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A new cross-talk between Rap1 and Rac that regulates sAPPα secretion

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Abstract

Cyclic AMP (cAMP) production results from the activation of Gs protein-coupled receptors, and regulates many physiological processes through activation of protein kinase A (PKA). Here we report that a Rho GTPase, Rac, is regulated by cAMP in a PKA-independent manner. We also show that Rac activation results from activation of a Ras GTPase, Rap1, via the guanine-nucleotide exchange factor Epac1. Activation of the Gs-coupled 5-HT4 receptor initiates this signalling cascade. Finally, we demonstrate that this new cross-talk between Ras and Rho GTPase families is involved in the cAMP–dependent processing of the amyloid precursor protein, indicating that this signalling pathway may serve important cellular functions.

Cyclic AMP is a second messenger which is produced in response to a large variety of extracellular signals and is involved in a wide range of cell-type specific effects, such as cell growth, cell differentiation, exocytosis and memory^{1,2}. Classically, cAMP mediates its cellular effects through activation of the protein kinase A (PKA) which induces phosphorylation of cytoplasmic and nuclear proteins, including enzymes and transcription factors³. However, a large body of evidence indicates that cAMP regulates a number of specific cellular functions through PKA-independent pathways. For instance, cAMP directly regulates some neuronal functions underlying neurotransmitter signalling and memory^{4,5}. More recently, we have shown that activation of a canonical Gs-coupled-receptor, the serotonin 5-HT₄ receptor may influence the processing of a key protein implicated in Alzheimer's disease (AD)⁶, the amyloid precursor protein (APP), through a mechanism involving cAMP production but not PKA activation⁷. The resulting effect of this PKA-independent stimulatory process is the secretion of the non-amyloidogenic soluble form of APP (sAPP α)⁷. These findings suggest the existence of alternative pathways by which G protein-coupled receptors (GPCRs) may regulate cell functions via cAMP.

Members of the Rho family of small GTPases are involved in the regulation of endocytotic pathways and have been shown to regulate secretory trafficking⁸⁻¹⁰. Therefore, we focused our research on small GTPase proteins as possible partners in the intracellular signalling pathway that couples the serotonin 5-HT₄ receptor to APP processing. We found that the 5-HT₄ receptor stimulates the activity of the small GTPase, Rac in a cAMP-dependent but PKA-independent manner. Furthermore, we provide experimental evidence that the cAMP regulated guanine nucleotide exchange factor (GEF), Epac1^{11,12} regulates sAPP α secretion through Rap1, a small GTPase of the Ras family and Rac. These data reveal an unsuspected connection between two families of small GTPases, and imply that Rac can function downstream of cAMP/Epac1/Rap1 in a novel signal transduction secretory pathway.

Results

Recombinant and native 5-HT₄ receptors induce sAPPa secretion and activate Rac. The amyloid β -protein (A β) is one of the major component of the senile plaques in the brain of AD patients and derives from its precursor APP by the action of β - and γ -secretases⁶. On the other hand, the α -secretase pathway splits APP within the A β domain producing a large amino-terminal non-amyloidogenic soluble APP (sAPPa), which is secreted in the extracellular medium and has neurotrophic activities and potent memory-enhancing effects^{13,14}. The effect of the activation of the 5-HT₄ receptor on sAPP α release was analysed in CHO cells stably expressing the neuronal human 5-HT_{4(e)} receptor isoform $(h5-HT_{4(e)})^{15}$. Since endogenous levels of APP in non neuronal cell lines are relatively low, we stably transfected these CHO cells with the major isoform of the human neuronal APP, APP695. As previously reported on native rodent APP⁷, activation of the h5-HT_{4(e)} receptor by 1 µM 5-HT enhances secretion of human sAPP α (Fig. 1a). This process is independent of PKA since the PKA inhibitor, H89 (10 µM) failed to block the stimulating effect of 5-HT on human sAPPa release (Fig. 1a). The regulation by the 5-HT₄ receptor of APP processing was not restricted to CHO cells, since it was also observed in primary culture of embryonic mouse cortical neurons (Fig. 1b & c). Indeed, two selective 5-HT₄ agonists, ML10302 and prucalopride^{16,17}, enhanced sAPPa secretion in primary neurons maintained in culture for 14 days (Fig. 1b & c). The stimulatory effect of 1 µM ML10302 on sAPPa secretion was inhibited by the selective 5-HT₄ antagonist, GR113808 (1 µM) in these neuronal cells (Fig. 1b). To examine the involvement of cAMP in APP metabolism, primary mouse cortical neurons were treated by the adenylyl cyclase activator forskolin (FSK, 10 µM) or a membrane permeant cAMP analogue adenosine 3',5' monophosphothioate (Rp-cAMPS, 10 µM) which does not activate PKA⁴⁶(Fig. 1c). Both drugs mimicked the effects of 5-HT₄ ligands on sAPPa secretion indicating that cAMP but not PKA is involved in APP metabolism in primary cortical neurons (Fig. 1c). Among the Rho family of GTPases which includes Rho, Cdc42 and Rac, RhoA and Cdc42 have been shown to be regulated by cAMP in a PKA-dependent manner¹⁸⁻²¹. Therefore, we focused our interest on Rac and examined its potential involvement in the above PKA-independent effects of 5-HT₄ receptor on APP processing.

We directly assayed Rac GTP-loading using a glutathione S-transferase (GST) fusion protein containing the Cdc42-Rac interactive binding domain (CRIB) of p21-activated kinase (PAK). When the GST-CRIB fusion protein coupled to glutathione beads was incubated with cell lysates from quiescent CHO cells that had been previously treated with 1 µM 5-HT for different periods of time, we found that 5-HT induced Rac activation (Fig. 2a). Increasing concentrations of 5-HT significantly enhanced Rac GTP-loading compared to untreated cells (Fig. 2b). As shown in Fig. 2c, 1 μ M of the selective 5-HT₄ receptor antagonist, GR113808, fully blocked 5-HT (1 µM)-enhanced Rac activation in CHO cells expressing the h5-HT_{4(e)} receptor indicating that the effect of 5-HT on Rac activation was indeed mediated by the h5-HT_{4(e)} receptor. GR113808 (1 µM) also reduced the basal level of Rac-GTP indicating a constitutive activity of the expressed h5-HT_{4(e)} receptor (Fig. 2c). To test whether native 5-HT₄ receptors can also activated Rac, we repeated these experiments in primary mouse cortical neurons and in the human IMR32 neuroblastoma cell line. In both systems, the two selective 5-HT₄ agonists, prucalopride (1 µM) and ML10302 (1 nM to 1 µM), induced a strong activation of Rac and their effects were inhibited by the selective 5-HT₄ antagonist, SB204070 (1 µM) (Fig. 2d & e). Furthermore, we observed that the increase in Rac activation upon treatment of the cells with the 5-HT₄ agonists was accompanied by an increase in the intracellular level of cAMP (data not shown). Altogether, these results demonstrate that recombinant and native 5-HT₄ receptors specifically induce the conversion of Rac into an activated state.

cAMP activates Rac. To examine the involvement of cAMP in Rac activity, CHO cells stably transfected with the h5-HT_{4(e)} receptor isoform were treated with FSK (10 μ M) or the phosphodiesterase inhibitor isobutylmethylxanthine (IBMX, 100 μ M) for 10 min. The amount of Rac-GTP was determined as above (Fig. 3a). Both compounds mimicked the effects of 5-HT since they caused a strong increase in the amount of Rac-GTP (Fig. 3a). The maximal amount of Rac-GTP was about 7-fold (range 5.6 ± 1.9 to 8 ± 1.5) above its basal level (Fig. 3a, lower panel). A 10 min exposure of CHO cells to cell-permeable cAMP analogues, 8-bromo–cyclic AMP (8-Br-cAMP, 10 μ M) or Rp-cAMPS (10 μ M), activated Rac to the same extent as did treatment with forskolin and IBMX (Fig. 3b). A similar effect of FSK (10 μ M), 8-Br-cAMP (100 μ M) or Rp-cAMPS (10 μ M) on Rac activation was found in IMR32 neuroblastoma cells (Fig. 3c) and primary cortical neurons (Fig. 3d).

cAMP regulates Rac activation in a PKA-independent manner. Functional consequences of a rise in cAMP upon activation of Gs protein-coupled receptors usually involves activation of PKA and phosphorylation of substrate proteins^{1,3}. Thus, we examined whether the effects of the 5-HT₄ receptor on Rac activation was PKA-dependent. First, CHO cells transfected with the 5-HT₄(e) receptor were pre-treated with 5 μ M of the potent PKA inhibitor, H89, and Rac activation in response to 5-HT (1 μ M) or FSK (10 μ M) was measured by the affinity precipitation assay (Fig. 4a & b). Although H89 (5 μ M) completely abolished PKA activity (Fig. 4a & b, lower panels), it failed to block 5-HT- or FSK-induced Rac activation (Fig. 4a, middle and upper panels and 4b, upper panel). These data are consistent with the results reported in Fig. 3b, c & d. Indeed, the cAMP analogue and PKA inhibitor, Rp-cAMPS (10 μ M), greatly enhanced the amount of the GTP bound form of Rac in CHO and neuronal cells (Fig. 3). Second, in a mutant CHO 10260 cell line that has been characterized as exhibiting

more than 95% reduction in both type I and II PKA activities²³, FSK still induced Rac activation (Fig. 4c). The increase in the amount of Rac-GTP in CHO 10260 cells was not significantly different from that observed in the parental CHO 10001 cell line (Fig. 4c). In contrast, basal and stimulated levels of PKA activities were dramatically lower in the CHO mutants cells than in the parental cells (Fig. 4c lower panel right). We therefore conclude that Rac is activated by cAMP independently of PKA.

Epac1 and Rap1 are involved in cAMP-dependent Rac activation. Rho GTPases are activated by a family of GEFs known as Dbl family proteins. These proteins share tandem Dbl homology (DH) and pleckstrin homology (PH) domains. DH domain catalyses nucleotide exchange on Rho GTPases²⁴. We have searched databases for GEFs containing both DH domain and cAMP-binding site but did not identify any sequence with the characteristics of a gene encoding such a protein. A family of GEFs that bind cAMP with affinities similar to that of the regulatory subunit of type I PKA and are directly activated by cAMP have been recently identified^{11,12}. These proteins, called Epac or cAMP-GEFs, have been shown to function as exchange factors for Rap GTPases which belong to the Ras superfamily¹¹. In particular, it has been clearly established that cAMP-dependent activation of Rap1 involves Epac²⁵. Therefore, we searched for a potential involvement of Epac-Rap1 in 5-HT₄ receptordependent Rac activation. To do this, we first investigated the effects of an activation of the 5-HT₄ receptor on Rap1 activation. Pull down assays using GST-Ral-GDS-Rap1-binding domain showed that 5-HT was able to activate Rap1 in CHO cells stably expressing the 5-HT_{4(e)} receptor (Fig. 5a). Interestingly, the PKA inhibitor and cAMP analogue, Rp-cAMPS efficiently activated Rap1 (Fig. 5a). 5-HT-induced Rap1 activation was blocked by the selective 5-HT₄ antagonist SB204070 indicating that the reaction specifically involved the 5-HT_{4(e)} receptor (Fig. 5a). 5-HT₄ receptor activation also enhanced Rap1 activation in primary neuronal culture (data not shown) and in IMR32 neuroblastoma cells (see supplemental data, Sl Fig. 1). Thus, we conclude that Rap1 is involved in the 5-HT₄ receptor signalling pathway.

Next, we examined the involvement of Epac1 in this process. To do this, we transiently transfected CHO cells with a plasmid encoding a mutated form of Epac1 (Epac- $\Delta cAMP$) which lacks the cAMP-binding domain¹¹. Epac- Δ cAMP therefore behaves as a constitutive activated form of Epac1 and cannot be regulated by cAMP²⁵. Transfection of Epac- Δ cAMP induced Rac activation in a concentration dependent manner (Fig. 5b). Furthermore, a novel Epac-specific cAMP analogue which does not activate PKA, 8CPT-2Me-cAMP²⁶ also increased Rac activation in CHO and IMR32 cells transiently transfected with Epac1 (Fig. 5c). This compound also induced Rac activation in cells non transfected with Epac1 such as primary neurons (Fig. 5c) and CHO cells (data not shown). Similarly, CHO cells transfection with a positive dominant form of Rap1 (Q63E-Rap1) strongly increased the amount of Rac GTP (Fig. 5d). Transfection of Epac- $\Delta cAMP$ or Q63E-Rap1 also enhanced Rac activation in IMR32 cells (see supplemental data, Sl Fig. 1). In a similar approach, we examined the effects of a functional inactive form of Epac (Δ 1-148Epac), in which the first 148 amino acid residues have been removed thereby preventing its proper cellular localisation²⁷, or a Rap1 GTPase activating protein (RapGAP) which strongly lowers the level of Rap1-GTP. Δ 1-148Epac or RapGAP were co-transfected into CHO cells with Rac1 and their effects on the activation of transfected tagged Rac were determined by pull down experiments. As shown in Fig. 5e and 5f, cell transfection with Δ 1-148Epac or RapGAP strongly inhibited 5-HTinduced Rac activation. The stimulatory effect of FSK on Rac activation was also blocked by these dominant negative constructs in CHO cells (data not shown). Since Ras has also been shown to be activated by cAMP independently of PKA²⁸, we performed Rac pull down experiments upon cotransfection of Rac1 and a negative dominant form of Ras (RasN17). Transfected RasN17 did not influence the effect of the 5-HT₄ receptor on Rac activation indicating that cAMP-induced Rac activation is independent of Ras (Fig. 5g). Altogether, these data clearly indicate that cAMP-dependent Rac activation is mediated by Epac1 and Rap1.

Involvement of Rac in the h5-HT4(e) receptor induced-sAPPa secretion. To further validate this new signalling pathway, we analysed its possible involvement in sAPPa secretion. Transfection of constitutive active Epac-AcAMP in CHO cells increased the cellular release of sAPP α in the absence of stimulation and rendered it independent of 5-HT (Fig. 6a). Furthermore, 8CPT-2Me-cAMP also increased sAPPa secretion in CHO and IMR32 cells transfected with Epac1 (Fig. 6b). Activation of endogenous Epac with 8CPT-2Me-cAMP also increased sAPPa secretion in primary neurons and IMR32 neuroblastoma cells (Fig. 6b and supplemental data, Sl Fig. 1c). Consistent with this result, Q63E-Rap1 enhanced basal secretion of sAPP α whereas functional inactive form of Epac, Δ 1-148Epac, and RapGAP inhibited 5-HT-induced cellular release of sAPPa (Fig. 6c & d). Similar results were observed in IMR32 cells (see supplemental data, SI Fig. 1). Finally, a dominant negative Rac (N17Rac) inhibited basal and 5-HT-induced cAMP-dependent secretion of sAPPa when transfected in CHO cells (Fig. 6e). In contrast, constitutive activated form of Rac (V12Rac) mimicked the stimulating effect of 5-HT on sAPPa secretion in the absence of 5-HT₄ ligand (Fig. 6e). Transfected dominant negative and positive forms of Rac were effective in our system since they decreased and enhanced Rac activation, respectively (Fig. 6e). The lesser increase in Rac activation observed in control cells stimulated with 5-HT was due to the longer incubation period with the neurohormone. We indeed noticed that upon 30 min treatment with 1 µM 5-HT Rac activation decreased compared to the short period of activation (data not shown). Altogether, our results indicate that this novel signalling pathway involving Epac1, Rap1 and Rac is implicated in APP processing.

Discussion

On the basis of our observations, we propose a new signalling pathway (Fig. 7), initiated at a GPCR positively coupled to adenylyl cyclase (AC), in which Rac is activated by Rap1 following cAMP binding to Epac1. In the case of the h5-HT₄ receptor this novel cAMPdependent, PKA-independent signalling pathway is involved in APP processing. Serotonin 5-HT₄ receptors are distributed in a variety of mammalian tissues including brain, heart and digestive system, where they mediate many physiological effects of 5-HT. In addition, 5-HT₄ receptors are thought to be involved in various pathological conditions, such as atrial arrhythmia and neurodegenerative disease²⁹. Our results thus open new insights into the signalling pathways by which 5-HT through the 5-HT₄ receptor may mediate its biological effects. Besides its role in APP metabolism, this finding suggests that activation of the 5-HT₄ receptor may also influence cytoskeletal dynamic. Accordingly, activation of the recombinant h5-HT_{4(a)} receptor isoform in a murine neuronal cell line promotes RhoA-dependent neurite retraction³⁰.

In our study, we found that forskolin and three cAMP analogues, 8-Br-cAMP, Rp-cAMPS and 8CPT-2Me-cAMP mimicked the activation of Rac by the 5-HT₄ receptor. This indicates a cross talk between cAMP and Rac and is in agreement with a recent report showing that another GPCR positively linked to the heterotrimeric protein Gs, the adenosine A₂ receptor, activates the small GTPase Cdc42 by a mechanism involving cAMP²¹. In most cellular systems, cAMP effects have been associated with the activation of PKA. For instance, PKA phosphorylation of the GTP binding protein RhoA was reported to be a central event in mediating the morphological effects of cAMP in cytotoxic lymphocytes¹⁸. PKA has also been

recently involved in cAMP-dependent Cdc42 activation²¹. Here we showed that Rac activation is induced by cAMP in a PKA-independent manner in CHO and neuronal cells suggesting an alternative mode of cAMP action. The stimulating and inhibiting effects of dominant positive and negative forms of Epac on Rac activation, respectively, indicated that Epac is involved in this signalling pathway. In contrast to our study, Rac activation by chemoattractants, β 1 integrins and prostaglandin E2 requires PKA activity in MDA-MB-435 human breast cancer carcinoma cells and in human umbilical vein endothelial cells^{31,32}. As Rac does not contain any consensus PKA phosphorylation sites, it is suggested that PKA may regulate Rac indirectly via the phosphorylation of its specific GEFs such as Tiam-1 and Trio³¹. Therefore, depending on the level of expression of the GEF in a given cell, the effects of cAMP on Rac may be mediated by PKA-dependent or PKA-independent mechanisms.

Several studies have shown that Ras and Rho GTPases can be activated in cascade so that one small GTPase stimulates GTP loading on another one³³. For instance, Cdc42 has been shown to activate Rac in different cellular systems and Rac may downregulate or upregulate Rho activity^{34,35}. With respect to the Ras superfamily, Rap1 antagonizes Ras-induced activation of mitogenesis³⁶. Cross-talk can also occur between the two families of GTPases and, for instance, in neuroblastoma cells Ras–induced outgrowth is mediated by both Cdc42 and Rac activity³⁷. Our report is the first to show a cross-talk between Rap1 and Rac that involves the second messenger cAMP.

Interestingly, we found that the cAMP analogue and PKA inhibitor, Rp-cAMPS not only activated Rac but also strongly enhanced Rap1 activity in our cellular systems. Likewise, Rp-cAMPS also enhanced sAPPα release. In accordance with our results, Rp-cAMPS was shown to mimic the stimulating effect of calcitonin on H,K-ATPase in collecting duct cells, a process which is entirely dependent of Epac and Rap1⁴⁷. However, Rehmann and colleagues⁴⁸ have recently mentioned in their study the unpublished observation that Rp-cAMPS binds to but

does not activate Epac. Clearly, if this finding is confirmed *in vitro* and *in vivo*, a question arises on how Rp-cAMPS may activate Rap1? We propose that Rp-cAMPS may act indirectly, by increasing the intracellular pool of cAMP available for Epac activation. This could be done two ways: Rp-cAMPS, which is hydrolysis-resistant, may bind to but not activate cAMP phosphodiesterases (PDE), which might result in PDE inhibition and cAMP accumulation. In support of this hypothesis, Sp-cAMPS, another cAMP analogue related to Rp-cAMPS, functions as a competitive inhibitor of PDE3A⁴⁹. Alternatively, Rp-cAMPS, which inhibits PKA, might relieve an inhibitory effect of PKA on adenylyl cyclase⁵⁰ and/or alleviate the stimulatory effect of PKA on PDE3⁵⁰ and PDE4⁵⁰⁻⁵⁵ activities. Such a direct or indirect inhibitory effect of Rp-cAMPS on PDE could lead to comparable effects as those of IBMX (Fig. 3a).

Rap1 has been implicated in various biological processes such as cell differentiation and integrin-mediated cell adhesion³⁸. Interestingly, Rap1 may be associated with secretory vesicles and play a role in the regulation of secretion³⁹. In favour of this hypothesis, cAMP-GEFII (also called Epac2) which regulates Rap1 activity²⁵ is responsible for cAMP-dependent and PKA-independent exocytosis, particularly the Ca²⁺-dependent secretion of growth hormone⁴⁰. Members of the Rho family have also been shown to regulate exocytosis. Indeed, Cdc42 and Rac regulate secreting processes in different cell types such as neurones and mast cells^{9,10}. Here, we show for the first time that Rac is implicated in a precise cellular function which involves the 5-HT₄ receptor and APP metabolism. Rap1 activity is required in this process. It is unclear at the moment whether the effects of the small GTPases, Rap1 and Rac on APP metabolism are directly linked to an increase in the activity of the α -secretases or an effect on cellular trafficking of APP. Given the involvement of Rap1 and Rac in the regulation of secretory processes, it is reasonable to think that these GTPases may target APP to cellular compartments involved in α -secretase processing or are directly involved in the

secretion of sAPPα at the cell surface. Recent experiments performed in yeast supports such a connection between Rap1, Rho GTPases and secretory processes. Bud1 is the yeast homologue of Rap1 and interacts and activates Cdc24p, a GEF for the Rho-family GTPase. In *Saccharomyces cerevisiae*, Cdc24p activates Cdc42 which recruits a number of effectors to the budding site allowing polarization of the actin cytoskeleton and targeting secretion to that site^{41,42}. Further investigation is needed to test whether Rap1 GTPases may recruit and activate a specific Rho GEF which would switch on Rac in mammalian cells. Alternatively, a possible connection between Rap1 and Rac could involve integrins. Indeed, integrins induce local Rac effector coupling by targeting Rac to membranes and leading to the dissociation of Rho guanine nucleotide dissociation inhibitors⁴³. This observation combined with the recent data that Rap1A activates integrins may suggest that these molecules may be involved in cAMP dependent Rap-Rac activation⁴⁴.

In summary, we show here that Rac is regulated in a cAMP-dependent but PKAindependent manner through Epac1 and Rap1. The identification of Rap1 and Rac as two elements in a signalling pathway controlling a cAMP-dependent secretory process provides the molecular basis for a new cross-talk between Ras GTPases and Rho GTPases.

Methods

Reagents, cell culture and transfection

All media, sera, and antibiotics used in the cell culture were purchased from Life Technologies (Cergy Pontoise, France). GR113808, GR127935, SB204070 were gifts from Glaxo SmithKline Beecham (Harlow, UK). ML10302 and prucalopride were generously given by Sames Sicsic (BIOCIS UPRES A CNRS 8076, Châtenay-Malabry, France) and Janssen Research Foundation (Beerse, Belgium), respectively. 5-HT and IBMX were from Sigma Aldrich (L'Isle d'Abeau Chesmes, France). Forskolin, 8-Br-cAMP, Rp-cAMPS and H89 were purchased from Calbiochem (France Biochem, Meudon, France). 8-(4-chlorophenylthio)-2'-O-methyladenosine-3'-5'-cyclic monophosphate (8CPT-2Me-cAMP) was from Biolog Life Science Institute (Bremen, Germany). Pure neuronal cultures of mice containing less than 5% astrocytes were prepared as previously described⁴⁵. Dissociated cortical cells in MEM medium supplemented with 5% foetal bovine and 5% horse serum were plated in multiwell vessels that had been previously coated with poly-D-lysine and laminin and 10 µM cytosine arabinoside was added after 3 days in vitro. At 12 days in vitro, neuronal culture was switched to serum free condition to avoid any presence of 5-HT in the culture medium provided by the serum. Experiments were performed after 14-15 days in vitro in serum free medium. CHO and IMR32 cells were grown respectively in HamF12 and MEM media with 10% FCS and antibiotics. Before any treatment, IMR32 cells and CHO cells were serum starved for 48 h. In CHO cells stably expressing the h5-HT4(e) receptor, 1 µM GR127935 was used to block the endogenous 5-HT_{1(b)} receptors. Transient transfection experiments were performed with jetPEI[™] (Polyplus-transfection, Illkirch, France) according to the manufacturer instructions. Expression of transfected proteins was determined by immunoblotting with anti-Flag, anti-HA or anti-GFP antibodies.

Rap and Rac activation assay

Rap1 pull down experiments were exactly performed as previously described using the GST fusion protein containing the Rap1-binding domain of Ral-GDS¹¹. Rac activation was measured as follows. After stimulation, cells were lysed in RIPA buffer (50 mM Tris pH 7.5, 500 mM NaCl, 20 mM MgCl₂, 0.5% deoxycholic acid, 0.1% SDS, 1% Triton X100, 1 mM PMSF, 10 µg/ml leupeptin and aprotinin) and clarified by centrifugation at 15000g at 4°C for 20 min. Protein concentration was determined by BCA assay (Sigma Aldrich, L'Isle d'Abeau Chesnes, France). Two mg of proteins (1 mg for transient transfection) were incubated with 50 µg GST-CRIB of PAK coupled to gluthatione sepharose beads for 1 hour at 4°C. The beads were then washed 3 times in washing buffer (50 mM Tris pH 7.5, 150 mM NaCl, 20 mM MgCl₂, 1% Triton X-100, 0.1 mM PMSF, 10 µg/ml leupeptin and aprotinin) and protein extracted in Laemmli's buffer. Rac-GTP samples and total lysates (40 µg) were separated in a 15% SDS-PAGE and transfered onto a polyvinylidene difluoride (PVDF) membrane (Hybond-P, Amersham Pharmacia Biotech, Saclay, France). Membranes were hybridized with an anti-Rac monoclonal antibody (a gift from Toshifumi Azuma) and proteins were visualized by enhanced chemiluminescence (ECL⁺, Amersham Pharmacia Biotech, Saclay, France). Densitometric analysis was performed using a Biorad image acquisition system (Ivry-Sur-Seine, France, gel doc 1000) and fold induction of Rac activation compared to control was calculated.

In vitro kinase reactions and cAMP assays.

Protein kinase A activity and cAMP assays were exactly performed as previously described^{7,28}.

Measurement of sAPPa.

CHO cells expressing the h5-HT_{4(e)} receptor were stably transfected with APP695. Confluent CHO cells were cultured in 10% dialysed FCS-containing medium overnight. After drug treatment in serum free medium, media were centrifuged and concentrated by ultrafiltration (Vivaspin columns, Vivascience, Palaiseau, France). sAPP α was detected as previously described⁷ by Western blotting using the monoclonal antibody 6E10 (Signet pathology systems, Dedham, USA) or the rabbit polyclonal antiserum R1736 (provided by Dr Selkoe, Harvard Medical School, Boston) which are selective of the α secreted cleavage of APP.

Statistical analysis.

An unpaired Student's *t*-test was used to calculate differences between means; differences were considered significant when p was at least <0.05.

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Figure 1. The h5-HT₄ receptor and cAMP regulate APP processing in CHO and primary neuronal cultures. a, 5-HT (1 µM) increased sAPPa secretion in a PKAindependent manner in CHO cells. A representative immunoblot is shown. The cDNA coding for the human APP695 was stably cotransfected with the h5-HT_{4(e)} receptor. Stable cell clones were selected for their ability to bind 5-HT₄ ligands, stimulate cAMP production and regulate sAPPa secretion. H89 (10 µM) was preincubated 10 min before the 30 min treatment with 1 µM 5-HT. Immunoblots were performed with the monoclonal 6E10 antibody which is selective of sAPPa. **b**, Selective 5-HT₄ ligands regulate sAPPa secretion in primary neuronal cultures. Primary cortical neurons from 14 days in vitro were incubated overnight with 1 µM of the selective 5-HT₄ agonist, ML10302 in the absence or presence of 1 µM of the 5-HT₄ antagonist, GR113808. sAPP α was detected in the culture medium by immunoblotting using the R1736 antiserum. The lower panel shows the quantification of the signal which was expressed as fold activation of the control untreated cells in the same experiment. Results are means \pm S.E.M. for 3 independent experiments. p<0.01, compared with untreated control cells, GR113808 or ML10302+GR113808 (t-test). c, sAPPa secretion is increased by cAMP in primary neurons. Mouse embryonic cortical neurons maintained in culture for 14 days were stimulated with the selective 5-HT₄ agonist, prucalopride (1 µM), FSK (10 µM) or with RpcAMPS (10 µM) overnight. sAPPa secretion was determined as described in b. A 110 kDa molecular weight marker is indicated at the left. CT, control cells.

Figure 2. The 5-HT₄ receptor induces Rac activation. a, Time-dependent stimulation of Rac by 1 μ M 5-HT in CHO cells stably expressing the h5-HT_{4(e)}. Rac activation was analysed by pull down experiments using the GST-CRIB fusion protein. Precipitated proteins were

subjected to Western blot using a monoclonal antibody to Rac. **b**, CHO cells expressing the h5-HT_{4(e)} receptor isoform were incubated with increasing concentrations of 5-HT for 10 min and the amount of Rac-GTP was determined as in b. **c**, The specificity of the reaction was determined with the selective 5-HT₄ receptor antagonist GR113808 (1 μ M), used in the presence of 1 μ M 5-HT. **d**, Selective 5-HT₄ ligands regulate Rac activation in primary cortical neurons. Cells were treated with a 5-HT₄ agonist, prucalopride (1 μ M) for 10 min, in the absence or presence of the 5-HT₄ antagonist, SB204070 (1 μ M). Cells were then lysed and assayed for GTP-bound Rac. **e**, The left panel shows a concentration-dependent stimulation of Rac by the selective 5-HT₄ agonist, ML10302 in IMR32 neuroblastoma cells. Right panel, the selective 5-HT₄ antagonist, SB204070 (1 μ M) inhibited ML10302 (1 μ M, 5 min)–induced Rac activation in IMR32 cells.

Figure 3. cAMP-dependent activation of Rac. a, b, CHO cells expressing the 5-HT_{4(e)} receptor were treated with 5-HT (1 μ M), forskolin (FSK) (10 μ M), IMBX (100 μ M), 8-Br-cAMP (10 μ M) or Rp-cAMPS (10 μ M) for 10 min and the amount of Rac-GTP was determined by pull down experiments and Western blot using an anti-Rac antibody. Lower panels, Rac-GTP is expressed as fold activation of the control (CT) cells (means \pm s.e.m; n=4 to 9). **c, d**, IMR32 cells and primary cortical neurons were treated and analysed for Rac activation as in (a) and (b). Prucalopride was used at a final concentration of 1 μ M. CT, control untreated cells. p<0.05, p<0.01 compared with control cells (Student *t*-test).

Figure 4. 5-HT₄ receptor– and forskolin–induced Rac activation is independent of PKA. a, b, CHO cells were preincubated for 15 min with the PKA inhibitor H89 (5 μ M) and stimulated with 5-HT (1 μ M) and forskolin (FSK, 10 μ M) for 10 min. Amounts of Rac-GTP and PKA activity were determined. A control for total Rac expression (total lysates) is shown. The upper panels in (a) and (b) show typical immunoblots. The middle panel in (a) shows means \pm s.e.m. of 4 independent experiments. Results are expressed as fold activation of the control (CT) cells. The lower panels in (a) and (b) show the measurements of PKA activity obtained from 3 to 5 independent experiments (means \pm s.e.m., n=9 to 15). p<0.05, p<0.01 or p<0.001 compared with CT (Student *t*-test) **c**, CHO 10001 and CHO 10260 cells were stimulated with increasing concentrations of FSK for 10 min and the amounts of Rac-GTP (upper panel and lower panel left) and PKA activity (lower panel right) were determined. Lower panels, values are means \pm s.e.m (n=9 to 12). ns, non significant; CT, control untreated cells. p<0.001 compared with control CHO 10001 cells (Student *t*-test).

Figure 5. Epac1 and Rap1 regulate Rac activation. a, The 5-HT₄ receptor and the cAMP analogue and PKA inhibitor, Rp-cAMPS activate Rap1. CHO cells stably expressing the 5-HT₄(e) receptor were treated with 1 μ M 5-HT in the absence or presence of the selective 5-HT₄ antagonist, SB204070 (1 μ M) or 10 μ M Rp-cAMPS for 10 min. After cell lysates were prepared, Rap1-GTP was assessed by pull down assays using the GST fusion protein containing the Rap1-binding domain of Ral-GDS followed by Western blot using an anti-Rap antibody. **b**, **d**, Constitutive activated forms of Epac and Rap1 induced Rac activation. CHO cells were transiently transfected with increasing amounts (from 2 to 8 μ g) of cDNA encoding an activated version of Epac (Epac- Δ cAMP) (b), Q63E-Rap1 (d). In each condition, the total amount of cDNA was kept constant to 8 μ g by transfecting the corresponding empty vector. One day after transfection, cells were treated or not with forskolin (FSK, 10 μ M, 10 min) and Rac activation was assessed by pull down experiments followed by Western blot using an anti-Rac antibody. **c**, 8-CPT-2Me-cAMP induced Rac activation.. Upper panel, CHO cells were transfected with 4 μ g of Epac WT, then stimulated with 15 μ M 8-CPT-2Me-cAMP for

the indicated period of time and Rac activation was determined. Lower panel, mouse cortical neurons maintained in culture for 14 days were stimulated with the indicated concentration of 8-CPT-2Me-cAMP for 10 min and then Rac pull-down assays were performed. **e**, **f**, 5-HT-induced Rac activation is inhibited by a dominant negative form of Epac or RapGAP. CHO cells were cotransfected with the indicated amounts of empty vector, AU5-tagged Rac1 wild type and Δ 1-148Epac or RapGAP. One day after transfection, cells were treated or not with 5-HT (1 µM, 10 min) and Rac activation was determined as above. Upper panels, representative western blot are shown. Lower panels, AU5-Rac-GTP was normalized to total AU5-Rac. p<0.01, p<0.001 compared with either control cells (CT) (f) or 5-HT treated cells (e) (means ± s.e.m; n=3 to 4) (Student *t*-test). **g**, Ras is not involved in cAMP-dependent Rac activation. Cotranfection and Rac pull down experiments were performed as in (d) and (e) using the indicated amounts of empty vector, AU5-tagged Rac1 WT and RasN17. A Western blot of the same lysates was carried out using anti-Flag, anti-HA and anti-GFP antibodies to monitor the level of expression of the transfected cDNA constructs.

Figure 6. Epac1, Rap1 and Rac are involved in PKA-independent sAPP α secretion. a, CHO cells were transiently transfected with increasing amounts (from 2 to 8 µg, total amount of transfected DNA was kept constant to 8 µg with the empty plasmid vector) of Epac- Δ cAMP cDNA. One day later secreted sAPP α was determined by immunoblot upon 30 min treatment with 1 µM 5-HT. **b**, The selective activator of Epac, 8CPT-2Me-cAMP enhanced sAPP α secretion. CHO and IMR32 cells were transiently transfected with 4 µg of Epac WT. One day later secreted sAPP α was determined by immunoblot upon treatment with 8CPT-2Me-cAMP (10 or 100 µM). The expression of the transfected construct was monitored using an antibody directed against GFP. CT, control untreated cells. Primary cortical neurons from 14 days *in vitro* were incubated overnight with 8CPT-2Me-cAMP (10 or 100 µM) and secreted sAPP α was determined as above. **c**, CHO cells were transfected with 8 µg of empty vector or Q63E-Rap1. Cell treatment and immunoblots were performed as in (a). **d**, A dominant negative form of Epac (Δ 1-148Epac) and RapGAP block sAPP α secretion. CHO cells were cotransfected with the empty vector (4 µg), or AU5-tagged Rac1 WT (4 µg) and Δ 1-148Epac (4 µg) or RapGAP (4 µg). One day after, cells were stimulated with 1 µM 5-HT for 30 min and sAPP α secretion was determined as in (a). The expression of the transfected constructs were monitored using antibodies directed against the different epitopes. **e**, Rac regulates sAPP α secretion. Cells were transiently transfected with 8 µg of RacN17 or RacV12, or vector alone. Cell treatment and immunoblots were performed as above. Corresponding cell lysates were assayed for Rac-GTP by affinity precipitation.

Figure 7. Model of Rac activation by cAMP. Stimulation of Gs coupled receptor such as the 5-HT₄ receptor activates the adenylyl cyclase which increases intracellular cAMP levels Subsequent activation of Epac1 increases Rac activation via Rap1. In the case of the 5-HT₄ receptor this novel signalling pathway is involved in APP processing.

SI Figure 1. The 5-HT₄ receptor regulates Rap1 and Rac activities and sAPP α secretion in IMR32 neuroblastoma cells. a, The 5-HT₄ receptor activates Rap1. IMR32 cells were treated with a selective 5-HT₄ agonist, prucalopride (1 µM) or forskolin (FSK, 10 µM) for 10 min and the amount of Rap-GTP was determined by pull down experiments followed by a Western blot using an anti-Rap antibody. b, d, Constitutive activated forms of Epac and Rap1 induced Rac activation and sAPP α secretion. IMR32 cells were transiently transfected with 8 µg of cDNA encoding an activated version of Epac (Epac- Δ cAMP), Q63E-Rap1, or V12Rac. The total amount of cDNA was kept constant to 8 µg by transfecting the corresponding empty vector. One day after transfection, cells were treated or not with prucalopride (1 μ M) and Rac activation or sAPP α release were assessed by pull down experiments and immunoblots, respectively. **c**, 8CPT-2Me-cAMP induced sAPP α secretion in IMR32 cells. Cells were treated with prucalopride (1 μ M) or 8CPT-2Me-cAMP (10 or 100 μ M) for 3 h and secreted sAPP α was determined as above.



Fig. 1

а

Fig. 2



Fig. 4

а

b

GFP-Epac WT

Q63E-Rap1

Fig. 5

– HA-

f

g

е

d

Supplementary information

IMR32 neuroblastoma cells

Sl. Fig. 1