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**INTERACTION OF THE INSULIN RECEPTOR WITH THE RECEPTOR-
LIKE PROTEIN TYROSINE-PHOSPHATASES PTP α and PTP ϵ IN
LIVING CELLS*.**

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Abbreviations:

PTPases, Protein Tyrosine Phosphatases; PTP α , Protein Tyrosine Phosphatase α ;
PTP ϵ , Protein Tyrosine Phosphatase ϵ ; PTP1B, Protein Tyrosine Phosphatase 1B;
BRET; Bioluminescence Resonance Energy Transfer; mBU, milliBioluminescence
Resonance Energy Transfer Unit; GFP or YFP, Green or Yellow Fluorescent Protein;
Tpz: Topaz fluorescent protein; IR-Rluc, Insulin Receptor-Renilla luciferase; WGL,
Wheat-Germ Lectin; DSSTP, Dithiobis(sulfosuccinimidylpropionate); bpV(phen),
bisperoxo(1,10-phenanthroline)-oxovanadate; SDS-PAGE, sodium dodecyl sulfate
polyacrylamide gel electrophoresis.

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ABSTRACT

The interactions between the insulin receptor and the two highly homologous receptor-like protein-tyrosine-phosphatases PTP α and PTP ϵ were studied in living cells by using bioluminescence resonance energy transfer. In HEK293 cells expressing the insulin receptor fused to luciferase and substrate-trapping mutants of PTP α or PTP ϵ fused to the fluorescent protein Topaz, insulin induces an increase in resonance energy transfer that could be followed in real time in living cells. Insulin effect could be detected at very early time-points and was maximal less than 2 min after insulin addition. Bioluminescence resonance energy transfer saturation experiments indicate that insulin does not stimulate the recruitment of protein tyrosine-phosphatases molecules to the insulin receptor but rather induces conformational changes within pre-associated insulin receptor / protein tyrosine-phosphatases complexes. Physical pre-association of the insulin receptor with these protein tyrosine-phosphatases at the plasma membrane, in the absence of insulin, was also demonstrated by chemical cross-linking with a non-cell permeable agent. These data provide the first evidence that PTP α and PTP ϵ associate with the insulin receptor in the basal state and suggest that these protein tyrosine-phosphatases may constitute important negative regulators of the insulin receptor tyrosine-kinase activity, by acting rapidly at the plasma membrane level.

Introduction

Reversible tyrosine-phosphorylation is an important mechanism for the regulation of cellular functions. This process is controlled by opposite activities of protein tyrosine kinases and protein tyrosine phosphatases (PTPases). PTPases are divided in two classes, cytosolic PTPases and receptor-like PTPases. Receptor like-PTPases, including PTP α and PTP ϵ , are widely expressed. These PTPases are composed of a highly glycosylated extracellular domain, a single membrane-spanning domain and two tandem catalytic intracellular domains, designated as D1 and D2 (Sap et al., 1990). Extracellular ligands for these two PTPases have not as yet been found (Tonks and Neel, 2001). Dimerization involving multiple domains of the proteins can lead to inactivation of PTP α and PTP ϵ (Jiang et al., 1999; Tertoolen et al., 2001; Toledano-Katchalski et al., 2003). Furthermore, oxidative stress was recently shown to inhibit PTP α by stabilization of PTP α dimers (Blanchetot et al., 2002).

PTP α is involved in several cellular processes such as integrin-mediated responses by dephosphorylation/activation of c-src (Su et al., 1999) and cell adhesion by interaction with contactin (Zeng et al., 1999). The PTP ϵ subfamily comprises four proteins produced by a single gene. The cytosolic form of PTP ϵ is implicated in inhibition of Jak/STAT signaling by cytokines of the IL-6 family (Tanuma et al., 2001). PTP ϵ , the expression of which is elevated in mammary tumors, is also a physiological activator of Src and supports the transformed cell phenotype (Gil-Henn and Elson, 2003).

Insulin exerts its biological effects through a plasma membrane receptor that possesses a tyrosine-kinase activity (Combettes-Souverain and Issad, 1998). PTP α and PTP ϵ have also been implicated in the regulation of insulin signaling. Indeed, PTP α has been shown to inhibit insulin-induced prolactin gene expression (Jacob et

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al., 1998) and translocation of the glucose transporter GLUT4 (Cong et al., 1999). In the adipocyte, an important insulin target cell, PTP α exhibits a subcellular distribution at the plasma membrane similar to that of the insulin receptor. Importantly, in these cells, insulin induces a redistribution of both PTP α and the insulin receptor from the plasma membrane to heavy microsomes (Calera et al., 2000). Moreover, by using cells overexpressing the insulin receptor, PTP α and PTP ϵ were found to be efficient negative regulators of the insulin receptor tyrosine kinase activity (Andersen et al., 2001; Lammers et al., 1998; Moller et al., 1995). However, although these data indicate that PTP α and PTP ϵ can regulate the activity of the insulin receptor, the dynamics of the interaction of the IR with these PTPases in living cells has never been investigated.

Bioluminescence Resonance Energy Transfer (BRET) is a recently described methodology allowing the study of protein-protein interactions in intact living cells (Angers et al., 2000; Xu et al., 1999). To study the interaction between two partners, one of the partners is fused to Renilla Luciferase (Rluc) and the other to a GFP (i.e. YFP or Topaz). The luciferase is excited by addition of its cell permeable substrate, coelenterazine. If the two proteins are close enough for resonance energy transfer to occur, part of the energy of the excited luciferase can be transferred to the GFP, which then emits a fluorescent signal. As previously shown by our laboratory, this BRET methodology can be used to monitor the activation state of the insulin receptor (Boute et al., 2001) and the dynamics of interaction of this receptor with PTP1B in living cells (Boute et al., 2003).

In the present study, we have used this methodology to study the interaction of the insulin receptor with PTP α and PTP ϵ in living cells. We demonstrate that these PTPases physically interact with the insulin receptor, even in the absence of insulin.

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Insulin stimulation induces a rapid and dose-dependent increase in BRET signal. BRET saturation experiments revealed that this increase in BRET signal reflects a conformational change between the insulin receptor and the PTPases that are pre-associated with the receptor in the basal state, and not an insulin-induced recruitment of the PTPases to the activated receptor.

Materials and Methods

Materials.

All materials have been described previously (Boute et al., 2003; Boute et al., 2001), except anti-GFP monoclonal antibody (clones 7.1 and 13.1, Roche Applied Science, Indianapolis, IN), bisperoxo(1,10-phenanthroline)-oxovanadate anion (bpV(phen), Calbiochem, La Jolla, CA) and the thiol-cleavable cross-linker 3,3'-dithiobis [sulfosuccinimidylpropionate] (DTSSP, Pierce, Rockford, IL).

Expression vectors.

The cDNA encoding IR-RIuc has been described previously (Boute et al., 2001). Wild-type PTP α and PTP ϵ were subcloned in frame with the coding sequence of the fluorescent protein Topaz (Tpz) in the cytomegalovirus-Topaz (pGFPtpz-N1) vector (Packard, Meriden, CT). The substrate-trapping mutant forms of PTP α and PTP ϵ were generated by converting Asp to Ala at position 401 in PTP α (PTP α -D401A) and at position 302 in PTP ϵ (PTP ϵ -D302A), using a site-directed mutagenesis kit (Quick Change, Stratagene, La Jolla, CA). In some experiments (covalent cross-linking), a non-tagged insulin receptor cDNA was used (pcDNA3-IR).

Cell culture and transfection.

Cell cultures were performed as described previously (Boute et al., 2003). Briefly, HEK-293 cells were seeded at a density of 2×10^5 cells per 35-mm dish. One day later, the cells were transfected with 600 ng of IR-RIuc cDNA and 300 ng of PTP α -D401A-Tpz cDNA or 200 ng of PTP ϵ -D302A-Tpz cDNA per dish, unless otherwise specified in the figure legends. One day after transfection, cells were transferred into 96-well microplates (white culture plate-96, Perkin Elmer, Foster City, CA) at a density

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of 3×10^4 cells per well. BRET measurements were carried out in these microplates on the following day.

BRET measurements

BRET measurements were performed as previously described (Boute et al., 2003). Briefly, cells were preincubated for 20 min in PBS in the presence of $5 \mu\text{M}$ coelenterazine. The dynamics of interaction of IR-Rluc with PTP α -Tpz or PTP ϵ -Tpz was monitored for more than 20 min after insulin addition, by light-emission acquisition at 485 nm and 530 nm. In some experiments, the dynamics of the interaction of IR-Rluc with PTP α -D401A-Tpz or PTP ϵ -D302A-Tpz was monitored at very early time points as described in (Boute et al., 2003). BRET signal was expressed in milliBRET units (mBU). The BRET unit has been defined previously as the ratio 530 nm / 485nm obtained when the two partners are present, corrected by the ratio 530 nm / 485 nm obtained under the same experimental conditions, when only the partner fused to *Renilla* luciferase is present in the assay (Angers et al., 2000; Boute et al., 2003; Boute et al., 2002; Boute et al., 2001). Each measurement corresponded to the signal emitted by the whole population of cells present in a well.

Dephosphorylation of IR-Rluc by PTP α and PTP ϵ in intact cells.

HEK-293 cells were cotransfected with IR-Rluc and PTP α -D401A-Tpz, PTP α -wt-Tpz, PTP ϵ -D302A-Tpz or PTP ϵ -wt-Tpz. 48h after transfection, cells were incubated with 100 nM insulin for 5 min. Cell extracts were prepared as described in (Boute et al., 2001). After normalization according to luciferase activity, soluble extracts were incubated for 2h at 4°C with 50 μl of Wheat Germ Lectin-sepharose (Issad et al., 1995) and partially purified proteins were submitted to western-blotting.

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Immunoprobings were performed with anti-phosphotyrosine (4G10), anti-IR β and anti-GFP antibodies.

Cross-linking experiments:

48 h after transfection, HEK cells expressing the insulin receptor and either PTP α -D401A-Tpz or PTP ϵ -D302A-Tpz were incubated with 100 nM insulin for 5 min. Then, chemical cross-linking with the thiol-cleavable cross-linker DTSSP was performed at the final concentration of 1 mg/ml for 2h at 4°C. After cell lysis and normalization according to Topaz fluorescence, Tpz-fused proteins were immunoprecipitated using an anti-GFP antibody. Immunoblotting was performed using anti-IR β , anti-phosphotyrosine (4G10) and anti-GFP antibodies.

Statistical analysis

All results are expressed as means \pm SEM of at least three individual experiments. Comparisons between experimental groups were made using a Student *t* test for unpaired values or a Scheffe's test, as indicated in the legends of the figures.

Results

Functional expression of wild-type and substrate-trapping mutants of PTP α -Tpz and PTP ϵ -Tpz in HEK cells.

PTP α and PTP ϵ contain two tandem catalytic domains: a membrane proximal domain (D1) and a membrane distal domain (D2). The majority of the catalytic activity resides within D1, while D2 displays little or no activity. Inactivation of D1 abolishes the biological activity of PTP α (den Hertog et al., 1993). In a previous study on the interaction of the insulin receptor with PTP1B, we observed that no insulin-induced BRET signal could be detected with wild-type PTP1B (Boute et al., 2003). This probably reflects the fact that PTPases are enzymes with very high turnover rates, rendering the interaction between the phosphorylated IR and PTP1B too transitory to be detected by BRET. In contrast, a robust insulin-induced BRET signal was observed with a substrate-trapping mutant of PTP1B, which bind to but cannot dephosphorylate tyrosine-phosphorylated proteins. Substrate-trapping mutants of PTP α and PTP ϵ have also been previously designed to identify substrates of these enzymes (Buist et al., 2000; Gil-Henn and Elson, 2003). In the present work, we have generated substrate-trapping mutants of PTP α and PTP ϵ , by replacing aspartates 401 in PTP α and 302 in PTP ϵ by alanines. The carboxy-terminal regions of either wild-type or substrate-trapping mutant versions of PTP α and PTP ϵ were fused to the Tpz fluorescent protein (PTP α -wt-Tpz, PTP α -D401A-Tpz, PTP ϵ -wt-Tpz, PTP ϵ -D302A-Tpz). As expected, when expressed in HEK-293 cells, the fusion proteins were localized at the plasma membrane (Figure 1A).

We first studied the effect of expression of Topaz-tagged PTPases and their trapping mutants on the phosphotyrosine content of the β -subunit of the insulin receptor. As shown in Figure 1B and C, the insulin receptor was dephosphorylated by the wild-

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type versions of PTP α -Tpz and PTP ϵ -Tpz, whereas it was not dephosphorylated by the mutated versions of these enzymes.

Study of the interaction of the insulin receptor with PTP α and PTP ϵ by BRET

BRET measurements were performed using HEK-293 cells expressing the fusion proteins IR-Rluc and either PTP α -Tpz (wild-type or D401A mutant) or PTP ϵ -Tpz (wild-type or D302A mutant). BRET measurements were performed 10 min after insulin addition. As shown in Figure 2, a basal BRET signal could be detected with both fusion proteins. This signal was higher with PTP ϵ -wt-Tpz than with PTP α -wt-Tpz, despite similar PTPases and IR expression levels, determined by measuring Topaz fluorescence and luciferase activity, respectively (data not shown). Insulin treatment for 10 min at 100 nM had no effect on the BRET signal in cells expressing wild-type PTPases. In contrast, insulin increases the BRET signal in cells expressing PTP α -D401A-Tpz and PTP ϵ -D302A-Tpz. This increase was more pronounced with PTP ϵ than with PTP α .

In another set of experiments, BRET measurements were started immediately after insulin addition, and the BRET signal was monitored during more than 20 min. For all fusion proteins, the basal BRET signal remained stable throughout the experiment (Figures 3A and B). As observed previously (Figure 2), an insulin effect on BRET signal could only be detected with the substrate-trapping mutant versions of PTP α and PTP ϵ (Figures 3A and B). Interestingly, the insulin-induced BRET signal had already reached its maximal value at the first BRET measurement (2 min), and remained stable throughout the experiment. We then studied insulin effect on BRET signal at earlier time-points (Figures 3C and D). The effect of insulin on BRET signal

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could be detected as early as 20 sec, and the maximal value was reached between 1 and 2 min after insulin addition.

Dose-response experiments were performed by measuring the effect of different concentrations of insulin on BRET signal at time 10 min. Figure 4 shows dose-response curves of insulin-induced BRET signal (BRET signal above basal). We observed that, for both PTPases, insulin dose-dependently increases the BRET signal. The maximal response to insulin was higher for PTP ϵ -D302A-Tpz than for PTP α -D401A-Tpz (62.3 ± 2.5 mBU versus 22.2 ± 0.8 mBU, $p < 0.0002$). However, the half-maximal effect was obtained for similar insulin concentrations (5.9 ± 0.7 nM insulin for PTP α -D401A-Tpz and 4.0 ± 0.5 nM insulin for PTP ϵ -D302A-Tpz). These concentrations are in the same range as those that give half-maximal autophosphorylation of the insulin receptor (Boute et al., 2001).

Insulin-induced BRET signal was indeed tightly dependent on the autophosphorylation of the insulin receptor, as shown in experiments using an inhibitor of the tyrosine kinase activity of the receptor, the tyrphostin AG1024. We observed that, for both PTPases (Figure 5A and B), insulin-induced BRET signal was markedly inhibited in cells treated with AG1024. In addition, basal BRET signal was also significantly inhibited by AG1024, suggesting that part of the basal BRET signal depends on an insulin-independent autophosphorylation of the insulin receptor.

The BRET methodology also allows for the study of the effect of PTPase inhibitors (Boute et al., 2003). Peroxovanadium compounds are general inhibitors of PTPases that act by oxidizing the catalytic cysteine common to all PTPases (Huyer et al., 1997; Posner et al., 1994). We observed, for both PTPases, that basal and insulin-induced BRET signals were markedly inhibited by the bpV(phen) compound (Figure 6A and B). This could be due to an impaired accessibility of the phosphorylated tyrosine to

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the oxidized active site, or to a conformational change induced by oxidation of the PTPases, resulting in a decrease in energy transfer (Persson et al., 2004).

BRET saturation experiments

We have observed that insulin induces an increase in the BRET signal, both with PTP α and PTP ϵ . The BRET signal not only depends on the distance between the luminescent and fluorescent proteins, but also on their relative orientation. Therefore, an insulin-induced BRET signal could either reflect the recruitment of additional PTPases molecules to insulin receptors upon their activation, or a conformational change within complexes containing insulin receptors pre-associated with PTPases.

Quantitative BRET analysis has been used to assess GPCR homo- and heterodimerization (Mercier et al., 2002) as well as leptin receptor oligomerization (Couturier and Jockers, 2003). This analysis is based on the principle that the level of energy transfer detected for a constant donor concentration should rise with increasing concentrations of acceptor. When all donor molecules are engaged by an acceptor molecule, the energy transfer reaches a plateau. Thus, saturation curves can be constructed where the maximal level reached reflects the total number of acceptor fusion/donor fusion protein complexes. The relative amount of acceptor giving 50% of maximal energy transfer (BRET₅₀) should reflect the relative affinity of the acceptor fusion protein for the donor fusion protein.

Here, we applied this approach to determine whether insulin induces an increase in the relative affinity of PTP α or PTP ϵ for the IR (i.e., an increase in the proportion of PTPases recruited to the IR at a given PTPase/IR ratio). BRET saturation experiments were performed in HEK-293 cells co-transfected with a constant amount of IR-Rluc and increasing concentrations of PTP α -D401A-Tpz or PTP ϵ -D302A-Tpz. As shown in Figure 7A and B, the curves behaved as hyperbolic functions reaching a

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saturation level. The $BRET_{max}$ values obtained for PTP α -D401A-Tpz were 68.7 ± 8.3 mBU in the absence of insulin and 99.2 ± 11.6 mBU in the presence of 100 nM insulin. For PTP ϵ -D302A-Tpz, $BRET_{max}$ values were 158.1 ± 16 mBU in the absence and 207 ± 15 mBU in the presence of 100 nM insulin. However, $BRET_{50}$ values were not affected by insulin, neither for PTP α (0.77 ± 0.37 in the absence and 0.83 ± 0.36 in the presence of 100 nM of insulin) nor for PTP ϵ (2.14 ± 0.80 in the absence and 1.25 ± 0.44 in the presence of insulin). These saturation curves were also represented as percentage of $BRET_{max}$ for PTP α -D401A-Tpz and PTP ϵ -D302A-Tpz. As shown in Figure 7C and D, the shape of these curves is identical in the absence and presence of insulin, clearly indicating that insulin does not modify the relative affinities of the insulin receptor for the PTPases. Therefore, for both PTPases, insulin-induced BRET signal reflects a conformational change between insulin receptors and PTPases molecules that are pre-associated in the basal state.

Covalent cross-linking of the insulin receptor with PTP α and PTP ϵ in transfected HEK -293cells.

HEK-293 cells co-transfected with the insulin receptor and either PTP α -D401A-Tpz or PTP ϵ -D302A-Tpz were incubated for 5 min in the absence or presence of 100 nM insulin. The water-soluble DTSSP cross-linking agent was then added to the cells and cross-linking was carried-out for 2 h at 4°C. After cell lysis, the Tpz-fused proteins were immunoprecipitated with anti-GFP antibody and immunoblotting was performed with anti-IR β antibody. Figure 8 shows that in the absence of insulin, substantial amounts of insulin receptors could be co-precipitated with PTP α or PTP ϵ . Insulin did not significantly increase the amount of receptor precipitated with the PTPases. This strongly suggest that the insulin receptor indeed interacts with these PTPases in the basal state, and that insulin does not further increase this interaction.

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This is in agreement with the results obtained in BRET saturation experiments indicating that insulin-induced BRET signal reflects a conformational change between pre-associated partners rather than an additional recruitment of the PTPases to the activated insulin receptors.

Discussion

Among the different PTPases that control the insulin receptor, the intracellular protein tyrosine-phosphatase PTP1B appears to play a major role (Cicirelli et al., 1990; Ahmad et al., 1995; Kenner et al., 1996; Elchebly et al., 1999; Klaman et al., 2000)

However, several lines of evidence suggest that other PTPases may also be involved in the control of IR activity (Cheng et al., 2002). Notably, in PTP1B knockout mice, insulin sensitivity was increased in liver and muscles, but not in adipose tissue, revealing important tissue-specific differences in the regulation of the IR by PTPases (Elchebly et al., 1999). In major contrast with PTP1B, which is targeted to the endoplasmic reticulum by means of its C-terminal sequence (Frangioni et al., 1992), PTP α and PTP ϵ are localized, like the insulin receptor, at the plasma membrane level. Therefore, the dynamics of interaction of the insulin receptor with these two types of PTPases is expected to be very different. Using the BRET methodology, we have shown previously (Boute et al., 2003) that insulin-induced interaction between the insulin receptor and PTP1B necessitates internalization of the insulin receptor. In these experiments, although an insulin-induced BRET signal could be detected at very early time points (less than one minute), it reached its maximal value only 20 min after insulin addition (Boute et al., 2003). In contrast, for PTP α and PTP ϵ , insulin-induced BRET signal was already maximal 2 min after insulin stimulation (Figure 3). Because the insulin receptor and the two receptor-like PTPases are localized in the same membrane, at the surface of the cell, it is conceivable that insulin stimulation rapidly induces the association of the PTPases to the insulin receptor. However, BRET saturation experiments suggest that insulin receptors and these plasma membrane PTPases are pre-associated in the absence of insulin, and insulin does not further recruit additional PTPase molecules to the activated receptor (Figure 7).

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This notion is strongly supported by covalent cross-linking experiments (Figure 8). This important observation suggests that, in the basal state, the insulin receptor and the PTPases are present in a pre-formed complex. Upon activation of the receptor, a conformational change occurs that can be detected as an increase in BRET signal. This increase is inhibited by the tyrphostin AG1024, indicating that it depends on the autophosphorylation of the receptor (Figure 5). Therefore, the increase in BRET signal could reflect a conformational change that occurs within the insulin receptor itself upon autophosphorylation, a conformational change within the PTPase upon docking of a phosphorylated tyrosine of the insulin receptor into the active site of the PTPase, or both. Interestingly, basal BRET signal was also significantly reduced by AG1024, indicating that part of the basal BRET signal may reflect the control, by these plasma membrane PTPases, of an autonomous autophosphorylation activity of the insulin receptor.

We have observed that insulin-induced BRET signal is higher with PTP ϵ than with PTP α (Figure 4). However, since BRET saturation experiments indicate that insulin effect on BRET only reflects a conformational change between already interacting partners, it is likely that the higher insulin-induced BRET signal observed with PTP ϵ only reflects a more favorable orientation of luciferase and topaze for resonance energy transfer in PTP ϵ /IR complexes than in PTP α /IR complexes.

PTPases activity can be regulated by oxidation of the cystein residue located in their active site. Interestingly, in the case of PTP α , it has been shown that the second phosphatase domain (D2) is more susceptible to oxidation than D1 (Persson et al., 2004). Moreover, oxidation of D2 results in a conformational change in PTP α (van der Wijk et al., 2003). We found that bpV(phen) treatment inhibited basal and insulin stimulated BRET signal (Figure 6). This could reflect a conformational change of the

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PTPases induced by oxidation of D2, resulting in a lower efficiency of energy transfer between the luciferase and Tpz within the insulin receptor/PTPases complex. It has also been shown that a substantial proportion of PTP α at the plasma membrane is in the form of pre-existing dimers (Jiang et al., 2000; Tertoolen et al., 2001), and that oxidation of D2 results in the stabilization of these dimers into an inactive conformation (Blanchetot et al., 2002). It is unclear at the present time whether PTP α and PTP ϵ pre-associated with the insulin receptor at the plasma membrane are in the form of monomers or dimers, and whether insulin-induced conformational changes in these complexes imply modification in the dimerization state of these PTPases.

One of the most important findings of this work is the observation of pre-existing complexes between the insulin receptor and PTP α or PTP ϵ at the cell surface. Pre-association of the insulin receptor with plasma membrane tyrosine-phosphatases may have important consequences on cell physiology. For instance, it has been shown recently that tyrosine kinase receptors such as EGF, PDGF and insulin receptors, stimulate the production of phosphatidyl inositol-3 phosphate by endomembranes after endocytosis, indicating that these endocytosed receptors are engaged in the activation of signalling pathways before being inactivated by PTP1B (Sato et al., 2003). Therefore, depending on the relative levels of expression of plasma membrane PTPases (e.g. PTP α or PTP ϵ) versus endomembrane PTPases (e.g. PTP1B) in a given cell type, the insulin signal could be interrupted either at the plasma membrane or at the endomembrane level, and this may induce very different cellular responses.

In summary, our work provides a new model for the mechanism of interaction of the insulin receptor with plasma membrane protein tyrosine-phosphatases. In this model, the insulin receptor, in the basal state, is engaged in molecular complexes with the

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PTPases. Insulin treatment induces a conformational change within these complexes, without modifying the number of receptor interacting with the PTPases. This information may be important for the development of therapeutic agents specially designed for the treatment of insulin resistance and diabetes.

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Footnotes

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Legends for Figures

Figure 1: (A) Expression of topaz fluorescent protein (Tpz) fusion proteins in human HEK293 cells. Transfection of cDNAs coding for PTP α -wt-Tpz, PTP α -D401A-Tpz, PTP ϵ -wt-Tpz and PTP ϵ -D302A-Tpz constructs results in localization of the fluorescence at the plasma membrane. (B, C) Dephosphorylation of the insulin receptor by PTP α -Tpz and PTP ϵ -Tpz fusion proteins. HEK cells were co-transfected with IR-Rluc and PTP α -D401A-Tpz or PTP α -wt-Tpz (B) and PTP ϵ -D302A-Tpz or PTP ϵ -wt-Tpz (C). Cells were stimulated with 100nM insulin for 5 min and then lysed. After purification on wheat-germ lectin sepharose beads, proteins were analysed by SDS-PAGE followed by immunoblotting with anti-phosphotyrosine antibody (anti PY), anti-IR β antibody and anti-GFP antibody. The lower panel shows the results of densitometric analysis of the anti-phosphotyrosine signal corrected by the anti-IR β signal. The data presented are means \pm sem of 4 to 5 independent experiments (**, ***, $P < 0.01$, $P < 0.001$ respectively, when compared to cells transfected with IR-Rluc alone using a Scheffe's test).

Figure 2: Effect of insulin on BRET signal in HEK-293 cells expressing IR-Rluc and wild-type or substrate-trapping mutant of PTP α -Tpz or PTP ϵ -Tpz fusion proteins. BRET measurements were performed 48h after transfection in cells incubated for 10 min in absence or presence of 100 nM insulin. Results are means \pm SEM of 3-6 independent experiments (**, ***, $P < 0.01$, $P < 0.001$ respectively when compared to basal condition using a Student's t test).

Figure 3: Dynamics of the interaction between the insulin receptor and PTP α or PTP ϵ . BRET signal was monitored during 20 min after insulin addition in cells

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expressing IR-Rluc and wild-type or the substrate-trapping versions of PTP α -Tpz (A) or PTP ϵ -Tpz (B). Early effects of insulin on BRET between insulin receptor and the substrate-trapping versions of PTP α -Tpz (C) or PTP ϵ -Tpz (D) were also studied. The data presented are representative of 3 independent experiments.

Figure 4: Dose-dependent effect of insulin on BRET signal. HEK cells were co-transfected with IR-Rluc and either PTP α -D401A-Tpz or PTP ϵ -D302A-Tpz. 48h after transfection, cells were incubated with increasing concentrations of insulin and BRET signal was measured. Insulin-induced BRET signal corresponds to the increase in BRET above basal in each experimental condition. The results are means \pm SEM of 6 separate experiments.

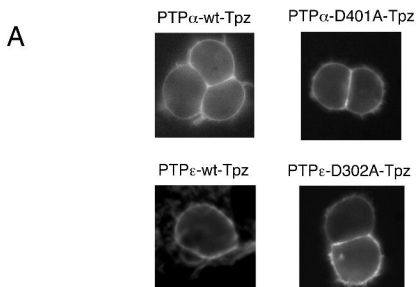
Figure 5: Effect of tyrphostin AG 1024 on BRET signal. HEK cells co-transfected with IR-Rluc and PTP α -D401A-Tpz (A) or PTP ϵ -D302A-Tpz (B) were preincubated for 1h in absence or presence of 100 μ M AG 1024. Cells were then stimulated with 100 nM insulin and BRET measurements were performed. Results are means \pm SEM of 3-4 separate experiments. (*, ***, $P < 0.05$, $P < 0.001$ respectively when compared to AG1024-untreated controls using a Scheffe's test).

Figure 6: Effect of bpV(phen) on BRET signal. HEK cells co-transfected with IR-Rluc and PTP α -D401A-Tpz (A) or PTP ϵ -D302A-Tpz (B) were preincubated in the absence or the presence of 1 mM bpV(phen) for 30 min. Cells were then stimulated with 100 nM insulin and BRET measurements were performed. Results are means \pm SEM of 3-4 independent experiments. (*, **, ***, $P < 0.05$, $P < 0.01$, $P < 0.001$ respectively when compared to bpV(phen)-untreated controls using a Scheffe's test).

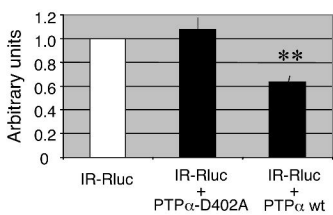
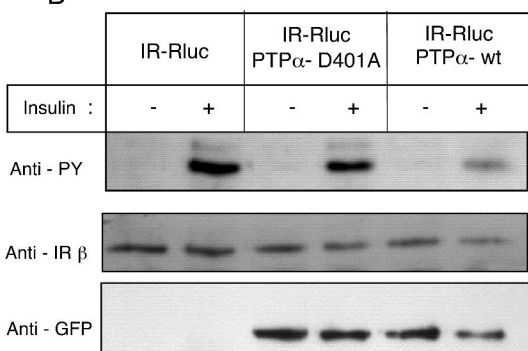
Figure 7: Saturation curves of BRET between insulin receptor and PTP α or PTP ϵ . HEK-293 cells were co-transfected with a constant amount of cDNA coding for IR-Rluc (600 ng/dish) and increasing amounts of PTP α -D401A-Tpz (150-600 ng/dish) (A) or PTP ϵ -D302A-Tpz (100-600ng/dish) (B). 48h after transfection, BRET signal, luciferase and fluorescence levels were measured in each experimental condition. BRET signals were plotted as a function of the ratio of PTP α -D401A-Tpz fluorescence or PTP ϵ -D302A-Tpz fluorescence to IR-Rluc luminescence. The results presented are from at least 6 independent experiments. The curves were fitted using a non-linear regression equation assuming a single binding site (GraphPad Prism). Data from A and B are presented in C and D as percent of BRET max.

Figure 8: Chemical cross-linking of the insulin receptor with PTP α -D401A-Tpz or PTP ϵ -D302A-Tpz. HEK-293 cells co-transfected with the insulin receptor and PTP α -D401A-Tpz or PTP ϵ -D302A-Tpz were incubated for 5 min in absence or presence of 100 nM insulin. Cells were then incubated with the non-cell permeable cross-linker DTSSP. Immunoprecipitation was performed using anti-GFP antibodies. Immune complexes were isolated on protein G sepharose and submitted to SDS-PAGE followed by western-blot. Immunodetection was performed using anti-IR β , anti-phosphotyrosine or anti-GFP antibody. The data presented are representative of 2 independent experiments.

Figure 1



B



C

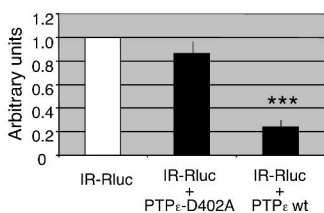
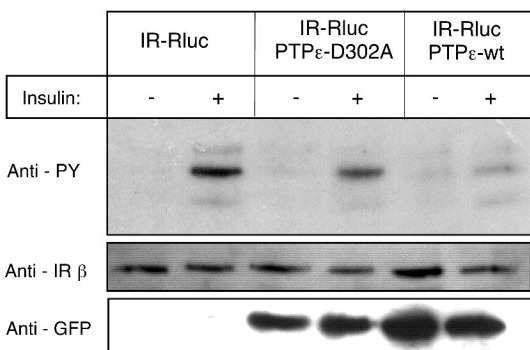


Figure 2

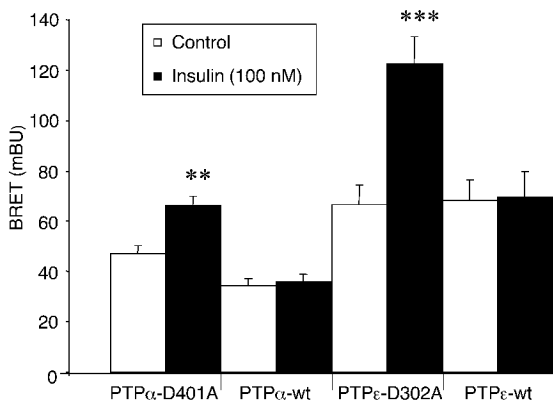


Figure 3

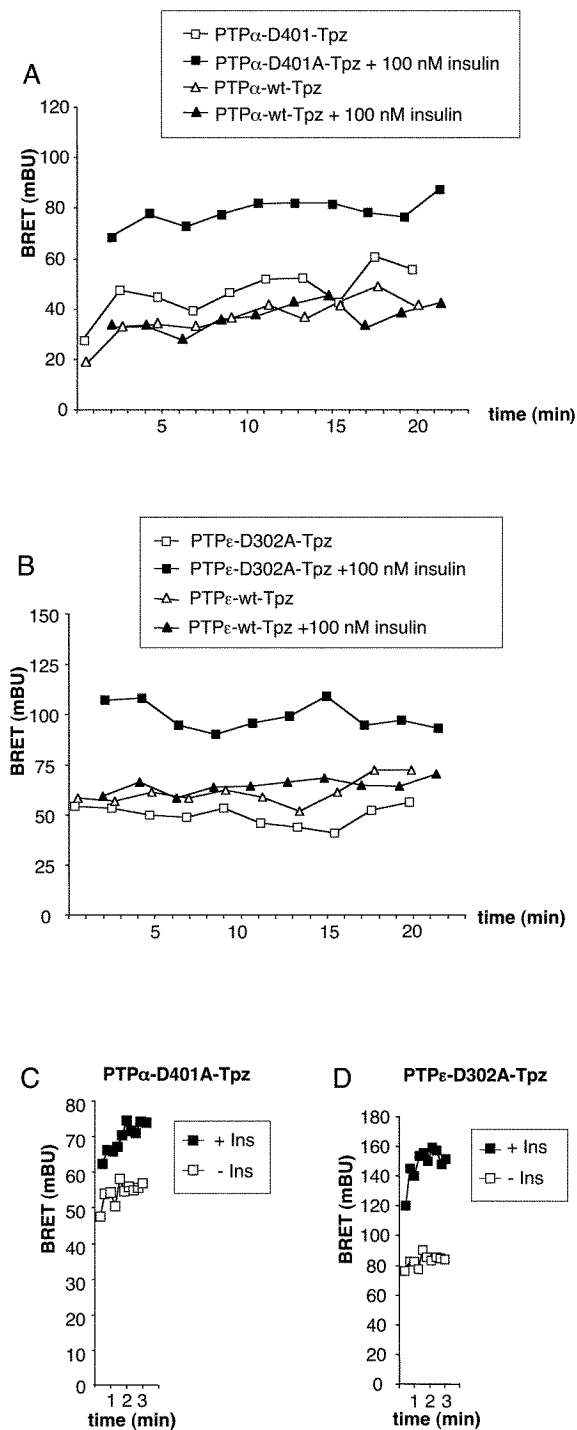


Figure 4

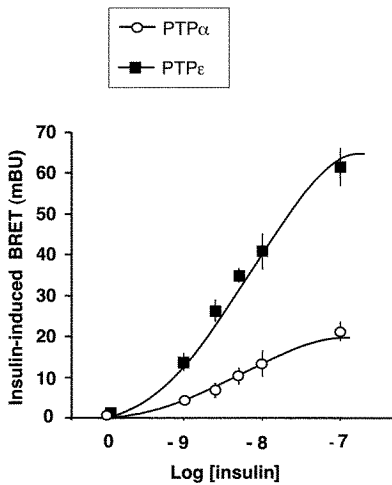


Figure 5

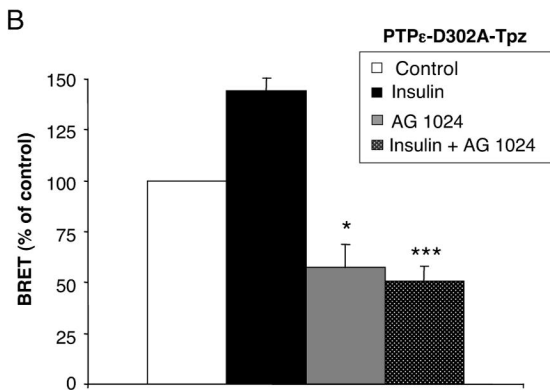
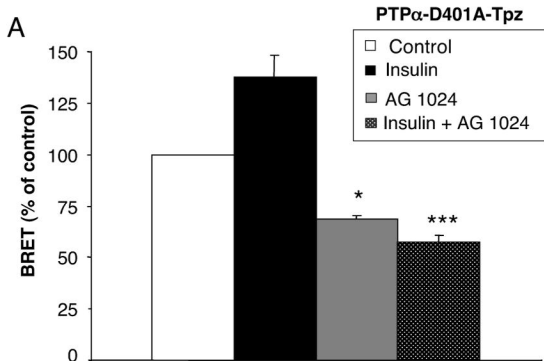


Figure 6

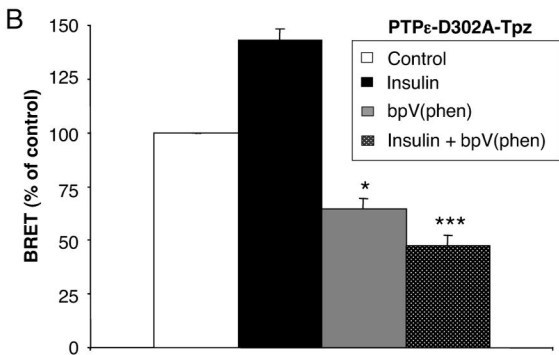
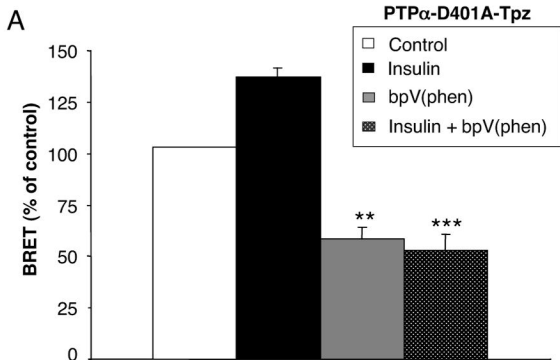
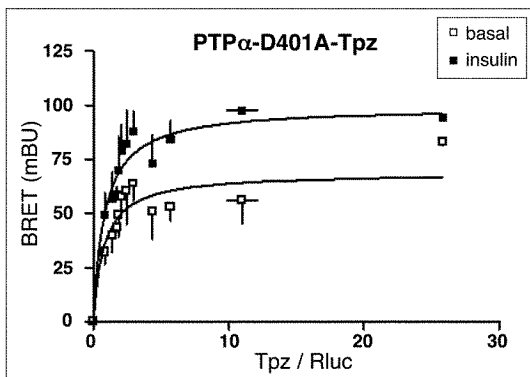
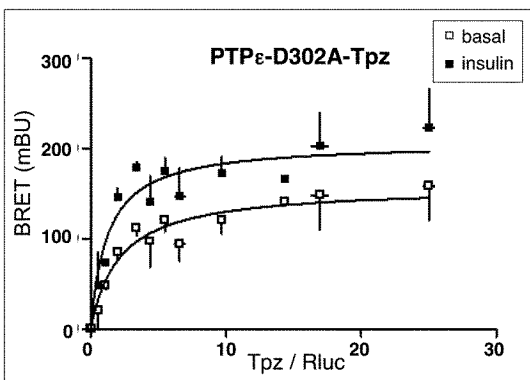


Figure 7

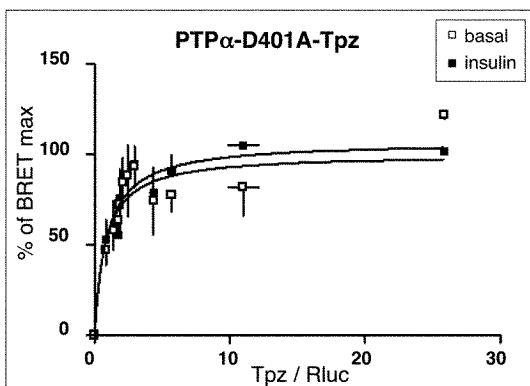
A



B



C



D

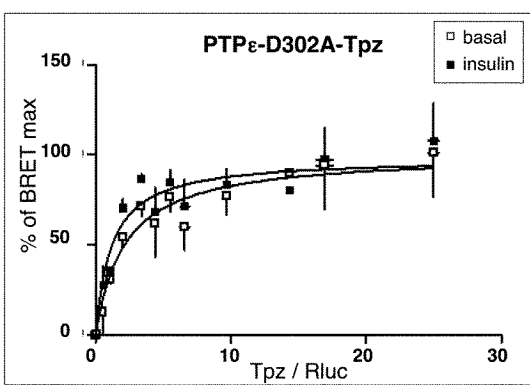


Figure 8

