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## The biology of mammalian multi-copper ferroxidases

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### Abstract

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The mammalian multicopper ferroxidases (MCFs) ceruloplasmin (CP), hephaestin (HEPH) and zyklopen (ZP) comprise a family of conserved enzymes that are essential for body iron homeostasis. Each of these enzymes contains six biosynthetically incorporated copper atoms which act as intermediate electron acceptors, and the oxidation of iron is associated with the four electron reduction of dioxygen to generate two water molecules. CP occurs in both a secreted and GPI-linked (membrane-bound) form, while HEPH and ZP each contain a single C-terminal transmembrane domain. These enzymes function to ensure the efficient oxidation of iron so that it can be effectively released from tissues via the iron export protein ferroportin and subsequently bound to the iron carrier protein transferrin in the blood. CP is particularly important in facilitating iron release from the liver and central nervous system, HEPH is the major MCF in the small intestine and is critical for dietary iron absorption, and ZP is important for normal hair development. CP and HEPH (and possibly ZP) function in multiple tissues. These proteins also play other (non-iron-related) physiological roles, but many of these are ill-defined. In addition to disrupting iron homeostasis, MCF dysfunction perturbs neurological and immune function, alters cancer susceptibility, and causes hair loss, but, despite their importance, how MCFs co-ordinately maintain body iron homeostasis and perform other functions remains incompletely understood.

### Keywords

Hephaestin; Ceruloplasmin; Zyklopen; Ferroxidase; Copper; Iron

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### Introduction

The MCFs are representatives of the large and diverse blue copper protein family (Rydén and Hunt 1993; Nakamura and Go 2005). The blue copper proteins vary greatly in size and complexity, and contain up to 9 copper atoms. Their functions are also broad, and include proteins involved in iron homeostasis (CP, HEPH, ZP), electron transfer (plastocyanin), blood clotting (coagulation factors V and VIII), ascorbate metabolism (ascorbate oxidase), oxidation of poly-phenols (laccase), and oxidative phosphorylation (cytochrome c oxidase subunit II). The relatively complex mammalian members of this family evolved from an ancestral single-domain, cupredoxin-fold protein, and the MCFs are the only known members of the blue copper protein family in mammals that contain multiple copper atoms (Nakamura and Go 2005).

CP is the most studied MCF and is the major ferroxidase in the blood (Hellman and Gitlin 2002). It was first identified in 1948 by Holmberg and Laurell as an abundant plasma protein with a distinctive sky blue color, and was subsequently shown to be synthesized and secreted, mainly by hepatocytes (Hellman and Gitlin 2002). Other cell types, including astrocytes and neurons in the brain, Sertoli cells in the testes and cells in the retina, lung and pancreas can also synthesize CP. While most CP is secreted into the extracellular fluids, alternative splicing of the CP transcript leads to production of a GPI-linked form of the protein which tethers it to the plasma membrane (Fig. 1). GPI-CP is expressed in astrocytes (Patel and David 1997; Jeong and David 2003), Sertoli cells, hepatocytes, macrophages and retinal epithelial cells (Mostad and Prohaska 2011; Marques et al. 2012).

HEPH is a protein critical for intestinal iron absorption and was cloned by identifying the gene mutated in the sex-linked anemic (*sla*) mouse (Vulpe et al. 1999). Unlike the secreted form of CP, HEPH is tethered to membranes by a single C-terminal transmembrane domain (Fig. 1). HEPH is strongly expressed in the small intestine, and at low to moderate levels in multiple other tissues, most notably the CNS (Vulpe et al. 1999; Hudson et al. 2009; Jiang et al. 2015).

ZP is the least studied MCF. It was identified relatively recently as a possible placental iron oxidase (Chen et al. 2010), although this now seems unlikely as placental iron transfer is not impeded in mice in which the gene encoding ZP (*Heph1l*) is knocked out, at least under standard (unstressed) physiological conditions (Helman et al. 2021). The only confirmed role for ZP at present is a contribution to normal hair development (Eragene et al. 2019; Sharma et al. 2019; Kuca et al. 2021). Like HEPH, ZP contains a single C-terminal transmembrane domain. ZP is most strongly expressed in the placenta, mammary gland, kidney, brain and testis (Chen et al. 2010).

The MCFs are very similar structurally (Fig. 1). At the amino acid level, HEPH and ZP show 50% and 46% identity to CP respectively, while HEPH and ZP are 49% identical (Vulpe et al. 1999; Chen et al. 2010). The residues involved in copper binding and disulfide bond formation (the latter suggesting similar folding of the proteins) are highly conserved between the three proteins. In addition, both HEPH and ZP contain a single C-terminal transmembrane domain of 86 amino acids, although the sequences of these domains are only 23% conserved at the protein level (Chen et al. 2010).

Most of our current knowledge about the structure of the mammalian MCFs comes from the detailed characterization of CP (Zaitsev et al. 1999). It is a large protein consisting of 1046 amino acids that are arranged in three major homology units, each of which appears to have arisen from internal duplication of an ancient cupredoxin fold, to give a six-domain protein (Ortel et al. 1984; Nakamura and Go 2005). An X-ray crystallographic structure of CP is available (Zaitseva et al. 1996; Bento et al. 2007), and this confirmed the domain organization of the protein predicted by earlier sequence analysis (Ortel et al. 1984). The six domains are each made up of a plastocyanin-like fold and beta-barrel strands organized similar to those found in the cupredoxin family of metalloreductases (Bento et al. 2007). A key feature of CP's structure, and one that underlies its biological activity, is the presence of three types of copper-containing sites, each of which has distinct spectroscopic properties resulting from their specific coordination environments. Three of the six copper atoms in CP are distributed between three type I copper sites, in domains 2, 4, and 6, and the remaining three copper atoms are in a distinctive type II/type III trinuclear cluster at the interface of domains 1 and 6. Molecular modelling of HEPH and ZP reveals predicted structures that closely align with the known structure of CP (Syed et al. 2002; Chen et al. 2010).

Both CP and HEPH have been shown to possess a high affinity ferrous iron binding site near the type I copper atom in domain 6, and ZP is predicted to possess a similar site (Lindley et al. 1997; Brown et al. 2002; Vashchenko and MacGillivray 2012). Additional iron-binding sites have also been demonstrated in CP (a second site in domain 4) and HEPH (additional sites in domains 2 and 4), but they are of lower affinity (Lindley et al. 1997;

Vashchenko and MacGillivray 2012; Zaitsev and Lindley 2019). Following binding, ferrous iron predicted to deliver an electron to the nearest type I copper site. This is associated with the transfer of the oxidized iron to an adjacent ferric iron holding site. Electrons are subsequently transferred through the protein to the type I copper site in domain 6, and then to the closely associated type II/type III trinuclear cluster where molecular oxygen binds. The Type II/Type III site can then sequentially transfer a total of four electrons, originally donated from different ferrous iron atoms, to dioxygen to form water (Lindley et al. 1997; Vashchenko and MacGillivray 2012). The MCFs can also oxidize a range of substrates in addition to iron, and, in CP, many of these bind at sites different from the iron binding sites (Bielli and Calabrese 2002).

Some of the residues important for ferroxidase and amine oxidase activity in CP have been identified by site directed mutagenesis, (Brown et al. 2002) and most, but not all of these are conserved in HEPH and ZP (Chen et al. 2010). CP interestingly has a non-typical, permanently-reduced type I copper site in domain 2 that is believed to be non-functional, while all three type I copper sites are predicted to be functional in HEPH and ZP (Chen et al. 2010). In CP, the copper atoms in two of the three type I sites are coordinated in a trigonal pyramid arrangement, with three ligands (a cysteine and two histidines) arranged in a plane, and the fourth, a methionine, at the apex of the pyramid. In the third (non-functional) type I site, however, the apical methionine has been replaced by a leucine residue. In contrast, in both HEPH and ZP, the third type I site shows the 'typical' coordination arrangement with an apical methionine.

Not only is copper required for the ferroxidase activity of CP and the other MCFs, it is also structurally important and the *apo*-proteins (lacking copper) are inherently unstable (Gitlin et al. 1992; Reeves and Demars 2005; Chen et al. 2006). The half-life of *apo*-CP is approximately five hours whereas the *holo*-protein has a half-life of over five days (Hellman and Gitlin 2002). Although the precise details of the mechanisms responsible for the metalation of CP are not known, the necessary copper is transported into the *trans*-Golgi network by ATPase copper-transporting Beta (ATP7B) where the metal is inserted into the nascent protein (Sato and Gitlin 1991; Hellman et al. 2002; Lutsenko et al. 2007). In cases of ATP7B deficiency (Wilson Disease), this process is less efficient and much of the CP that is produced lacks both copper and oxidase activity (Członkowska et al. 2018). Consequently, measurements of serum ferroxidase activity are a useful diagnostic tool for Wilson Disease and for assessing moderate to severe copper deficiency (Sánchez-Monteagudo et al. 2021). It is likely that the biosynthetic pathways for HEPH and ZP are similar to that of CP, but these processes have yet to be studied in detail. There is certainly very strong evidence that the biological activities of HEPH and ZP are also copper-dependent (Chen et al. 2006; Chen et al. 2010).

### **The role of the MCFs in iron homeostasis**

In 1966, Frieden and Osaki demonstrated that plasma CP had the ability to oxidize ferrous iron (Osaki et al. 1966) and subsequently showed that the protein was important for mobilizing iron from the liver (Osaki et al. 1971). These pioneering studies, along with an important series of experiments in pigs showing that copper deficiency was associated

with anemia (Lee et al. 1968; Roeser et al. 1970), highlighted what is now known to be a very close physiological relationship between copper and iron (Gulec and Collins 2014).

Understanding how the MCFs function requires a basic understanding of iron homeostasis. Despite being essential, iron is also toxic when present in excess as it can catalyze the formation of reactive oxygen species. To meet cellular iron requirements while preventing iron accumulating to toxic levels, cellular and body iron concentrations must be kept within defined limits. To achieve this, both dietary iron absorption and internal iron trafficking are tightly regulated (Anderson and Frazer 2017). Most iron within the body is used for the production of hemoglobin in red blood cells. At the end of their functional life, senescent red cells are phagocytosed by macrophages, which can store iron or return it to the circulation, a process referred to as iron recycling. In addition, iron in excess of immediate metabolic needs may be stored within tissues, particularly in the liver, but it can be readily mobilized in times of metabolic need. Dietary iron absorption, internal iron recycling and mobilization of stored iron are all critically dependent on MCFs. This requirement is best illustrated by situations in which MCF dysfunction occurs. In the rare genetic disease aceruloplasminemia, CP is either absent or expressed at very low levels (Marchi et al. 2019). This condition is typified by extensive iron deposition in many tissues, illustrating the key role that CP plays in iron mobilization. Subsequent studies with *Cp* knockout mice, and analogous work with mouse models of HEPH deficiency, have supported and extended these early findings (Harris et al. 1999; Fuqua et al. 2014; Fuqua et al. 2018).

Iron is exported from cells as  $\text{Fe}^{2+}$  via the unique iron export protein ferroportin (FPN), however,  $\text{Fe}^{2+}$  must be oxidized before it can bind to transferrin, the major plasma iron transport protein (Anderson and Vulpe 2009). The MCFs likely facilitate this process, but how they contribute to cellular iron efflux is incompletely understood. The membrane-bound MCFs HEPH and GPI-CP have been demonstrated to interact physically with FPN (Jeong and David 2003; Yeh et al. 2009, 2011; Dlouhy et al. 2019), but there is not yet any convincing data that the soluble form of CP behaves similarly (Musci et al. 2014). FPN has been shown to co-localize with GPI-CP in astrocytes (Jeong and David 2003) and with HEPH in enterocytes (Yeh et al. 2009; Fuqua et al. 2018). It has been suggested that MCFs are able to oxidize  $\text{Fe}^{2+}$  that has been transported by FPN and that this oxidation is important for the release of the iron from the transporter (De Domenico et al. 2007). If iron is not released from FPN, the transporter is ubiquitinated, internalized and degraded, which suggests that the MCF is important in maintaining FPN stability. Although, this is an attractive hypothesis, the data have not yet been independently verified, and this work was mentioned in a scientific misconduct investigation (McCormack 2013). Nevertheless, there is some support for this concept. For example, brain microvascular endothelial cells express both HEPH and soluble CP, and when the levels of these proteins are depleted by copper chelation, FPN levels are also reduced (McCarthy and Kosman 2013). Similarly, knockdown of HEPH in neurons reduces FPN levels and iron export (Ji et al. 2018). Interestingly, however, loss of HEPH in the small intestine is not associated with any overt decline in FPN expression (Fuqua et al. 2018), suggesting that the MCFs may only stabilize FPN in some tissues. There is also some evidence that GPI-CP is able to interfere with the binding of hepcidin to FPN (Kono et al. 2010). Hepcidin is a liver-derived peptide hormone that is considered the master regulator of systemic iron homeostasis, and it acts by binding to FPN

and removing it from the cell surface, consequently blocking iron export. This interaction could be a second mechanism by which MCFs facilitate iron efflux, and is consistent with a close physical association between FPN and the MCF, but it requires further investigation.

Essentially all of the (ferrous) iron that is exported from cells via FPN and oxidized by an MCF is bound by transferrin (TF) in the plasma, and this protein can then deliver iron to the various sites in the body where it is required (Anderson and Vulpe 2009). TF binds ferric iron ( $\text{Fe}^{3+}$ ), thus oxidation of  $\text{Fe}^{2+}$  by MCFs could not only facilitate the release of iron from FPN (as discussed above), but also permit binding of  $\text{Fe}^{3+}$  to TF. Although  $\text{Fe}^{2+}$  will auto-oxidize in the plasma, this process is considered too slow for the trafficking of iron under physiological conditions (Osaki et al. 1966). While some studies have found evidence for a physical association between CP and TF (Ha-Duong et al. 2010), others have found no evidence that either CP or HEPH can stably interact with TF (Hudson et al. 2008). A strong interaction between CP and lactoferrin (LF), a close relative of TF, has been identified (Zakharova et al. 2000; Ha-Duong et al. 2010), although the physiological relevance of this interaction has yet to be established.

In addition to its well-described role in cellular iron efflux, CP has been shown to stimulate the uptake of iron by some mammalian cells in culture using a TF-independent pathway (Mukhopadhyay et al. 1998; Attieh et al. 1999). In this case, it is proposed that non-transferrin-bound iron (NTBI) is reduced by a ferrireductase and that the resulting ferrous iron is oxidized by CP to make it available for a trivalent cation-specific transporter (Attieh et al. 1999). However, the identity of this uptake pathway remains unknown, and further work is required to determine its physiological relevance in vivo.

### Other physiological roles of MCFs

Although their role in oxidizing iron is clear, MCFs have broader roles and can utilize a range of other substrates. Some of these are summarized in Table 1. The non-ferroxidase activities of the MCFs have only been extensively studied in the case of CP. In addition to iron, CP can oxidize a range of organic amines, including biogenic amines (such as catecholamines, serotonin and dopamine), low-density lipoprotein (LDL), and nitric oxide (NO) (Bielli and Calabrese 2002; Vashchenko and MacGillivray 2013). HEPH and ZP can also oxidize amines (e.g. *para*-phenylene diamine) (Vulpe et al. 1999; Chen et al. 2010; Deshpande et al. 2017), so it is quite possible that they also have a broad spectrum of catalytic actions. Studies with CP and recombinant HEPH have demonstrated that they share the ability to oxidise some amines, but not others (Vashchenko et al. 2011), so there are likely to be some differences in the range of activities that individual MCFs participate in.

The physiological significance of many of these non-iron-related activities has never been clearly defined. For example, in humans with aceruloplasminemia and *Cp* knockout mice, the dominant phenotypes are either directly or indirectly related to tissue iron accumulation (Harris et al. 1999; Ayton et al. 2013; Marchi et al. 2019), but this does not exclude more subtle effects on other aspects of body physiology. It has been estimated that no more than 5–10% of the CP in plasma is sufficient to perform all its iron-related functions (Roeser et al. 1970; Osaki et al. 1971), so is the excess simply 'reserve' iron oxidizing capacity, or does the protein perform other functions? Furthermore, HEPH is expressed at high

levels throughout the small intestine and colon, even though dietary iron absorption occurs mainly in the duodenum (Frazer et al. 2001). Its function in the regions of the gut that absorb little iron is unknown. There is also considerable evidence that CP can act as an anti-oxidant (Goldstein 1983; Song et al. 2010; Bakhautdin et al. 2013; Vasilyev 2019), and HEPH and ZP may play similar roles, but this function of the MCFs has never been investigated systematically. Interestingly, CP can also act as a pro-oxidant, causing damage to macromolecules such as LDL (Mukhopadhyay and Fox 1998; Fox et al. 2000), and thus may act as a risk factor for the development of atherosclerotic disease.

Although the main physiological roles of the MCFs most likely relate to iron homeostasis, they also are relevant to the biology of other metals. For example, CP possesses cuprous oxidase activity (Mukhopadhyay et al. 1997; Stoj and Kosman 2003) and may play a role in copper distribution in some circumstance (such as copper transport to the fetus and neonate) (Lee et al. 1993; Chu et al. 2012; Kenawi et al. 2019). Indeed, CP-bound copper represents the largest pool of copper in the circulation, and it has been suggested that it may provide an important pathway for delivering copper to a range of tissues (Linder 2016). Nevertheless, there is no evidence for altered copper homeostasis in aceruloplasminemia (Meyer et al. 2001). Studies with CP-deficient rodents have also shown that CP plays a role in the body distribution of manganese (Jursa and Smith 2009; Kenawi et al. 2019). Whether HEPH or ZP also contribute to the homeostasis of metals other than iron has yet to be determined.

In addition to its enzymatic roles, CP has been shown to bind to a range of other proteins, thereby affecting their activity (Vasilyev 2019). These include myeloperoxidase (MPO), matrix metalloproteinases 2 and 12, eosinophil peroxidase, 5-lipoxygenase and coagulation factor FIIa. CP binding to other iron-associated proteins such as ferritin (Van Eden and Aust 2000) and lactoferrin (Zakharova et al. 2000; Pulina et al. 2002) has also been reported. The binding of iron by LF is facilitated by CP, and it has been suggested that this is one mechanism by which the host might protect itself against the respiratory burst that occurs when neutrophils infiltrate a site of inflammation or infection (Sokolov et al. 2005). Neutrophils are a rich source of LF, and secreted LF has the capacity to bind a considerable amount of iron at the site of inflammation, particularly in the presence of CP. Another potential anti-inflammatory function of CP is its ability to bind to MPO and inhibit the production of hypochlorous acid (Sokolov et al. 2007; Chapman et al. 2013).

### Regulation of the MCFs

Given their important role in cellular iron efflux, it is not surprising that the expression of MCFs is increased under iron deficient conditions (Vulpe et al. 1999; Hellman and Gitlin 2002; Chen et al. 2010; Ranganathan et al. 2011). This response is likely an adaptation to support hemoglobin production in developing red cells when iron supply is limiting. The mechanisms responsible for this regulation have not been defined. The expression of many of the central proteins in iron homeostasis is regulated post-transcriptionally (either translationally or via mRNA stability) by the binding of iron regulatory proteins (IRPs) to iron regulatory elements (IREs) in target transcripts (Zhang et al. 2014), but none of the mRNAs encoding MCFs appears to contain an IRE. Thus, alternative transcriptional or post-transcriptional mechanisms must be involved. CP, at least, can be regulated by a combination



of transcriptional and translational mechanisms by factors such as hypoxia (Chepelev and Willmore 2011; Linder 2016), proinflammatory cytokines (Gitlin 1988; Sampath et al. 2003; Conley et al. 2005; Mazumder et al. 2006; Persichini et al. 2010; Linder 2016; Tisato et al. 2018) and oxidative stress (Das et al. 2007; Tapryal et al. 2009; Tisato et al. 2018). The other MCFs have not been extensively investigated, however, HEPH has been shown to be regulated by transcriptional and epigenetic mechanisms. In the small intestine and colon, the *HEPH* gene is a prominent target of the transcription factor CDX2 (Hinoi et al. 2005), which plays a major role in intestinal development and cancer. In addition, CDX2 itself was found to be regulated by iron, providing a transcriptional loop for the regulation of HEPH in the absorptive epithelium of the gut (Hinoi et al. 2005). The transcription of *HEPH* is also regulated by the H3K9 methyltransferase G9a (Wang et al. 2017), which represses *HEPH* expression, resulting in cellular iron accumulation and enhanced growth of breast cancer cells. Other epigenetic modes of regulation of the MCFs are likely to be uncovered upon further investigation.

Changes in the subcellular localization of membrane proteins represent another mode of regulation which is particularly useful as a rapid response mechanism. Several studies have demonstrated that HEPH will re-localize from an intracellular supra-nuclear compartment to the basolateral membrane (or a compartment in close proximity to the basolateral membrane) of polarized epithelial cells in response to iron treatment (Yeh et al. 2009; Lee et al. 2012). This likely represents an adaptation to reduce the risk of intracellular iron increasing to toxic levels by stimulating expression of components of the iron export pathway. This effect has also been observed in non-polarized cells (Dlouhy et al. 2019).

### Genetics and the pathology of MCF loss or dysfunction

Loss of MCF function leads to increased cellular iron accumulation and its associated pathological consequences (Hellman and Gitlin 2002; Musci et al. 2014; Linder 2016). The best characterised MCF disturbance in humans is aceruloplasminemia, in which iron accumulates in multiple tissues, particularly the liver, CNS and pancreas (Kono 2013; Piperno and Alessio 2018; Marchi et al. 2019). Liver injury, neurodegenerative disease, diabetes and retinal degeneration are common complications. Reduced CP expression has also been implicated in atherosclerosis (Ji et al. 2015), while increased expression has been noted in various cancers (Gangoda et al. 2017; Matsuoka et al. 2018; Dai et al. 2020; Zhang et al. 2021b). Fewer data are available on HEPH or ZP deficiency. Polymorphisms in HEPH have been associated with migraine (Maher et al. 2012; Albury et al. 2017), and reduced HEPH expression has been noted in atherosclerotic plaques (Ji et al. 2015) and is linked to the iron deficiency associated with a high-fat diet (Albury et al. 2017). Increased HEPH (and CP) levels have also been found in the placenta from a GRACILE syndrome fetus (Kotarsky et al. 2010). A patient carrying a missense mutation in *HEPHL1*, the gene which encodes ZP, had notable hair defects, including pili torti (flattened, twisted hair) as well as behavioural abnormalities (Sharma et al. 2019). A missense mutation in *HEPHL1* has also been identified in patients with Parkinson's disease (Al Yemni et al. 2019).

Rodent strains with disturbed MCF function have proven particularly important in revealing some of the physiological roles of these proteins. No natural or induced



Supporting this observation is the recent identification of *HEPHL1* as the gene affected in the Curly Whiskers mouse (Eragene et al. 2019), and demonstration that a mutation in *HEPHL1* is responsible for the hair abnormalities in Belted Galloway Cattle (Kuca et al. 2021). While the available data suggest that ZP plays an important role in hair development, whether this role reflects altered iron homeostasis or some other function of ZP has yet to be determined. Whether ZP plays roles in other tissues has not yet been determined.

More recently, mice with disrupted expression of both HEPH and CP have been generated and analysis of their phenotype has clearly shown that both of these MCFs play roles in multiple tissues. Such studies were first carried out by examining the progeny of crosses of *Cp* knockout and *sla* mice (Hahn et al. 2004), and more recently by generating animals in which each of these genes has been deleted (Fuqua et al. 2018). These studies have shown that the phenotype arising in the mice when both MCFs are disrupted is more severe than that observed in either of the single mutants. Double knockout mice were severely anemic and had defects in intestinal iron absorption, but they also loaded iron in a broad range of tissues, including the heart, kidney, pancreas, retina and adrenal gland. In these extra-intestinal tissues, little or no iron loading was observed in either single knockout, indicating that the MCFs work together to facilitate iron release from these tissues. Interestingly, even mice lacking both ferroxidases were still able to absorb orally administered iron, but that iron was distributed in an unusual pattern, with particularly high retention in the liver (Fuqua et al. 2018). It is possible that in these mice, the limited ferroxidase activity meant that iron loading onto transferrin was inefficient, and consequently that the level of NTBI was unusually high. NTBI is very efficiently taken up by the liver (Craven et al. 1987). Although there is no report in the literature describing the consequences of the combined deletion of ZP and CP, or ZP and HEPH, our unpublished studies have shown that the progeny of both crosses are viable.

Since an adequate iron supply is required to support cancer cell growth, there are numerous links between the expression of various proteins of iron homeostasis, including the MCFs, and tumor progression (Chen et al. 2019a). Much of the interest in this area in recent years has focused on HEPH. Reduced HEPH protein levels have been associated with more advanced disease in mesothelioma (Crovella et al. 2016), breast cancer (Wang et al. 2017) and colorectal cancer (Brookes et al. 2006), and high HEPH expression has been linked to improved prognosis in lung cancer (Zacchi et al. 2021) and renal clear cell carcinoma (Chen et al. 2019a). In breast cancer, both lower HEPH (Wang et al. 2017) and lower FPN (Pinnix et al. 2010) expression are associated with iron accumulation and more aggressive tumours. Together, these data indicate that reduced iron export favours tumour growth. The recent demonstration that Euchromatic histone-lysine N-methyltransferase 2 (EHMT2) can repress HEPH transcription (Wang et al. 2017) in breast cancer cells makes it a potential therapeutic target for modulating HEPH expression in tumours. Indeed, EHMT2 is frequently elevated in breast cancer (Wang et al. 2017). Reduced HEPH expression has also been linked to enhanced resistance of colon cancer cells to the chemotherapeutic drug methotrexate (Zhang et al. 2021a). In this case, the mechanism involves a regulatory path-way whereby the microRNA miR-24-3p released from cancer-associated fibroblasts acts on colon cancer cells to reduce the activity of the transcription factor CDX2, a major (positive) regulator of the *HEPH* gene. Although reduced HEPH levels show a clear link to a more severe

tumor phenotype, studies with CP have shown the opposite relationship, with increased CP expression being associated with poorer prognosis (Gangoda et al. 2017; Matsuoka et al. 2018; Zhang et al. 2021b). Much of this increase may reflect the fact that CP is an acute phase protein, but some studies have indicated that CP itself can drive cellular changes that promote tumorigenesis (Dai et al. 2020; Zhang et al. 2021b). Understanding how the cellular iron export machinery contributes to tumor progression and exploring the MCFs as therapeutic targets are areas that warrant further investigation.

### **The need for multiple ferroxidases: tissue and organ-specific roles of the MCFs**

As CP is secreted into the plasma, it might be expected that it could be involved in iron mobilization from most cells, but, surprisingly, it only appears to perform this role efficiently for some tissues. In other tissues, either other MCFs are more important, or MCFs work together to coordinate iron efflux. As noted above, the characterization of rare human diseases where MCF function is disturbed and a range of MCF knockout/mutant mouse strains has demonstrated that CP is particularly important for facilitating iron efflux from the liver, and certain cell types in the brain and pancreas (Hellman and Gitlin 2002; Jeong and David 2003; Linder 2016), whereas HEPH is the major MCF functioning in the small intestine (Fuqua et al. 2014). However, more recent studies with mice lacking both CP and HEPH have indicated a more complex picture, with both MCFs contributing to iron efflux from multiple peripheral tissues (Fuqua et al. 2018). Tissue-specific knockouts of particular proteins in mice can be a powerful tool for assessing their function in individual tissues, but very few tissue-specific MCF knockout mice have been generated to date (e.g. HEPH deletion in the intestine (Fuqua et al. 2014) and retina (Wolkow et al. 2012)).

The MCFs clearly function in multiple tissues, but do so with varying efficiencies and in varying combinations. This may reflect factors such as accessibility to circulating CP, varying MCF expression levels, differential subcellular distribution (particularly for membrane-bound MCFs), or even differences in their catalytic profile. A few remarks on the contributions of the MCFs to iron homeostasis in selected tissues are provided below and an overview is provided in Fig. 2.

**Liver**—The liver is quantitatively the major iron storage organ in the body, so it is critically important that iron is efficiently mobilized from the liver when required (Anderson and Frazer 2005). CP is clearly the major MCF involved in liver iron mobilization, as hepatic iron levels increase substantially when CP is missing or dysfunctional (Harris et al. 1999; Piperno and Alessio 2018; Kenawi et al. 2019; Marchi et al. 2019). Loss of HEPH either globally or specifically in the small intestine leads to a reduction in dietary iron absorption, and consequently hepatic iron levels are low (Fuqua et al. 2014). If HEPH played a significant role in hepatic iron release, liver iron levels might be expected to increase, or at least not decrease, but this is not the case. The low expression of HEPH in the liver is consistent with it playing a limited role in this organ (Vulpe et al. 1999; Frazer et al. 2001; Hudson et al. 2009). Nevertheless, studies with CP/HEPH double knockout mice have suggested that lack of HEPH in the intestine may influence the kinetics of iron flux through the liver (Fuqua et al. 2018), and this requires further investigation.

**Small intestine**—The best characterized role of HEPH is its contribution to dietary iron absorption, as it is required at the basolateral membrane of enterocytes to facilitate the efficient export of iron into the portal circulation (Fuqua et al. 2014; Fuqua et al. 2018; Doguer et al. 2017). Early studies with *sla* mice demonstrated this phenotype, even though at the time HEPH had not been identified. *Heph* knockout mice show a similar phenotype to *sla* animals, but loss of HEPH does not completely block iron absorption and the anemia observed in intestine-specific HEPH knockouts is relatively mild (Fuqua et al. 2014). Indeed, the anemia in these mice is milder than that of whole body HEPH knockout mice, indicating that HEPH makes contributions to iron supply to the plasma from tissues other than the gut. The fact that dietary iron absorption can continue even in the absence of HEPH indicates that either another iron oxidase performs this function or that ferroxidase activity is not essential for iron export from enterocytes. Under normal physiological conditions, CP appears to make little contribution to dietary iron absorption, but when body iron demands are particularly high, such as following hemolysis, CP can help to mobilize iron from enterocytes (Cherukuri et al. 2005). The apparent lower efficiency of CP in facilitating dietary iron absorption may indicate that the precise subcellular location of the ferroxidase activity is important. Although HEPH is generally considered to act on the basolateral membrane and some localization studies support this contention (Han and Kim 2007), other data suggest HEPH could be located in an intracellular compartment very close to the membrane, rather than on the basolateral membrane itself (Kuo et al. 2004). Intriguingly, a ferroxidase activity that cannot be attributed to CP or HEPH has been detected in rodent enterocytes (Ranganathan et al. 2012), but its identity is not yet known. It has also been demonstrated that HEPH appears to be important for iron absorption under normal physiological conditions, but is not limiting when dietary iron absorption is stimulated by iron deficiency or following haemolytic anemia (Doguer et al. 2017), a finding consistent with early studies using *sla* mice (Sorbie et al. 1974) and supporting the suggestion that other ferroxidase activities are present.

**Central nervous system**—The important role the MCFs, and particularly CP, play in the CNS has been extensively documented (Piperno and Alessio 2018). Progressive neurodegeneration and retinopathy are characteristic features of aceruloplasminemia and reflect iron deposition in these tissues associated with loss of CP activity (Kono 2013; Marchi et al. 2019). Thus, CP can be considered a neuroprotective molecule. It is the GPI-linked form of CP in astrocytes that appears to be particularly important in brain iron metabolism, as it facilitates the export of iron from astrocytes, which makes this iron available for other types of cells in the brain (Jeong and David 2006; Marchi et al. 2019). When CP is missing or defective, not only is this iron supply disrupted, but iron can accumulate within astrocytes to cytotoxic levels and this in turn may exacerbate the neurodegeneration (Kono 2013). HEPH has also been shown to play a significant role in maintaining iron homeostasis in the brain, and mice lacking both functional CP and HEPH show greater iron accumulation in multiple brain regions than either of the individual mutant strains (Zhao et al. 2015; Zheng et al. 2018b). The double knockout animals also showed significant learning and memory defects (Zheng et al. 2018b). Interestingly, while CP is predominantly expressed in astrocytes, HEPH is most strongly expressed in oligodendrocytes. Loss of CP function in astrocytes or HEPH function in oligodendrocytes

leads to iron accumulation in each of these cell types and concomitant oxidative stress (Chen et al. 2019b). Loss of HEPH leads to iron accumulation in oligodendrocytes in the gray matter, but not in the white matter, with a compensatory increase in CP expression in the white matter cells explaining the difference (Schulz et al. 2011). The retina is another component of the CNS where MCFs play important roles. Retinal pigment epithelial (RPE) cells express both CP and HEPH, and mutation or loss of these proteins is associated with iron accumulation in these cells and retinal degeneration (Hahn et al. 2004; Hadziahmetovic et al. 2008; Wolkow et al. 2012). These ferroxidases play a very limited role in photoreceptor cells, but since global MCF knockout mice show more severe retinal damage than mice lacking the CP and HEPH only in the RPE, these MCFs must be contributing to retinal iron homeostasis via their roles in some other cell type(s) (Wolkow et al. 2012).

**Pancreas**—The pancreas is an organ that is particularly prone to iron loading (Kimita and Petrov 2020), so ensuring it has an efficient mechanism for divesting itself of iron is critical for its normal function. Diabetes is a common feature of aceruloplasminemia (Piperno and Alessio 2018), but interestingly, there is no overt iron loading in the endocrine pancreas in this condition (Kato et al. 1997), suggesting the effects on the endocrine cells are secondary to iron loading in other cell types. In mice, loss of either CP or HEPH alone appears to have little effect on iron levels in the pancreas, but when both genes are ablated together, significant iron loading is observed in this organ (Chen et al. 2018; Fuqua et al. 2018). Most of this iron loading is observed in the exocrine pancreas (i.e. the acinar cells) rather than in the Islets (i.e. the endocrine pancreas). This iron loading causes oxidative damage and enhanced production of plasma lipase and trypsin, which are typically elevated when the exocrine pancreas is damaged, but there were no significant changes in insulin levels (Chen et al. 2018). These studies clearly show the important protective role the MCFs play in the pancreas.

**Placenta**—Given the importance of MCFs in iron export from other tissues (Vashchenko and MacGillivray 2013), it is likely that a placental ferroxidase is required for efficient transfer of iron from the mother to the fetus, particularly in the final weeks of pregnancy. Although newborn *sla* mice, which essentially lack functional HEPH, are anemic and have reduced total body iron (Bannerman et al. 1973), radioactive iron transport studies did not indicate a defect in placental iron transfer (Kingston et al. 1978). Fetal CP has also been suggested to facilitate the export of iron from the placenta based on studies in BeWo cells (Danzeisen et al. 2000, 2002), but studies with CP knockout mice argue against CP playing a major role in iron delivery to the fetus (Harris et al. 1999). When ZP was identified, it was initially considered to be a putative placental MCF based on its high expression in the placenta at embryonic day 18 and its major contribution to the ferroxidase activity of BeWo cells (Chen et al. 2010). However, highest expression was observed in the implantation chamber at embryonic day 7, before the definitive placenta has formed. A recent study has shown that when ZP was knocked out in the labyrinth, the nutrient transporting layer of the placenta, there was no change in hemoglobin or total fetal iron content compared with littermate controls (Helman et al. 2021). Dissection of late-gestation placentas to separate the various layers revealed that expression of *Heph11*, the gene encoding ZP, is high in the

maternal decidua, but undetectable in the labyrinth, making it unlikely to play a role in iron export to the fetus (Helman et al. 2021). Furthermore, no iron phenotype was found under normal conditions when ZP was knocked out in the maternal decidua, but the placentas were 26% heavier, suggesting that ZP may play a role in placental development (Helman et al. 2021). Thus, it appears that no one MCF plays a dominant role in placental iron transport, but the studies conducted to date do not exclude a combined role of multiple ferroxidases.

**Hair**—An association between iron deficiency and hair loss has been recognized for some time, but the mechanisms involved are poorly understood (Trost et al. 2006; Almohanna et al. 2019). The MCFs may play an important role in hair development as disruption of either HEPH or ZP has been associated with an abnormal hair phenotype. The importance of ZP for maintaining normal hair growth was noted above. A patient carrying a mutation in *HEPHL1*, as well as *Zp*<sup>-/-</sup> and *Zp*<sup>ew/cw</sup> mice, all exhibit loss of ferroxidase activity and develop similar hair abnormalities, with reduced lysyl oxidase activity in fibroblasts (Eragene et al. 2019; Sharma et al. 2019). A recent genome-wide association study in Belted Galloway cattle with hypotrichosis demonstrated the alopecia phenotype is caused by a mutant variant in *HEPHL1* (Kuca et al. 2021). Together, these studies suggest that the skin-expressed ZP may play an important role in facilitating hair growth and preventing hair disorders. HEPH may also play a role in normal hair development. Pre-weaning pups born to or nursed by *Heph* knockout dams developed truncal hair loss (the “mask” phenotype) and systemic iron deficiency (Wolkow et al. 2012; Zhou 2020). Transcriptomic studies have indicated that the WNT/β-catenin signaling pathway, the key regulatory pathway for hair growth and hair follicle development, was downregulated in the affected skin (Zhou 2020). Preliminary data suggest that decreased expression and activity of stearoyl coenzyme A desaturase, an iron-containing enzyme essential for WNT palmitoylation (Stoffel et al. 2017), may provide the link between disturbed iron homeostasis and hair loss in the *Heph* deficient mice (Zhou 2020). In addition, single nucleotide polymorphisms in the *HEPH* gene have been associated with male pattern baldness in humans (Hagenaaers et al. 2017), however, whether the association reflects HEPH itself or the nearby androgen receptor gene has yet to be established.

**Other tissues**—Loss of either CP or HEPH in mice does not lead to overt iron accumulation in the heart, kidney or in adipose tissue, but when both of these MCFs are deleted, iron loading is observed (Qian et al. 2007; Jiang et al. 2016b; Fuqua et al. 2018). This finding suggests that the combined actions of these two ferroxidases are important in maintaining iron homeostasis in these tissues. In the kidney, loss of CP and HEPH leads to iron loading in both the medulla and cortex, and is associated with proteinuria and elevated urinary iron, consistent with renal damage (Jiang et al. 2016a). Some histological changes, particularly in the medulla, were also observed. In the heart, the iron loading in CP/HEPH double knockout mice was particularly apparent in cardiac myocytes (Fuqua et al. 2018). The iron loading that is observed in *Cp*<sup>-/-</sup>/*Heph*<sup>-/-</sup> adipocytes was associated with decreased leptin and adiponectin mRNA expression (Zheng et al. 2018a). Fasting glucose levels were significantly increased in the double knockout mice, and both glucose and insulin tolerance were impaired. This finding is of clinical interest given the recent observation that adipocyte iron levels regulate intestinal lipid absorption (Zhang et al. 2021c), and raises the

possibility that pharmacologically increasing MCF levels could prove beneficial by limiting lipid intake.

### Future areas of research

Although CP has been studied for many decades, and HEPH and ZP have been investigated for more than 20 and 10 years respectively, there are still many aspects of their biology that are incompletely understood. The following list highlights some of the areas where further investigation is warranted.

1. Although we have some valuable clues as to how the MCFs function in conjunction with FPN to facilitate efficient iron export, the precise details of this process remain unclear, and it is possible that MCF/FPN interactions will vary depending on the tissue and the MCF involved. Some key questions include: Do the membrane-tethered MCFs act in the same way as soluble CP? Is the oxidation of recently transported ferrous iron necessary for its release from FPN? What is the link between MCF action and FPN stability?
2. Understanding the structure of the MCFs individually, and when complexed with FPN (and potentially other proteins) will be extremely helpful in elucidating the mechanisms of MCF action.
3. The nature of the interactions of the MCFs with transferrin has yet to be defined.
4. How the three MCFs work together to co-ordinately regulate body iron homeostasis is incompletely understood. Studies with single and multiple MCF knockout in specific tissues are beginning to answer these questions, but more extensive investigations are required.
5. Why MCF ferroxidase activity is apparently in gross excess in the plasma and some other tissues needs to be resolved.
6. Are there ferroxidases other than CP, HEPH and ZP that play important roles in body iron trafficking?
7. Does ZP have physiological roles in addition to its described function in hair development?
8. Which MCFs contribute to iron recycling through macrophages and what are the relative contributions of cell-specific and systemic mechanisms?
9. The cell biology of the MCFs, particularly the membrane bound forms, is not well understood and warrants further investigation. This is particularly important as changes in the subcellular localization of these proteins appears to be a significant mode of regulation, at least in response to altered iron availability.
10. It is known that each of the MCFs is able to oxidize substrates in addition to  $\text{Fe}^{2+}$ , but how broad this spectrum of substrates is (particularly for HEPH and ZP) has not been determined and the physiological relevance of many of these 'non-iron' activities has yet to be defined.



11. Why does the loss of ferroxidase activity in CP or HEPH knockout mice not lead to a phenotype as severe as might be expected, either in terms of tissue iron loading or the anemia that develops (Prohaska 2011)? While redundancy between the MCFs might explain this observation to some extent, other factors may also be involved.
12. How does lack of HEPH contribute to anemia beyond its role in intestinal iron absorption? For example, why are *Hephr<sup>int/int</sup>Cp<sup>-/-</sup>* mice (with knockout of HEPH only in the intestine) not more anemic than *Cp* knockout mice, while *Hephr<sup>-/-</sup>/Cp<sup>-/-</sup>* mice (with global deletion of both HEPH and CP) have a much more severe anemia?
13. The demonstration that only specific cells in various tissues of *Hephr/Cp* double knockout mice retain iron (Fuqua et al. 2018) suggests a mosaic expression of the MCFs. Understanding MCF function at the single-cell resolution is thus essential to fully understand their biology.
14. Do the MCFs have sex-specific roles and/or are there quantitative differences in their impact between the sexes? We have some understanding of their role in female specific tissues, but historically, most experimental studies have been carried out in males only, or in mixed sex groups.
15. What roles do the MCFs play in the pathogenesis of various diseases? Do they generally play a role as antioxidants as appears to be the case with CP? Understanding their role in CNS disorders is particularly significant. Given the association of iron accumulation in specific regions of the brain with multiple neurodegenerative disorders, including both Alzheimer's disease and Parkinson's disease, understanding the roles of the MCFs may provide important insight into the pathophysiology of these disorders.
16. Perhaps the most exciting future direction of MCF research is the exploration of whether modulating MCF function could provide a therapeutic avenue for human diseases. Either blocking or enhancing MCF expression could have application in different situations, and altering MCF functions could be relevant to either iron-related pathologies or to some of their non-iron roles. As an example, there is growing evidence that reduced HEPH expression plays a role in the progression of multiple cancers, so increasing its activity has therapeutic potential.

## Concluding remarks

Investigation of the mammalian MCFs over the past several decades has increased our collective understanding of the complex process of iron homeostasis. More recent work has demonstrated new, potentially important functions for these proteins, but additional experimentation is required to make definitive conclusions about possible other physiological roles. Collectively, these advances have led to postulates regarding MCF function and development of chronic diseases in humans, which has logically led to speculation that therapeutic targeting of these proteins could be a fruitful area for future research.

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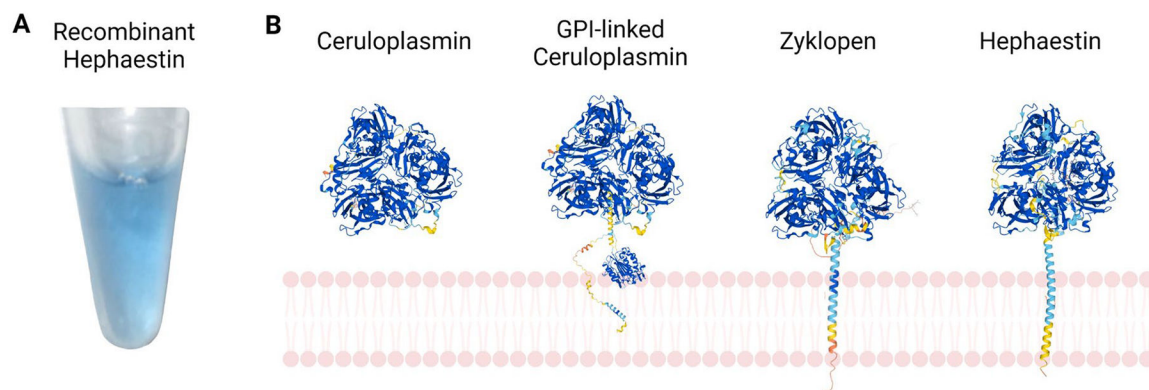
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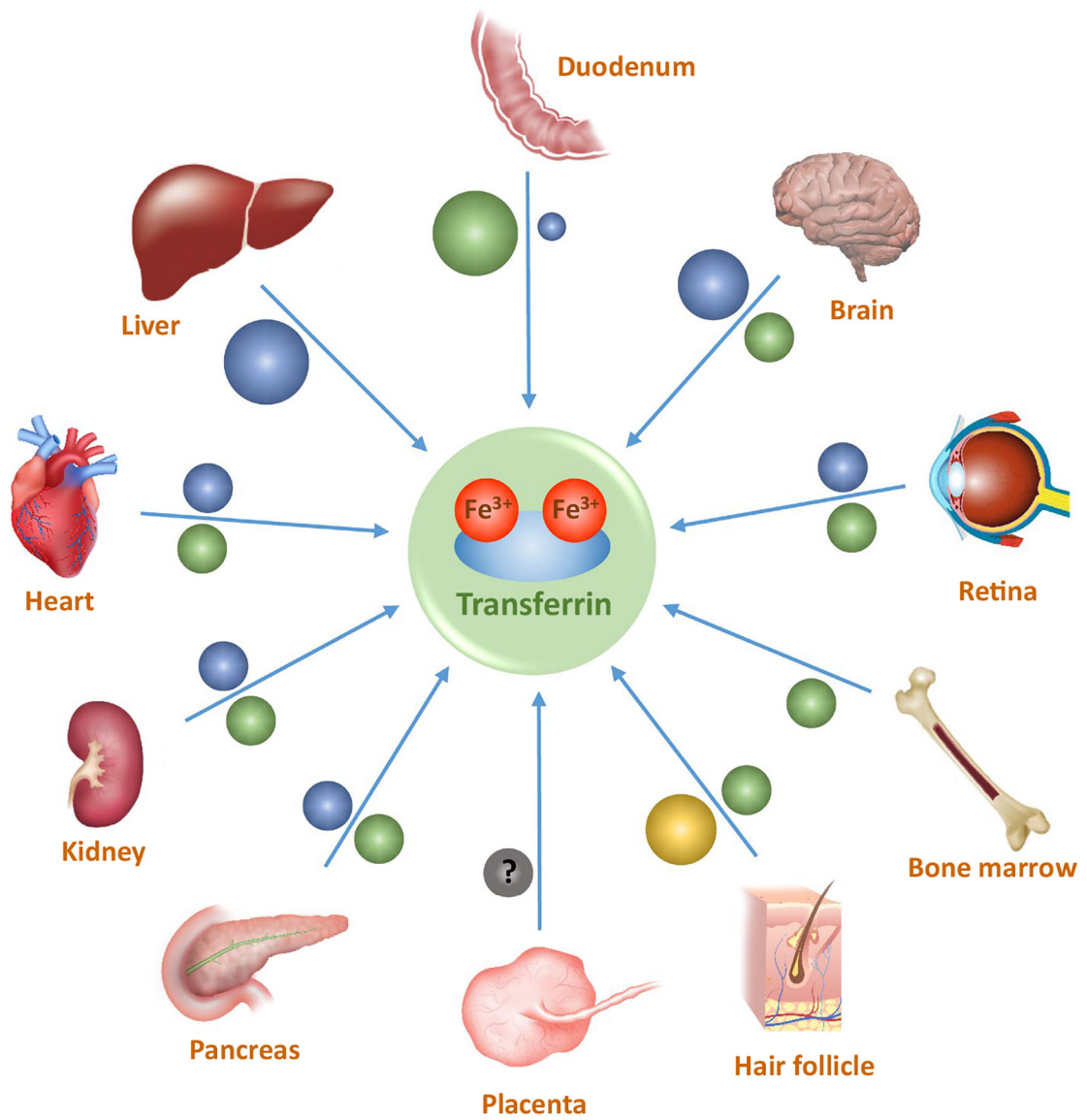


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**Fig. 1.**

Nature of the mammalian MCFs. **A** Recombinant mouse HEPH (without its transmembrane domain), generated as described by Deshpande et al. (2017), showing the intense blue color typical of the MCFs and related blue copper proteins. **B** General structure of the four forms of mammalian MCFs. From left to right: secreted CP; GPI-CP; and the two MCFs with single transmembrane domains, ZP and HEPH. The ribbon diagrams of each protein were generated using AlphaFold (<https://alphafold.ebi.ac.uk/entry/Q9BQS7>) (Jumper et al. 2021). Created with [BioRender.com](https://www.biorender.com/).



**Fig. 2.** Relative contributions of CP, HEPH and ZP to iron export from selected tissues. We now know that the MCFs contribute to iron export from multiple tissues, and often from specific cell types within those tissues. For example, in the brain, the MCFs are important for intra-organ iron trafficking. In this diagram, CP, HEPH and ZP are represented by blue, green and yellow balls respectively. For each tissue, the size of the ball represents the relative contribution of that MCF to iron transport from (or within) that tissue. For example, CP appears to be largely, if not solely, responsible for enhancing iron efflux from the liver, whereas in the duodenum, HEPH provides the majority of the ferroxidase activity, with CP contributing, albeit less efficiently, in some circumstances. In a number of tissues, both CP and HEPH appear to contribute significantly. The contributions of ZP to body iron homeostasis have yet to be firmly established, but its role in hair development is well supported. It is presumed a ferroxidase is required for efficient iron transport by the placenta

to the fetus, but the identity of that ferroxidase remains uncertain, and is represented here by a grey ball.

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Table 1

Some catalytic roles of the MCFs

MCF	Activity	Substrate	Reference
CP	Ferroxidase	Fe <sup>2+</sup>	Osaki (1966)
	NO-oxidase	NO	Shiva (2006)
	Glutathione peroxidase	Hydrogen peroxide, organic peroxides	Cha (1999), Park (1999)
	Cuprous oxidase	Cu <sup>+</sup>	Mukhopadhyay (1997)
	Amine oxidase	Organic amines (e.g. <i>p</i> -phenylene diamine, <i>o</i> -dianisidine)	Young (1972)
	Amine oxidase	Biogenic amines (adrenaline, noradrenaline, serotonin, dopamine)	Young (1972)
	Superoxide dismutase	Superoxide	Vasilyev (1988)
HEPH	Ferroxidase	Fe <sup>2+</sup>	Vulpe (1999), Deshpande (2017)
	Amine oxidase	Organic amines (e.g. <i>p</i> -phenylene diamine, <i>o</i> -dianisidine)	Vulpe (1999), Deshpande (2017)
ZP	Ferroxidase	Fe <sup>2+</sup>	Chen (2010)
	Amine oxidase	Organic amines (e.g. <i>p</i> -phenylene diamine)	Chen (2010)