAR α , β and γ (NR1C1, NR1C2, NR1C3)(1), which are encoded by separate genes and exhibit distinct tissue-distribution, belong to the nuclear receptor superfamily (2). This family includes receptors for sexual and adrenal steroids, retinoids, thyroid hormone, vitamin D, ecdysone, and a number of receptors whose ligands are still unknown and therefore are called orphan receptors (3, 4). PPARs were first shown to be activated by substances that induce peroxisome proliferation (5, 6). PPARs share a common structure of 5 domains named A/B, C, D, E (the F domain is absent from PPARs compared with other members of the superfamily) (7). Key functions have been assigned to each of these domains (for review, (2, 8)). The N-terminal A/B domain contains the ligand-independent transcription activation function 1 (AF-1) (9). The C domain has a characteristic helix-loop-helix structure stabilized by two zinc atoms and is responsible for the binding to peroxisome proliferator response elements (PPREs) in the promoter region of target genes. The D domain is a hinge region which can modulate the DNA binding ability of the receptor and which is involved in corepressors binding (10). The E domain has a ligand binding function and exhibits a strong ligand-dependent activation function (AF-2). In a classical manner, the ligand binding domain facilitates the heterodimerization of PPAR with the retinoid X receptor (RXR). In its active state, this heterodimer is able to associate with coactivators (11-13) and when bound to a PPRE to modulate the expression of target genes.

It was also recently demonstrated that phosphorylation can modulate PPAR activity (14-21), but there are no data available concerning the phosphorylation of PPAR by the protein kinase A (PKA) pathway. PKA activity is enhanced in the liver under conditions of stress, fasting or exercise (22, 23). Under these conditions, PPAR target genes such as AcylCoA Oxidase, are up-regulated (24-26).

The aim of this work was to investigate whether PKA activators, mimicking stress, fasting or exercise, could directly modulate the activity of PPARs. We performed a detailed analysis of the effects of PKA on the activity of the mouse PPAR α , β and γ isotypes. We observed a PKA-dependent enhancement of the activity of all three isotypes in a ligand-independent and ligand-dependent manner on two distinct promoters. Several but not all PKA activators were able to produce this effect. Moreover, PKA inhibitors were able to repress the action of PKA activators. Very interestingly, PKA inhibitors were able to repress the effect of PPAR ligands by 50 to 75%, suggesting that these ligands are acting in part by recruiting the PKA pathway. By using GST fusions of PPARs, we demonstrate that PKA is able to phosphorylate different subdomains of PPARs *in vitro*, the DBD being the main phosphorylation target. Finally, using hepatocytes isolated from WT and PPAR α KO mice, we show that PKA activators can enhance the expression of certain PPAR α target genes and that PKA inhibitors repress WY 14, 643 dependent stimulation of these genes. These findings highlight the involvement of the PKA pathway in PPAR action and furthermore suggest that it is essential for the regulation of gene activation by PPAR ligands.

RESULTS

PKA activators enhance PPAR α activity on PPRE containing promoters. To evaluate the effect of PKA on PPAR activity, we transfected HEK-293 cells with a mPPAR α expression vector along with the PPRE-containing 2CYPA6-TK-CAT reporter construct or the TK-CAT construct as a control (Fig 1). Cells were treated or not with WY 14,643 (a PPAR α ligand) and/or cholera toxin (CT) as a PKA activator. In the absence or the presence of cotransfected PPAR α , expression of the TK-CAT construct was not significantly affected by WY 14, 643 nor by CT. In the absence of PPAR α , the 2CYPA6-TK-CAT construct, which contains functional PPREs (PPAR response element), was not stimulated by WY 14,643 nor by CT. When the PPAR α expression vector was cotransfected, PPAR α stimulated the activity of the 2CYPA6-TK-CAT construct in the presence of WY 14,643 (up to 7 fold compared to PPAR α in the absence of ligand). Addition of CT to the WY 14,643 treatment produced a further enhancement of PPAR α activity of about 2 fold compared to WY 14,643 treatment alone. In the absence of WY 14,643, CT by itself had also a stimulatory effect of about 2 fold. These results demonstrate that PKA activators can enhance PPAR α activity.

Dose dependent activation of mouse PPARs by their ligands. To further evaluate the effect of PKA on mouse PPAR activity, we transfected mPPAR α , β and γ expression vectors along with the 2CYPA6-TK-CAT reporter construct (Fig 2). Cells were treated or not with PPAR ligands (WY 14,643 for mPPAR α , bromopalmitate for mPPAR β and BRL 49,653 for mPPAR γ) and/or CT. PPAR α stimulated the activity of the reporter construct in a WY 14,643 dose dependent manner. Addition of CT to the WY 14,643 treatment produced a further enhancement of PPAR α activity even at saturating concentrations of WY 14,643. The same enhancement of activity by CT in the presence of ligand was obtained with PPAR β and with PPAR γ . Since these observations were made with mouse PPARs, we tested whether

amphibian PPARs would behave similarly. We observed that amphibian PPAR activity was also enhanced by PKA in a ligand-independent and ligand-dependent manner (data not shown), thus suggesting that the PPAR response to the PKA pathway is well conserved throughout evolution.

PPAR activity enhancement by PKA can be obtained with PKA activators and mediated by different PPREs. To ensure that the results obtained above were not dependent on the 2CYPA6-TK-CAT reporter construct only, we tested the effect of PKA on the PPAR-dependent stimulation of the ACO-TK-CAT reporter construct (Fig. 3). This construct, which contains the PPRE containing region of the ACO gene promoter was inducible by mouse PPAR α , β and γ in the presence of exogenous ligands, even though the effect was less pronounced than with the 2CYPA6-TK-CAT reporter gene. Addition of CT was able to increase PPAR activity 2 to 3 times. Interestingly, the ligand-independent effect of CT on PPAR activity was more pronounced on the ACO-TK-CAT reporter than on the 2CYPA6-TK-CAT reporter. These data suggest that the enhancement by PKA of PPAR activity can be mediated by distinct PPREs.

To determine whether other PKA activators (see Fig. 4A) had similar effects as CT (a Gs protein activator), we tested 8-Br-cAMP (a cAMP analog), forskolin (an adenylate cyclase activator) and IBMX (a phosphodiesterase inhibitor) in transient transfection with the mouse PPAR α expression vector and the 2CYPA6-TK-CAT reporter construct (Fig. 4B). CT and forskolin had roughly the same activation ability, while 8 Br-cAMP and IBMX had no significant effect even when higher concentrations of these activators were used (data not shown). These results suggest that PKA activators acting directly on cAMP production are the major stimulators of PPAR activity.

Ligand activation of PPARs involves the PKA pathway. Next, we tested whether ligand activation of PPARs was also involving the PKA pathway, even in the absence of added PKA

activators. To answer to this question, we used the H89 compound (a specific inhibitor of PKA at low concentrations) (27) in transient transfection experiments (Fig 5A). We observed that H89 was able to repress not only CT-dependent induction of PPAR α , β and γ activity but also the ligand-dependent activation as well as the activation due to CT and ligand used together. In addition, the basal receptor activity, in the absence of exogenous ligand, was also repressed. These data suggest that PKA affects both the basal and ligand-regulated activities of the PPARs. To ensure that the effects observed with CT or H89 were not due a change in PPAR expression, we checked by western blot the levels of expression PPAR α after treatment with WY 14,643, CT or H89 (Fig. 5B). PPAR α protein was only detected after transfection of PPAR α expression vector. Moreover, PPAR α protein levels were unchanged following treatment with WY 14,643, CT or H89, alone or in combination. These data demonstrate that the effects observed in transfection assays are not due to changes in PPAR expression levels. We then checked whether these drugs could affect PPAR DNA binding. The same extracts as in Fig. 5B were used to perform a gel shift assay using the PPRE from the ACO gene (Fig. 5C). We observed a strong binding with PPAR α WCE in the absence of added drug. Surprisingly, WY 14,643 treatment decreased PPAR α DNA binding ability. CT also diminished PPAR α DNA binding ability but to a lesser extent. More interestingly, the combination of WY 14,643 and CT enabled PPAR to bind to DNA more strongly than WY 14,643 or CT treatment alone. This in agreement with the situation observed for ER α for which PKA inhibits the dimerization of the receptor in the absence of ligand (28). Finally H89 had the same ability to reduce PPAR DNA binding as did WY 14,643. These data suggest that CT acts by stabilizing the decreased DNA binding ability of the liganded receptor in this *in vitro* assay.

RXR contributes to PPAR activation by PKA. As RXR is an obligate heterodimerization partner of the PPARs for DNA binding and transactivation, we determined whether RXR

could be involved in the PKA activation of PPAR α (Fig 6A). In the absence of transfected RXR or PPAR, the activity of the 2CYPA6-TK-CAT construct was very low and not modulated by the WY 14,643 and CT. In the absence of transfected RXR, transfected PPAR was active and modulated by PKA in HEK-293 cells, as these cells express low levels of endogenous RXR. In contrast, transfection of RXR alone in these cells had almost no effect on the expression of the 2CYP4A6-TK-CAT reporter gene even in the presence of 9-cis-retinoic acid (9cRA, is a ligand of RXR). However, we observed an enhancement of PPAR α activity in the presence of 9cRA and CT both in the absence and in the presence of WY 14,643. Indeed, enhancement of the PPAR α activity was even more potent with CT + 9cRA than with WY 14,643 + 9cRA. On the contrary, in the presence of WY 14,643, 9c-RA had only a minor effect. By overexpressing simultaneously RXR and PPAR α , in the absence of 9cRA, we observed an increase by about 30% of PPAR α activation by WY 14,643, and a 2 fold activity enhancement in the presence of CT and without ligand compared to PPAR α without cotransfected RXR. RXR affected only moderately PPAR α activation (about 20%) by WY 14,643 + CT. In the presence of RXR and 9cRA and in the absence of WY 14,643 and CT, we observed a 3 fold enhancement of PPAR α activity compared to cells without 9cRA. In the presence of WY 14,643 or WY 14,643 + CT, 9cRA only increased by 30% the activity seen in the absence of 9cRA. Finally, 9cRA was unable to affect CT induction of PPAR α in the absence of WY 14,643. These data suggest that RXR cooperates with PPAR α in the absence of exogenous ligand to increase both the basal and CT-induced activity of PPAR α on PPRES. We next checked whether RXR was itself the target of PKA when bound to its preferred binding site (DR1). To do so, we used the DR1-TK-CAT construct containing a strong RXR binding site (Fig. 6B). We observed a strong activation of the construct by RXR in the presence of RA. CT treatment increased both ligand-independent and ligand-dependent

activity of RXR. Thus, RXR by being itself the target of PKA can enhance PPAR activity on PPREs.

AF-1 is dispensable for PPAR stimulation by PKA activators. In order to explore which domain of PPAR is the target of the PKA pathway, we created 3 truncated versions of PPAR α (Fig 7A): one was deleted of the entire AB region (lacking AF-1, Δ AB mPPAR α), and the 2 others lacked the AF-2 domain as the entire LBD (Δ LBD mPPAR α) or the last 13 C-terminal residues (Δ AF2 mPPAR α) were deleted (Fig. 7A). Surprisingly, the Δ AB mPPAR α construct had roughly the same transactivation ability and exhibited the same enhancement of activity by CT as the wild-type mPPAR α . On the contrary, the Δ LBD and Δ AF2 constructs were totally insensitive to WY 14,643 or CT treatments. These data suggest that the AF-2 region is the major mediator of the effects of PKA activators and that the AF-1 region is not essential. As a control, we checked the DNA binding ability of the mutants by performing a gel shift assay using ACO PPRE (Fig. 7B). We *in vitro* translated the different mutants and checked first that they were produced in similar amounts (data not shown). We observed that Δ AB had the same ability to bind to DNA as wild-type PPAR α , whereas Δ AF2 and Δ LBD had a weak or not detectable binding ability respectively. This lack of binding of Δ LBD mutant is in agreement with previous reports (29). However, the weaker DNA binding ability of the Δ AF2 mutant cannot explain totally the lack of responsiveness to CT and we rather hypothesized that the AF2 function was involved in PKA stimulation. To confirm this hypothesis, we constructed GAL4 chimera comprising either the AB domain or the LBD of PPAR α fused to the GAL4 DNA binding domain (Fig. 7C). In transfection assays in HEK-293 cells, the GAL-AF1 chimera exhibited a strong ligand independent activity. This product was insensitive to WY 14,643 and CT alone or in combination, whereas the activity of the GAL-AF2 chimera was synergistically enhanced by WY 14,643 and CT treatments.

Interestingly, the GAL-AF2 chimera was not significantly affected by CT treatment alone, suggesting that other domains are involved in the effects of CT.

The main phosphorylation target of PKA activators is the DBD. To investigate which regions of PPARs were phosphorylated by PKA, we constructed a set of different GST fusion proteins corresponding either to the AB, DNA-binding (DBD) or ligand-binding (LBD) domains of the mouse PPAR α and β . The fusion proteins produced in bacteria were then purified on GSH columns and submitted to PKA treatment with recombinant enzyme in the presence of γ -[32 P]-ATP (Fig. 8A). We observed a strong phosphorylation of the DBD and a weaker labelling of the LBD from PPAR α and PPAR β , suggesting that the DBD is the main phosphorylation target. These data are in agreement with the effects of CT on DNA binding of PPAR α (see above). Interestingly, the AB domain of mPPAR α was also phosphorylated at a low level. We then mapped the putative sites of PKA phosphorylation (Fig. 8B). The most likely sites were mapped in the AB, C and E domain confirming that several domains of PPAR are potential targets of PKA phosphorylation.

PKA modulates PPAR α target genes. It was of interest to determine the effect of the PKA pathway on endogenous genes regulated by PPAR α *in vivo*, and particularly in liver which is one of the main sites of PPAR α action. To this end, isolated hepatocytes either from wild-type mice or from PPAR α KO mice (30) were cultured *in vitro* and treated with PKA activators or inhibitors in the absence or the presence of WY 14,643. The expression of the ACO (peroxisomal acylCoA oxidase) and the FABP (fatty acid binding protein) genes, two PPAR α target genes was analysed by northern blot (Fig. 9A). Under the conditions used, the ACO gene was not induced by WY 14,643 alone in the cultured wild-type hepatocytes, possibly because of an unidentified limiting factor was lost, due to *in vitro* partial dedifferentiation of the hepatocytes (31). However, CT was able to weakly stimulate ACO gene expression in the

absence of WY 14,643, but CT was a strong activator in the presence of WY 14,643. These data underline the ability of PKA activators to potentiate PPAR α ligand-dependent activation. In contrast, PPAR α KO hepatocytes were not sensitive to CT nor to WY 14,643 treatment, demonstrating that PPAR α was required for the WY 14,643 and CT synergism in the stimulation of the ACO gene. The expression of the FABP gene was strongly enhanced by WY 14,643 in wild-type hepatocytes but not in PPAR α KO hepatocytes, suggesting that distinct factors are required for ACO and FABP stimulation by PPAR α . In wild-type hepatocytes, addition of CT did not significantly affect FABP expression in the absence and in the presence of WY 14,643. On the contrary, H89 reduced by about 90% the WY 14,643 induction of FABP expression, which is in agreement with our transfection experiments. In PPAR α KO hepatocytes, the expression of the FABP gene was not significantly affected by WY 14,643, CT or H89, demonstrating that PPAR α was required for FABP induction. In conclusion, our data suggest that ligand and PKA activation of PPAR α converge in the stimulation of the PPAR target genes in hepatocytes.

DISCUSSION

Among the different stimuli known to phosphorylate and modulate nuclear receptor activity, the PKA pathway is certainly one of the best studied. However, no data are yet available concerning the potential effect of PKA on PPAR activity. This is in contrast with several studies focusing on PPAR phosphorylation by other stimuli. The scope of this work was to evaluate the role of PKA in modulating PPAR activity.

Early work from Shalev *et al.* (14) has shown that insulin treatment can phosphorylate PPAR α . Insulin can also increase PPAR α , and PPAR γ 2 activity in transient transfections. Insulin stimulation of PPAR γ involves MAP kinases (15, 19). On the contrary, other pathways stimulated by EGF (epidermal growth factor) and PDGF (platelet-derived growth factor) also involving MAP kinase have a negative effect on mPPAR γ 1 activity by phosphorylating serine 82 in the AB domain, which corresponds to serine 112 of mPPAR γ 2 (16, 17, 20). Further studies have demonstrated that this negative effect of MAP kinase was due to the inhibition of ligand binding consequent to an alteration of the three-dimensional structure of the receptor (32).

Here we demonstrate by transient transfection experiments that PKA activators can stimulate PPAR activity in an exogenous ligand-independent manner. Moreover, a combination of PKA activators and PPAR ligands leads to an increased activation of PPAR target genes.

Interestingly, this effect was obtained even at saturating concentrations of PPAR ligands. Moreover, we show that these effects are not due to change in PPAR expression. This stimulatory effect of PKA is in agreement with the results obtained with other nuclear receptors. Most studies report an activation of nuclear receptors such as estrogen receptor (ER α), glucocorticoid receptor (GR), mineralocorticoid receptor (MR), progesterone receptor (PR), androgen receptor (AR) or steroidogenic factor-1 (SF-1) by PKA (28, 33-38), whereas HNF4 has been shown to be down-regulated by PKA (39). Moreover, PKA activated PPARs

were able to stimulate two types of PPREs, even though the amplitude of response was different. Interestingly, PKA activators were nearly as effective as PPAR ligands to activate the ACO PPRE, whereas they could only activate the CYPA6 PPRE to levels corresponding to 25-30% of those obtained with PPAR ligands. Such differences were also found with ER α according to the cell-type and the promoter used (40-42).

Using different PKA pathway activators, we observed that the most potent ones were those affecting adenylate cyclase activity and not those leading to a direct increase of cAMP such as supplementation with cAMP analogs or inhibition of phosphodiesterase activity. To address the question whether ligands by themselves would activate the PKA pathway, we treated cells with H89, an inhibitor of PKA (27). Interestingly, we observed that H89 could reduce not only the CT induction of PPARs but also their induction by ligands such as WY 14,643, bromopalmitate and BRL 49,653, suggesting that these ligands act in part through the PKA pathway. This result was not due to a change in PPAR levels as shown by western blot. An attractive hypothesis would be that the ligands can also act indirectly by modulating intracellular cAMP levels. Such observations have indeed been reported for estrogens, which are able to increase intracellular cAMP levels, which in turn activate PKA and increases ER α activity (34).

As PPARs act essentially as heterodimers, it was of particular interest to determine whether its heterodimer partner (RXR) could be involved in PPAR activation by PKA. Surprisingly, in the absence of 9-cis-retinoic acid (RXR ligand), cotransfection of RXR moderately affected PPAR α activity in the presence of WY 14,643, but increased PPAR α activity by about 2 fold without any ligand, leading to an activation by CT equal to the one with WY 14,643. In the presence of RA, RXR conferred an even stronger ligand-independent activity to PPAR α . This could be explained by the fact that RXR is itself the target of PKA as shown clearly on DR1-TK-CAT construct. RAR and RXR activation by PKA have been previously reported (43-45).

Thus, in the absence of WY 14,643 but in the presence of RA, it seems that the PKA action on RXR heterodimerized with PPAR leads to a major enhancement of the activity of PPAR/RXR heterodimers. However, we cannot exclude the possibility that other factors such as coactivators could also be the target of PKA.

Using truncated PPAR α constructs, we determined that the AF-2 domain was most important for transactivation by PKA. Deletion of the AB domain from PPAR α only slightly affected its basal and ligand induced activity, which is in agreement with previous data (29). Interestingly, CT was still able to potentiate WY 14,643 induction of the truncated PPAR α , demonstrating that the AF-1 function was not involved in the potentiation by PKA. On the contrary, AF-2 deletion completely abolished WY 14,643 as well as CT activation of PPAR α . Our data obtained with the GAL4 chimeras clearly confirm that AF2 but not AF1 is the target of PKA action. The demonstration that AF2 is the target of PKA is in agreement with the results found for PKA activation of ER α (42, 46) or SF-1 (38). However, for AR (37) and MR (47), the N-terminal portion seemed to be involved in PKA effects. Interestingly, CT had no effect on the ligand-independent activity of the GAL-AF2 chimera suggesting that other domains of the receptors are necessary. To better characterize the domains involved, we analysed the phosphorylation of different domains of PPAR by using GST-PPAR fusions. We observed a very strong phosphorylation of the DBD, a weaker one for the AB domain and a faint one for the LBD, again suggesting that several domains are involved in the activation of PPARs by the PKA pathway. This result is confirmed by the mapping of the most conserved putative PKA sites which are present in the A/B, DBD and LBD domains. As the main phosphorylation site is present in the DBD, we analysed whether the drugs used could modulate the DNA binding ability of PPAR α *in vitro*. To our surprise, WY 14,643 and CT strongly inhibited PPAR α DNA binding. However, cotreatment with WY 14,643 and CT led only to a limited decreased binding. We thus propose that CT acts in part by preventing the

decreased binding of PPAR liganded with WY 14,643. This stabilization would in turn increase PPAR activity. This in agreement with a previous report (28), which shows that ER α DNA binding is inhibited by PKA only in the absence of estradiol. This report also shows that the target of PKA is in the DBD of ER α . Rangarajan et al. (33) have also demonstrated that PKA enhancement of liganded GR required specific residues of the DBD. In this case, however, they observed an enhancement of GR binding in the presence of PKA. This suggested that depending on the receptors, the mechanisms of enhancement of the activity by PKA requires different functions of the receptor.

A crucial question was whether PKA does modulate the expression of PPAR target genes? To answer this question, we focused on the role of PPAR α in the liver and took advantage of PPAR α KO mice. We analysed the expression of two target genes, ACO and FABP (48, 49). In wild-type mice hepatocytes cultured *in vitro*, we demonstrated that cotreatment with WY 14,643 and CT led to a synergistic activation of the ACO gene expression. On the other hand, the FABP gene was only weakly affected by addition of exogenous PKA activators, but addition of PKA inhibitors strongly diminished its induction by WY 14,643, confirming the results obtained in transient transfection experiments. In PPAR α KO mice, ACO was not subjected to stimulation by WY 14,643 or CT alone or in combination, confirming that PPAR α was essential to PKA activation of the ACO gene. In these mice, FABP induction by WY 14,643 was completely abolished. FABP which is involved in fatty acid (FA) binding in hepatocytes and ACO in β -oxidation of FA could therefore be integrated in the following model involving the PKA pathway (Fig 9B): Under conditions of stress, fasting or exercise, the limiting factor for brain and muscles rely on increased energy fuel availability, essentially glucose and ketone bodies. One way for the organism to meet these needs is to stimulate gluconeogenesis and ketogenesis. The adipose tissue hydrolyses triglycerides (TG) to liberate free non esterified fatty acids, which are released into the blood circulation and are then

rapidly taken up by the liver to be transformed into ketone bodies. PPAR α and PPAR γ are directly involved in the regulation of several key enzymes of these pathways. Stress, fasting or exercise are also associated with an increased glucagon production (one of the key factors increasing cAMP levels in cells and thus activating PKA) by the adrenal gland (50). PKA increases PPAR α activity in liver, which in turn stimulates the β -oxidation and in particular the conversion of FA into acetyl-CoA used in the production of ketone bodies (51). Results from our laboratory (52) have also demonstrated that fasting did not affect FABP expression in wild-type mice. On the contrary, ACO gene expression has been shown to be up-regulated by fasting in wild-type mice but not in PPAR α KO mice (24, 26), confirming the scheme of regulation we propose. In addition, PPAR α KO mice exhibited an increased accumulation of FA in liver, due to impaired β -oxidation (53).

In conclusion, our results suggest that under conditions of stress, fasting and exercise, PPAR α activity is increased by the PKA pathway and leads to an enhancement of β -oxidation, production of glucose and ketone bodies, which serve as fuel for muscles and brain.

MATERIALS AND METHODS

Chemicals.

Bromopalmitate, CT (cholera toxin), Forskolin, 8-Br-cAMP (8-Bromoadenosine 3':5'-cyclic monophosphate), IBMX (isobutyl methylxanthine) were from Sigma (St Louis, MO). WY 14,643 was from Chemsyn Science Laboratories (Kansas, USA). BRL 49,653 was a kind gift from Parke Davis. H89 (N-[2-((p-Bromocinnamyl)amino)ethyl]-5-isoquinolinesulfonamide, 2HCl) was from Calbiochem (La Jolla, CA). PKA catalytic subunit was from Promega (Madison, WI).

Oligonucleotide Sequences.

ACO1: GCCACCGCCTATGCCTTCCACTTT

ACO2: CGGCTTGCACGGCTCTGTCTTGA

LPL1: CCTGCGGGCCCTATGTTTG

LPL2: CTCGCCGATGTCTTTGTCCAGT

mFABP1: CAATTGCAGAGCCAGGAGAACTTT

mFABP2: CAATGTCGCCCAATGTCA

Plasmids.

The reporter plasmid 2CYP-TK-CAT contains two copies of the CYP4A6 PPRE cloned in opposite orientation upstream of the minimal herpes simplex virus thymidine kinase promoter in the pBLCAT8+ plasmid (54). ACO-TK-CAT plasmid corresponds to the Acyl-CoA oxidase promoter PPRE cloned in pBLCAT8+ as described by Dreyer et al. (6). DR1-TK-CAT corresponds to the perfect DR1 sequence (AGCTTCATTCTAGGTCAAAGGTCATCCCCT) cloned in the pBLCAT8+ plasmid. pG5CAT reporter plasmid (Clontech) corresponds to 5 Gal4 binding sites upstream of the E1b minimal promoter. Mouse and Xenopus PPAR α , β and γ cDNAs were cloned into the BamHI site of the pSG5 mammalian expression vector. For PKA *in vitro* assays, portions of mouse

PPAR cDNAs were amplified by PCR and then subcloned into the BamHI site of the prokaryotic expression vector pGEX1. mPPAR α AB (aa 1-101) was cloned into the SmaI/BamHI sites of pGEX1. mPPAR α DBD domain (aa 98-203), mPPAR β DBD domain (aa 68-129), mPPAR α LBD (aa 202-468) and mPPAR β LBD (aa 136-396) were cloned into the BamHI site of pGEX1. pSG5-mPPAR α Δ AB was obtained by removing aa 1 to 101 from WT mPPAR α by PCR and pSG5-mPPAR α Δ LBD by removing sequences downstream from residue 247. pSG5-mPPAR α Δ AF2 was obtained by removing the last 13 residues from WT mPPAR α . GAL-AF1 and GAL-AF2 expression plasmids correspond to AB domain (aa 1 to 100) LBD domain (165-468) respectively cloned in Gal4 DBD pM vector (Clontech).

***In vitro* Translation**

In vitro translation was performed using the TNT Promega kit. Briefly, 1 μ g of expression vector was mixed to 25 μ l of TNT rabbit reticulocyte lysate, 2 μ l of TNT buffer, 1 μ l of mix containing all amino acids, 1 μ l of RNasin (50 U/ μ l), 1 μ l of T7 RNA polymerase (20 U/ μ l). A control reaction was performed under the same conditions but [35 S]-methionine (15 μ Ci/ μ l) was used to label the protein produced. The final reaction volume was 50 μ l. The reaction was performed for 1.5 h at 30 $^{\circ}$ C. The translation efficiency was checked by loading 1 μ l of labelled lysate on an SDS-PAGE gel.

Gel Mobility Shift Assays.

Gel mobility shift assays were carried out as previously described (55). Briefly, [32 P]-labeled ACoA (GATCCCGAACGTGACCTTTGTCCTGGTCCCGATC) double strand oligonucleotide, (56) was combined with *in vitro* translated PPAR or HEK-293 WCE and when indicated mouse RXR β_2 Sf9 cellular extract. Protein-DNA complexes were separated from the free probe by non-denaturing gel electrophoresis with 4% polyacrylamide (29/1) gels in 0.5 X TBE.

Cell Culture and Transient Transfection.

HEK-293 cells (human embryonic kidney cells) were cultured in 10% FCS (foetal calf serum) DMEM-F12 with 5% CO₂. Cells were plated in 24-well plates in 10% CDFCS, phenol-free DMEM 24 h before transfection. Transfections were performed by lipofection (lipofectamine, Life Technologies, Rockville, MA) using 200 ng of CAT reporter construct, 400 ng of the internal reference β -galactosidase reporter plasmid (pCH110) and 100 ng of pSG5-PPAR or pSG5-hRXR α expression vectors per well. After lipofection, the cells were grown in 10% CDFCS, DMEM in the presence of different ligands for 36 h. Transactivation ability was determined by CAT activity on the whole cell extract as previously described (55).

Hepatocyte Isolation and Culture

Hepatocytes were isolated from liver of adult male wild-type (SV129) or PPAR α KO mice using a two-step *in situ* portal vein collagenase A (Boehringer Mannheim) perfusion method (57). Freshly isolated hepatocytes were filtered through nylon membrane to remove tissue debris and cell clumps. The cell suspension was washed in Leibovitz's L-15 medium and resuspended twice after centrifugation. The isolated hepatocytes were suspended in William's medium E supplemented with 10% FCS, 100 μ g/ml streptomycin and 100 μ g/ml penicillin and seeded in dishes at the density of $5 \cdot 10^5$ cells/ml medium. The medium was renewed 4 hr later to remove dead cells. Cells were then treated with or without WY 14,643 (10 μ M) and cholera toxin (1 μ g/ml), in presence or not of H89 (10 μ M). Cultures were maintained at 37°C in a humidified air / CO₂ incubator (5% CO₂, 95% air) for 24 h.

Whole cell extract preparation and western blot.

HEK-293 cells were harvested, washed in PBS, and resuspended in TEG (10 mM Tris-HCl, pH 7.4, 1.5 mM EDTA, and 10% glycerol)/ 0.4 M KCl containing 5 μ g/ml aprotinin, leupeptin and pepstatin A and 0.1 mM phenylmethylsulfonyl fluoride. Then, cells were sonicated and the cellular debris were pelleted by centrifugation at 14000 rpm for 20 minutes

in microfuge tubes. Thirty μg of whole cell extract proteins were subjected to SDS-PAGE followed by electrotransfer onto a nitrocellulose membrane. The blot was probed with anti PPAR α AB antibody (1/1000) (polyclonal rabbit antibody produced in our laboratory and directed against mouse PPAR α AB region) and then incubated with rabbit anti-rabbit IgG horseradish peroxidase conjugated antibody (1 $\mu\text{g}/\text{ml}$). ECL kit from Amersham (Arlington, IL, USA) was used for protein detection.

RNA Isolation and Northern Blot.

Total RNA was isolated from isolated hepatocytes using the Trizol reagent from Life Technologies (Rockville, MA) as described by the manufacturer. ACO and FABP probes were amplified by RT-PCR. The amplifying primers were: ACO1 and ACO2 primers for mouse peroxisomal acylCoA oxidase (ACO) probe (1036-1690), mFABP1 and mFABP2 for mouse fatty acid binding protein (FABP) probe (61-394) (see above).

For northern blot analysis, 20 μg of total RNA was electrophorized in a 2.2 M formaldehyde-1% agarose gel in MOPS buffer and then hybridized with the different probes as previously described (58).

Production of GST Fusion Proteins

Production of GST fusion proteins was performed as previously described (13). Protein concentration was estimated by the Bradford method. The levels of expressed fusion proteins were determined by an *in vitro* binding assay followed by a SDS-PAGE gel and a Coomassie Blue staining.

***In vitro* PKA Assays with Glutathione Sepharose**

Glutathione Sepharose (Pharmacia Biotech, Uppsala, Sweden) was equilibrated with NET binding buffer (150 mM NaCl, 50 mM Tris-HCl (pH 7.4), 5 mM EDTA). Crude bacterial extract containing GST fusion proteins was incubated at 4 $^{\circ}\text{C}$ with 25 μl of beads for 2.5h . After 2 washes with NETN (NET + 0.5% NP40), the beads were washed 2 times with PKA

buffer (50 mM Tris-HCl (pH 7.5), 10 mM NaCl, 1 mM DTT, 10 mM MgCl₂, 10% glycerol). The beads were then incubated in a mix containing 50 µl of PKA buffer, 45U of PKA catalytic subunit, 0.5 µl γ-[³²P]-ATP and 0.5 µl ATP 2.5 mM for 45 min at 30 °C. After 2 washes with NETN, beads were boiled in SDS loading buffer, and a quarter of the proteins were run on SDS-PAGE. The gel was then stained with coomassie blue. After extensive washes with a solution containing 20% methanol and 10% acetic acid, the gel was submitted to autoradiography.

FIGURE LEGENDS

Fig 1: CT increases mPPAR α activity on a PPRE containing promoter

Mouse PPAR α was transfected along with TK-CAT or 2CYPA6-TK-CAT constructs in HEK-293 cells. Transfections were with 200 ng of TK-CAT or 2CYPA6-TK-CAT reporter construct and without or with 100 ng of pSG5-mPPAR α expression vector per well. Cells were grown for 36 h in the presence of 1 μ M of WY 14,643 (WY), with or without 1 μ g/ml CT. Results are shown as the mean \pm SD (n = 6) of CAT activity after normalization for β -galactosidase activity.

Fig 2: CT increases mPPAR activity at various concentrations of PPAR ligands

Mouse PPAR α , β and γ were tested for their ability to respond to CT in HEK-293 cells. Transfections were with 200 ng of 2CYPA6-TK-CAT reporter construct and 100 ng of pSG5-mPPAR α , β and γ expression vectors per well. Cells were grown for 36 h in the presence of various concentrations of WY 14,643 (WY), Bromopalmitate (Bro) and BRL 49,653 (BRL), with or without 1 μ g/ml CT. Results are shown as the mean \pm SD (n = 6) of CAT activity after normalization for β -galactosidase activity.

Fig 3: PKA enhancement of PPAR activity via the ACO PPRE

Mouse PPAR α , β and γ cDNAs were cotransfected in HEK-293 cells with ACO-TK-CAT reporter construct instead of the 2CYPA6-TK-CAT reporter construct in the same conditions as in fig 1. Cells were grown for 36 h in the presence of 1 μ M WY 14,643 (WY), 50 μ M Bromopalmitate (Bro) and 5 μ M BRL 49,653 (BRL), with 1 μ g/ml CT when indicated. Results are shown as the mean \pm SD (n = 6) of CAT activity after normalization for β -galactosidase activity.

Fig 4: various PKA activators stimulate PPAR activity

A. Schematic representation of the PKA pathway. cAMP is produced from ATP by a membrane-bound adenylate cyclase (Ac). Transmembrane receptors (R) for numerous hormones, neurotransmitters and other stimuli (H) are coupled to adenylate cyclase via heterotrimeric G-proteins (α , β and γ subunits). This interaction promotes the exchange of GDP, bound to the α -subunit, for GTP and the subsequent dissociation of the α subunit from the $\beta\gamma$ heterodimer. The $G\alpha$ -GTP then binds to adenylate cyclase and modulates its activity. PDE: phosphodiesterase; FSK: forskolin; CT: cholera toxin; . **B.** 100 ng of pSG5-mPPAR α was cotransfected in HEK-293 cells with 2CYPA6-TK-CAT reporter construct under the same conditions as in Fig 1. Cells were grown for 36 h in the presence of 1 μ M of WY 14,643 (WY), with 1 μ g/ml Cholera toxin (CT), 50 mM 8-BrcAMP, 200 μ M forskolin (FSK) or 10 μ M IBMX when indicated. Results are shown as the mean \pm SD (n = 3) of CAT activity after normalization for β -galactosidase activity.

Fig 5: PKA inhibitors can repress PPAR activity

A. Mouse PPAR α , β and γ were tested for their ability to respond to CT in HEK-293 cells using 200 ng of 2CYPA6-TK-CAT reporter construct and 100 ng of pSG5-PPAR expression vectors. After lipofection, 10 μ M of H89 was added to the medium 1 h before ligands. Cells were grown for 36 h in the presence of 1 μ M WY 14,643 (WY), 50 μ M Bromopalmitate (Bro) and 5 μ M BRL 49,653 (BRL), with 1 μ g/ml CT when indicated. Results are shown as the mean \pm SD (n = 6) of CAT activity after normalization for β -galactosidase activity. **B.** 30 μ g of whole cell extracts from transfected cells were loaded on SDS-PAGE and probed by western blot using mPPAR α antibody. The first lane corresponds to HEK-293 cells transfected with the empty pSG5 vector and the remaining lanes correspond to 293 cells

transfected with pSG5-mPPAR α vector and treated or not (-) with WY, CT or H89 under the same conditions as in Fig. 5A. **C.** 5 μ g of the same WCE were used in gel shift assays using the ACoA probe.

Fig 6: RXR modulates PPAR activity in the presence of PKA activators

A. 100 ng of pSG5, pSG5-mPPAR α and pSG5-mRXR β 2 expression vectors per well in combination or alone were cotransfected in HEK-293 cells with 200 ng of 2CYPA6-TK-CAT reporter construct. After lipofection, cells were grown for 36 h with or without 1 μ M of WY 14,643 (WY), and 1 μ M 9-cis retinoic acid (RA), with 1 μ g/ml CT when indicated. Results are shown as the mean \pm SD (n = 6) of CAT activity after normalization for β -galactosidase activity. **B.** 100 ng of pSG5 or pSG5-mRXR β 2 expression vectors were cotransfected in HEK-293 cells with 200 ng of DR1-TK-CAT reporter construct per well. After lipofection, cells were grown for 36 h with or without 1 μ M 9-cis retinoic acid (RA) and 1 μ g/ml CT when indicated. Results are shown as the mean \pm SD (n = 6) of CAT activity after normalization for β -galactosidase activity.

Fig 7: The AB domain is dispensable for PPAR response to PKA

A. 100 ng of pSG5-mPPAR α WT, or pSG5-mPPAR α Δ AB, pSG5-mPPAR α Δ LBD, or pSG5-mPPAR α Δ AF2 constructs were transfected in HEK-293 cells with 200 ng of 2CYPA6-TK-CAT reporter construct per well. After lipofection, cells were grown for 36 h with or without 1 μ M of WY 14,643 (WY) with or without 1 μ g/ml CT. Results are shown as the mean \pm SD (n = 6) of CAT activity after normalization for β -galactosidase activity. **B.** Equivalent amounts of *in vitro* translated mPPAR α WT, mPPAR α Δ AB, mPPAR α Δ LBD, or mPPAR α Δ AF2 receptors were used in gel shift assays in combination with the ACoA probe.

Lane 1 corresponds the probe alone and lane C to mock lysate. In addition, mouse RXR β_2 Sf9 cellular extract was eventually added (RXR). The arrow indicated the position of RXR retarded complex and the star the position of a non specific complexes. C. 100 ng of GAL, GAL-AF1, or GAL-AF2 constructs were transfected in HEK-293 cells with 200 ng of pG5CAT reporter construct. After lipofection, cells were grown for 36 h with or without 1 μ M of WY 14,643 (WY) with or without 1 μ g/ml CT. Results are shown as the mean \pm SD (n = 6) of CAT activity after normalization for β -galactosidase activity.

Fig 8: PPAR phosphorylation in response to PKA occurs mainly in the DBD

A. Regions encoding AB (AB α), DBD (DBD α) and LBD (LBD α) domains of mPPAR α as well as DBD (DBD β) and LBD (LBD β) of mPPAR β were cloned in pGEX1 prokaryotic expression vector as GST fusions. *In vitro* PKA assay was performed as described in the Materials and Methods. The upper panel corresponds to the gel autoradiogram and the lower panel to the protein expression levels after coomassie blue staining of the gel. The arrow labelled NS corresponds to a non specific band. **B.** Mapping of the main putative PKA sites of phosphorylation on mPPAR α .

Fig 9: PKA modulates some PPAR target genes

A. Hepatocytes were isolated from wild-type and PPAR α KO mice and cultured in Williams medium supplemented with 10% CDFCS. Cells were then treated for 24 h with or without 10 μ M WY 14,643 (WY) and 1 μ g/ml cholera toxin (CT). When H89 was used (H: 10 μ M), it was added to the medium 1 h before treatment with WY or CT. After RNA extraction, 10 μ g of total RNA were used for northern blotting and then hybridized sequentially with ACO, FABP and 28S probes. The figure shows a representative experiment. **B.** General model of

cross-talk between fatty acids and PKA signalling involving PPARs. TG: triglycerides, FA: fatty acids, Glucocor: glucocorticoids, β OX: β oxidation, LPL: lipoprotein lipase.

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REFERENCES

1. [letter] 1999 A unified nomenclature system for the nuclear receptor superfamily. *Cell* 97:161-3
2. Desvergne B, Wahli W 1999 Peroxisome proliferator-activated receptors: nuclear control of metabolism. *Endocr Rev* 20:649-88
3. Beato M, Herrlich P, Schutz G 1995 Steroid hormone receptors: many actors in search of a plot. *Cell* 83:851-7
4. Mangelsdorf DJ, Thummel C, Beato M, Herrlich P, Schutz G, Umesono K, Blumberg B, Kastner P, Mark M, Chambon P, et al. 1995 The nuclear receptor superfamily: the second decade. *Cell* 83:835-9
5. Issemann I, Green S 1990 Activation of a member of the steroid hormone receptor superfamily by peroxisome proliferators. *Nature* 347:645-50
6. Dreyer C, Krey G, Keller H, Givel F, Helftenbein G, Wahli W 1992 Control of the peroxisomal beta-oxidation pathway by a novel family of nuclear hormone receptors. *Cell* 68:879-87
7. Krust A, Green S, Argos P, Kumar V, Walter P, Bornert JM, Chambon P 1986 The chicken oestrogen receptor sequence: homology with v-erbA and the human oestrogen and glucocorticoid receptors. *EMBO J.* 5:891-897
8. Schoonjans K, Martin G, Staels B, Auwerx J 1997 Peroxisome proliferator-activated receptors, orphans with ligands and functions. *Curr Opin Lipidol* 8:159-66
9. Hi R, Osada S, Yumoto N, Osumi T 1999 Characterization of the amino-terminal activation domain of peroxisome proliferator-activated receptor alpha. Importance of alpha-helical structure in the transactivating function. *J Biol Chem* 274:35152-8
10. Kumar R, Thompson EB 1999 The structure of the nuclear hormone receptors. *Steroids* 64:310-9

11. McKenna NJ, Lanz RB, O'Malley BW 1999 Nuclear receptor coregulators: cellular and molecular biology. *Endocr Rev* 20:321-44
12. Krey G, Braissant O, L'Horsset F, Kalkhoven E, Perroud M, Parker MG, Wahli W 1997 Fatty acids, eicosanoids, and hypolipidemic agents identified as ligands of peroxisome proliferator-activated receptors by coactivator-dependent receptor ligand assay. *Mol Endocrinol* 11:779-91
13. Lazennec G, Ediger TR, Petz LN, Nardulli AM, Katzenellenbogen BS 1997 Mechanistic aspects of estrogen receptor activation probed with constitutively active estrogen receptors - correlations with dna and coregulator interactions and receptor conformational changes. *Mol Endocrinol* 11:1375-1386
14. Shalev A, Siegrist KC, Yen PM, Wahli W, Burger AG, Chin WW, Meier CA 1996 The peroxisome proliferator-activated receptor alpha is a phosphoprotein: regulation by insulin. *Endocrinology* 137:4499-502
15. Zhang B, Berger J, Zhou GC, Elbrecht A, Biswas S, White-Carrington S, Szalkowski D, Moller DE 1996 Insulin- and mitogen-activated protein kinase-mediated phosphorylation and activation of peroxisome proliferator-activated receptor gamma. *J Biol Chem* 271:31771-31774
16. Adams M, Reginato MJ, Shao D, Lazar MA, Chatterjee VK 1997 Transcriptional activation by peroxisome proliferator-activated receptor gamma is inhibited by phosphorylation at a consensus mitogen- activated protein kinase site. *J Biol Chem* 272:5128-32
17. Camp HS, Tafuri SR 1997 Regulation of peroxisome proliferator-activated receptor gamma activity by mitogen-activated protein kinase. *J Biol Chem* 272:10811-10816

18. Passilly P, Schohn H, Jannin B, Malki MC, Boscoboinik D, Dauca M, Latruffe N 1999 Phosphorylation of peroxisome proliferator-activated receptor alpha in rat Fao cells and stimulation by ciprofibrate. *Biochem Pharmacol* 58:1001-8
19. Juge-Aubry CE, Hammar E, Siegrist KC, Pernin A, Takeshita A, Chin WW, Burger AG, Meier CA 1999 Regulation of the Transcriptional Activity of the Peroxisome Proliferator-activated Receptor alpha by Phosphorylation of a Ligand- independent trans-Activating Domain. *J Biol Chem* 274:10505-10510
20. Hu ED, Kim JB, Sarraf P, Spiegelman BM 1996 Inhibition of adipogenesis through map kinase-mediated phosphorylation of ppar-gamma. *Science* 274:2100-2103
21. Werman A, Hollenberg A, Solanes G, Bjorbaek C, Vidalpuig AJ, Flier JS 1997 Ligand-independent activation domain in the n terminus of peroxisome proliferator-activated receptor gamma (ppar-gamma) - differential activity of ppar-gamma-1 and -2 isoforms and influence of insulin. *J Biol Chem* 272:20230-20235
22. Winder WW, Duan C 1992 Control of fructose 2,6-diphosphate in muscle of exercising fasted rats. *Am J Physiol* 262:E919-24
23. Begum N, Graham AL, Sussman KE, Draznin B 1992 Role of cAMP in mediating effects of fasting on dephosphorylation of insulin receptor. *Am J Physiol* 262:E142-9
24. Kroetz DL, Yook P, Costet P, Bianchi P, Pineau T 1998 Peroxisome Proliferator-activated Receptor alpha Controls the Hepatic CYP4A Induction Adaptive Response to Starvation and Diabetes. *J Biol Chem* 273:31581-31589
25. Lemberger T, Saladin R, Vazquez M, Assimacopoulos F, Staels B, Desvergne B, Wahli W, Auwerx J 1996 Expression of the peroxisome proliferator-activated receptor alpha gene is stimulated by stress and follows a diurnal rhythm. *J Biol Chem* 271:1764-1769

26. Leone TC, Weinheimer CJ, Kelly DP 1999 A critical role for the peroxisome proliferator-activated receptor alpha (PPARalpha) in the cellular fasting response: The PPARalpha-null mouse as a model of fatty acid oxidation disorders. *PNAS* 96:7473-7478
27. Chijiwa T, Mishima A, Hagiwara M, Sano M, Hayashi K, Inoue T, Naito K, Toshioka T, Hidaka H 1990 Inhibition of forskolin-induced neurite outgrowth and protein phosphorylation by a newly synthesized selective inhibitor of cyclic AMP-dependent protein kinase, N-[2-(p-bromocinnamylamino)ethyl]-5-isoquinolinesulfonamide (H-89), of PC12D pheochromocytoma cells. *J Biol Chem* 265:5267-72
28. Chen DS, Pace PE, Coombes RC, Ali S 1999 Phosphorylation of human estrogen receptor alpha by protein kinase A regulates dimerization. *Mol Cell Biol* 19:1002-1015
29. Hsu MH, Palmer CNA, Song W, Griffin KJ, Johnson EF 1998 A carboxyl-terminal extension of the zinc finger domain contributes to the specificity and polarity of peroxisome proliferator-activated receptor DNA binding. *J Biol Chem* 273:27988-27997
30. Lee SS, Pineau T, Drago J, Lee EJ, Owens JW, Kroetz DL, Fernandez-Salguero PM, Westphal H, Gonzalez FJ 1995 Targeted disruption of the alpha isoform of the peroxisome proliferator- activated receptor gene in mice results in abolishment of the pleiotropic effects of peroxisome proliferators. *Mol Cell Biol* 15:3012-22
31. Menjo M, Yamaguchi S, Murata Y, Hayashi Y, Nagaya T, Ohmori S, Refetoff S, Seo H 1999 Responsiveness to thyroid hormone is enhanced in rat hepatocytes cultured as spheroids compared with that in monolayers: altered responsiveness to thyroid

hormone possibly involves complex formed on thyroid hormone response elements.

Thyroid 9:959-67

32. Shao DL, Rangwala SM, Bailey ST, Krakow SL, Reginato MJ, Lazar MA 1998 Interdomain communication regulating ligand binding by ppar-gamma. Nature 396:377-380
33. Rangarajan PN, Umesono K, Evans RM 1992 Modulation of glucocorticoid receptor function by protein kinase A. Mol Endocrinol 6:1451-7
34. Aronica SM, Kraus WL, Katzenellenbogen BS 1994 Estrogen action via the cAMP signaling pathway: stimulation of adenylate cyclase and cAMP-regulated gene transcription. PNAS 91:8517-8521
35. Aronica SM, Katzenellenbogen BS 1991 Progesterone receptor regulation in uterine cells: stimulation by estrogen, cyclic adenosine 3',5'-monophosphate, and insulin-like growth factor I and suppression by antiestrogens and protein kinase inhibitors. Endocrinology 128:2045-52
36. Kahmann S, Vassen L, Klein-hitpass L 1998 Synergistic enhancement of prb-mediated ru486 and r5020 agonist activities through cyclic adenosine 3',5'-monophosphate represents a delayed primary response. Mol Endocrinol 12:278-289
37. Sadar MD 1999 Androgen-independent induction of prostate-specific antigen gene expression via cross-talk between the androgen receptor and protein kinase A signal transduction pathways. J Biol Chem 274:7777-83
38. Jacob AL, Lund J 1998 Mutations in the activation function-2 core domain of steroidogenic factor-1 dominantly suppresses pka-dependent transactivation of the bovine cyp17 gene. J Biol Chem 273:13391-13394

39. Viollet B, Kahn A, Raymondjean M 1997 Protein kinase A-dependent phosphorylation modulates DNA-binding activity of hepatocyte nuclear factor 4. *Mol Cell Biol* 17:4208-19
40. Ince BA, Montano MM, Katzenellenbogen BS 1994 Activation of transcriptionally inactive human estrogen receptors by cyclic adenosine 3',5'-monophosphate and ligands including antiestrogens. *Mol Endocrinol* 8:1397-406
41. Fujimoto N, Katzenellenbogen BS 1994 Alteration in the agonist/antagonist balance of antiestrogens by activation of protein kinase A signaling pathways in breast cancer cells: antiestrogen selectivity and promoter dependence. *Mol Endocrinol* 8:296-304
42. Cho H, Katzenellenbogen BS 1993 Synergistic activation of estrogen receptor-mediated transcription by estradiol and protein kinase activators. *Mol Endocrinol* 7:441-52
43. Dowhan DH, Muscat GE 1996 Characterization of the AB (AF-1) region in the muscle-specific retinoid X receptor-gamma: evidence that the AF-1 region functions in a cell-specific manner. *Nucleic Acids Res* 24:264-71
44. Rochette-Egly C, Oulad-Abdelghani M, Staub A, Pfister V, Scheuer I, Chambon P, Gaub MP 1995 Phosphorylation of the retinoic acid receptor-alpha by protein kinase A. *Mol Endocrinol* 9:860-71
45. Taneja R, Rochette-Egly C, Plassat JL, Penna L, Gaub MP, Chambon P 1997 Phosphorylation of activation functions AF-1 and AF-2 of RAR alpha and RAR gamma is indispensable for differentiation of F9 cells upon retinoic acid and cAMP treatment. *Embo J* 16:6452-65
46. El Tanani M, Green CD 1997 Two separate mechanisms for ligand-independent activation of the estrogen receptor. *Mol Endocrinol* 11:928-37

47. Massaad C, Houard N, Lombes M, Barouki R 1999 Modulation of human mineralocorticoid receptor function by protein kinase A. *Mol Endocrinol* 13:57-65
48. Belury MA, Moya CS, Sun H, Snyder E, Davis JW, Cunningham ML, Vanden Heuvel J 1998 Comparison of dose-response relationships for induction of lipid metabolizing and growth regulatory genes by peroxisome proliferators in rat liver. *Toxicol Appl Pharmacol* 151:254-61
49. Aoyama T, Peters JM, Iritani N, Nakajima T, Furihata K, Hashimoto T, Gonzalez FJ 1998 Altered constitutive expression of fatty acid-metabolizing enzymes in mice lacking the peroxisome proliferator-activated receptor alpha (ppar-alpha). *J Biol Chem* 273:5678-5684
50. Bankir L, Martin H, Dechaux M, Ahloulay M 1997 Plasma cAMP: a hepatorenal link influencing proximal reabsorption and renal hemodynamics? *Kidney Int Suppl* 59:S50-6
51. Bahnsen M, Burrin JM, Johnston DG, Pernet A, Walker M, Alberti KG 1984 Mechanisms of catecholamine effects on ketogenesis. *Am J Physiol* 247:E173-80
52. Kersten S, Seydoux J, Peters JM, Gonzalez FJ, Desvergne B, Wahli W 1999 Peroxisome proliferator-activated receptor alpha mediates the adaptive response to fasting. *J Clin Investig* 103:1489-98
53. Costet P, Legendre C, More J, Edgar A, Galtier P, Pineau T 1998 Peroxisome Proliferator-activated Receptor alpha-Isoform Deficiency Leads to Progressive Dyslipidemia with Sexually Dimorphic Obesity and Steatosis. *J Biol Chem* 273:29577-29585
54. Devchand PR, Keller H, Peters JM, Vazquez M, Gonzalez FJ, Wahli W 1996 The PPARalpha-leukotriene B4 pathway to inflammation control. *Nature* 384:39-43

55. Lazennec G, Alcorn JL, Katzenellenbogen BS 1999 Adenovirus-mediated delivery of a dominant negative estrogen receptor gene abrogates estrogen-stimulated gene expression and breast cancer cell proliferation. *Mol Endocrinol* 13:969-80
56. IJpenberg A, Jeannin E, Wahli W, Desvergne B 1997 Polarity and specific sequence requirements of peroxisome proliferator- activated receptor (PPAR)/retinoid X receptor heterodimer binding to DNA. A functional analysis of the malic enzyme gene PPAR response element. *J Biol Chem* 272:20108-17
57. Seglen PO 1976 Preparation of isolated rat liver cells. *Methods Cell Biol* 13:29-83
58. Sambrook J., Fritsch E.F., Maniatis T (ed.). 1989. *Molecular cloning: a laboratory Manual*, Ed. 2 ed. Cold Spring Harbor Laboratory Press, Cold Spring Harbor.