Monocarboxylate transporter 8 expression in the human placenta: the effects of severe intrauterine growth restriction


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Abstract

Thyroid hormones (THs) are essential for normal fetal development, with even mild perturbation in maternal thyroid status in early pregnancy being associated with neurodevelopmental delay in children. Transplacental transfer of maternal THs is critical, with increasing evidence suggesting a role for 3,3',5-tri-iodothyronine (T3) in development and function of the placenta itself, as well as in development of the central nervous and other organ systems. Intrauterine growth restriction (IUGR) is associated with fetal hypothyroxinaemia, a factor that may contribute to neurodevelopmental delay. The recent description of monocarboxylate transporter 8 (MCT8) as a powerful and specific TH membrane transporter, and the association of MCT8 mutations with profound neurodevelopmental delay, led us to explore MCT8 expression in placenta. We describe the expression of MCT8 in normal human placenta throughout gestation, and in normal third-trimester placenta compared with that associated with IUGR using quantitative reverse transcriptase PCR. MCT8 mRNA was detected in placenta from early first trimester, with a significant increase with advancing gestation (P=0.007). In the early third trimester, MCT8 mRNA was increased in IUGR placenta compared with normal samples matched for gestational age (P<0.05), but there was no difference between IUGR and normal placenta in the late third trimester. Western immunoblotting findings in IUGR and normal placentae were in accord with mRNA data. MCT8 immunostaining was demonstrated in villous cytotrophoblast and syncytiotrophoblast as well as extravillous trophoblast cells from the first trimester onwards with increasingly widespread immunoreactivity seen with advancing gestation. In conclusion, expression of MCT8 in placenta from early gestation is compatible with an important role in TH transport during fetal development and a specific role in placental development. Altered expression in placenta associated with IUGR may reflect a compensatory mechanism attempting to increase T3 uptake by trophoblast cells.


Introduction

Thyroid hormones (THs) are essential for the normal growth and development of the fetus with even minor perturbations in maternal thyroid status during early pregnancy being associated with neurodevelopmental abnormalities in children in later life (Haddow et al. 1999, Pop et al. 1999, Chan & Kilby 2000, Kilby 2003, Pop et al. 2003). Since effects of maternal thyroid status impact on outcome at a time before development of the thyroid in the fetus, it is clear that supply of maternal THs across the placenta is critical to normal fetal development during pregnancy. Significant transport of maternal THs across the placenta is also evidenced by the finding of significant concentrations of serum thyroxine (T4) and 3,3’,5-tri-iodothyronine (T3) in neonates with congenital hypothyroidism and absent endogenous thyroid function (Vulsm et al. 1989).

Babies born with intrauterine growth restriction (IUGR) are major contributors to perinatal and neonatal mortality and up to 10% display neurodevelopmental delay (Gaffney et al. 1994). Thyroid status is one of several factors believed to play a role in this morbidity, especially in development and function of the central nervous system and placenta (Kilby 2003). We and others have used cordocentesis to perform fetal blood sampling in babies affected by severe IUGR and have reported a statistically significant reduction in circulating concentrations of free T4 and free T3 (Thorpe-Beeston et al. 1991, Kilby et al. 1998). Furthermore, the areas of the
central nervous system affected by IUGR in both human and rodent models are those that express T3-regulated genes (e.g. PCP-2 in the cerebellum) and are affected similarly in terms of histopathological changes by overt thyroid hormone deficiency (Schwartz 1983).

While circulating concentrations of T4 and T3 are considered the major determinants of cellular supply of these hormones, it is clear that other factors potentially modulate thyroid hormone action at a tissue level. These include deiodination of the pro-hormone T4 to the receptor-binding ligand T3, and the expression of functional thyroid hormone receptor isoforms. The availability of T3 for local receptor binding, and indeed the transplacental passage of maternal THs, is determined in part by the action of three selenocysteine monooxygenase subtypes (D1, D2 and D3), as well as by sulphatotransferases (Koopdonk-Kool et al. 1996, Stanley et al. 2001, Hernandez & St Germain 2003).

However, much less is known about the mechanisms of entry of THs into thyroid-responsive tissues. It has been considered that passive diffusion represents a major mode of entry into cells (Ekins 1992), although active uptake has been increasingly recognized to play a part (Hennemann et al. 2001, Abe et al. 2002). Interest has again focused on active uptake of THs since the recent description of the novel, specific, highly potent and active thyroid hormone transporter protein monocarboxylate transporter 8 (MCT8; Friesema et al. 2003). Human MCT8 was cloned in 1994 (Laforenere et al. 1994); the gene is located at Xq13.2 and comprises six exons encoding a protein with 12 putative transmembrane domains.

We hypothesize that MCT8 plays a critical role in the cellular uptake of T4 and T3 by the placenta for both thyroid hormone action within this organ and for transport of THs to the fetus. To define the role of MCT8 in human development we have determined the expression of MCT8 mRNA in human placenta across gestation. We have demonstrated the presence of MCT8 mRNA by reverse transcriptase (RT)-PCR and protein by Western immunoblotting, and have described the distribution of MCT8 protein in human placenta (both villous and extravillous) by immunohistochemistry. To investigate the relevance of MCT8 to normal and abnormal placental development and function, we have also compared the expression of MCT8 mRNA encoding MCT8 in placenta of uncomplicated pregnancies and those complicated by early- and late-onset IUGR.

Materials and Methods

Collection of placental samples

The study had the approval of the hospital ethics committee (South Birmingham) and the Research and Development Committee of the Birmingham Women’s Hospital. Placental samples were collected either at surgical termination of pregnancy (in accordance with the Polkinghorne Report) or after elective Caesarean sections. Normal placentae from the early third trimester were obtained following emergency Caesarean section for placenta praevia, maternal tumours or prelabour rupture of membranes (with no infection) with a breech presentation. Placental samples were grouped as follows: early first trimester (6–8 weeks gestation; n = 17), late first trimester (9–12 weeks gestation; n = 19), second trimester (13–20 weeks gestation; n = 13), early third trimester (28–34 weeks gestation; n = 6), and term (37–41 weeks gestation; n = 16). The gestational ages of the pregnancies were determined by ultrasound scan in the first trimester (using crown–rump length) and confirmed post-natally by feet-length/femur-length ratios (Hern 1984).

Ten cases of severe IUGR were diagnosed prospectively using ultrasound, having at least three of the following four characteristics: (i) ultrasound measurement of the fetal abdominal circumference of less than equal to the third centile for gestation; (ii) abdominal circumference growth velocity of less than 1.5 S.D. over 14 days; (iii) severe oligohydramnios (amniotic fluid index <10th centile for gestation); (iv) absent end-diastolic flow velocity in the umbilical artery Doppler velocity waveform measurements.

These criteria selected a relatively homogenous, but severe, phenotype and all were delivered by Caesarean section under maternal spinal analgesia. The IUGR group was sub-divided into early- and late-gestation groups, six being at 26–32 weeks gestation and four being at 37–38 weeks gestation. The placentae from the early IUGR group were compared with placentae from 28–34 weeks gestation with appropriately grown fetuses for gestational age, and the late IUGR group with normal term placenta.

Immediately after delivery, multiple full-thickness placental biopsies were obtained from the central (peri-umbilical) region, which has previously been demonstrated to be representative of the entire placenta (Bacon et al. 1986). Placental tissue were dissected off the chorionic plate and washed in iced 0·9% saline solution as previously described (Chan et al. 2003). No membrane was left adherent to the placental samples and these were snap-frozen in liquid nitrogen and stored at −70 °C until analysis or fixed in 4% formaldehyde solution. A representative sample from these formalin-fixed placental samples was taken for examination by immunohistochemistry.

In addition, placental bed biopsies from women following elective pregnancy termination in the first (8–12 weeks; n = 3) and early second trimester (14–15 weeks; n = 5) of pregnancy were obtained transcervically as previously described (Robson et al. 2002). All placental bed biopsies included in the study were selected to include interstitial extravillous trophoblast and ‘transformed’ decidual and/or myometrial spiral arteries containing endovascular and/or intramural extravillous trophoblast.

RNA extraction

Total RNA was extracted from approximately 100 mg tissue after homogenization, utilizing the Sigma Trisol kit: this used a single-step acid guanidinium/phenol/chloroform extraction procedure following the manufacturer’s guidelines, as previously described (Chan et al. 2002). RNA was reverse transcribed using avian myeloblastosis virus (AMV) RT (Promega) in a total reaction volume of 20 µl with 1 µg total RNA, 30 pmol random hexamer primers, 4 µl 5 × AMV RT buffer (Promega), 2 µl dNTP mix (200 µM each; Boehringer Mannheim), 20 units ribonuclease inhibitor (RNasin; Promega) and 15 units AMV RT (Promega).

Quantitative TaqMan PCR

Expression of mRNA encoding MCT8 was determined using the ABI PRISM 7700 Sequence Detection System as described previously by our group (Boelaert et al. 2003, 2004, Chan et al. 2003). RT-PCR was carried out in 25 µl volumes on 96-well plates in a reaction buffer containing 1 × TaqMan Universal PCR Master Mix, 100–200 nmol TaqMan probe and 900 nmol primers, as described below. All reactions were multiplexed with a pre-optimized mixture of control probe and primers for 18 S rRNA (PE Biosystems, Warrington, Cheshire, UK), enabling data to be expressed in relation to an internal reference to allow for differences in RT efficiency. All samples were measured in triplicate.

Primer and probe sequences for MCT8 were as follows: forward primer, 5′-CAA CGC ACT TAC CGC ATC TG-3′; reverse primer, 5′-GTA GCC CCA ATA CAC ACC AAG AG-3′; probe, 5′-TCC ACA TAC TTC ATC AGG TGT ACA TAG GGA ACA AA-3′.

The TaqMan primers yielded amplicons of 150 bp. The MCT8 probe was labelled with flurochrome 6-carboxy-fluorescein (FAM) and the 18 S probe with fluorophore VIC. Reactions were as follows: 50 °C for 2 min, 95 °C for 10 min, followed by 44 cycles of 95 °C for 15 s and 60 °C for 1 min.

As per the manufacturer’s guidelines, data were expressed as Ct values (the cycle number at which logarithmic PCR plots cross a calculated threshold line) and used to determine ΔCt values (ΔCt = Ct of the target gene (e.g. MCT8) minus Ct of the control housekeeping gene, 18 S) for each sample. Validation studies of the MCT8 TaqMan primers and probe demonstrated parallel changes in Ct values for MCT8 and 18 S across a wide range of specific concentrations of RNA (data not shown). The mean ΔCt for each experimental group was compared with the mean ΔCt for the control group (ΔΔCt) and transformed through the equation 2−ΔΔCt to calculate the fold change in gene expression.

Western immunoblotting

Placental proteins were prepared in lysis buffer (100 mmol/l NaCl, 0–1% Triton X-100 and 50 mmol/l Tris, pH 8·3) containing enzyme inhibitors (1 mmol/l PMSF, 0·3 µmol/l aprotinin and 0·4 mmol/l leupeptin), and denatured (2 min, 100 °C) in loading buffer. Protein concentration prior to loading was measured by the Bradford assay with BSA as standard. Western-blot analyses were performed as we have described previously (Friesema et al. 2003, Boelaert et al. 2003). Briefly, soluble proteins (30 µg) were separated by electrophoresis in 12·5% SDS/polyacrylamide gels, transferred to PVDF membranes, incubated in 5% non-fat milk in PBS with 0·1% Tween prior to incubation with a specific, validated rabbit polyclonal MCT8 antibody used at 1:1000 as described previously (Friesema et al. 2003). After washing, blots were incubated with appropriate secondary antibodies conjugated to horseradish peroxidase for 1 h at room temperature. After further washes, antigen–antibody complexes were visualized by the ECL chemiluminescence detection system on Hyperfilm ECL. Actin expression (monoclonal anti-β-actin clone AC-15 mouse ascites fluid; Sigma-Aldrich; used at 1:10 000) was used to assess protein loading.

Immunohistochemistry

Formalin-fixed paraffin-embedded sections of representative placental samples and placental bed biopsies were immunostained using an avidin–biotin peroxidase technique (Vectastain Elite; Vector Laboratories, Peterborough, Cambs, UK) as described in detail previously (Barber et al. 2005). All reagents were prepared as per the kit instructions. Briefly, after dewaxing and rehydration the sections were processed in 0·1 mol/l sodium citrate buffer (pH 6·0) in a pressure cooker at 103 kPa for 60 s for antigen retrieval. Slides were then incubated with 1–6% H2O2 in methanol to block endogenous peroxidase activity. After washing in 0·1 mol/l Tris/0·05 mol/l saline (pH 7·6; Tris-buffered saline, TBS), the slides were overlain with the supplied blocking serum in TBS for 10 min, before incubation with a validated rabbit polyclonal antibody to MCT8 (1:1000) for 1 h at room temperature, as reported before (Friesema et al. 2003). For negative controls the primary antibody was replaced by non-immune serum. After three 5-min washes in TBS the sections were incubated with biotinylated secondary antibody for 30 min, followed, after further TBS washes, by addition of the avidin–biotin–peroxidase complex. The reaction was developed by incubation in 3,3′-diaminobenzidine (Sigma Chemical Co) for 5–10 min. Sections were lightly counterstained with Mayer’s haematoxylin, dehydrated, cleared and mounted in synthetic resin.

Statistical analyses

Data from the quantitative RT-PCR were expressed as ΔGt values upon which all statistics were performed.
The means ± S.E. of ΔCt values for each group are shown in Figs 1 and 2. Data were analysed using GraphPad software (San Diego, CA, USA). Unpaired two-tailed t-test was used for comparison between two groups and ANOVA was used for between-group comparisons of more than two groups with Dunnett’s (post-hoc) multiple comparisons test to isolate groups that differed from others. Significance was taken as $P<0.05$.

## Results

### Expression of mRNA encoding MCT8

mRNA encoding MCT8 was present from the early first trimester and throughout gestation. Quantification of MCT8 mRNA indicated a significant increase with advancing gestation (Fig. 1; ANOVA, $P=0.007$). With post-hoc testing, a significant reduction in mRNA expression compared with term placenta was evident for placental samples collected at 9–12 weeks gestation (0.26-fold, $P<0.01$) and at 13–20 weeks gestation (0.37-fold, $P<0.05$). However, placenta from the 6–8 and 28–34 weeks gestation groups did not demonstrate statistically significant reductions in MCT8 mRNA expression compared with term placenta. There were also no statistically significant differences between the gestational groups prior to term.

In severe IUGR resulting in delivery in ‘early’ gestation (26–32 weeks), there was a significant increase of 7.5-fold in the expression of mRNA encoding MCT8 in placenta from IUGR pregnancies compared with gestationally matched placental samples from normal controls ($P<0.05$). However, the difference was no longer statistically significant when comparing ‘late’ IUGR placenta (37–38 weeks gestation) with normal term placenta (Fig. 2).

### Western immunoblots

Protein was extracted from placental homogenates of early (29 weeks) and late (38 weeks) IUGR pregnancy and compared with gestationally matched samples from uncomplicated pregnancies. The MCT8 antibody utilized demonstrated a 50 kDa band on Western immunoblotting, with weaker bands at 63 kDa, as previously described in other tissues (Friesema et al. 2003; Fig 3). The early IUGR sample expressed more MCT8 protein than the gestationally matched control while the late IUGR sample showed
no obvious difference compared with control in accord with mRNA data (Fig. 2). Immunoblotting for β-actin confirmed even protein loading across samples (Fig. 3).

**Immunohistochemistry**

Immunohistochemistry was used to localize MCT8 protein in first-, second- and third-trimester placental and placental bed biopsies (Fig. 4). MCT8 was localized to both cytoplasm and nucleus of trophoblast cells. In the first trimester (Fig. 4A1 and A2) there was focal strong reactivity with villous syncytiotrophoblast, with clear positive cells adjacent to negative syncytiotrophoblast. Immunostaining of villous cytotrophoblast was also focal, with intense staining of both cytoplasm and nucleus, often adjacent to negative syncytiotrophoblast. Samples from the early second trimester (14–18 weeks gestation; Fig. 4B1 and B2) showed intense reactivity with villous syncytiotrophoblast and cytotrophoblast, which was again focal, although more widespread than with first-trimester placenta. Again clear positive cells were present adjacent to non-reactive cells of the same type. In normal term placental tissues (Fig. 4C1 and C2) villous syncytiotrophoblast showed consistent strong reactivity for MCT8 with weaker staining of the occasional residual cytotrophoblast cells. At all gestational ages there was weaker reactivity with mesenchymal cells within the villous stroma, including endothelial cells of fetal vessels.

Stromal cells in human decidua from first-trimester pregnancies showed moderate to strong immunostaining.
(Fig. 4E), although there was no reactivity with lymphocytes within the decidual stroma. In second-trimester placental bed biopsies (Fig. 4F) the majority of interstitial trophoblast showed strong staining, although a small proportion of cells remained negative. Where present, endovascular trophoblast was also positive for MCT8 (Fig. 4F).

**Discussion**

We have demonstrated that the potent iodothyronine transporter MCT8 is present in the human placenta from early gestation. mRNA encoding MCT8 was evident in normal placenta from 6 weeks gestation, with evidence for increasing expression as gestation advanced. The early presence of MCT8 protein in placenta from the first trimester of pregnancy was also confirmed by immunohistochemistry findings. These results suggest both an early and an increasing role for MCT8 in determining T3 supply to the fetus during ontogeny, as well as a potential role in trophoblast development. Furthermore, significantly higher expression of MCT8 mRNA was evident in placenta affected by severe IUGR, where delivery was required in the early third trimester.

Immunohistochemistry demonstrated that MCT8 protein is expressed in villous trophoblast cells across gestation. In addition, in late first- and second-trimester placental bed biopsies, MCT8 protein was demonstrated in the endovascular extravillous trophoblast. This localization of MCT8 in both villous and extravillous trophoblast is similar to that of thyroid hormone receptor isoform localization described previously (Kilby et al. 1998, Barber et al. 2005). Furthermore, the pattern of localization of MCT8 in focal areas of villous placenta is similar to that of D2 (the enzyme which converts the prohormone, T4, to the active ligand, T3) in the first trimester, as we have previously reported (Chan et al. 2003), with most intense staining in the cytotrophoblast layers and weaker staining in the syncytiotrophoblast. The coordinated expression of MCT8, D2 and thyroid hormone receptors in the relatively undifferentiated villous cytotrophoblast (trophoblast stem cell line) in the first trimester suggests a role for T3 in early placentation. Indeed, T3 has been shown to modulate epidermal growth factor-stimulated changes in first-trimester primary cytotrophoblast proliferation and function in vitro, confirming it to be a T3-responsive tissue (Barber et al. 2005). A role for T3 in modulating uterine invasion by extravillous trophoblast in early pregnancy has also been reported (Laoag-Fernandez et al. 2004), an effect recently shown to be associated with upregulation by T3 of matrix metalloproteinases (Oki et al. 2004). We have shown that in SGHPL-4 cells (derived by transformation of primary extravillous trophoblasts) T3 interacts with epidermal growth factor to modulate trophoblast proliferation, migration and invasion in vitro (Barber et al. 2005). We postulate that MCT8 expression in trophoblast in the first trimester plays a crucial role in facilitating the T3-regulation of placental development, which would have a bearing on optimal placental function and fetal growth in later gestation.

The recent description (Dumitrescu et al. 2004, Friesema et al. 2004) of association between mutations of the MCT8 gene on the X chromosome and a syndrome of severe neurodevelopmental delay with abnormal circulating thyroid hormone concentrations in boys supports the view that fetal development and function are critically dependent upon an intact MCT8 transporter system within the brain. Their heterozygote mothers do not display an abnormal neurological phenotype, but are characterized by mild perturbation of tests for thyroid function. Since development prior to function of the fetal thyroid is dependent on effective transfer of maternal THs across the placenta it is plausible that alterations in both maternal circulating concentrations of TH as well as the expression of mutant MCT8 in the maternal uteroplacental bed and the fetoplacental unit, with consequent influences on transport of THs across the placenta, may exacerbate the observed phenotype in affected boys.

A focus of the present study of expression of MCT8 in human placenta is investigation of the effects of severe IUGR. This study has revealed an increase in mRNA encoding MCT8 in human villous placenta affected by severe IUGR requiring preterm delivery when compared with normal placenta. We and others have previously reported that IUGR is associated with relative fetal hypothyroxinaemia (Thorpe-Beeston et al. 1991, Kilby et al. 1998). We have also reported upregulation of thyroid receptor proteins in placenta affected by IUGR and postulated that this may reflect a compensatory mechanism enhancing thyroid hormone action in the hypothyroxinaemic environment of IUGR (Kilby et al. 1998, Kilby 2003). Enhanced expression of MCT8 may represent an additional compensatory mechanism active in the fetoplacental unit attempting to increase T3 uptake in trophoblast cells or to increase the transplacental passage of THs in these compromised pregnancies.

It would be important to establish a functional role for MCT8 within the placenta which could partly be addressed through in vitro studies in primary trophoblast cultures and choriocarcinoma cell lines. The factors affecting the regulation of MCT8 expression at a molecular level in normal physiology, and in syndromes of malplacentation such as IUGR, are as yet unknown and are the subject of further investigation.

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