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Preferential formation of MT_1/MT_2 melatonin receptor heterodimers

with distinct ligand interaction properties compared to MT_2 homodimers

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Abbreviations : BRET, bioluminescence resonance energy transfer; CFP, cyan fluorescent

protein; FRET, fluorescence resonance energy transfer; GPCR, G protein-coupled receptor;

Rluc, *Renilla* luciferase; YFP, yellow fluorescent protein, ^{125}I-MLT, 2(^{125}I)-iodomelatonin.

Abstract

Heterodimerization has been documented for several members of the G protein-coupled receptor (GPCR) super-family, including the closely related MT_1 and MT_2 melatonin receptors. However, the relative abundance of hetero- versus homodimers and the specific properties, which can be attributed to each form, are difficult to determine. Using a bioluminescence resonance energy transfer (BRET) donor saturation assay, we show that half maximal MT_1/MT_2 heterodimer formation is reached for expression levels as low as ~4000 receptors/ cell. The relative propensity of MT_1 homodimer and MT_1/MT_2 heterodimer formation are similar, whereas that for the MT_2 homodimer formation is 3-4 fold lower. These data indicate that both the relative expression level of each receptor isoform and the affinities between monomers may determine the actual proportion of homo- and heterodimers. The specific interaction of ligands with the MT_1/MT_2 heterodimer was studied using a BRET-based assay as a read out for the conformational changes of the heterodimer. A MT_1/MT_2 heterodimer-specific profile as well as ligands selective for the MT_1/MT_2 heterodimer compared to the MT_2 homodimer could be identified. Classical radioligand binding and BRET studies suggest that heterodimers contain two functional ligand binding sites that maintain their respective selectivity for MT_1 and MT_2 ligands. Occupation of either binding site is sufficient to induce a conformational change within the heterodimer. Taken together, we show that the probability of GPCR heterodimer formation may be equal or even higher than that of the corresponding homodimers and that specific properties of heterodimers can be revealed using a BRET-based ligand/receptor interaction assay.

INTRODUCTION

A growing number of observations suggest that G protein-coupled receptors (GPCR) form homodimers and heterodimerize with other members of the same receptor super-family. Heterodimerization may have important consequences in terms of receptor function, as significant changes in ligand binding, signaling or trafficking were observed for several heterodimers (Gazi et al., 2002). Considering that multiple receptors are expressed simultaneously in tissues and cells, it is reasonable to assume that most cells co-express several different GPCR that may be engaged into heterodimeric complexes. So far, little is known about the rules that govern homo- and heterodimer formation. The homo-/heterodimer ratio is expected to depend on the relative affinity of receptor subtypes for each other and on the expression level of the interacting partners. Most studies on GPCR heterodimerization did not examine the proportion of heterodimers versus homodimers. In addition, these studies were principally performed in cells expressing supra-physiological levels of recombinant receptors, in which the formation of GPCR heterodimers might be over-estimated, compared to normal tissues that express endogenous levels of receptors. Recent quantitative BRET-based studies in transfected cells expressing high concentrations (1-10 pmol/mg of protein) of β_1 and β_2-adrenergic receptors (Mercier et al., 2002) or oxytocin and vasopressin receptors (Terrillon et al., 2003) suggested that the probability to form homo- and heterodimers is similar.

Ligand binding properties of receptors may change when they are engaged in heterodimeric complexes (Jordan and Devi, 1999). Because GPCRs are major pharmacological targets, the discovery of specific ligand binding profiles for heterodimers may have important implications for the development and screening of new drugs. However,

the determination of a specific binding profile for heterodimers is difficult to establish using classical radioligand competition binding assays. This limitation is particularly true for heterodimers composed of two receptors, which display similar affinities for the same radioligands.

The interaction between ligand and receptor can be studied with alternative approaches, which measure the conformational changes of ligand-bound receptors such as fluorescence and electron paramagnetic resonance spectroscopy (Farrens et al., 1996; Ghanouni et al., 2001; Lee et al., 1997). Indeed, the efficiency to promote specific ligand-induced conformations (EC_{50}) is correlated, in theory, with the binding affinities of the ligands (Kenakin and Onaran, 2002). Recently, resonance energy transfer techniques such as fluorescence and bioluminescence resonance energy transfer (FRET, BRET), have also emerged as sensitive approaches to monitor conformational changes of a wide range of proteins in living cells, including membrane receptors (Heyduk, 2002; Truong and Ikura, 2001). Energy transfer occurs if the energy donor is in close proximity (10 - 100 Å) to the energy acceptor and if the respective orientation of donor and acceptor is appropriate (Heyduk, 2002; Truong and Ikura, 2001). The extreme sensitivity to relatively small perturbations makes this technique an attractive approach to detect receptor conformational changes. Two strategies have been used to study a protein of interest with this approach. Both, energy donor and acceptor may be fused to the same protein to be studied (intramolecular energy transfer), as reported previously to monitor calcium- and cAMP-dependent signaling, phosphorylation ((Heyduk, 2002; Truong and Ikura, 2001) for review) or the activation of parathyroid hormone and alpha$_{2A}$ adrenergic receptors (Vilardaga et al., 2003). The second strategy takes advantage of the fact that most receptors exist as dimers. The co-expression of two receptors, one fused to the energy donor and the second to the acceptor allows to monitor ligand-induced conformational changes within constitutive receptor dimers (intermolecular

energy transfer) ((Heyduk, 2002; Truong and Ikura, 2001) for review). Such an approach was used to study the insulin receptor (Boute et al., 2001), the leptin receptor (Couturier and Jockers, 2003) and several GPCRs (Angers et al., 2000; Kroeger et al., 2001; Rocheville et al., 2000a). Stimulation of these receptors with the appropriate hormones modified the constitutive energy transfer in a dose-dependent manner, supporting the idea that the conformational changes modify the distance and/or the orientation between the two BRET partners.

Using a BRET-based approach, we have shown recently that MT_1 and MT_2 melatonin receptors, which share 70 % sequence homology, form both homo- and heterodimers (Ayoub et al., 2002). Here we report that MT_1/MT_2 heterodimers constitute a significant proportion of overall dimers, which can be distinguished from homodimers in living cells on the basis of their ligand-receptor interaction profile determined by a proximity-based BRET-assay.

EXPERIMENTAL PROCEDURE

Materials. Compounds were obtained from the following sources : melatonin were from Sigma, St Louis, MO, S20098 (N-[2-(7-methoxynapht-1-yl)ethyl]acetamide), S20928 (N-[2-(1-naphtyl)ethyl]cyclobutanecarboxamide), S22153 (N-[2-(5-ethylbenzo[b]thiophen-3-yl)ethyl]acetamide), S24773 (N-{2-[3-(3-aminophenyl)-7-methoxy-1-naphtyl]ethyl}acetamide) and S26284 (N-(2-{7-[4-({8-[2-acetylamino)ethyl]-2-naphtyl}oxy)butoxy]-1-naphtyl}ethyl)acetamide) were from the Institut de Recherche Servier (Audinot et al., 2003), 2-iodomelatonin from RBI, Natick, MA and luzindole (2-benzyl *N*-acetyltryptamine) and 4P-PDOT (4-phenyl-2-proprionamidotetraline) from Tocris, Ellisville, MO.

Plasmid constructions, transfections and cell culture. Construction of Rluc and YFP fusion proteins and Flag-MT1 and Myc- MT_2 constructs have been described elsewhere (Ayoub et al., 2002). HEK 293 cells were grown in complete medium (DMEM supplemented with 10 % (v/v) FBS, 4.5 g/liter glucose, 100 U/ml penicillin, 0.1 mg/ml streptomycin, 1 mM glutamine) (all from Life Technologies (Gaithersburg, MD)). Transient and stable transfections were performed using the transfection reagent FuGene 6 (Roche, Basel, Switzerland) according to supplier instructions.

Radioligand Binding Experiments. Whole-cell radioligand binding assays were performed as described (Brydon et al., 1999b). Radioligand binding assays were performed in PBS (pH 7.4) using the lipophilic $2(^{125}I)$-iodomelatonin (^{125}I-MLT) (NEN, Boston, MA) at 25 - 1000 pM as radioligand in saturation experiments. Specific binding was defined as binding displaced by 10 µM melatonin (Sigma, St Louis, MO). Competition binding assays were carried out at 100-200 pM ^{125}I-MLT and increasing concentrations of different compounds.

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Assays were carried out for 60 min at 37°C, terminated by rapid filtration through Whatman

GF/F glass fiber filters previously soaked in PBS and filters counted in a γ-counter.

Competition curves were fitted using a one or two site non-linear regression (GraphPad

Prism). IC_{50} values were transformed into K_i values using the Cheng-Prussof formula : $K_i = IC_{50}/ [1+(L/ K_d)]$ where L corresponds to the ^{125}I-MLT concentration and K_d corresponds to

the respective values obtained in ^{125}I-MLT saturation binding assays.

Crude membrane preparation, solubilization and immunoprecipitation. Crude

membranes were prepared, solubilized with 1 % digitonin, a detergent known to maintain

MTR in a native conformation and immunoprecipitated as described recently (Brydon et al.,

1999a; Roka et al., 1999) with 2 µg/ml of the monoclonal anti c-Myc 9E10 antibody (Santa

Cruz Biotechnology, CA).

SDS-PAGE / Immunoblotting. Immunoprecipitates were denatured over night in 62.5

mM Tris/HCl (pH 6.8), 5 % SDS, 10 % glycerol, 0.05 % bromophenol blue at room

temperature. Proteins were separated by 10 % SDS-PAGE and transferred to nitrocellulose.

Immunoblot analysis was carried out with the polyclonal anti-Flag antibody (2 µg/ml)

(Sigma, St Louis, MO). Immunoreactivity was revealed using appropriate secondary antibody

coupled to horseradish peroxidase and the ECL chemi-luminescent reagent (Amersham,

Aylesbury, UK).

Microplate BRET assay. Forty-eight hours post-transfection, HEK 293 cells were

detached and washed with PBS. $1-2 \times 10^5$ intact cells were distributed in a 96-well microplate

and incubated for 10 minutes at 25°C in the absence or presence of the indicated ligands.

Coelenterazine h substrate (Molecular Probes, Eugene, OR) was added at a final

concentration of 5 µM and readings were performed with a lumino/fluorometer (Fusion$^{\text{TM}}$,

Packard Instrument Company, Meriden, CT) that allows the sequential integration of

luminescence signals detected with two filter settings (Rluc filter : 485 ± 10 nm; YFP filter :

530 ± 12.5 nm) as described previously (Ayoub et al., 2002). The EC_{50} was defined as the

ligand concentration necessary to promote 50 % of the maximal ligand-induced BRET signal.

Correlation of fluorescence and luminescence levels of receptor fusion proteins to

^{125}I-MLT binding sites. Luminescence and fluorescence levels of several luciferase and GFP

receptor fusion proteins have been shown to be linearly correlated to receptor numbers

(Ayoub et al., 2002; Couturier and Jockers, 2003; McVey et al., 2001; Mercier et al., 2002).

Since this correlation is an intrinsic characteristic of each fusion protein, correlation curves

have to be established for each construct. HEK 293 cells were transfected with increasing

DNA concentrations of the melatonin receptor Rluc or YFP fusion protein constructs.

Maximal luminescence was determined at 485 ± 10 nm (Gain 4, PMT 1100 V, 1.0 sec) in 96-

well optiplates using coelenterazine h (5 µM) as substrate in Rluc-expressing cells and

fluorescence obtained upon exogenous YFP excitation (Gain 4, PMT 1100 V, 1.0 sec) was

measured in 96-well HTRF plates (Packard Instrument Company, Meriden, CT) in YFP-

expressing cells with a lumino/fluorometer Fusion[TM]. Background luminescence and

fluorescence determined in wells containing untransfected cells was substracted. To correlate

the luminescence and the fluorescence values with relative receptor numbers, the total number

of ^{125}I-MLT binding sites was determined in the same cells as described under "Radioligand

Binding Experiments". Luminescence and fluorescence were plotted against binding sites and

linear regression curves were generated (Fig. 1 of supplemental material). To determine the

expression level of YFP versus Rluc fusion proteins in cells co-expressing both proteins, the

maximal luciferase activity and fluorescence were determined using the same parameters as

described above and the YFP/Rluc ratio was calculated using the corresponding standard

curves. Reliable quantification of luciferase activity was possible under conditions of energy

transfer between YFP and Rluc fusion proteins since the amount of energy transfer observed

in the presence of YFP fusion receptors was negligible compared to the luciferase signal.

Luciferase activity remained constant under conditions where the basal energy transfer

increased 1.5 - 3 fold in the presence of melatonin (see Fig. 2 of supplemental material).

RESULTS

**Evidence for MT$_1$/MT$_2$ Heterodimerization based on Co-immunoprecipitation and

BRET experiments.** In a previous study we reported, using a BRET-based approach, that

both MT$_1$ and MT$_2$ melatonin receptors form homodimers in living HEK 293 cells (Ayoub et

al., 2002). In addition, our data suggested that these receptors may also form heterodimers. To

extend these observations, co-immunoprecipitation experiments were carried out with

epitope-tagged receptors. The human MT$_1$ receptor tagged with a Flag at its N terminus was

transiently expressed in HEK 293 cells, which stably express 25 fmol per mg of total protein

of MT$_2$ receptors tagged with a Myc epitope at their N-terminus. In accordance with previous

observations (Ayoub et al., 2002), western blot analysis of membranes prepared from these

cells with an anti-flag antibody revealed two major immunoreactive forms with apparent

molecular weights of 55 kDa and 110 kDa, likely corresponding to the monomeric and the

dimeric forms of the Flag-MT$_1$ (Fig. 1a, mb). The Flag-MT$_1$ was also pulled down by the

immunoprecipitation of the Myc-tagged MT$_2$ receptor, confirming the existence of MT$_1$/MT$_2$

heterodimers in cell lysates (Fig. 1a, IP). MT$_1$/MT$_2$ heterodimerization was further studied in

intact cells by BRET. Wild-type human MT$_1$ and MT$_2$ receptors were tagged at their C

terminus with either Rluc (BRET donor) or YFP (BRET acceptor). Fusion proteins retained

both their ligand binding (Tab. 1) and signaling properties (Ayoub et al., 2002). Two different

combinations of MT$_1$ and MT$_2$ fused to either BRET donor or acceptor were studied. A

significant BRET signal was observed for both combinations (Fig. 1b). The specificity of

these signals is illustrated by the absence of significant transfer between MT_1-Rluc or MT_2-

Rluc and a control β_2-adrenergic receptor YFP fusion protein expressed at comparable levels.

Taken together, these data confirm our previous observation that MT_1/MT_2 heterodimers are

formed in intact HEK 293 cells.

Evaluation of the proportion of MT_1 and MT_2 homo- and heterodimers in living

cells. Although it is clear that MT_1 and MT_2 homo- and heterodimers are formed in HEK 293

cells, the actual proportion of homo- and heterodimers present in these cells remains

unknown. To address this question, we measured the relative tendency of forming homo- and

heterodimers in living cells with a BRET donor saturation assay (Couturier and Jockers, 2003;

Mercier et al., 2002). Cells were co-transfected with constant amounts of cDNA coding for

the BRET donor (receptor fused to Rluc) and increasing quantities of cDNA for the BRET

acceptor (receptor fused to YFP). The amount of each receptor species effectively expressed

in transfected cells was determined, for each individual experiment, by correlating both

luminescence and fluorescence signals with ^{125}I-MLT binding sites (see Fig. 1 of the

supplemental material). As shown in Fig. 2, BRET signals increased as an hyperbolic function

of the ratio between the BRET acceptor and the BRET donor reaching an asymptote, which

corresponds to the saturation of all BRET donor molecules by acceptor molecules. Assuming

that the association of interacting proteins, fused to the BRET donor and the BRET acceptor

respectively, is random, the amount of acceptor required to obtain the half-maximal BRET

($BRET_{50}$) for a given amount of donor reflects the relative affinity of the two partners

(Mercier et al., 2002; Terrillon et al., 2003). Comparable $BRET_{50}$ values were observed for

the MT_1-homodimer and MT_1/MT_2 heterodimer formation, whereas the $BRET_{50}$ value for

MT_2 homodimers was 3 - 4 times higher (Tab. 2). In all cases, total receptor densities at the

$BRET_{50}$ were in the range of 3000 – 10000 receptors/cell or 30 – 100 fmol/mg of protein

(Tab. 1), which corresponds to physiological values for melatonin receptors in tissues
(Dubocovich and Takahashi, 1987; Morgan et al., 1994; Paul et al., 1999). Estimating an
average cell surface of 240 μm^2 for HEK 293 cells, the density of dimers at the cell surface
would be comprised between 10 and 50 receptors/ μm^2, a value that is at least 100 times
lower than that promoting non-specific BRET in HEK 293 cells (Mercier et al., 2002). Taken
together, these data support the hypothesis that MT_1/MT_2 heterodimers may form at low
expression levels, and indicate, in addition, that in cells, which co-express both receptor
isoforms, the formation of MT_1/MT_2 heterodimers is even more probable than that of MT_2
homodimers.

Pharmacological Properties of Co-expressed MT_1 and MT_2 Receptors. To identify
unambiguously MT_1/MT_2 heterodimers in tissues, it is necessary to characterize the specific
binding properties of heterodimers versus those of homodimers. Such specific
pharmacological properties have been documented for some GPCR heterodimers but not for
others (Jordan and Devi, 1999; Pfeiffer et al., 2001; Rocheville et al., 2000a). Binding
experiments with ^{125}I-MLT as radioligand were performed on cells expressing MT_1-Rluc and
MT_2-YFP receptors either separately or in combination at a 1:1 protein ratio. The expression
level of these receptors was monitored by measuring either luciferase activity or YFP
fluorescence using calibration curves, which correlate luminescence and fluorescence signals
to the number of ligand binding sites (Fig. 1 of supplemental material). Both MT_1 and MT_2
receptors bound the specific agonist ^{125}I-MLT with high affinity (K_d = 115 ± 22 pM and 250
± 60 pM for MT_1-Rluc and MT_2-YFP, respectively). When receptors were co-expressed, the
apparent K_d was similar (200 pM ± 21). The pharmacological profile of melatonin receptors
was then determined in ^{125}I-MLT competition binding experiments. When expressed
individually, MT_1-Rluc and MT_2-YFP displayed K_i values and pharmacological profiles very

similar to those reported for the corresponding wild-type receptors (Audinot et al., 2003;

Dubocovich et al., 1997; Petit et al., 1999) indicating that Rluc and YFP moieties did not

significantly affect receptor binding properties (Tab. 1 and Fig. 3 of supplemental material).

In cells co-expressing MT_1-Rluc and MT_2-YFP at a 1:1 protein ratio competition curves for

melatonin, S20098, S22153, S20928 and luzindole were monophasic with K_i values

comparable to those observed for cells expressing each receptor separately. The competition

profiles of the MT_2-selective ligands 4P-PDOT and S24773 were biphasic with K_i values

consistent with the binding to MT_1 and MT_2 binding sites. These data may be interpreted in

different ways. According to the results obtained with the BRET donor saturation assay (see

Table 2) the absence of MT_1/MT_2 heterodimers can be excluded since this receptor species

represents a major receptor fraction in cells co-expressing MT_1 and MT_2 receptors at a 1:1

protein ratio. We can also exclude that MT_1/MT_2 heterodimers are unable to bind ^{125}I-MLT

and that the ligand binding profile observed in cells co-expressing both receptors would

correspond to the sum of competition profiles of co-existing MT_1 and MT_2 homodimers.

Indeed, no decrease in ^{125}I-MLT binding has been observed in cells co-expressing both

receptors compared to cells expressing equivalent amounts of both receptors individually

(quantified by fluorescence/ luminescence measurements), as would be expected if the

heterodimer is unable to bind ^{125}I-MLT (not shown). In addition, the effect of ligands on the

BRET signal presented below shows that MT_1/MT_2 heterodimers are ligand binding

competent. Having excluded these possibilities, the competition profiles in cells co-expressing

MT_1 and MT_2 receptors, may be either explained by the fact that the affinity of MT_1 and MT_2

binding sites for the ligand is identical whether they are part of an homodimer or of an

heterodimer or by the fact that existing differences in ligand binding properties are not

revealed in this assay due to the superposition of multiple competition profiles caused by the

different coexisting receptor species (monomers, homo- and heterodimers). To discriminate

between these possibilities and to identify unambiguously the ligand binding properties of

MT_1/MT_2 heterodimers, we developed an alternative approach.

Correlation between ligand affinity and ligand-induced changes of BRET. A direct

consequence of ligand binding to receptors is the induction of conformational changes within

the core of the helical transmembrane domain that may be monitored with the BRET assay.

For the MT_2 homodimer and the MT_1/MT_2 heterodimer, ligand-promoted modifications of

BRET signals can indeed be observed in the presence of agonists and inverse agonists.

Changes of the BRET signal are most likely induced by the conformational change of the

receptor and does not results from dimer recruitment, receptor redistribution or alterations in

local pH (a parameter that could influence energy transfer efficacy) (Ayoub et al., 2002).

Importantly, the change of BRET signals upon ligand binding can be attributed to a specific

receptor dimer since the energy transfer occurs only between BRET-competent receptors. We

first verified whether the efficiency to promote ligand-induced BRET signals (EC_{50}),

correlates with binding affinities of the ligands. A good correlation would be expected for

receptor homodimers such as the MT_2 homodimer. To test this prediction, MT_2-Rluc and

MT_2-YFP fusion proteins were co-expressed at a 1:3 ratio that corresponds to the optimized

condition for BRET measurements (Ayoub et al., 2002). K_i values were determined in [125]I-

MLT competition binding experiments for selected ligands and were shown to be similar to

those observed in cells expressing MT_2-YFP alone (compare Tab. 1 and Tab. 3). The same

compounds increased the BRET signal in cells expressing MT_2 homodimers in a dose-

dependent manner with maximal BRET values ranging between 115 and 175 % of the basal

BRET (Fig. 3a). The rank order of potency of the ligands was the similar in the BRET assay

and the ^{125}I-MLT competition binding assay (K_i : 2-iodomelatonin = 20098 > melatonin = 4P-

PDOT > S22153 = S24773 = luzindole > S20928; EC_{50} : 2-iodomelatonin ≥ S20098 =

melatonin = S24773 = 4P-PDOT = S22153 > luzindole > S20928). A good correlation was

obtained when EC_{50} values were plotted against the corresponding K_i values (linear

regression, R^2=0.74) (Fig. 3b) indicating that the efficiency of a ligand to promote BRET

changes within dimers is correlated with its affinity for the receptor.

Assessment of ligand-promoted BRET changes of the MT_1/MT_2 heterodimer.

Similar experiments were conducted in cells expressing MT_1-Rluc/ MT_2-YFP heterodimers

(at a 1:3 protein ratio) for a panel of ligands (Tab. 4). Competition binding curves were

monophasic and K_i values close to those measured in cells expressing MT_2-Rluc and MT_2-

YFP. Again a dose-dependent ligand-induced BRET was observed for all compounds tested

with maximal values ranging from 130 to 140 % of the basal BRET (Fig. 4). However, no

correlation could be established between EC_{50} and K_i values (R^2=0.02) (Fig. 5a) indicating

that the efficiency of a ligand to promote BRET changes specifically within the MT_1/MT_2

heterodimer does not correlate with the apparent affinity constant measured in cells co-

expressing MT_1 and MT_2 receptors. Similarly no correlation was observed when EC_{50} values

of non-selective and MT_1-selective ligands were plotted against K_i values of the MT_1 receptor

(here shown for MT_1-Rluc) (Fig. 5b) or when EC_{50} values of non-selective and MT_2-selective

ligands were plotted against K_i values of the MT_2 receptor (here shown for MT_2-YFP)

(Fig.5c). This indicates that the binding properties of the MT_1 and MT_2 binding site in the

heterodimer are different from those detected in the corresponding homodimers. Further

evidence for this hypothesis comes from the comparison between BRET EC_{50} values of MT_2

homodimers and MT_1/ MT_2 heterodimers (Fig. 5d). The absence of correlation suggests that

ligand-promoted conformational changes of MT_1/MT_2 heterodimers differ from those elicited

in MT_2 homodimers and indicate the existence of MT_1/MT_2 heterodimer-specific ligand

binding properties. Heterodimer selectivity of ligands can be estimated by the ratio of EC_{50}

values for homo- and heterodimers (Tab. 4). Whereas melatonin and S20098 are equally

potent, EC_{50} values for S22153 and S24773 are eight times lower for the heterodimer. S20928

and luzindole are clearly more potent on MT_1/MT_2 heterodimers compared to MT_2 homodimers (26 and 126 times, respectively) and EC_{50} values for 4P-PDOT are 5 times lower for the MT_2 homodimer. Taken together these results show that melatonin receptor-specific ligands are binding to MT_1/MT_2 heterodimers and that the potency of these ligands to induce conformational changes is similar for some ligands (melatonin, S20098) but clearly different for others (S20928, luzindole) compared to those measured for MT_2 homodimers.

Both ligand binding sites are functional within the MT_1/MT_2 heterodimer. In cells co-expressing MT_1 and MT_2 receptors, ^{125}I-MLT competition binding curves for the MT_2– selective compounds S24773 and 4P-PDOT were biphasic with a MT_2 binding site of high affinity and a MT_1 binding site with 30-100 times lower affinity (see Tab. 1 and Fig. 3 of supplemental material). Accordingly, concentration-response curves of the ligand-induced BRET are expected to be biphasic for these compounds in cells expressing MT_1/MT_2 heterodimers. However, experimental BRET curves were monophasic for these compounds and EC_{50} values corresponded to the affinity for MT_2 (see Fig. 4) indicating that the ligand-promoted BRET change is due to binding to the MT_2 binding site at this ligand concentration. The absence of the second (low-affinity) component of the BRET curves might be explained by the absence of the second (MT_1-like) functional binding site in the heterodimer. To address this point, we studied the effect of the MT_1-selective ligand S26284 (Audinot et al., 2003). The BRET dose-response curve of this compound for the MT_1/MT_2 heterodimer was monophasic with an EC_{50} of 48 ± 3 nM (Fig. 6), which is close to the K_i value measured for MT_1-Rluc (47 ± 5 nM, n=2) compared to the K_i value for MT_2-YFP (605 ± 420 nM, n=2). Thus S26284 bound to the MT_1-like binding site within the heterodimer and induced conformational changes that decrease of the basal BRET signal by either increasing the distance between the BRET partners or promoting a less favorable orientation of the two partners. Binding of S26284 to the MT_1-like binding site within the heterodimer is further

supported by the fact that no BRET change can be observed upon S26284 stimulation in cells expressing MT_2-YFP (Fig. 6). Taken together these data indicate that both the MT_1- and the MT_2-like binding site within the heterodimer are functional and shows that ligand binding to either of the two binding sites of the heterodimer is sufficient to induce a conformational change within the heterodimer.

DISCUSSION

In the present study, we have shown that the probability of MT_1/MT_2 heterodimer formation is similar or even higher than those of the corresponding homodimers in cells expressing low levels of receptor and that heterodimers are competent for binding ligands. Both, the MT_1- and MT_2-binding site are functional within the heterodimer. We have also shown that the two binding sites maintain their respective selectivity for MT_1- and MT_2 selective ligands and that the ligand interaction profile of the MT_1/MT_2 heterodimer determined by BRET is not identical to that of the MT_2 homodimer.

The homo-/ heterodimer ratio of MT_1 and MT_2 receptors has been determined with the BRET donor saturation assay. The engagement of MT_1 receptors into MT_1 homodimers or MT_1/MT_2 heterodimers appears to be governed exclusively by the relative expression levels of MT_1 and MT_2 receptors since similar relative affinities were observed in BRET saturation assays. In contrast, MT_2 receptors have a higher tendency to form heterodimers with MT_1 receptors than to form homodimers suggesting that MT_2 receptors may be preferentially engaged into heterodimers in cells co-expressing both receptors. The documented co-expression of MT_1 and MT_2 receptors in many melatonin sensitive tissues, such as the hypothalamic suprachiasmatic nuclei (Reppert et al., 1988), the retina (Dubocovich, 1983), arteries (Krause et al., 1995) and adipose tissue (Brydon et al., 2001) suggests that

heterodimerization could indeed occur in native mammalian tissues assuming simultaneous expression of both receptors in the same cells. Our results indicate, for the first time, that homo- and heterodimer formation may not only depend on the relative expression levels of receptor subtypes but also on the relative affinity of the monomers for each other. For the γ-aminobutyric acid (GABA) receptor B (GABA$_B$R), heterodimerization between GABA$_{B1}$ and GABA$_{B2}$ subunits was shown to be obligatory for the formation of functional receptors (Jones et al., 1998). Expression of each subunit alone does not form functional receptors. However, most GPCR clearly form functional homodimeric receptors when expressed alone. As shown for the MT$_2$ receptor, some receptors may form homodimers but preferentially engage into heterodimers. This may also be the case for the α_{1D}-adrenergic receptor (α_{1D}-AR) (Hague et al., 2004). This receptor forms homodimers that accumulate intracellularly when expressed alone. Co-expression of α_{1D}-AR with α_{1B}-AR caused heterodimer formation and the quantitative translocation of the α_{1D}-AR to the cell surface. A large spectrum of affinities is likely to exist for the formation of different GPCR heterodimers. Such an affinity spectrum may provide a framework for a better understanding of the formation of homo- and heterodimeric complexes in cells, which naturally co-express several different GPCRs.

The determination of the specific pharmacological profile of heterodimers is difficult to achieve using classical radioligand competition binding assays, in particular for heterodimers composed of two receptor subtypes (MT$_1$ and MT$_2$), which display similar affinities for the same radioligand (^{125}I-MLT). The major difficulty resides in the simultaneous detection of all ligand binding-competent receptor species (monomers, homo- and heterodimers). In contrast, energy transfer assays such as the BRET assay, have the unique feature to focus on one given combination of receptors (those competent for BRET). We have shown that the efficiency for a ligand to induce conformational changes in a homodimer reflects its affinity for the receptor

as observed in cells co-expressing MT_2-Rluc and MT_2-YFP fusion proteins (MT_2

homodimer). Then, the efficiency of ligands to induce conformational changes was compared

between MT_2 homodimers and MT_1/MT_2 heterodimers. MT_1 homodimers could not be

studied directly because the ligand-induced conformational change does not translate into

alterations of the BRET signal for this specific subtype (Ayoub et al., 2002). Some ligands,

including the natural hormone melatonin, showed similar efficiencies to induce BRET

changes in MT_2 homodimers and heterodimers, whereas several synthetic compounds

(S20928, luzindole, S26284) caused clearly different effects on homo- and heterodimers

showing that the efficiency to promote ligand-induced conformational changes of MT_1/MT_2

heterodimers differs from that of MT_2 homodimers.

Subtype selective ligands are frequently used to define the specific melatonin receptor

subtype involved in the physiological effects of melatonin (Masana and Dubocovich, 2001).

We have shown that both the MT_2-selective 4P-PDOT and S24773, and the MT_1-selective

S26284 bind also with high affinity to MT_1/MT_2 heterodimers. This may have important

implications for the interpretation of data obtained in cells co-expressing the two melatonin

receptor subtypes since these compounds will not only bind with high affinity to the selected

homodimer but also to the MT_1/MT_2 heterodimer.

GPCR dimers are potentially composed of two functional ligand binding sites. Whether

both sites are indeed functional and whether ligand binding to both sites is necessary for

receptor activation are critical questions to understand the activation mechanism of GPCRs.

^{125}I-MLT binding and BRET experiments with subtype-selective ligands in cells co-

expressing MT_1 and MT_2 receptors suggested that MT_1/MT_2 heterodimers are composed of

two functional ligand binding sites with distinct properties, a MT_1-like binding site and a

MT_2-like binding site. The conservation of two ligand binding sites within GPCR dimers

showing variable changes of the pharmacological properties, have also been observed for

other heterodimers (δ and κ opioid (Jordan and Devi, 1999), μ and δ opioid (George et al.,

2000), somatostatin sst1 and sst5 receptors (Patel et al., 2002), adenosine A1 and dopamine

D1 (Ferre et al., 1998), somatostatin sst5 and dopamine D2 (Rocheville et al., 2000a)).

Whether ligand binding to both sites is necessary for receptor activation has been a

difficult issue to address. Early studies on the $GABA_BR$, which forms obligatory heterodimers

between the $GABA_{B1}$ and $GABA_{B2}$ subunits, showed that ligand binding to the $GABA_{B1}$

subunit is sufficient to promote G protein trans-activation through the $GABA_{B2}$ subunit

(Kniazeff et al., 2002). Further evidence comes from internalization studies of somatostatin

receptor sst1/ sst5 heterodimers. Whereas sst5 receptors can be internalized, sst1 receptors are

unable to be internalized when expressed alone. However, following binding of a sst1

selective ligand to the sst1/ sst5 heterodimer, the sst1 receptor was reported to be internalized

(Rocheville et al., 2000b) indicating that ligand binding to one monomeric unit (sst1) of the

heterodimer is sufficient to induce the conformational change and receptor internalization of

the dimer. Similar observations were made for the internalization of the V1a/V2 vasopressin

receptor heterodimer where the fate of the internalized heterodimer depends on the activation

of the specific monomeric unit within the heterodimer (Terrillon et al., 2004). Similarly,

induction of conformational changes upon binding of subtype-selective ligands to either MT_1-

or MT_2-like binding sites of the MT_1/MT_2 heterodimer is also consistent with the model that

occupation of only one ligand binding site within the dimer may be sufficient for receptor

activation.

Although ligand-induced conformational changes are supposed to be a general

phenomenon, ligand-induced energy transfer has not been observed for all receptors studied

so far (Issafras et al., 2002; Terrillon et al., 2003) indicating that conformational changes do

not always result into energy transfer variations. Indeed, the ligand-induced conformational change within the receptor moiety may have little effect on the position and the orientation of the energy transfer partners. Data available so for indicate that the development of energy transfer assays as a conformational sensors needs some optimization of the assay conditions and of the fusion protein design (Boute et al., 2001; Couturier and Jockers, 2003). In this respect, the introduction of energy donors and acceptors at sites other than the C terminus of GPCRs may represent an interesting alternative as reported for the receptor "chameleon" constructs carrying CFP and YFP respectively, in the third intracellular loop and after the C terminus of the parathyroid hormone and alpha$_{2A}$ adrenergic receptors. These constructs were still functional and highly sensitive to ligand-induced conformational changes, in agreement with the predicted movement of the third intracellular loop away from the C terminus (Vilardaga et al., 2003). Ligand-promoted BRET changes were also observed for tyrosine kinase receptors (Boute et al., 2001) and cytokine receptors (Couturier and Jockers, 2003) demonstrating the general interest of BRET/FRET approaches to monitor ligand-induced conformational changes.

In conclusion, we investigated melatonin receptor heterodimerization using the BRET technology. The relative propensity for melatonin receptor homo- and heterodimer formation was determined in a BRET donor saturation assay and showed that MT$_1$/MT$_2$ heterodimers are formed at low expression levels and at equal or higher probability than the corresponding homodimers. We have shown that the efficiency to promote ligand-induced variations of the BRET signal correlates with the binding affinities of ligands to the receptor. Based on this correlation, we developed a BRET-based approach to study the specific ligand binding properties of MT$_1$/MT$_2$ heterodimers. This approach may be potentially applied to a wide range of ligand-regulated receptors.

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REFERENCES

Angers, S., Salahpour, A., Joly, E., Hilairet, S., Chelsky, D., Dennis, M., and Bouvier, M. (2000). Detection of b2-adrenergic receptor dimerization in living cells using bioluminescence resonance energy transfert (BRET). Proc Natl Acad Sci U S A *97*, 3684-3689.

Audinot, V., Mailliet, F., Lahaye-Brasseur, C., Bonnaud, A., Le Gall, A., Amosse, C., Dromaint, S., Rodriguez, M., Nagel, N., Galizzi, J. P., Malpaux, B., Guillaumet, G., Lesieur, D., Lefoulon, F., Renard, P., Delagrange, P., and Boutin, J. A. (2003). New selective ligands of human cloned melatonin MT1 and MT2 receptors. Naunyn Schmiedebergs Arch Pharmacol *367*, 553-561.

Ayoub, M. A., Couturier, C., Lucas-Meunier, E., Angers, S., Fossier, P., Bouvier, M., and Jockers, R. (2002). Monitoring of ligand-independent dimerization and ligand-induced conformational changes of melatonin receptors in living cells by bioluminescence resonance energy transfer. J Biol Chem *277*, 21522-21528.

Boute, N., Pernet, K., and Issad, T. (2001). Monitoring the activation state of the insulin receptor using bioluminescence resonance energy transfer. Mol Pharmacol *60*, 640-645.

Brydon, L., Barrett, P., Morgan, P. J., Strosberg, A. D., and Jockers, R. (1999a). Investigation of the human Mel 1a melatonin receptor using anti-receptor antibodies. Adv Exp Med Biol *460*, 215-220.

Brydon, L., Petit, L., Delagrange, P., Strosberg, A. D., and Jockers, R. (2001). Functional expression of MT2 (Mel1b) melatonin receptors in human PAZ6 adipocytes. Endocrinology *142*, 4264-4271.

Brydon, L., Roka, F., Petit, L., de Coppet, P., Tissot, M., Barrett, P., Morgan, P. J., Nanoff, C., Strosberg, A. D., and Jockers, R. (1999b). Dual signaling of human Mel1a melatonin receptors via G(i2), G(i3), and G(q/11) proteins. Mol Endocrinol *13*, 2025-2038.

Couturier, C., and Jockers, R. (2003). Activation of the Leptin Receptor by a Ligand-induced Conformational Change of Constitutive Receptor Dimers. J Biol Chem *278*, 26604-26611.

Dubocovich, M. L. (1983). Nature *306*, 782-784.

Dubocovich, M. L., Masana, M. I., Iacob, S., and Sauri, D. M. (1997). Melatonin receptor antagonists that differentiate between the human Mel1a and Mel1b recombinant subtypes are used to assess the pharmacological profile of the rabbit retina ML1 presynaptic heteroreceptor. Naunyn Schmiedebergs Arch Pharmacol *355*, 365-375.

Dubocovich, M. L., and Takahashi, J. S. (1987). Use of 2-[125I]iodomelatonin to characterize melatonin binding sites in chicken retina. Proc Natl Acad Sci U S A *84*, 3916-3920.

Farrens, D. L., Altenbach, C., Yang, K., Hubbell, W. L., and Khorana, H. G. (1996). Requirement of rigid-body motion of transmembrane helices for light activation of rhodopsin. Science *274*, 768-770.

Ferre, S., Torvinen, M., Antoniou, K., Irenius, E., Civelli, O., Arenas, E., Fredholm, B. B., and Fuxe, K. (1998). Adenosine A1 receptor-mediated modulation of dopamine D1 receptors in stably cotransfected fibroblast cells. J Biol Chem *273*, 4718-4724.

Gazi, L., Lopez-Gimenez, J. F., and Strange, P. G. (2002). Formation of oligomers by G protein-coupled receptors. Curr Opin Drug Discov Devel *5*, 756-763.

George, S. R., Fan, T., Xie, Z., Tse, R., Tam, V., Varghese, G., and O'Dowd, B. F. (2000). Oligomerization of mu- and delta-opioid receptors. Generation of novel functional properties. J Biol Chem *275*, 26128-26135.

Ghanouni, P., Steenhuis, J. J., Farrens, D. L., and Kobilka, B. K. (2001). Agonist-induced conformational changes in the G-protein-coupling domain of the beta 2 adrenergic receptor. Proc Natl Acad Sci U S A *98*, 5997-6002.

Hague, C., Uberti, M. A., Chen, Z., Hall, R. A., and Minneman, K. P. (2004). Cell surface expression of alpha 1D-adrenergic receptors is controlled by heterodimerization with alpha 1B-adrenergic receptors. J Biol Chem.

Heyduk, T. (2002). Measuring protein conformational changes by FRET/LRET. Curr Opin Biotechnol *13*, 292-296.

Issafras, H., Angers, S., Bulenger, S., Blanpain, C., Parmentier, M., Labbe-Jullie, C., Bouvier, M., and Marullo, S. (2002). Constitutive agonist-independent CCR5 oligomerization and antibody-mediated clustering occurring at physiological levels of receptors. J Biol Chem *277*, 34666-34673.

Jones, K. A., Borowsky, B., Tamm, J. A., and Gerald, C. (1998). GABAb receptors function as a heterotrimeric assembly of the subunits GABAbR1 and GAGbR2. Nature *396*, 674-679.

Jordan, B. A., and Devi, L. A. (1999). G-protein-coupled receptor heterodimerization modulates receptor function. Nature *399*, 697-700.

Kenakin, T., and Onaran, O. (2002). The ligand paradox between affinity and efficacy: can you be there and not make a difference? Trends Pharmacol Sci *23*, 275-280.

Kniazeff, J., Galvez, T., Labesse, G., and Pin, J. P. (2002). No ligand binding in the GB2 subunit of the GABA(B) receptor is required for activation and allosteric interaction between the subunits. J Neurosci *22*, 7352-7361.

Krause, D. N., Barrios, V. E., and Duckles, S. P. (1995). Melatonin receptors mediate potentiation of contractile responses to adrenergic nerve stimulation in rat caudal artery. Eur J Pharmacol *276*, 207-213.

Kroeger, K. M., Hanyaloglu, A. C., Seeber, R. M., Miles, L. E., and Eidne, K. A. (2001). Constitutive and agonist-dependent homo-oligomerization of the thyrotropin-releasing hormone receptor. Detection in living cells using bioluminescence resonance energy transfer. J Biol Chem *276*, 12736-12743.

Lee, J., Pilch, P. F., Shoelson, S. E., and Scarlata, S. F. (1997). Conformational changes of the insulin receptor upon insulin binding and activation as monitored by fluorescence spectroscopy. Biochemistry *36*, 2701-2708.

Masana, M. I., and Dubocovich, M. L. (2001). Melatonin receptor signaling: finding the path through the dark. Sci STKE *2001*, E39.

McVey, M., Ramsay, D., Kellett, E., Rees, S., Wilson, S., Pope, A. J., and Milligan, G. (2001). Monitoring receptor oligomerization using time-resolved fluorescence resonance energy transfer and bioluminescence resonance energy transfer. The human delta -opioid receptor displays constitutive oligomerization at the cell surface, which is not regulated by receptor occupancy. J Biol Chem *276*, 14092-14099.

Mercier, J. F., Salahpour, A., Angers, S., Breit, A., and Bouvier, M. (2002). Quantitative assessment of beta 1- and beta 2-adrenergic receptor homo- and heterodimerization by bioluminescence resonance energy transfer. J Biol Chem *277*, 44925-44931.

Morgan, P. J., Barrett, P., Howell, H. E., and Helliwell, R. (1994). Melatonin receptors: localization, molecular pharmacology and physiological significance. Neurochem Int *24*, 101-146.

Patel, R. C., Kumar, U., Lamb, D. C., Eid, J. S., Rocheville, M., Grant, M., Rani, A., Hazlett, T., Patel, S. C., Gratton, E., and Patel, Y. C. (2002). Ligand binding to somatostatin receptors induces receptor-specific oligomer formation in live cells. Proc Natl Acad Sci U S A *99*, 3294-3299.

Paul, P., Lahaye, C., Delagrange, P., Nicolas, J. P., Canet, E., and Boutin, J. A. (1999). Characterization of 2-[125I]iodomelatonin binding sites in Syrian hamster peripheral organs. J Pharmacol Exp Ther *290*, 334-340.

Petit, L., Lacroix, I., de Coppet, P., Strosberg, A. D., and Jockers, R. (1999). Differential signaling of human Mel1a and Mel1b melatonin receptors through the cyclic guanosine 3'-5'-monophosphate pathway. Biochem Pharmacol *58*, 633-639.

Pfeiffer, M., Koch, T., Schroder, H., Klutzny, M., Kirscht, S., Kreienkamp, H. J., Hollt, V., and Schulz, S. (2001). Homo- and heterodimerization of somatostatin receptor subtypes. Inactivation of sst(3) receptor function by heterodimerization with sst(2A). J Biol Chem *276*, 14027-14036.

Reppert, S. M., Weaver, D. R., Rivkees, S. A., and Stopa, E. G. (1988). Putative melatonin receptors in human biological clock. Science *242*, 78-81.

Rocheville, M., Lange, D. C., Kumar, U., Patel, S. C., Patel, R. C., and Patel, Y. C. (2000a). Receptors for Dopamine and Somatostatin: Formation of Hetero-Oligomers with Enhanced Functional Activity. Science *288*, 154-157.

Rocheville, M., Lange, D. C., Kumar, U., Sasi, R., Patel, R. C., and Patel, Y. C. (2000b). Subtypes of the Somatostatin Receptor Assemble as Functional Homo- and Heterodimers. J Biol Chem *275*, 7862-7869.

Roka, F., Brydon, L., Waldhoer, M., Strosberg, A. D., Freissmuth, M., Jockers, R., and Nanoff, C. (1999). Tight association of the human Mel(1a)-melatonin receptor and G(i): precoupling and constitutive activity. Mol Pharmacol *56*, 1014-1024.

Terrillon, S., Barberis, C., and Bouvier, M. (2004). Heterodimerization of V1a and V2 vasopressin receptors determines the interaction with {beta}-arrestin and their trafficking patterns. Proc Natl Acad Sci U S A.

Terrillon, S., Durroux, T., Mouillac, B., Breit, A., Ayoub, M. A., Taulan, M., Jockers, R., Barberis, C., and Bouvier, M. (2003). Oxytocin and Vasopressin V1a and V2 Receptors Form Constitutive Homo- and Heterodimers during Biosynthesis. Mol Endocrinol *17*, 677-691.

Truong, K., and Ikura, M. (2001). The use of FRET imaging microscopy to detect protein-

protein interactions and protein conformational changes in vivo. Curr Opin Struct Biol *11*,

573-578.

Vilardaga, J. P., Bunemann, M., Krasel, C., Castro, M., and Lohse, M. J. (2003).

Measurement of the millisecond activation switch of G protein-coupled receptors in living

cells. Nat Biotechnol *21*, 807-812.

Footnotes

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Legends to figures:

Figure 1 : Detection of MT$_1$/MT$_2$ heterodimers. *A,* HEK 293 cells stably expressing Myc-MT$_2$ were transiently transfected or not with the Flag-MT$_1$ construct and crude membranes prepared. Receptors were immunoprecipitated with a monoclonal anti-Myc antibody as described in "Experimental Procedure". Membranes and immunoprecipitates were then submitted to SDS-PAGE and revealed by Western blot analysis using a polyclonal anti-Flag antibody. ***B,*** The indicated Rluc and YFP fusion proteins were expressed at a 1:10 protein ratio in HEK 293 cells as determined using standard curves correlating ^{125}I-MLT binding sites with luminescence or YFP fluorescence (Fig. 1 of the supplemental material). Energy transfer measurements were performed in living cells by adding coelenterazine and measuring light emission in a luminometer with Rluc and YFP filter settings as described in the "Experimental Procedure" section. Data are means ± S.E.M. of at least three independent experiments each performed in duplicate.

Figure 2 : BRET donor saturation curves of MT$_1$ and MT$_2$ homo- and heterodimers.
BRET measurements were performed with HEK 293 cells co-expressing 12 fmol/mg of protein of the indicated Rluc fusion proteins and increasing amounts of the indicated YFP fusion proteins. BRET values were plotted as a function of the ratio of YFP/Rluc fusion proteins (as determined by transforming luminescence and fluorescence values measured for each data point into receptor numbers by using the correlation curves shown in Fig. 1 of the supplemental material). The curves represent 7 - 10 individual normalized saturation curves that were fitted using a non-linear regression equation assuming a single binding site (GraphPad Prism).

Figure 3 : Dose-response curves of the ligand-induced BRET in MT$_2$ homodimers

A, HEK 293 cells co-expressing MT$_2$-Rluc and MT$_2$-YFP receptors at a 1:3 protein ratio (as determined using the correlation curves shown in Fig. 1 of the supplemental material) were incubated in the presence of increasing concentrations of the indicated ligands and BRET measurements performed according to the "Microplate BRET assay" protocol. Data are represented as % of energy transfer in the absence of ligand and are representative of at least three independent experiments each performed in triplicate. Curves were analyzed by non-linear regression (GraphPad Prism). **B,** Correlation between K$_i$ and EC$_{50}$ values of the MT$_2$ homodimer. Data were fitted using a linear regression equation (GraphPad Prism). Ligands have been numbered according to the numbers used in Table 3.

Figure 4 : Dose-response curves of the ligand-induced BRET in MT$_1$/MT$_2$ heterodimers

A, HEK 293 cells co-expressing MT$_1$-Rluc and MT$_2$-YFP receptors at a 1:3 protein ratio (as determined using the correlation curves shown in Fig. 1 of the supplemental material) were incubated in the presence of increasing concentrations of the indicated ligands and BRET measurements performed according to the "Microplate BRET assay" protocol. Data are represented as % of maximally induced BRET and are representative of at least three experiments each performed in triplicate. Curves were analyzed by non-linear regression (GraphPad Prism).

Figure 5 : Correlations between EC$_{50}$ values of MT$_1$/MT$_2$ heterodimers and K$_i$ values of MT$_1$ and MT$_2$ receptors. EC$_{50}$ values of the MT$_1$/MT$_2$ heterodimer shown in Tab. 4 were plotted against K$_i$ values determined in cells co-expressing MT$_1$ and MT$_2$ receptors (**A**) or expressing MT$_1$-Rluc (**B**) or MT2-YFP (**C**) receptors individually. **D,** Correlation between EC$_{50}$ values of the MT$_2$ homodimer and the MT$_1$/MT$_2$ heterodimer. Ligands have been

numbered according to the numbers used in Table 3. In panel B and C, non-selective and

selective compounds are represented by filled and open circles, respectively. Ligand 9 shown

in panel B corresponds to the MT_1-selective S26284 compound. Data were fitted using a

linear regression equation (GraphPad Prism).

Figure 6 : Ligand-dependent BRET changes of S26284 in cells expressing MT_1 and MT_2

homo- and heterodimers. HEK 293 cells co-expressing MT_1-Rluc and MT_2-YFP (□) or

MT_2-Rluc and MT_2-YFP (▲) receptors at a 1:3 protein ratio were incubated in the presence

of increasing concentrations of S26284 and BRET experiments performed as described under

"Experimental Procedure". Data representative of at least three independent experiments each

performed in triplicate. Curves were analyzed by non-linear regression (GraphPad Prism).

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TABLE 1

Binding affinities measured in HEK 293 cells expressing MT_1 and MT_2 receptors.

HEK 293 cells expressing MT_1-Rluc or MT_2-YFP or both together at a 1:1 ratio were incubated with ^{125}I-MLT and various concentrations of the indicated compounds. K_i values were calculated as described under "Experimental procedures ". Data are means ± S.E. of three independent experiments each performed in duplicate.

Ligands	K_i (nM)			Ratio K_i		
	MT_1-Rluc	MT_2-YFP	MT_1-Rluc + MT_2-YFP (Ratio 1:1)	$\dfrac{MT_1}{MT_2}$	$\dfrac{MT_1}{MT_1+MT_2}$	$\dfrac{MT_2}{MT_1+MT_2}$
1) - Melatonin	0.36 ± 0.1	0.48 ± 0.1	1.13 ± 0.5	0.7	0.3	0.4
2) - S20098	0.78 ± 0.2	0.08 ± 0.05	1.64 ± 1.5	9.7	0.5	0.05
3) - S22153	39.4 ± 9.0	13 ± 3	31.7 ± 8.9	3	1.2	0.4
4) - S20928	244 ± 105	210 ± 81	281 ± 114	1.2	0.9	0.7
5) - 4P-PDOT	53.5 ± 13	2 ± 0.3	1.40 ± 0.4	26	75	1.4
			70.3 ± 15 (2nd site)	-	0.7	0.02
6) - Luzindole	31.6 ± 7.0	12.3 ± 4.3	10 ± 1.7	2.6	3.2	1.2
7) - S24773	295 ± 26	3.70 ± 1.7	0.60 ± 0.1	80	491	6.2
			192 ± 23 (2nd site)	1.5	-	0.01

MOL # 398

TABLE 2

Relative affinities between two BRET partners of melatonin receptors.

The $BRET_{50}$ represents the acceptor/donnor ratio required to reach half-maximal BRET in BRET-donnor saturation experiments. Results are the mean + S.E. of 7-10 independent saturation curves (see Fig. 2). Receptor densities at $BRET_{50}$ are determined by averaging BRET values recorded close to the calculated $BRET_{50}$ of 7-10 individual saturation experiments. The luciferase activity and YFP fluorescence were used to calculated the number of ^{125}I-MLT binding sites according to standard curves shown in the Fig. 1 of the supplemental material. 7500 HEK 293 cells correspond to 1 µg of total protein. (** : $p < 0.02$, compared to conditions 1, 2 and 3).

Dimer	$BRET_{50}$	Receptor density at $BRET_{50}$	
		(Receptor/cell)	(fmoles/mg of protein)
1- MT_1-Rluc/MT_1-YFP	3.7 ± 0.8	3080 ± 520	33 ± 4
2- MT_1-Rluc/MT_2-YFP	3.0 ± 1.6	3434 ± 436	43 ± 5
3- MT_2-Rluc/MT_1-YFP	4.2 ± 1.6	4554 ± 974	51 ± 12
4- MT_2-Rluc/MT_2-YFP	$12.5 \pm 2.2^{**}$	$11318 \pm 2176^{**}$	$142 \pm 27^{**}$

TABLE 3

Binding affinities (K_i) and EC_{50} values of ligand-induced BRET for MT_2 homodimers in HEK 293 cells.

MT_2-Rluc and MT_2-YFP receptors were expressed at a 1:3 ratio (~80 fmol/mg of protein) and ^{125}I-MLT competition binding experiments and BRET measurements were performed as described under "Experimental procedures " and in Fig. 3. Data are means ± S.E. of at least three independent experiments each performed in triplicate.

Ligands	K_i (nM)	EC_{50} (nM)	Emax (%)
1) - Melatonin	0.27 ± 0.01	26 ± 7.75	173 ± 20
2) - S20098	0.06 ± 0.05	14.2 ± 13.6	143 ± 9
3) - S22153	3 ± 1.90	47 ± 24	147 ± 18
4) - S20928	60 ± 0.10	368 ± 234	126 ± 6
5) - 4P-PDOT	0.35 ± 0.31	40.4 ± 12	146 ± 3
6) - Luzindole	7.23 ± 4.5	126 ± 57.2	150 ± 8
7) - S24773	5.70 ± 1.7	32.7 ± 16.4	161 ± 6
8) - 2-iodomelatonin	0.16 ± 0.03	4.4 ± 1.95	113 ± 3

TABLE 4

Binding affinities (Ki) measured in HEK 293 cells co-expressing MT_1 and MT_2 receptors and EC50 values of ligand-induced BRET for MT_1/MT_2 heterodimers.

MT_1-Rluc and MT_2-YFP receptors were expressed at a 1:3 ratio (~80 fmol/mg of protein) and ^{125}I-MLT competition binding experiments and BRET measurements were performed as described under "Experimental procedures " and in Fig. 4. Data are means ± S.E. of at least three independent experiments each performed in triplicate.

Ligands	K_i (nM)	EC_{50} (nM)	Emax (%)	Ratio EC_{50} MT_2 homodimer / heterodimer
1) - Melatonin	0.40 ± 0.16	16.8 ± 9.7	142 ± 9	1.5
2) - S20098	0.23 ± 0.17	7.7 ± 4.6	136 ± 6	1.8
3) - S22153	8.87 ± 2.34	6.4 ± 5.6	132 ± 10	7.3
4) - S20928	116.6 ± 1.75	14 ± 9.5	136 ± 7	26.3
5) - 4P-PDOT	8.60 ± 7.40	211 ± 159	141 ± 4	0.2
6) - Luzindole	32.4 ± 9.20	1.0 ± 0.1	134 ± 13	126
7) - S24773	4.5 ± 1.0	4.1 ± 2.2	137 ± 4	8

Figure 1 :   **MOL # 398**

A)

IP : anti-Myc
Blot : anti-Flag

B)
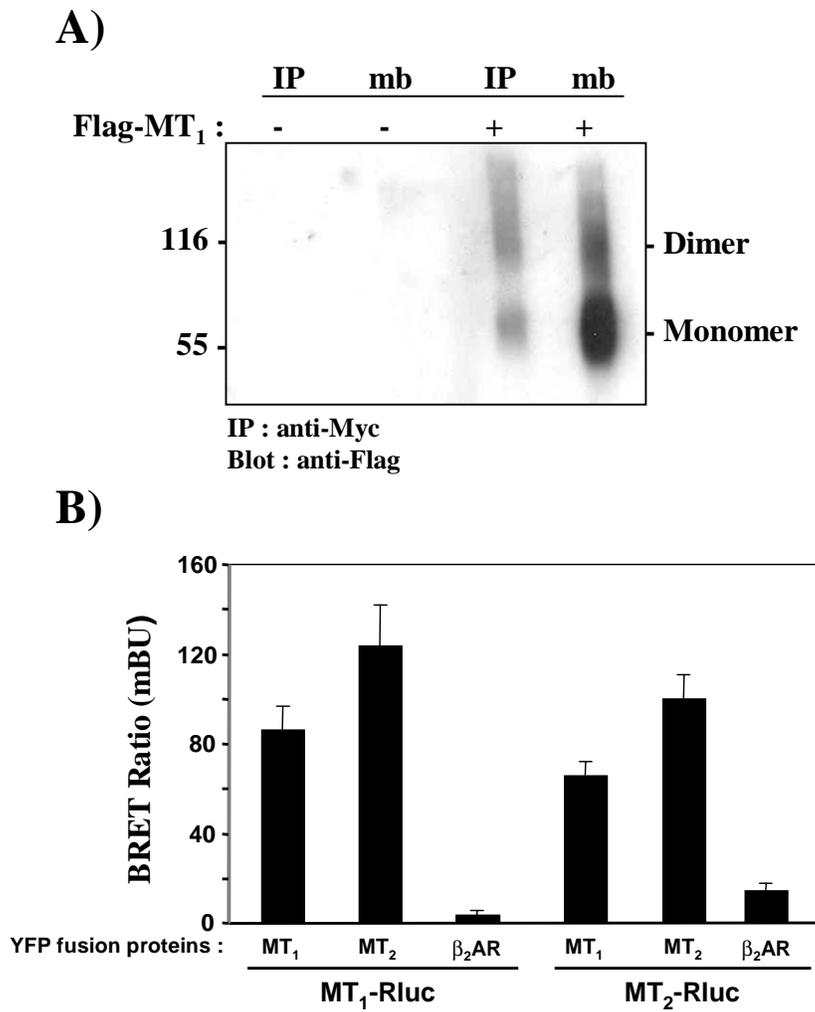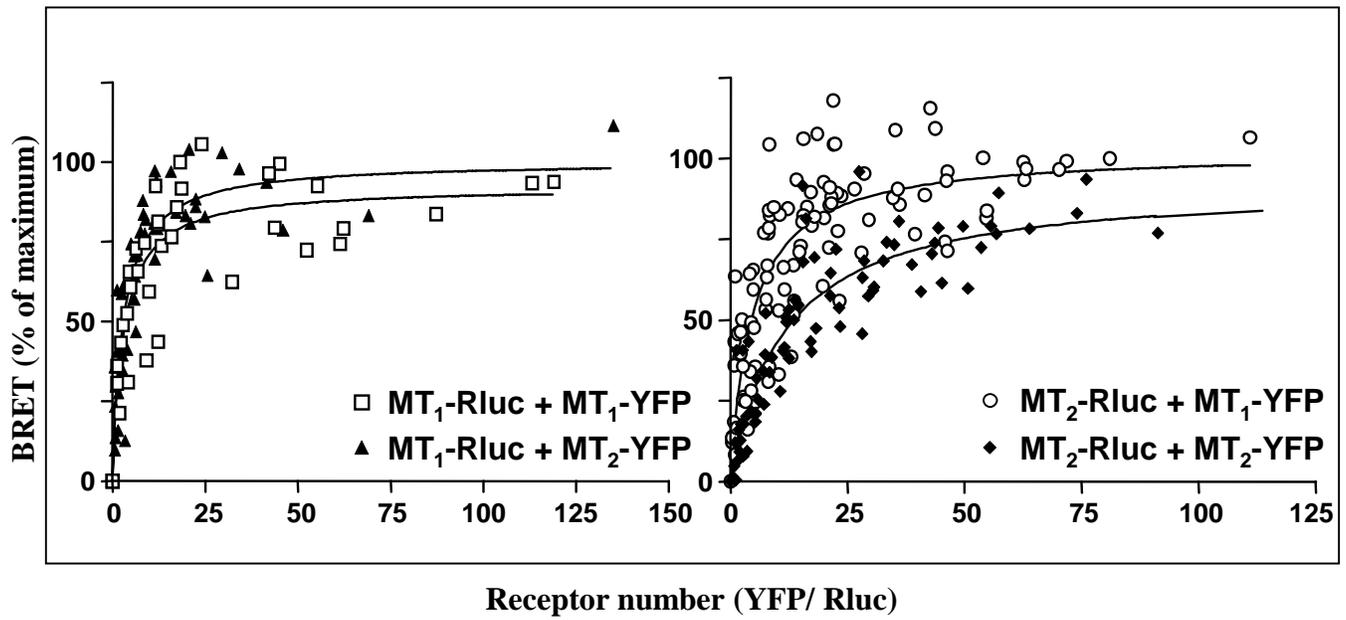
Figure 2 :

MOL # 398

Figure 3 :

MOL # 398

A)
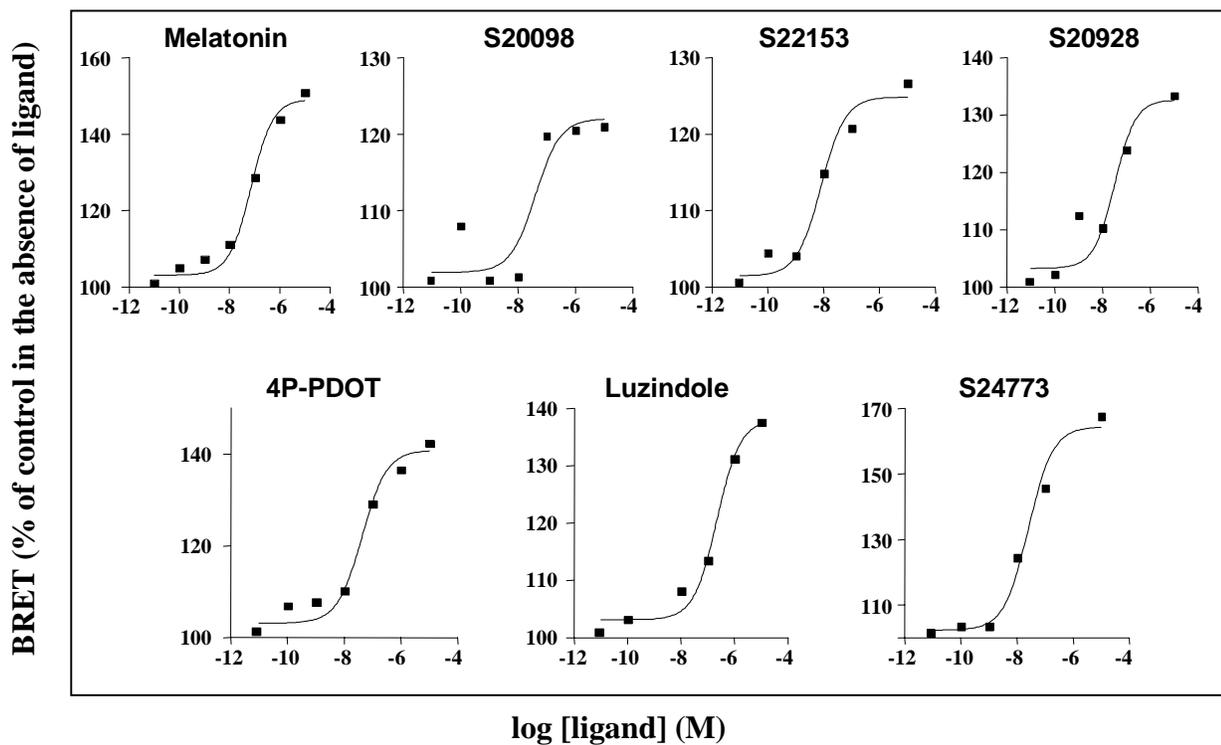
B)
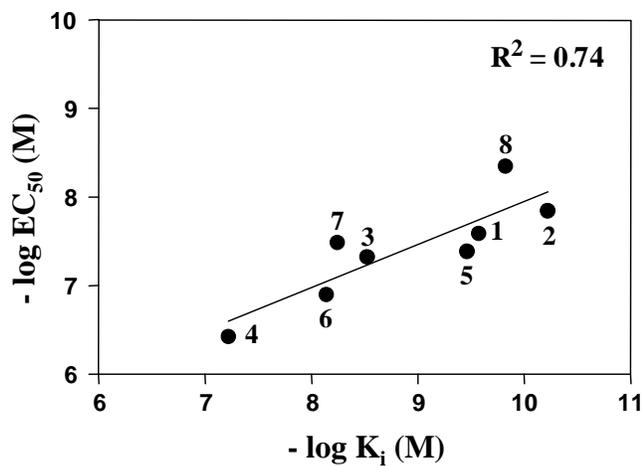
Figure 4 :

MOL # 398

Figure 5 :   **MOL # 398**

A)

B)

C)

D)

Figure 6 :   **MOL # 398**