Preferential formation of MT₁/MT₂ melatonin receptor heterodimers

with distinct ligand interaction properties compared to MT₂ 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 β₁ and β₂-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 alpha2A 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]cyclobutane-carboxamide), 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-([3-(3-aminophenyl)-7-methoxy-1-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-propionamidotetraline) from Tocris, Ellisville, MO.

Plasmid constructions, transfections and cell culture. Construction of Rluc and YFP fusion proteins and Flag-MT1 and Myc-MT2 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 $^2(125I)$-iodomelatonin ($^{125I}$-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 $^{125I}$-MLT and increasing concentrations of different compounds.
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. 1-2x10$^5$ 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™).
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\textsubscript{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
\textsuperscript{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 \mu M) 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™. Background luminescence and
fluorescence determined in wells containing untransfected cells was subtracted. To correlate
the luminescence and the fluorescence values with relative receptor numbers, the total number
of \textsuperscript{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 MT₁/MT₂ Heterodimerization based on Co-immunoprecipitation and BRET experiments. In a previous study we reported, using a BRET-based approach, that both MT₁ and MT₂ 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 MT₁ 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 MT₂ 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-MT₁ (Fig. 1a, mb). The Flag-MT₁ was also pulled down by the immunoprecipitation of the Myc-tagged MT₂ receptor, confirming the existence of MT₁/MT₂ heterodimers in cell lysates (Fig. 1a, IP). MT₁/MT₂ heterodimerization was further studied in intact cells by BRET. Wild-type human MT₁ and MT₂ 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 MT₁ and MT₂ 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 MT1-Rluc or MT2-Rluc and a control β2-adrenergic receptor YFP fusion protein expressed at comparable levels. Taken together, these data confirm our previous observation that MT1/MT2 heterodimers are formed in intact HEK 293 cells.

**Evaluation of the proportion of MT1 and MT2 homo- and heterodimers in living cells.** Although it is clear that MT1 and MT2 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 125I-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 MT1-homodimer and MT1/MT2 heterodimer formation, whereas the \(BRET_{50}\) value for MT2 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 \( \mu \text{m}^2 \) for HEK 293 cells, the density of dimers at the cell surface would be comprised between 10 and 50 receptors/\( \mu \text{m}^2 \), 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 \(^{125}\text{I}-\text{MLT} \) 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 \(^{125}\text{I}-\text{MLT} \) with high affinity (\( K_d = 115 \pm 22 \text{ pM} \) and \( 250 \pm 60 \text{ pM} \) for MT\(_1\)-Rluc and MT\(_2\)-YFP, respectively). When receptors were co-expressed, the apparent \( K_d \) was similar (200 pM \( \pm \) 21). The pharmacological profile of melatonin receptors was then determined in \(^{125}\text{I}-\text{MLT} \) 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\textsubscript{1}-Rluc and MT\textsubscript{2}-YFP at a 1:1 protein ratio competition curves for melatonin, S20098, S22153, S20928 and luzindole were monophasic with \textit{K}_i values comparable to those observed for cells expressing each receptor separately. The competition profiles of the MT\textsubscript{2}-selective ligands 4P-PDOT and S24773 were biphasic with \textit{K}_i values consistent with the binding to MT\textsubscript{1} and MT\textsubscript{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\textsubscript{1}/MT\textsubscript{2} heterodimers can be excluded since this receptor species represents a major receptor fraction in cells co-expressing MT\textsubscript{1} and MT\textsubscript{2} receptors at a 1:1 protein ratio. We can also exclude that MT\textsubscript{1}/MT\textsubscript{2} heterodimers are unable to bind \textsuperscript{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\textsubscript{1} and MT\textsubscript{2} homodimers. Indeed, no decrease in \textsuperscript{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 \textsuperscript{125}I-MLT (not shown). In addition, the effect of ligands on the BRET signal presented below shows that MT\textsubscript{1}/MT\textsubscript{2} heterodimers are ligand binding competent. Having excluded these possibilities, the competition profiles in cells co-expressing MT\textsubscript{1} and MT\textsubscript{2} receptors, may be either explained by the fact that the affinity of MT\textsubscript{1} and MT\textsubscript{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
MT₁/MT₂ 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 MT₂ homodimer and the MT₁/MT₂ 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 result 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 (EC₅₀),
correlates with binding affinities of the ligands. A good correlation would be expected for
receptor homodimers such as the MT₂ homodimer. To test this prediction, MT₂-Rluc and
MT₂-YFP fusion proteins were co-expressed at a 1:3 ratio that corresponds to the optimized
condition for BRET measurements (Ayoub et al., 2002). Kᵢ values were determined in
¹²⁵I-MLT competition binding experiments for selected ligands and were shown to be similar to
those observed in cells expressing MT₂-YFP alone (compare Tab. 1 and Tab. 3). The same
compounds increased the BRET signal in cells expressing MT₂ 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 ¹²⁵I-MLT competition binding assay (Kᵢ : 2-iodomelatonin = 20098 > melatonin = 4P-
PDOT > S22153 = S24773 = luzindole > S20928; EC₅₀ : 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 MT₁/MT₂ heterodimers compared to MT₂ homodimers (26 and 126 times, respectively) and EC₅₀ values for 4P-PDOT are 5 times lower for the MT₂ homodimer. Taken together these results show that melatonin receptor-specific ligands are binding to MT₁/MT₂ 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 MT₂ homodimers.

**Both ligand binding sites are functional within the MT₁/MT₂ heterodimer.** In cells co-expressing MT₁ and MT₂ receptors,¹²⁵I-MLT competition binding curves for the MT₂-selective compounds S24773 and 4P-PDOT were biphasic with a MT₂ binding site of high affinity and a MT₁ 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 MT₁/MT₂ heterodimers. However, experimental BRET curves were monophasic for these compounds and EC₅₀ values corresponded to the affinity for MT₂ (see Fig. 4) indicating that the ligand-promoted BRET change is due to binding to the MT₂ 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 (MT₁-like) functional binding site in the heterodimer. To address this point, we studied the effect of the MT₁-selective ligand S26284 (Audinot et al., 2003). The BRET dose-response curve of this compound for the MT₁/MT₂ heterodimer was monophasic with an EC₅₀ of 48 ± 3 nM (Fig. 6), which is close to the Kᵢ value measured for MT₁-Rluc (47 ± 5 nM, n=2) compared to the Kᵢ value for MT₂-YFP (605 ± 420 nM, n=2). Thus S26284 bound to the MT₁-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 MT₁-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 MT2-YFP (Fig. 6). Taken together these data indicate that both the MT1- and the MT2-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 MT1/MT2 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 MT1- and MT2-binding site are functional within the heterodimer. We have also shown that the two binding sites maintain their respective selectivity for MT1- and MT2 selective ligands and that the ligand interaction profile of the MT1/MT2 heterodimer determined by BRET is not identical to that of the MT2 homodimer.

The homo-/ heterodimer ratio of MT1 and MT2 receptors has been determined with the BRET donor saturation assay. The engagement of MT1 receptors into MT1 homodimers or MT1/MT2 heterodimers appears to be governed exclusively by the relative expression levels of MT1 and MT2 receptors since similar relative affinities were observed in BRET saturation assays. In contrast, MT2 receptors have a higher tendency to form heterodimers with MT1 receptors than to form homodimers suggesting that MT2 receptors may be preferentially engaged into heterodimers in cells co-expressing both receptors. The documented co-expression of MT1 and MT2 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 (GABABR), heterodimerization between GABAB1 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 MT2 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 (MT1 and MT2), 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 MT1- or MT2-like binding sites of the MT1/MT2 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 far 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 alpha2A 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 MT1/MT2 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 MT1/MT2 heterodimers. This approach may be potentially applied to a wide range of ligand-regulated receptors.
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Footnotes

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Legends to figures:

Figure 1: Detection of MT₁/MT₂ heterodimers. A, HEK 293 cells stably expressing Myc-MT₂ were transiently transfected or not with the Flag-MT₁ 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 ¹²⁵I-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 MT₁ and MT₂ 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 MT₂ homodimers

A, HEK 293 cells co-expressing MT₂-Rluc and MT₂-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 Ki and EC₅₀ values of the MT₂ 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 MT₁/MT₂ heterodimers

A, HEK 293 cells co-expressing MT₁-Rluc and MT₂-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 EC₅₀ values of MT₁/MT₂ heterodimers and Ki values of MT₁ and MT₂ receptors. EC₅₀ values of the MT₁/MT₂ heterodimer shown in Tab. 4 were plotted against Ki values determined in cells co-expressing MT₁ and MT₂ receptors (A) or expressing MT₁-Rluc (B) or MT₂-YFP (C) receptors individually. D, Correlation between EC₅₀ values of the MT₂ homodimer and the MT₁/MT₂ 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 MT₁-selective S26284 compound. Data were fitted using a linear regression equation (GraphPad Prism).

**Figure 6 : Ligand-dependent BRET changes of S26284 in cells expressing MT₁ and MT₂ homo- and heterodimers.** HEK 293 cells co-expressing MT₁-Rluc and MT₂-YFP (□) or MT₂-Rluc and MT₂-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).
TABLE 1

*Binding affinities measured in HEK 293 cells expressing MT₁ and MT₂ receptors.*

HEK 293 cells expressing MT₁-Rluc or MT₂-YFP or both together at a 1:1 ratio were incubated with ¹²⁵I-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.

<table>
<thead>
<tr>
<th>Ligands</th>
<th>Kᵢ (nM)</th>
<th>Ratio Kᵢ</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>MT₁-Rluc</td>
<td>MT₂-YFP</td>
</tr>
<tr>
<td>1) - Melatonin</td>
<td>0.36 ± 0.1</td>
<td>0.48 ± 0.1</td>
</tr>
<tr>
<td>2) - S20098</td>
<td>0.78 ± 0.2</td>
<td>0.08 ± 0.05</td>
</tr>
<tr>
<td>3) - S22153</td>
<td>39.4 ± 9.0</td>
<td>13 ± 3</td>
</tr>
<tr>
<td>4) - S20928</td>
<td>244 ± 105</td>
<td>210 ± 81</td>
</tr>
<tr>
<td>5) - 4P-PDOT</td>
<td>53.5 ± 13</td>
<td>2 ± 0.3</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>6) - Luzindole</td>
<td>31.6 ± 7.0</td>
<td>12.3 ± 4.3</td>
</tr>
<tr>
<td>7) - S24773</td>
<td>295 ± 26</td>
<td>3.70 ± 1.7</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
**TABLE 2**

*Relative affinities between two BRET partners of melatonin receptors.*

The BRET$_{50}$ represents the acceptor/donorr ratio required to reach half-maximal BRET in BRET-donorr 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).

<table>
<thead>
<tr>
<th>Dimer</th>
<th>BRET$_{50}$</th>
<th>(Receptor/cell)</th>
<th>(fmoles/mg of protein)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1- MT$_1$-Rluc/MT$_1$-YFP</td>
<td>3.7 ± 0.8</td>
<td>3080 ± 520</td>
<td>33 ± 4</td>
</tr>
<tr>
<td>2- MT$_1$-Rluc/MT$_2$-YFP</td>
<td>3.0 ± 1.6</td>
<td>3434 ± 436</td>
<td>43 ± 5</td>
</tr>
<tr>
<td>3- MT$_2$-Rluc/MT$_1$-YFP</td>
<td>4.2 ± 1.6</td>
<td>4554 ± 974</td>
<td>51 ± 12</td>
</tr>
<tr>
<td>4- MT$_2$-Rluc/MT$_2$-YFP</td>
<td>12.5 ± 2.2**</td>
<td>11318 ± 2176**</td>
<td>142 ± 27**</td>
</tr>
</tbody>
</table>
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.

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

Binding affinities (Ki) measured in HEK 293 cells co-expressing MT₁ and MT₂ receptors and EC50 values of ligand-induced BRET for MT₁/MT₂ heterodimers.

MT₁-Rluc and MT₂-YFP receptors were expressed at a 1:3 ratio (~80 fmol/mg of protein) and ¹²⁵I-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.

<table>
<thead>
<tr>
<th>Ligands</th>
<th>Kᵢ (nM)</th>
<th>EC₅₀ (nM)</th>
<th>Emax (%)</th>
<th>MT₂ homodimer / heterodimer</th>
<th>Ratio EC₅₀</th>
</tr>
</thead>
<tbody>
<tr>
<td>1) - Melatonin</td>
<td>0.40 ± 0.16</td>
<td>16.8 ± 9.7</td>
<td>142 ± 9</td>
<td>1.5</td>
<td></td>
</tr>
<tr>
<td>2) - S20098</td>
<td>0.23 ± 0.17</td>
<td>7.7 ± 4.6</td>
<td>136 ± 6</td>
<td>1.8</td>
<td></td>
</tr>
<tr>
<td>3) - S22153</td>
<td>8.87 ± 2.34</td>
<td>6.4 ± 5.6</td>
<td>132 ± 10</td>
<td>7.3</td>
<td></td>
</tr>
<tr>
<td>4) - S20928</td>
<td>116.6 ± 1.75</td>
<td>14 ± 9.5</td>
<td>136 ± 7</td>
<td>26.3</td>
<td></td>
</tr>
<tr>
<td>5) - 4P-PDOT</td>
<td>8.60 ± 7.40</td>
<td>211 ± 159</td>
<td>141 ± 4</td>
<td>0.2</td>
<td></td>
</tr>
<tr>
<td>6) - Luzindole</td>
<td>32.4 ± 9.20</td>
<td>1.0 ± 0.1</td>
<td>134 ± 13</td>
<td>126</td>
<td></td>
</tr>
<tr>
<td>7) - S24773</td>
<td>4.5 ± 1.0</td>
<td>4.1 ± 2.2</td>
<td>137 ± 4</td>
<td>8</td>
<td></td>
</tr>
</tbody>
</table>
**Figure 1:**

**A)**

<table>
<thead>
<tr>
<th></th>
<th>IP</th>
<th>mb</th>
<th>IP</th>
<th>mb</th>
</tr>
</thead>
<tbody>
<tr>
<td>Flag-MT1</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>+</td>
</tr>
</tbody>
</table>

**Blot:** anti-Flag

**IP:** anti-Myc

- Mono mer
- Dimer

**B)**

<table>
<thead>
<tr>
<th>YFP fusion proteins</th>
<th>MT₁</th>
<th>MT₂</th>
<th>β₂AR</th>
</tr>
</thead>
<tbody>
<tr>
<td>MT₁-Rluc</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>MT₂-Rluc</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**BRET Ratio (mBU):**

- MT₁-Rluc: 120 ± 10 mBU
- MT₂-Rluc: 90 ± 10 mBU
- β₂AR: 5 ± 1 mBU
Figure 3:

A)

BRET (% of control in the absence of ligand)

log [ligand] (M)

Melatonin

S20098

S22153

S20928

4P-PDOT

Luzindole

S24773

B)

-log EC50 (M)

-log K1 (M)

R^2 = 0.74
Figure 4:

BRET (% of control in the absence of ligand) vs. \( \log [\text{ligand}] \) (M) for various ligands:

- **Melatonin**
- **S20098**
- **S22153**
- **S20928**
- **4P-PDOT**
- **Luzindole**
- **S24773**
Figure 6:

![Graph showing BRET (% of control in the absence of ligand) vs. log [S26284] (M).]

- ▲ ▲ ▲ ▲ MT₂-Rluc/MT₂-YFP
- □ □ □ □ MT₁-Rluc/MT₂-YFP