

# Melanocortin receptor accessory proteins in adrenal gland physiology and beyond

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## Abstract

The melanocortin receptor (MCR) family consists of five G-protein-coupled receptors (MC1R–MC5R) with diverse physiological roles. MC1R controls pigmentation, MC2R is a critical component of the hypothalamic–pituitary–adrenal axis, MC3R and MC4R have a vital role in energy homeostasis and MC5R is involved in exocrine function. The melanocortin receptor accessory protein (MRAP) and its paralogue MRAP2 are small single-pass transmembrane proteins that have been shown to regulate MCR expression and function. In the adrenal gland, MRAP is an essential accessory factor for the functional expression of the MC2R/ACTH receptor. The importance of MRAP in adrenal gland physiology is demonstrated by the clinical condition familial glucocorticoid deficiency, where inactivating MRAP mutations account for ~20% of cases. MRAP is highly expressed in both the zona fasciculata and the undifferentiated zone. Expression in the undifferentiated zone suggests that MRAP could also be important in adrenal cell differentiation and/or maintenance. In contrast, the role of adrenal MRAP2, which is highly expressed in the foetal gland, is unclear. The expression of MRAPs outside the adrenal gland is suggestive of a wider physiological purpose, beyond MC2R-mediated adrenal steroidogenesis. *In vitro*, MRAPs have been shown to reduce surface expression and signalling of all the other MCRs (MC1,3,4,5R). MRAP2 is predominantly expressed in the hypothalamus, a site that also expresses a high level of MC3R and MC4R. This raises the intriguing possibility of a CNS role for the MRAPs.

## Key Words

- ▶ MRAPs
- ▶ Melanocortin receptor
- ▶ HPA axis
- ▶ Adrenal cortex

*Journal of Endocrinology*  
(2013) 217, R1–R11

## Hypothalamic–pituitary–adrenal axis

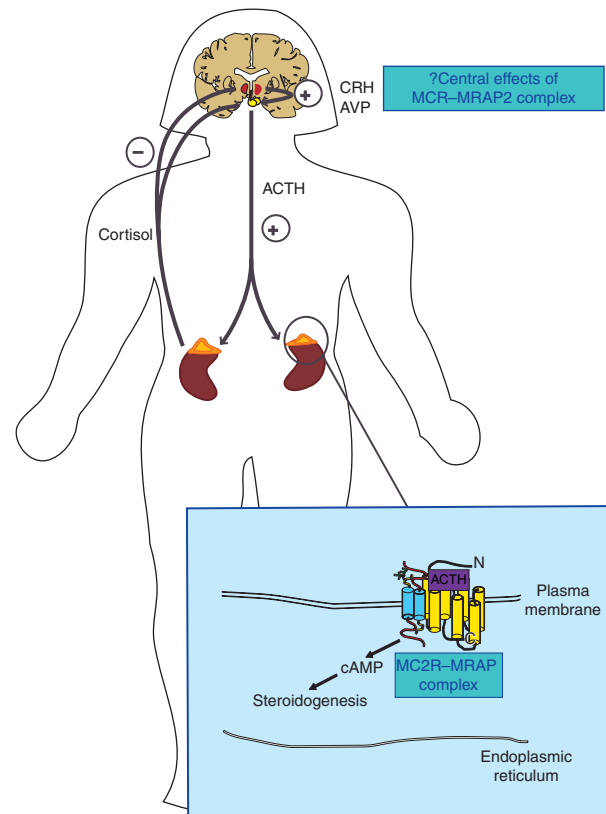
The adrenal glands are responsible for releasing different classes of hormones. The inner part of the glands, the medulla, secretes catecholamines under the control of the sympathetic nervous system. The outer part, the cortex, has three functionally distinct layers – the outer zona glomerulosa that secretes aldosterone, the middle zona fasciculata that produces glucocorticoids and the inner zona reticularis that is responsible for the production of adrenal androgens. Production and release of

glucocorticoids by the zona fasciculata are tightly regulated by the hypothalamus and pituitary to control diurnal cortisol levels and the rapid increase of cortisol secretion during emotional or physical stress. Stress triggers a number of brain circuits such as brainstem catecholamine-producing pathways that cause excitation of the parvocellular neurons of the hypothalamic paraventricular nucleus (PVN; reviewed in Herman & Cullinan (1997) and Arnsten (2009)). In response to stress, these neurons start secreting corticotropin-releasing hormone

(CRH) and arginine vasopressin (AVP) (Sawchenko & Swanson 1985) into the hypophyseal portal circulation. Both CRH and AVP trigger secretion of ACTH by cells of the anterior pituitary gland (Vale *et al.* 1981, Aguilera *et al.* 1983). ACTH is produced by cleavage from its precursor proopiomelanocortin (POMC), as are  $\alpha$ -,  $\beta$ - and  $\gamma$ -melanocyte-stimulating hormones (MSHs). ACTH is released into the circulation to act on peripheral sites, mainly the adrenal glands, to stimulate glucocorticoid hormone production. Glucocorticoids negatively feedback to regulate the release of CRH and AVP at the hypothalamus and ACTH at the pituitary, thus providing tight regulation of cortisol production (Fig. 1). This regulatory system is termed the hypothalamic–pituitary–adrenal (HPA) axis, and perturbations of this system have been shown to be involved in a wide range of physiological processes and diseases such as Cushing's syndrome, adrenal insufficiency, depression and sepsis.

### Melanocortin 2 receptor is a critical component of the HPA axis

Melanocortin receptors (MCRs) are a subfamily of seven-transmembrane (TM)-domain G-protein-coupled receptors (GPCRs) that mediate signalling of hormones derived from POMC such as ACTH and MSHs (Mountjoy *et al.* 1992, Cone *et al.* 1993). To date, five MCRs have been described, and they have been shown to have a diverse range of functions. MC1R controls skin pigmentation (Valverde *et al.* 1995), MC3R and MC4R are major contributors to the regulation of food intake and energy homeostasis (Marsh *et al.* 1999, Butler *et al.* 2000, Chen *et al.* 2000) and MC5R is highly expressed during embryogenesis but in adults is believed to be involved in exocrine function (Labbe *et al.* 1994, Chagnon *et al.* 1997, Ogawa *et al.* 2004). MC1,3,4,5R respond to all POMC-derived hormones (ACTH and  $\alpha$ -,  $\beta$ - and  $\gamma$ -MSH) but differ in ligand preference, whereas MC2R responds to ACTH only (Cone *et al.* 1993). Binding of ACTH to MC2R results in increase of cAMP and activation of protein kinase A, which is essential for the expression of steroidogenic enzymes and cortisol production. MC2R appears to be positively regulated by its ligand (Hofland *et al.* 2012), and some studies have suggested a rare involvement of MC2R in adrenal cortisol-secreting tumours (Latronico *et al.* 1995, Arnaldi *et al.* 1998). MC2R is predominantly expressed in all zones of the adrenal cortex, but ACTH mostly acts on the zona fasciculata (Mountjoy *et al.* 1992). ACTH activation in zona reticularis stimulates the production of adrenal androgens (Weber *et al.* 1997).



**Figure 1**

Schematic overview of the HPA axis and the sites of predominant MRAP/MRAP2 action. In short, CRH and AVP at the hypothalamus stimulate ACTH production in the anterior pituitary gland. ACTH released into the circulation acts on the adrenal cortex to produce glucocorticoids (cortisol in humans and corticosterone in rodents). Glucocorticoids negatively feedback to regulate the release of CRH and AVP at the hypothalamus and ACTH at the pituitary. MC2R and MRAP act predominantly in the adrenal cortex as demonstrated by patients with isolated cortisol deficiency caused by deleterious mutations in MC2R and MRAP. The function of MRAP2 is yet unclear, although it is thought to have a predominant central effect due to its expression in the hypothalamus.

Failure of MC2R to activate in response to ACTH causes familial glucocorticoid deficiency (FGD; Clark *et al.* 1993), a rare autosomal recessive disorder characterised by severe cortisol deficiency with high plasma ACTH levels and normal mineralocorticoid levels (Shepard *et al.* 1959, Clark & Weber 1998). Untreated, FGD is lethal in early childhood, usually due to profound hypoglycaemia or overwhelming infection. MC2R mutations resulting in effective loss of the receptor function are responsible for FGD type 1, which accounts for up to 25% of all FGD cases (Clark *et al.* 2005). MC2R activating mutations are extremely rare, and to date only one (F278C) has been reported, which was found in a patient with cyclical Cushing's syndrome (Swords *et al.* 2002). The same

mutation was found in conjunction with a non-activating mutation presenting with FGD (Chan *et al.* 2009a). The majority of MC2R mutations result in defective trafficking of the receptor to the plasma membrane (Chung *et al.* 2008).

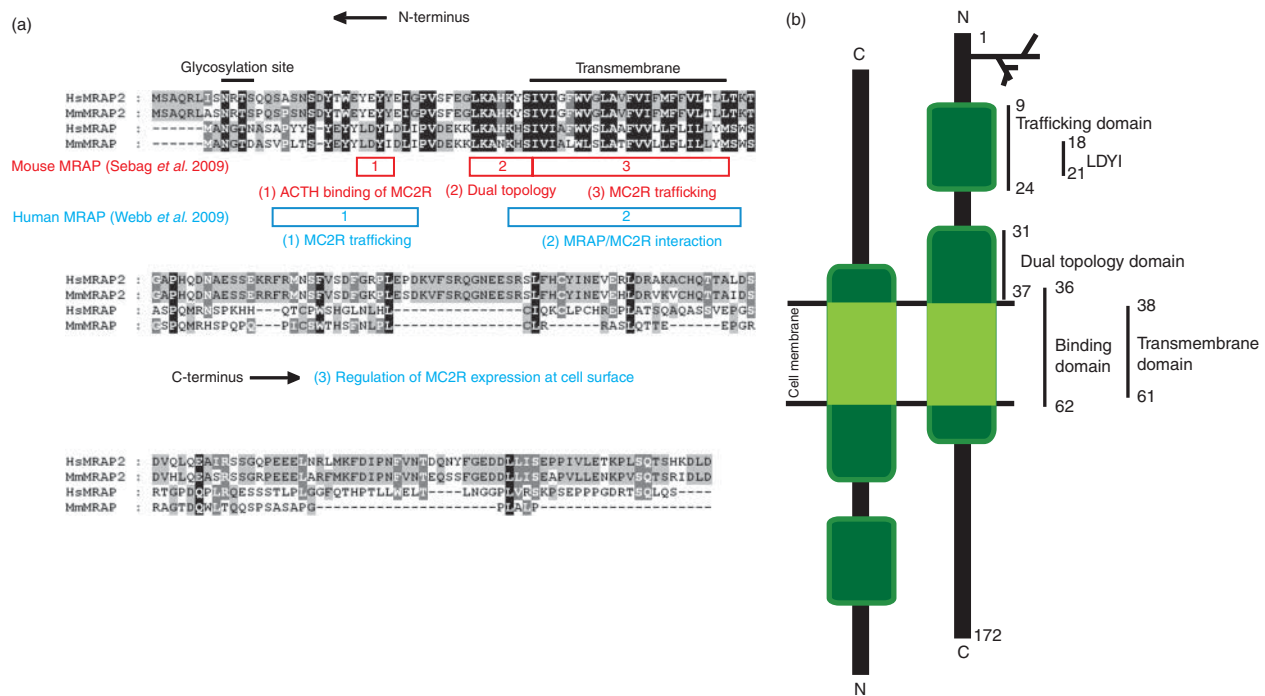
Functional expression of MC2R in cell lines of non-adrenal origin (heterologous cells) is notoriously difficult and results in retention of MC2R in the endoplasmic reticulum (ER) and unresponsiveness of such cells to ACTH (Schimmer *et al.* 1995). It was suggested that a specific factor, present in the adrenal gland, is essential for human MC2R function (Noon *et al.* 2002). This phenomenon is observed with all described MC2Rs in vertebrates apart from cartilaginous fish (elephant shark) where the MC2R can reach the plasma membrane and function in non-adrenal cells and responds not only to ACTH but to MSHs as well. It is suggested that the unique properties of MC2R such as selectivity to ACTH and requirement of the adrenal specific factor emerged after the divergence of the ancestral cartilaginous and bony fishes more than 400 million years ago (Reinick *et al.* 2012).

### Melanocortin 2 receptor accessory protein and adrenal gland physiology

A genetic screening of patients with FGD but with normal MC2R identified a number of nonsense and splice site mutations in the *C21ORF61* gene that would result in a complete absence or a short truncated protein thought to undergo rapid degradation if translated at all (Metherell *et al.* 2005). The gene product had previously been named fat tissue-specific low molecular weight protein due to its expression in differentiating adipocyte cells, but its function had been unknown (Xu *et al.* 2002). Further expression analysis revealed that this protein was highly expressed in the adrenal gland with lower expression levels in other tissues such as brain, thyroid, ovary, testis and breast (Metherell *et al.* 2005). From finding that mutations of this gene underlie ~20% cases of FGD, now known as FGD type 2, and that the protein was predominantly expressed in the adrenal gland, it was hypothesised that it could be the adrenal specific factor required for MC2R functional expression. It was shown that this protein was able to interact with MC2R and thus, the new protein was named melanocortin receptor 2 accessory protein (MRAP; Metherell *et al.* 2005). MRAP was shown to localise to the ER and plasma membrane, and its co-expression with MC2R in heterologous cells resulted in a significant proportion of the receptor being delivered to the plasma membrane. Moreover, such cells became ACTH

responsive, thus showing that MRAP was the adrenal specific factor required for MC2R function (Metherell *et al.* 2005). Later studies confirmed that MRAP is essential for trafficking of MC2R to the plasma membrane and its function. In the mouse adrenocortical Y1 cell line that endogenously expresses both MC2R and MRAP and responds to ACTH stimulation, knockdown of MRAP by shRNA resulted in loss of response to ACTH, which was rescued by overexpression of the shRNA-insensitive human MRAP (Cooray *et al.* 2008). Both MRAP and MC2R are highly expressed in the zona fasciculata in glucocorticoid-producing cells, with the highest expression levels in undifferentiated zone (Gorrigan *et al.* 2011), which is directly adjacent to zona fasciculata and is believed to contain stem cells that contribute to steroidogenic cell maintenance (Mitani *et al.* 2003). These data suggest that MC2R and MRAP are important for adrenal gland development, and the adrenal histology of FGD supports this hypothesis as the glands from deceased patients demonstrate disorganisation of glomerulosa cells with almost complete absence of fasciculata and reticularis cells (Clark & Weber 1998). It has also been demonstrated that *in vivo*, mRNA levels of both MC2R and MRAP in the adrenal gland are regulated by ACTH (Xing *et al.* 2010, Hofland *et al.* 2012). However, at times of acute stress, the immediate steroidogenic response would depend on adequate levels of MRAP and MC2R at the surface and that can be readily trafficked to the plasma membrane. MRAP and MC2R that can be readily trafficked to the plasma membrane. Furthermore, pulsatility of MRAP transcription induced by pulsatile administration of ACTH in rats leads to the suggestion that adrenocortical responsiveness and expression of steroidogenic enzymes are ultimately determined by the availability of MC2R at the cell surface (Spiga *et al.* 2011).

The MRAP protein comprises 172 amino acids and consists of a highly evolutionarily conserved N-terminus and a single TM domain followed by a less conserved C-terminus (reviewed in Webb & Clark (2010)). MRAP has a unique previously uncharacterised topology (Sebag & Hinkle 2007). It forms a stable antiparallel homodimer at the plasma membrane and in the ER. The seven amino acids flanking the TM domain from the N-terminus are essential for this dual topology (Sebag & Hinkle 2007, 2009a,b, Cooray *et al.* 2008; Fig. 2). Importantly, MRAP forms a stable heterodimer with MC2R and this heterodimer is present at the plasma membrane (Sebag & Hinkle 2009b). Co-immunoprecipitation studies using MRAP fragments mapped the interaction domain to between amino acids 36 and 62; this region includes the entire TM

**Figure 2**

Regions of MRAP action. (A) Protein alignment of mouse and human MRAP proteins with schematic representation of identified functional domains. Work on human MRAP by Webb *et al.* identified two functional domains involved in MC2R trafficking and MRAP/MC2R interaction. There was also a possible suggestion that the C-terminus could regulate MC2R surface expression (Webb *et al.* 2009). In comparison, work on mouse MRAP has

revealed three possible essential domains involved with ACTH binding of MC2R, dual topology and MC2R trafficking (Sebag & Hinkle 2009a,b). Hs, *Homo sapiens*; Mm, *Mus musculus*. (B) Schematic representation of human MRAP adopting an antiparallel form. Dark green boxes denote amino acid regions of interest whilst the light green area demonstrates the transmembrane domain.

domain and several flanking amino acids (Webb *et al.* 2009). It is suggested that the dual topology of MRAP is important for trafficking of MC2R as deletion of the residues responsible for antiparallel dimerisation resulted in failure to traffic the receptor to the plasma membrane (Sebag & Hinkle 2009b). However, the deletion of the dimerisation region also removes a residue in the MC2R binding domain, which may affect MRAP/receptor binding and hence trafficking. Interestingly, a tyrosine-rich region in the N-terminus (amino acids 9–24) is required for MC2R trafficking to the plasma membrane and this region is similar to those in other GPCR accessory proteins, such as odorant receptor accessory protein REEP1 (Saito *et al.* 2004, Webb *et al.* 2009). Moreover, upon deletion within this region, at residues 18–21 (LDYI), MRAP was able to traffic MC2R to the plasma membrane surface but MC2R failed to respond to ACTH, indicating that probably these four amino acids facilitate receptor–ligand binding (Sebag & Hinkle 2009b).

Two MRAP isoforms with differing C-termini have been described in humans – MRAP $\alpha$  (19 kDa) and MRAP $\beta$

(14 kDa), a shorter isoform produced by alternative splicing. The precise function of the variable C-terminal domain of MRAP remains to be fully elucidated. The isoforms are expressed in the adrenal gland at similar levels, but the precise role of each isoform remains unclear (Metherell *et al.* 2005, Webb *et al.* 2009). Deletion of the whole C-terminus causes a decrease in the ACTH response of MC2R-expressing cells (Roy *et al.* 2012). Some studies have demonstrated that expression of the shorter isoform, MRAP $\beta$ , leads to higher MC2R surface expression, suggesting it may regulate MC2R internalisation rate (Roy *et al.* 2007, Webb *et al.* 2009). Recently, another study (Roy *et al.* 2012) demonstrated that MRAP $\alpha$  localised mainly in the ER, whereas MRAP $\beta$  exhibited a strong localisation at the plasma membrane. Although both MRAP isoforms and the C-terminally truncated construct were able to facilitate surface expression of MC2R, it was co-expression of MRAP $\beta$  and MC2R that resulted in the highest cAMP production upon stimulation with ACTH (Roy *et al.* 2012). These data suggest that the C-terminus may determine intracellular

localisation of MRAP and potentially increase MC2R affinity to its ligand or facilitate MC2R signalling, but this hypothesis needs further validation.

### MRAP2 in the adrenal gland

Recently, a homologue of MRAP highly conserved through vertebrates was identified having 39% amino acid sequence identity within the N-terminus, again with a single-pass TM domain, and it was named MRAP2 (Chan *et al.* 2009b). The degree of conservation of MRAP2 across species is much higher compared with MRAP, suggesting that MRAP2 is the ancestral gene (Webb & Clark 2010). It has been shown that in rodents, MRAP2 is expressed in developing adrenals, but the expression levels are very low in adulthood (Gorrigan *et al.* 2011). Expression has also been demonstrated in brain, mainly in the hypothalamus (Lein *et al.* 2007, Chan *et al.* 2009b, Gorrigan *et al.* 2011). In contrast, a more ubiquitous pattern of expression has been shown in zebrafish MRAP2 (Agulleiro *et al.* 2010). MRAP2 is slightly bigger than MRAP, comprising 205 amino acids (23.5 kDa), and has a glycosylation site NRTS at positions 9–12. It is likely to form antiparallel homodimers, and intracellular localisation of MRAP2 also matches the intracellular distribution of its homologue MRAP (Chan *et al.* 2009b). Cell expression studies showed that MRAP2 was able to form a complex with MC2R, facilitate its trafficking, and enable receptor signalling in response to ACTH. MRAP2 is also able to form stable heterodimers with MRAP when co-expressed in cell lines (Chan *et al.* 2009b). However, the role of MRAP/MRAP2 heterodimers remains to be elucidated as one study demonstrated negative regulation of the MRAP/MC2R complex by MRAP2 (Sebag & Hinkle 2010), whereas other studies demonstrated either no effect on the signalling properties of the MC2R/MRAP complex (Gorrigan *et al.* 2011) or that MRAP2 augments MC2R signalling (Agulleiro *et al.* 2010). A study of human adrenocortical tissues demonstrated an association between patient ACTH and cortisol levels and adrenal expression of MRAP and MC2R, but not MRAP2 (Hofland *et al.* 2012). Mutation of the glycosylation site of MRAP2 abolishes MC2R response to ACTH but does not affect facilitation of MC2R surface expression, suggesting that MRAP2 might have two separate accessory roles (Chan *et al.* 2009b). Interestingly, MC2R–MRAP2 complex requires 1000 times higher ACTH concentration than the ACTH amount needed for the MC2R–MRAP complex to trigger the same cAMP production (Gorrigan *et al.* 2011). This difference can be reversed by inserting residues LDYI, not conserved

between MRAP and MRAP2, into the MRAP2 protein sequence (Sebag & Hinkle 2009b). Based on all the data described above, it would be tempting to hypothesise that both MRAPs regulate the dose response of MC2R to ACTH. However, patients with FGD type 2 have plasma ACTH levels elevated hundreds of times above normal, and according to this hypothesis, such levels would enable signalling via the MC2R–MRAP2 complex in the absence of MRAP, whereas in fact intact MRAP2 is unable to compensate for the loss of MRAP, suggesting that interaction with MC2R is not the main function of MRAP2 *in vivo*. However, an MRAP2 adrenal role cannot be excluded, as MRAP2 is expressed in the developing adrenal gland, which suggests its involvement in the adrenal development (Gorrigan *et al.* 2011). Thus, the function of MRAP2 interaction with MC2R and its relationships with MRAP require further investigation using *in vivo* models.

### MRAP and MRAP2 function beyond the adrenal gland

The relatively wide expression patterns of both MRAP and MRAP2 suggest that they may have other functions. Although MC2R expression is mostly restricted to the adrenal glands, low levels are also detected in the skin and lymphocytes (Mountjoy *et al.* 1992, Slominski *et al.* 1996, Andersen *et al.* 2005), and the role of MC2R and MRAPs in these tissues is unknown. The importance of MC2R expression in lymphocytes such as natural killer and T helper (CD4<sup>+</sup>) cells (Andersen *et al.* 2005) is difficult to assess as lack of glucocorticoids in FGD patients compromises the immune system via a number of mechanisms (Clark *et al.* 1993, Lowenberg *et al.* 2008).

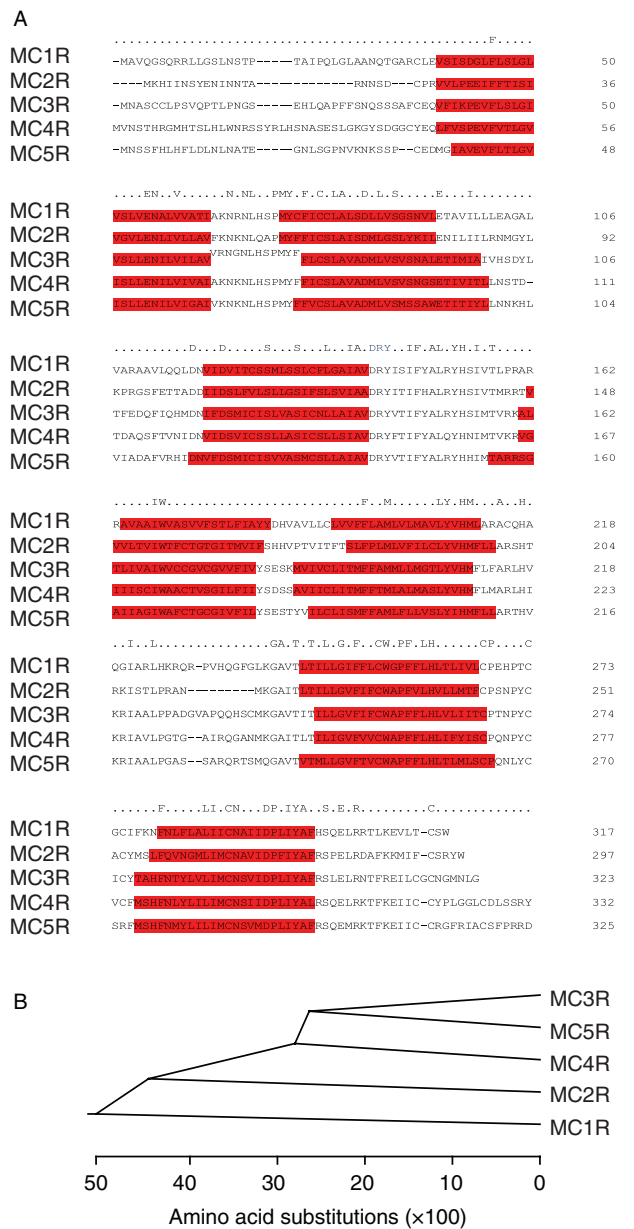
MC2R is also expressed in adipose tissue together with MC5R where they mediate the potent lipolytic effects of ACTH and  $\alpha$ -MSH (reviewed in Boston (1999) and Wikberg (1999)). At the same time, there is evidence that signalling mediated by MC2R and MC5R in adipocytes inhibits production of leptin, a hormone that completes a nutritional feedback loop to the hypothalamus (Zhang *et al.* 1994, Maffei *et al.* 1995), indicating that possibly there is a control mechanism for modulation of adipose tissue function via a melanocortin–leptin axis (Norman *et al.* 2003). MRAP was initially identified as a small protein upregulated in the differentiating mouse adipocyte 3T3-L1 cell line (Xu *et al.* 2002) and more recently was shown to be highly upregulated in the differentiation of mesenchymal stem cells into adipocytes, pointing to a role in adipogenesis (Menssen *et al.* 2011).

Interestingly, MC2R has a wide expression pattern during embryonic development and is also present in placenta in rodents (Izumi *et al.* 2004, Nimura *et al.* 2006). Placenta produces ACTH along with other pituitary-like peptides, and human foetal plasma ACTH levels are higher than those in maternal plasma (Krieger 1982, Petraglia *et al.* 1987), indicating the importance of ACTH during embryogenesis. In rodents, MC2R is highly expressed in a developing adrenal gland and is also found in lung where its function may be important for glucocorticoid-mediated lung maturation (Simard *et al.* 2010). MC2R is detected in developing testes where it may have a role in testosterone production by Leydig cells at an early stage of embryogenesis (O'Shaughnessy *et al.* 2003). MC2R is also found at different stages in the foetal brain, choroid plexus, genital ridge, mesonephros, metanephros and dorsal root and trigeminal ganglia, but its role in these tissues has not been investigated (Nimura *et al.* 2006). However, FGD type 1 patients with MC2R mutations are not known to have any congenital abnormalities other than glucocorticoid deficiency. Detailed study of MRAP and MRAP2 expression patterns during development would probably shed light on MC2R function regulation in the foetus.

Importantly, MRAP and MRAP2 are able to interact with all other MCRs, which broadens the potential range of MRAP function (Chan *et al.* 2009b; Fig. 3). For instance, *in vitro*, both MRAPs interact with MC4R but act in the opposite way to their effect on MC2R. Co-expression of MRAP and/or MRAP2 with MC4R in heterologous cells results in decreased surface expression of the receptor and reduction of signalling in response to NDP-MSH (Chan *et al.* 2009b). MC4R is highly expressed in PVN of the hypothalamus and nucleus accumbens and is involved in food intake regulation. Mutations in the *MC4R* gene are associated with severe obesity in early childhood (Vaisse *et al.* 1998, Yeo *et al.* 1998, Lee *et al.* 2008), and studies using knockout *Mc4r* mice showed that this obesity is due to hyperphagia (Huszar *et al.* 1997, Marsh *et al.* 1999). The majority of these mutations lead to intracellular retention of the receptor and lack of MC4R signalling (Farooqi *et al.* 2003, Lubrano-Berthelie *et al.* 2003) and have been reported in ~2–6% of the population. Polymorphisms V103I and I251L are thought to increase MC4R signalling, and affected individuals exhibit low body mass index (Stutzmann *et al.* 2007). MC4R signalling mediates some pathways of the stress response, and MC4R activation has been shown to result in anhedonia with weight loss (Lim *et al.* 2012). Hence, future drug targeting of MRAPs/MC4R could potentially alter appetite intake and body weight.

Both MRAPs interact with MC3R, which is involved in body weight and energy metabolism regulation. MRAPs regulate MC3R signalling but not surface expression, supporting the hypothesis of a dual functional role for MRAP proteins (Chan *et al.* 2009b). MC3R is highly expressed in ventromedial and arcuate nuclei of the hypothalamus. In knockout mice lacking the *Mc3r* gene, nutrients are preferentially partitioned into fat at the expense of lean mass, resulting in a reduction in locomotor activity and fine movements (Chen *et al.* 2000). Importantly, increased feed efficiency, not hyperphagia, causes weight increase in these mice (Chen *et al.* 2000). In humans, several polymorphisms of the *MC3R* gene are associated with high insulin levels and obesity in children (Feng *et al.* 2005). Although both MRAPs reduce MC3R signalling, it is more likely that MRAP2 regulates MC3R *in vivo* as both are mostly expressed in the hypothalamus. With the exception of one family, altered body weight has not been observed in patients with MRAP mutations, suggesting that MRAP may not be the key hypothalamic player (Rumie *et al.* 2007). However, this does not exclude the possibility of MC3R regulation by MRAP in other tissues where MC3R is also expressed such as in adipose tissue, heart (Chagnon *et al.* 1997), skeletal muscle, kidney (Chhajlani 1996), stomach, duodenum, placenta, pancreas (Gantz *et al.* 1993) and immunocompetent cells (Getting *et al.* 1999, Lindberg *et al.* 2005).

Another interesting interaction is with MC5R where both MRAPs have a significant negative effect on this receptor's surface expression and signalling. Despite a very wide expression pattern, the function of MC5R is mostly associated with exocrine regulation, and *Mc5r* knockout mice exhibit severe defects in water repulsion and thermoregulation due to decreased production of sebaceous lipids (Chen *et al.* 1997). Interestingly, MC5R mediates a pheromonal signal for aggression in mice (Morgan & Cone 2006). Previous work has indicated the requirement of  $\alpha$ -MSH in aldosterone production from the zona glomerulosa (Vinson *et al.* 1980, Shenker *et al.* 1985, Costa *et al.* 2011). As MC5R is expressed in the zona fasciculata, it has been suggested that  $\alpha$ -MSH may act through the MC5R, although this has yet to be verified. Along with MC2R, MC5R is also believed to play an important part in embryogenesis, having a similar tissue distribution. However, MC5R unlike MC2R is not found in placenta, is expressed at different stages, and therefore probably has a different role from MC2R (Nimura *et al.* 2006, Simamura *et al.* 2010). Mouse embryonic stem cells express MC5R instead of other MCRs and MC5R is suggested to be involved in the regulation of stem cell propagation

**Figure 3**

Conservation of the melanocortin receptors. (A) Alignment (ClustalW) of the protein sequences of human melanocortin receptors (NCBI, MC1R NP\_002377.4, MC2R NP\_000520.1, MC3R NP\_063941.3, MC4R NP\_005903.2 and MC5R NP\_005904.1). The classical feature of the GPCR family, seven predicted TM regions, is highlighted in red. The amino acids identical between all five receptors are shown above the alignment. Because MRAPs interact with all the melanocortin receptors, it is tempting to suggest that via a single TM domain, MRAPs bind MCRs within the regions of highest similarity between the receptors such as TM2, TM5 and TM6. (B) Phylogenetic tree of the melanocortin receptors showing evolutionary relationships within this GPCR family.

(Ogawa *et al.* 2004). In cultured B-lymphocytes,  $\alpha$ -MSH binding to MC5R activates the Jak/Stat pathway that is normally activated by cytokines, indicating that MC5R might have some function during immune response (Buggy 1998). It would be interesting to investigate whether regulation of MC5R by MRAPs has any role in the above functions of MC5R, perhaps revealing another level of GPCR regulation *in vivo*.

There is little suggestion of a role for MRAP in MC1R regulation as there is no significant effect on its surface expression and signalling. MRAP2 has a slight inhibitory effect on MC1R signalling without affecting the receptor presence on the plasma membrane (Chan *et al.* 2009b). There is no evidence for MRAP2 being expressed in the skin where MC1R is believed to function (Robbins *et al.* 1993). Using immunohistochemistry, it was shown that MC1R might be expressed in the placenta (Thornwall *et al.* 1997), but whether MRAP2 is expressed in this tissue or whether it has a regulatory role there is not known.

Interestingly, patients with FGD exhibit a dark skin colour, which is thought to be due to elevated ACTH levels acting on MC1R receptors expressed in melanocytes (Chung *et al.* 2010). This suggestion is supported by the recent case of a patient with FGD with mutations in both MC2R and MC1R who exhibited normal skin colour (Turan *et al.* 2012). It has also been proposed that changes in MC1R function can be associated with poor tanning response and melanoma (Valverde *et al.* 1995, 1996). There is some evidence that skin has an equivalent of the HPA regulated by MC1R and MC2R and is able to secrete cortisol (Ito *et al.* 2005), and MRAPs may potentially have a regulatory function on MCRs here.

Another potential role of MRAPs in signalling mediated by MCRs could be in manipulating their ligand preference. It is known that only MC2R responds to ACTH alone, whereas all the other MCRs also bind other POMC-derived peptides but with different affinities (Cone *et al.* 1993). The effect of MRAPs on the other MCRs' signalling has mostly been studied using a synthetic  $\alpha$ -MSH (Chan *et al.* 2009b, Sebag & Hinkle 2009b, 2010). However, it is known that MC3R is activated by  $\gamma$ -MSH and ACTH rather than  $\alpha$ -MSH, and MRAPs possibly have a different effect on MC3R signalling in response to these two hormones rather than the suppression shown for  $\alpha$ -MSH (Roselli-Reh fuss *et al.* 1993, Chan *et al.* 2009b). MC4R responds to  $\alpha$ -MSH with the same affinity as for ACTH, but the binding to these different ligands may result in distinct pathways being triggered within the cell. How accessory proteins regulate the affinity of these receptors for different POMC peptides needs further elucidation. The interaction of

MRAP2 with MC3R and MC4R is particularly intriguing as they all have hypothalamic expression; future studies will uncover physiological functions of MRAP2. The interplay between MRAP and MRAP2 in MCR regulation during embryonic development and into adulthood requires further investigation.

## Conclusions

The melanocortin family of proteins has diverse roles in the regulation of physiological processes. Identifying mutations in MC2R as causative of FGD triggered years of intensive research on MC2R regulation. Finding its accessory protein MRAP, where mutations result in FGD type 2, provided further insight into the regulation of MC2R function. MRAP regulates MC2R surface expression and facilitates its signalling. The role of the MRAP homologue, MRAP2, in relation to MC2R regulation needs exploration. However, it is more likely that the function of MRAP2 lies beyond MC2R and adrenal physiology. It is likely that MRAP2 is involved in the functioning of other MCRs such as MC4R and/or MC3R, thereby in the central regulation of energy metabolism and food intake. MRAP and MRAP2 may also have a role as regulators of MCRs during embryonic development. Further studies using animal transgenic models and genetic analysis are needed to elucidate *in vivo* functions of MRAP and MRAP2 and their importance. Understanding these functions will allow future research to develop new therapies for diseases caused by MCR dysfunction.

### Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the review reported.

### Funding

Research on MRAPs is supported by an MRC/Academy of Medical Sciences Clinician Scientist Fellowship to L F C (grant number G0802796).

### Acknowledgements

This article is based on the presentation by Dr L J Chan for the JOE/JME Prize Lecture 2012 at ICE/ECE 2012, Florence, Italy.

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Received in final form 30 January 2013

Accepted 11 February 2013

Accepted Preprint published online 11 February 2013