Positive Newborn Screen for Methylmalonic Aciduria Identifies the First Mutation in TCblR/CD320, the Gene for Cellular Uptake of Transcobalamin-bound Vitamin B$_{12}$

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Communicated by William S. Sly

Received 22 April 2010; accepted revised manuscript 18 May 2010.
Published online 3 June 2010 in Wiley InterScience (www.interscience.wiley.com). DOI 10.1002/humu.21297

ABSTRACT: Elevated methylmalonic acid in five asymptomatic newborns whose fibroblasts showed decreased uptake of transcobalamin-bound cobalamin (holo-TC), suggested a defect in the cellular uptake of cobalamin. Analysis of TCblR/CD320, the gene for the receptor for cellular uptake of holo-TC, identified a homozygous single codon deletion, c.262_264GAG (p.E88del), resulting in the loss of a glutamic acid residue in the low-density lipoprotein receptor type A-like domain. Inserting the codon by site-directed mutagenesis fully restored TCblR function. Hum Mutat 31:924–929, 2010. © 2010 Wiley-Liss, Inc.

KEY WORDS: methylmalonic acid; homocysteine; transcobalamin-receptor; cobalamin; vitamin B$_{12}$; TCblR; CD320

Introduction

Although genetic defects affecting the absorption, transport and metabolism of vitamin B$_{12}$ (cobalamin, Cbl) are now known [Fowler et al., 2008], no defects in the receptor TCblR (CD320; MIM# 606475) for cellular uptake of Cbl have been identified to date. TCblR expressed on the plasma membrane binds transcobalamin (TC) saturated with Cbl (holo-TC) and mediates cellular uptake of Cbl. TCblR expression is coupled to the cell cycle with highest expression in actively proliferating cells and down-regulation in quiescent cells [Amagasaki et al., 1990]. The receptor has a 29-fold higher affinity for holo-TC than for apo TC [Quadros et al., 2005]. The only route for physiologic transport of TC-bound Cbl into cells is via TCblR. The existence of a receptor for holo-TC was inferred from Ca$^{2+}$-dependent binding and uptake of holo-TC by cells [DiGirolamo and Huennekens 1975], but the identity of the protein and the gene encoding this receptor had eluded researchers for decades. Definitive purification of the receptor protein from human placenta provided the protein sequence for the identification of the CD320 gene on chromosome 19.p13.2 as the gene that encodes the receptor [Quadros et al., 2009]. This 282aa glycoprotein with a single transmembrane domain and a cytoplasmic tail is ubiquitously expressed. This report describes the first gene defect of a single amino acid deletion in the receptor leading to elevated methylmalonic acid (MMA) in newborns.

Case Report

The index case was born at term after a pregnancy (gravida1, para1) complicated by mild anemia and gestational diabetes, controlled by diet. The parents are of mixed European background and are nonconsanguineous. Blood for a newborn screen was collected on the second day of life and revealed an elevated C3-acylcarnitine of 6.22 μmol/l (cutoff 5.71). The C3/C2 ratio was 0.21 (cutoff 0.2). Urine organic acid analysis showed moderately elevated MMA and trace methylcitric acid without other propionate metabolites. The plasma vitamin B$_{12}$ level was 269 ng/l (reference range, RR; 180–914), and total plasma homocysteine was 8.6 μmol/l (RR 0–11). The hematocrit was 44% (RR 42–70), and the mean corpuscular volume (MCV) was 96.9 FL (RR 90–115). A repeat newborn screen, done on day 14 of life was normal; the C3-acylcarnitine was below cutoff at 3.2 μmol/l, the C3/C2 ratio was 0.17. Lactate, ammonia, electrolytes, liver function tests, coagulation studies, blood gas, CBC, urinalysis and a carnitine panel were normal. Repeat MMA levels were obtained during this period and ranged from 5.3 to 7.7 μmol/l. One milligram of hydroxocobalamin was administered intramuscularly (i.m.) at 21 days of age; the MMA level 24 hr later was normal at 0.82 μmol/l and remained within the reference range for the patient’s age. When last tested at 9 months of age, MMA was 1.1 μmol/l and the vitamin B$_{12}$ level, without additional supplementation, was elevated at >1,400 ng/l. Maternal vitamin B$_{12}$ level 2 weeks after delivery was 312 ng/l (RR 180–914); homocysteine was 7.8 μmol/l (RR 5.1–13.9) and MMA was 0.58 μmol/l (RR <0.89). All hematological parameters were within the normal range with no indication of vitamin B$_{12}$ deficiency. Following the positive newborn screen, a skin biopsy was obtained and a fibroblast culture was established.
Identification of the TCblR gene defect in the index case prompted the review of previous undiagnosed cases that were referred for genetic testing due to elevated levels of MMA in newborn screens.

The index case and the additional cases reported here were all referred to the medical genetics laboratory at McGill University as a result of an abnormal newborn screen in the form of an elevated C3-acylcarnitine and methylmalonic aciduria. This cell line and four additional lines with the identical deletion are available from the Repository for Mutant Human Cell Strains, Montreal Children’s Hospital, Montreal, Canada (http://www.cellbank.mcgill.ca/). The protocol was approved by the Royal Victoria Hospital, Research Ethics Board.

Methods

Propionate Incorporation in Cells

Fibroblasts were plated in 35-mm tissue culture dishes at a density of 400,000 cells per dish. After cells had attached, medium was removed and replaced with Puck’s F medium supplemented with 15% (v/v) fetal bovine serum and 1–[^14C]propionate (GE Health Sciences, Piscataway, NY) diluted with cold propionate to a final specific activity of 10 μCi/μmol. Cultures were incubated with labeled medium for 18 hr. At the end of this period, medium was removed and cellular macromolecules were precipitated by incubation in 5% trichloroacetic acid. The precipitate was dissolved in 0.2 N NaOH and radioactivity was determined by liquid scintillation counting. This assay was done in the absence of propionate to a final specific activity of 10 μCi/μmol. Cultures were incubated with labeled medium for 18 hr.

Methyltetrahydrofolate Incorporation in Cells

This procedure is identical to the propionate incorporation test except for the use of methionine-free minimal essential medium supplemented with 100 μM homocysteine thiolactone, 50 μg/ml sodium ascorbate, and 10% (v/v) dialyzed fetal bovine serum containing 0.5 μCi/ml (60 mCi/mmol) 5-[^14C]methyltetrahydrofolate (GE Health Sciences). This assay provides a measure of methionine synthase function in intact cells [Wyllard et al., 1978].

Synthesis of Cobalamin Coenzymes

Confluent cultures of fibroblasts in 175 cm² culture flasks were incubated for 96 hr in MEM supplemented with 25 pg/ml [^57Co]cyanocobalamin bound to TC in human serum. Cells were harvested by trypsinization and cobalamins were extracted in hot ethanol and separated by high-performance liquid chromatography (HPLC) using a modification of the method of Jacobsen et al. [1982].

Holo-TC Uptake Studies

Skin fibroblasts were cultured in Dulbecco’s modified Eagle’s minimal essential medium (DMEM) with 10% fetal bovine serum. Binding and uptake of Cbl was determined by incubating cells in fresh medium containing recombinant TC saturated with [^57Co]cyanocobalamin.

Homocysteine and Methylmalonic Acid in Fibroblast Cultures

Cells were cultured with varying amounts of holo-TC. Medium was analyzed for Hcy by an HPLC-based method with fluorescence detection [Jacobsen et al., 1994] and MMA by gas chromatography-mass spectrometry [Rasmussen, 1989].

Analysis of the TCblR Gene

Total RNA and genomic DNA were prepared from cultured fibroblasts using Trizol reagent (Invitrogen, Carlsbad, CA). The mRNA level was measured by SYBR GreenER (ABI PRISM)-based real-time PCR (qPCR) (Primer set: Forward [F]-AAGTTCCAGTGCGCACGACTGAGCAGTGA; Reverse [R]-AGTCAGTTGCAGGCCCTGAG; 95°C, 10 min, followed by 40 cycles of 95°C, 15 sec, and 60°C, 1 min). The size and sequence of the transcript was determined by reverse transcription using random primers followed by PCR amplification of the cDNA (Primer set: F-ACAGCATGACGGGCGCTCGA/R-GCTACGCCCCAGGCTGAGGT; 94°C, 2 min, followed by 30 cycles of 94°C, 40 sec, 60°C, 40 sec, 72°C, 40 sec, and one cycle at 72°C, 10 min). The nucleotide sequence encoding the mRNA for TCblR was determined by sequencing the PCR amplified cDNA fragment and confirmed by amplification (Primer set: F-ATATGCCGGCTGCCTGTCCTC/RTATACCCAGGCGAGCTGTCGG; 94°C, 2 min, followed by 30 cycles of 94°C, 40 sec, 58°C, 40 sec, 72°C, 40 sec, and one cycle at 72°C, 10 min), and sequencing the region of the gene corresponding to the mutation in the genomic DNA. The missing codon was reinserted into the patient’s cDNA using the QuikChange Lightning site-directed mutagenesis kit (Stratagene, La Jolla, CA). The patient’s cDNA in pCDNA3.1 vector was amplified by PCR using overlapping mutagenic primers (Primer set: F-GGGCATTGAGGAGGTGACGCTGAC/GGCTCAATCTCTGACTCCTCCTCGTGC). The PCR amplification, cDNA isolation and cloning was carried out as per instructions provided with the kit and the insertion of the missing codon confirmed by sequencing. Nucleotide numbering reflects cDNA numbering with +1 corresponding to the A of the ATG translation initiation codon in the reference sequence (GenBank NM_016579.3). The initiation codon is codon 1.

Results

Cellular Vitamin B₁₂ Metabolism

Studies in the index case with fibroblasts cultured for 18 hr with and without 3.7 μM hydroxocobalamin showed normal[^14C]propionate (9.5 and 12.2 nmol/mg protein) and[^14C]methyltetrahydrofolate (79 and 960 nmol/mg protein) incorporation. Initial testing in cultured fibroblasts revealed low uptake of[^57Co]cyanocobalamin (1.6 pg/10⁶ cells) bound to TC in human serum but conversion to adenosylcobalamin (11%) and methylcobalamin (52%) was normal. The decreased Cbl uptake suggested a potential defect in TC-receptor mediated cellular uptake of holo-TC and a detailed study to identify the genetic abnormality was initiated.

Binding and Uptake of TC-Cbl

Uptake of[^57Co]cyanocobalamin bound to human recombinant TC in fibroblast cultures showed consistently lower binding and uptake throughout the 3-hr time period tested (Fig. 1A). On average, the uptake in the patient’s cells was about half that
observed in two control cell lines. This difference in uptake appeared to be due to a twofold decrease in the affinity of the receptor for holo-TC (Kₐ = 4 nM⁻¹) compared to the control fibroblasts (Kₐ = 2 nM⁻¹), and a fourfold decrease in maximum binding (B_max) (Fig. 1B). The fibroblast cultures from the four additional cases also showed decreased uptake of TC-Cbl (Fig. 2).

Homocysteine and Methylmalonate in Fibroblast Cultures

The fibroblasts from the index case accumulated significantly higher Hcy and MMA in the medium compared to controls during 8 days in culture (Fig. 3). Both the normal and the TCblR mutant cell line showed less Hcy and MMA with increasing concentrations of holo-TC. Production of Hcy in the presence of 0 or 148 pmol/l holo-TC was significantly higher for the TCblR mutant than for the control cell line (Fig. 3A). The concentration of Hcy produced by the TCblR mutant cells decreased to normal levels when the concentration of holo-TC was increased to >1.4 nmol/l. In contrast, although a substantial decrease in the production of MMA was observed with increasing concentrations of holo-TC, the TCblR mutant cell line produced significantly higher concentration of MMA at all concentrations tested (Fig. 3B).

Analysis of the TCblR Gene

Genomic DNA and mRNA were extracted from cultured skin fibroblasts. The size of the transcript determined by PCR amplification of the mRNA and agarose gel electrophoresis was consistent with the full-length cDNA. In addition, the amount of the transcript as determined by qPCR was similar to that in control fibroblasts (Fig. 1C). However, the nucleotide sequence showed deletion of three base pairs (c.262_264delGAG) (Fig. 4A), resulting in loss of codon 88 (p.E88del) and deletion of a single glutamic acid residue in the 3’ end of the first LDLR type A domain (Fig. 4B). The patient was presumed to be homozygous for this deletion, because direct sequencing of the PCR amplified cDNA as well as the genomic DNA failed to show the presence of a normal allele. DNA from parents was not available. Other changes in the nucleotide sequence noted were c.489A>G (p.S142G), which is not a known polymorphism, c.723G>A (rs2336573; p.G220R), and the silent polymorphisms c.512G>T (rs2232783) and c.548C>T (rs2232784). The phenotype was confirmed by transfecting HEK 293 cells with the patient’s cDNA containing the
nant holo-TC was added to 2 ml culture medium containing 0.2 µmol/l holo-TC. Bovine recombinant long chain VLCFA (MMA) ([Kozyraki and Gofflot, 2007]). In the case of TCblR, the two LDLR-type A domains with consensus sequence for Ca$$^{++}$$ binding are involved in holo-TC/TCblR interaction. Epitope mapping studies [Fedosov et al., 2005] and the crystal structure of holo-TG [Wuerges et al., 2006] have identified the positively charged heparin-binding region of TC as the likely site of interaction with the receptor. This is further supported by the observation that a monoclonal antibody either to the second LDLR-A domain of TCblR (unpublished data) or to sequences near the heparin binding domain of TC [Fedosov et al., 2005], blocks holo-TG/TCblR interaction. The identification of a single amino acid deletion in the first LDLR-A domain of TCblR, which is involved in Ca$$^{++}$$ and TC-Cbl binding is consistent with the observation of decreased uptake contributing to intracellular Cbl deficiency and elevated MMA. This conclusion is further supported by the observation of decreased Hcy and MMA in fibroblast cultures when 10–100-fold excess holo-TC is available for Cbl uptake. Compensating for the decreased affinity by providing adequate TC-Cbl facilitates additional binding and uptake to lower the intracellular production of Hcy and MMA.

Discussion

Following an unremarkable pregnancy, a newborn screen positive for elevated C3-acylcarnitine led to the identification of methylmalonic academia, prompting additional studies to determine its cause. Somatic cell studies failed to identify any abnormalities in the synthesis of Cbl coenzymes or Cbl-dependent enzymes. However, a substantial decrease in the accumulation of Cbl in fibroblast cultures prompted additional studies to identify the underlying cause.

Both acquired and inherited causes of functional Cbl deficiency have been described. Juvenile and adult onset pernicious anemia due to intrinsic factor deficiency is primarily an autoimmune disease and mutations in the genes encoding intrinsic factor, cubulin, and amnionless result in disorders of Cbl absorption [Fyfe et al., 2004; Tanner et al., 2005; Yassin et al., 2004]. Inherited TC deficiency leads to Cbl insufficiency after birth, typically manifesting as decreased absorption and cellular uptake of the vitamin [Burman et al., 1979] due to the dual role of TC in the translocation of the absorbed Cbl in the gut as well as its delivery to tissue cells [Quadros et al., 1999]. Mutations in the genes for the two enzymes requiring Cbl or in the enzymes involved in the pathways leading to the synthesis of adenosyl- and methyl-Cbl affect intracellular Cbl metabolism [Goelho et al., 2008; Hannibal et al., 2009; Rutsch et al., 2009]. Depending on the location of the defect, these disorders can present with elevated Hcy and/or MMA, with disorders of Cbl transport typically presenting with lower levels of these metabolites when compared to disorders affecting intracellular metabolism. Early detection and treatment with pharmacologic doses of vitamin B$$_{12}$$ can prevent or reverse neuropathologic changes [Hall, 1990]. Therefore, newborn screening for Hcy and MMA, two sensitive indicators of intracellular Cbl deficiency, is recommended for early detection. Although testing for Hcy is not routine, testing for MMA through elevated C3-acylcarnitine is becoming widespread.

In the index case reported, elevated MMA was identified after an abnormal newborn screen. Failure to associate the high MMA with known genetic defects prompted additional studies, among them measurement of cellular TC-Cbl uptake to detect a possible defect in cellular uptake and processing of TC-Cbl. Preliminary studies indicated a potential defect in the binding of TC-Cbl to the cell surface receptor. The receptor belongs to a class of membrane receptors that share structural features such as the LDL receptor type A domains found in the LDL receptor family of proteins, separated by a CUB-like domain found in a number of multiligand binding receptors such as cubulin, megalin, and epidermal growth factor receptor; these domains are involved in ligand binding [Kozyraki and Gofflot, 2007]. The discovery of four additional cases with the identical gene deletion corrected the cell line’s decreased Cbl uptake (Fig. 4C).

![Figure 3. Accumulation of homocysteine (Hcy) (A) and methylmalonic acid (MMA) (B) in culture medium of fibroblasts grown in the presence of increasing concentrations of holo-TC. Bovine recombinant holo-TC was added to 2 ml culture medium containing 0.2 x 10$^6$ cells.](image-url)
previously reported in cases of Cbl deficiency in infants [Campbell et al., 2005; Monsen et al., 2003]. Thus, the MMA level appears to be a more sensitive indicator of subclinical Cbl deficiency. From the cases reported here, it is also evident that hematological changes are not apparent at this deficiency level in newborns and infants, and therefore, screening for acylcarnitine and MMA would be required. Earlier studies have also reported a decrease in Cbl level during the first weeks of life with corresponding benign and transient increase in MMA and Hcy that either remain elevated without clinical symptoms or revert to normal levels with time [Ledley et al., 1984; Shih et al., 1976]. The elevated Hcy observed in the in vitro studies done with fibroblast cultures from the index case contrasts with the in vivo data of normal Hcy. Although unconfirmed, it is possible that the metabolism of Hcy via the alternate betaine–homocysteine methyltransferase (BHMT) pathway could account for the normal plasma Hcy. Skin fibroblasts do not express BHMT [Wang et al., 1991], and therefore, reflect the effect of cellular Cbl deficiency on both the MMA and Hcy pathways. It should be noted, however, that serum Hcy levels were elevated in two of the additional cases of TcBlR deficiency. In all cases, indirect measures of propionate catabolism and conversion of homocysteine to methionine, with and without added Cbl, as well as synthesis of Cbl cofactors from exogenous cyanocobalamin was normal. The only defect identified in all five cases with the single codon deletion was decreased uptake of TC-Cbl.

Many inborn errors of Cbl metabolism cause severe functional blocks and present early in life. Patients with genetic abnormalities of TC, the ligand for TcBlR, develop severe cellular Cbl deficiency, pancytopenia, megaloblastic anemia, and neurological sequelae within the first 2 months of life due to complete loss of TC function [Prasad et al., 2008; Ratschmann et al., 2009]. The moderate MMA and the absence of an early clinical phenotype in these cases is likely due to the defect producing a mild but sustained intracellular Cbl deficiency. The elevated plasma vitamin B12 level 9 months after a single injection of vitamin B12 in the index case further attests to the defect in cellular uptake. The long-term effects of this condition are yet to be unraveled, but TcBlR defects should be considered in individuals with elevated blood and urine MMA who do not have other defects in Cbl metabolism.

Acknowledgments
We thank Lydia Vezina, Lara Reichman, and Laura Dempsey-Nunez for excellent technical support and discussions as part of their undergraduate research project. This study was supported by NIH Grant DK 064732 (E.V.), HL71907 and HL52234 (D.W.J.), and CIHR Grant 160439 (D.S.R.).

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