Menkes disease: recent advances and new aspects

Zeynep Tümér, Nina Horn

Copper is the third most abundant trace element in the body, after iron and zinc, and it is required for the normal function of several important copper enzymes. However, the same element in excess is highly toxic and has detrimental effects. Fine regulation of intracellular copper homeostasis is therefore vitally important and disturbance of this balance is reflected in two hereditary disorders, Menkes disease and Wilson disease. In recent years, remarkable progress has been made in this field following the isolation of the defective gene in Menkes disease (MD), which will be the main focus of this review.

Progressive neurodegeneration and connective tissue disturbances are the main manifestations of X-linked recessive Menkes disease and most of the clinical features can be explained by malfunction of one or more copper enzymes. The disease locus has been mapped to Xq13.3 and the gene (MNK) defective in Menkes disease has been isolated by positional cloning. The protein product is predicted to be a copper binding P type ATPase (ATP7A), the first intracellular copper transporter described in eukaryotes. Identification of MNK led to a series of advances in a very short time. The mouse homologue of MNK has been isolated and the allelic relationship between MD and the occipital horn syndrome confirmed. Most importantly, the gene defective in Wilson disease was isolated using sequences specific to MNK and the predicted protein product showed high homology to ATP7A. In this review we will outline these recent advances and their consequences with special reference to Menkes disease. Wilson disease will be described briefly, and the defective copper metabolism in both diseases will be summarised in the light of the new insights. Finally, copper translocating ATPases identified in other species and their significance in understanding copper metabolism will be discussed.

Keywords: Menkes disease; Wilson disease; copper metabolism.

Menkes disease

Menkes disease (MD) is a multisystemic lethal disorder, dominated by neurodegenerative symptoms and connective tissue manifestations. A striking and pathognomonic feature is the sparse, coarse, and depigmented hair, the reason why the disease has also been called "kinky or steely hair disease" (fig 1). Though most patients (90-95%) have a severe clinical course (table 1), various forms of the disease exhibiting different degrees of nervous system or connective tissue involvement can be distinguished. The occipital horn syndrome (OHS), mainly characterised by connective tissue manifestations, has been suggested to be a very mild allelic form of MD\(^\text{4}\) (table 1). Besides OHS, a late onset mild form\(^\text{7}\) and a moderate form can also be distinguished from the severe classical form. Variation also exists within these groups and there are, for example, a significant number of patients surviving more than six years despite the severe clinical symptoms. It is thus conceivable that MD covers a clinical continuum from the severe classical form to the mild OHS. Variation in copper metabolism was thought to be the primary cause, but later investigations showing copper accumulation in extrahepatic tissues, apart from the brain, indicated a multisystemic involvement. Most of

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Table 1  Main symptoms of patients with the classical form of Menkes disease and OHS

<table>
<thead>
<tr>
<th></th>
<th>MD</th>
<th>OHS</th>
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<tr>
<td>Neurological</td>
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<td>+/-</td>
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<tr>
<td>retardation</td>
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<td>Convulsions</td>
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<td>Hypothermia</td>
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<td>+</td>
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<tr>
<td>Feeding difficulties</td>
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<td>+</td>
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<td>Muscle tone changes</td>
<td>+</td>
<td>ND</td>
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<td>Connective tissue symptoms</td>
<td>+</td>
<td>+</td>
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<tr>
<td>Torcular vessels</td>
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<tr>
<td>Skeletal changes</td>
<td>+</td>
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<tr>
<td>Bladder diverticulce</td>
<td>+</td>
<td>++</td>
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<td>Loose skin</td>
<td>+</td>
<td>+</td>
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<tr>
<td>Loose joints</td>
<td>+</td>
<td>+</td>
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<tr>
<td>Other symptoms</td>
<td>+</td>
<td>+</td>
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<tr>
<td>Facial dysmorphism</td>
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<tr>
<td>Abnormal hair, pili torti</td>
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<tr>
<td>Hypopigmentation</td>
<td>+</td>
<td>+</td>
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<tr>
<td>Laboratory findings</td>
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<tr>
<td>Serum copper</td>
<td>↓</td>
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<td>Serum ceruloplasmin</td>
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<td>↓</td>
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<tr>
<td>Intracellular Cu accumulation</td>
<td>*</td>
<td>*</td>
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<tr>
<td>Life span</td>
<td>Death before</td>
<td>3 years</td>
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</table>

*Except for four patients, all the known OHS patients (about 22) are still alive, the oldest patient being 52 years old. Two patients have died in car accidents (I Katila and P Byers, personal communication), one patient has died of an unknown cause at 40 years of age (I Katila, personal communication), and one patient has died aged 49 years because of a bladder rupture (D Weaver, personal communication). ND, not determined.

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the clinical features of MD are attributable to deficiency of one or more important copper requiring enzymes, such as cytochrome c oxidase (electron transport), superoxide dismutase (free radical detoxication), dopamine β hydroxylase (catecholamine production), lysyl oxidase (cross linking of collagen and elastin), and peptidyl-glycine α-amidating mono-oxygenase (PAM, bioactivation of peptide hormones).

Genetics of Menkes disease

LOCALISATION OF THE MENKES LOCUS

Menkes disease was recognised to segregate as an X linked recessive trait when it was first described in 1962 by Menkes et al. Cell culture studies in phenotypically normal female Menkes carriers indicated that the Menkes disease gene (MNK) was subject to random X inactivation, also supporting an X linked mode of inheritance. Later, MNK was linked to the centromeric region of the X chromosome by C banding polymorphism studies and further linkage analyses, exclusion mapping, and comparative gene mapping between man and mouse suggested proximal Xq as the candidate region.

The first physical evidence for the location of MNK was the finding of a female Menkes patient with a balanced X;2 translocation and the X chromosome breakpoint was localised to Xq13.2-q13.3. This finding suggested that the X chromosome breakpoint was at or very near the Menkes locus, directly affecting the function of the gene. Description of another Menkes patient with an X chromosome abnormality involving Xq13 not only supported this region as the candidate locus for MNK, but also finely localised it to Xq13.3. This patient was a male carrying a unique intrachromosomal rearrangement where the segment Xq13.3-q21.2 was inserted into the short arm.

ISOLATION OF THE GENE DEFECTIVE IN MENKES DISEASE

The finding of two Menkes patients with X chromosome breakpoints within the candidate region expedited the efforts in cloning MNK. Three groups, including ours, isolated MNK using slightly different positional cloning strategies. The X chromosome breakpoint of the X;2 translocation patient was the starting point for all three groups, though our group also investigated the patient with the intrachromosomal rearrangement. The common step was to construct a physical map of the genomic region encompassing the site of the breakpoints and to isolate YACs (yeast artificial chromosomes) covering this region. Screening cDNA libraries with different kinds of probes enabled all three groups to isolate candidate cDNA sequences which were mapping to the breakpoint region and were identical to each other. Southern blot analysis of 16 unrelated Menkes patients referred to our institute showed partial gene deletions of different sizes and locations, indicating that this candidate gene was defective in MD, and hence was “the” Menkes disease gene (MNK).

Furthermore, in line with the disturbed copper metabolism, the predicted protein sequence showed a P type ATPase with striking sequence homology to a bacterial heavy metal binding protein. The predicted protein is now designated ATP7A and the MNK gene is occasionally called ATP7A.

The mRNA transcript of MNK is 8.5 kb long and the 3' end contains a 3.8 kb untranslated region. An open reading frame of 4500 bp is identified encoding a predicted protein of 1500 amino acids. The mRNA transcript is expressed in heart, brain, lung, liver, skeletal muscle, kidney, and placenta, but the level is low in pancreas and hardly detectable in liver. MNK is organised in 23 exons spanning about a 150 kb genomic region (fig 2). The first exon is a leader exon containing only untranslated sequences and the ATG start codon is in the second exon. The exons in general correspond to the predicted functional/structural domains of the protein product with a few exceptions.

About the protein (ATP7A)

Comparison of the predicted protein sequence (ATP7A) with other sequences in protein databases indicated a close similarity to cation transporting P type ATPases, especially a heavy metal ion ATPase, involved in copper homeostasis in Enterococcus hirae. It was also closely related to the cadmium resistance ATPases found in Staphylococcus aureus.

P type ATPases are energy using membrane proteins, functioning as cation pumps either for uptake, efflux, or cation exchange. These enzymes are called “P type” ATPases, because...
A

\[
\begin{array}{cccccccccccc}
\hline
\text{Cu} & \text{Cu} & \text{Cu} & \text{Cu} & \text{Cu} & \text{Cu} & \text{PD} & \text{CPC} & \text{D} & \text{ATP} \\
\end{array}
\]

B

\[
\begin{array}{c}
\text{NH}_2 - \text{Cu} \\
\text{Cytoplasm} \\
\text{Membrane} \\
\text{Transmembrane domains} \\
\text{COOH} \\
\text{ATP binding domain (ATP)} \\
\text{Phosphatase domain (PD)} \\
\text{TGE} \\
\text{DKTGT} \\
\end{array}
\]

**Figure 2** (A) Exon structure of MNK and corresponding protein domains.\(^\text{26}\) The vertical lines indicate the positions of the introns and the exons are indicated by numbers. (B) Predicted protein structure of ATP7A.\(^\text{22-23}\) The structure of ATP7B is also predicted to be similar to ATP7A.\(^\text{22-23}\) Transmembrane domain is a common structural feature of P type ATPases and anchors the protein into the membrane. ATP7A is predicted to have eight transmembrane domains (indicated with white bars in fig 2A). ATP binding domain (ATP) is an extramembranous segment, responsible for ATP binding in P type ATPases and it is one of the most conserved sites. Phosphorylation domain (D) contains an invariant cytoplasmic DKTGT motif. The aspartate residue (D) is crucial for the enzyme activity and it is phosphorylated with the terminal phosphate of ATP in the cation transport cycle. CPC is the predicted cation channel with a proline residue (P) highly conserved among P type ATPases. This residue is proposed to participate in the transduction of energy from the phosphorylation site to cation transport. In ATP7A proline is surrounded by cysteine residues, which may provide specificity for heavy metals. Phosphatase domain (PD) contains the TGE motif, which may have a role in removing the phosphate from the phosphorylated aspartic acid (D) as part of the cation transport. The amino-termini of P type ATPases is one of the most divergent domains. The GMXCXXX motif with a pair of conserved cysteine residues is repeated six times in ATP7A and this motif is suggested to be binding copper (Cu).

of a conserved aspartate residue (D) that is transiently phosphorylated with the terminal phosphate of ATP during the transport of the cations across a membrane. The overall sequence similarity among prokaryotic and eukaryotic cation transporting ATPases suggest that these proteins have been modified throughout evolution in response to the need for translocation of various cations. ATP7A has all the features common to P type ATPases\(^\text{22-23}\) (fig 2B). A remarkable feature of this protein is the presence of six successive repeats at the amino terminal. These repeats contain the consensus GMXCXXX motif and the presence of paired cysteine residues suggests that these domains are the copper binding sites.

**The mottled mouse**

Soon after the isolation of MNK the status of mottled mouse, which was long thought to be the murine model for Menkes disease, was established. In mouse, 23 known mutations (Y Boyd, personal communication) at the same X linked mottled locus (Mo) lead to a mottled coat pigmentation in the female heterozygotes and the affected males show neurological and connective tissue abnormalities, differing greatly in severity.\(^\text{34}\) Phenotypic and biochemical similarities between MD patients and mottled mutants,\(^\text{34,35}\) along with the conserved map positions of MNK and Mo,\(^\text{35,36}\) have long suggested that Mo phenotypes were caused by mutations in Mnk, the mouse homologue of MNK. Following the isolation of MNK, Mnk
was cloned using the human sequences. The predicted protein product was also a P type ATPase (atp7a), showing 89% sequence homology to ATP7A, and the tissue expression profile was also similar to the human counterpart. In the two mottled mutants, mottled blotchy (Mo⁰) and mottled dappled (Mo⁶), altered levels of the Mnk transcript were detected. Recently a mutation in Mnk was detected in Mo⁰ confirming the status of the Mo mouse as an animal model for MD (see below).

The occipital horn syndrome
Cloning of MNK provided the opportunity to analyse the occipital horn syndrome (OHS) patients for an allelic mutation. These two X linked recessive disorders of copper metabolism were suggested to be allelic based on their biochemical and clinical resemblances (table 1), and their homology with the possibly allelic forms of the mottled mouse. Two well characterised mottled mutants, the mottled brindled (Mo⁰) and the mottled blotchy (Mo⁶), were suggested as the murine models for the classical form of MD and OHS, respectively. The Mo⁰ phenotype and classical MD have similarly disturbed copper homeostasis leading to severe neurological impairment and death at an early age. The Mo⁶ phenotype resembles OHS showing predominantly connective tissue manifestations. Both in MD and OHS, serum copper and ceruloplasmin are low, though usually lower in MD and occasionally normal in OHS. Cultured fibroblasts of OHS and MD patients show increased copper accumulation and markedly low lysyl oxidase activity. In OHS the major effect is on lysyl oxidase, the copper dependent enzyme that initiates cross linking of collagen and elastin in connective tissues. However, the possibility of a primary defect in the lysyl oxidase gene was excluded by the assignment of the gene to an autosome.

Following the isolation of MNK, Levinson et al detected markedly reduced levels of the mRNA transcript in two unrelated OHS patients, suggesting that MNK had a role in OHS. Recently, impairment of MNK in three patients with OHS has been reported, giving direct molecular evidence for the allelic relationship between these two diseases (unpublished data). These mutations are base pair substitutions affecting the normal mRNA splicing and the Mo⁰ also has a similar splicing defect. Further studies, elucidating the effects of the gene mutations on the structure and function of the protein, are required to understand the cellular pathology resulting in different phenotypes in mice and humans.

Females with Menkes disease
A total of eight females affected with Menkes disease have been described (Tsukahara, personal communication). In two patients, diagnosis was suggested clinically, and cytogenetic and biochemical analyses were not performed. In three cases, diagnosis was confirmed by significantly increased copper accumulation in cultured fibroblasts. Two of these patients had normal karyotypes and the most likely explanation for the expression of the disease was odd X inactivation. The third patient had a 46,XX/45,X(31.5%) mosaicism in cultured fibroblasts and expression of the disease could be explained by monosomy of the X chromosome harbouring the defective gene or by odd X inactivation. Three female patients had balanced translocations, t(X;21)°, t(X;1)°, and t(X;21)° (Tsukahara, personal communication) and in each case the X chromosome breakpoint was shown to be at the MNK locus by fluorescence in situ hybridisation analyses (unpublished data). As expected, the normal X chromosomes were preferentially inactivated in these patients, resulting in the expression of the disease, which was also confirmed by copper uptake studies.

Mutation spectrum in Menkes disease
Menkes disease is one of the so called "new mutation disorders". These are usually X linked diseases, which are prone to new mutations, and almost every affected family shows a different alteration. Results to date indicate that the mutations leading to MD show a wide variety from cytogenetic abnormalities to single base pair changes.

Cytogenetically visible chromosome abnormalities comprise about 1% of the underlying genetic defect in MD. Three of these patients are females with balanced X;autosomal translocations as described above (fig 3). The only male patient with a cytogenetic aberration was detected during a systematic screening of 180 unrelated Menkes disease patients. We have recently localised the Xq13.3 breakpoint of the male patient and one of the female patients within MNK using Southern blot hybridisation (unpublished data).

In about 15-20% of patients the mutations are gross deletions or rearrangements. With Southern blot analysis we have identified rearrangements or partial gene deletions in about 45 unrelated Menkes patients (unpublished data) (fig 4). The boundaries of 30 partial gene

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**Figure 3** Chromosome painting of the metaphase and interphase chromosomes of the female Menkes patient with X;2 translocation, using hamster cell hybrid containing a single human X chromosome (CL-20, kindly provided by T Kruse). The normal X chromosome is indicated with an arrow and the derivative chromosomes with arrow heads. (The photograph was taken by T Hindkjær on a confocal laser microscope.)
Menkes disease:

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...exon coding for the ATG start codon and the first metal binding domain (fig 2A). The small-

deletions were further delimited by PCR amplification of the individual exons with intron specific primers (unpublished data). The deletions show great variability in size and location. The largest defect observed is the deletion of the whole gene except for the first two exons, the leader exon, and the second exon coding for the ATG start codon and the first metal binding domain (fig 2A). The small-
est mutation shown by this approach is the deletion of the leader exon (unpublished data).

The remaining genetic defects found in MNK are thus base pair changes or very small rearrangements which can be analysed by various PCR based methodologies. Das et al have characterised the mutations in 10 patients by RTPCR (reverse transcription-PCR) and chemical cleavage mismatch detection using RNA. To screen the mutations in our vast patient population, we first determined the genomic organisation of MNK25 and constructed primers to analyse each exon using genomic DNA. Using SSCP analysis (single strand conformation polymorphism) and subsequent direct sequencing of the exons (fig 5) we identified the genetic defect in 41 unrelated patients. All these point mutations show great variety, including missense and nonsense mutations, deletions and insertions of a single or more base pairs resulting in frameshifts and splice site mutations.

**Diagnosis of Menkes disease**

Initial diagnosis of Menkes disease is suggested by the clinical features (especially the typical hair changes) and supported by the reduced levels of serum copper and ceruloplasmin. However, interpretation of these markers may be difficult in the first months of life, as serum copper and ceruloplasmin levels may also be low in normal infants in this period. A definitive biochemical diagnosis exists and is based on the intracellular accumulation of copper owing to impaired efflux. Accumulation is evaluated in cultured cells by measuring radioactive copper ("Cu" retention after a 20 hour pulse, and impaired efflux is directly determined after a 24 hour pulse chase. However, these analyses demand expertise and are carried out only in a few specialised centres in the world.4,6,7 Postnatal diagnosis is performed on cultured fibroblasts and though a clear discrimination between affected and unaffected males is possible, milder phenotypes cannot be distinguished from the classical form.

Isolation of MNK and characterisation of the genomic organisation now opens up a new diagnostic possibility, mutation analysis. Demonstration of a defect in MNK will be the ultimate diagnostic proof and as our knowledge about the mutations in the different forms of MD increases, it may also be possible to distinguish the different clinical forms genotypically. However, mutation detection in Menkes disease is quite challenging; the large MNK transcript is organised in 23 exons, the genetic defect shows great variety, and each family has its own unique mutation.

**Prenatal Diagnosis**

**Aminocentesis**

Prenatal diagnosis of MD was initiated in 1974 by measuring "Cu accumulation in cultured amniotic fluid cells. However, copper accumulation in these cells is less pronounced than in fibroblasts and may occasionally give rise to diagnostic difficulties. Standardisation of the cell culture conditions is therefore crucial and needs expertise. Amniocentesis should be...
performed before the 18th week of gestation as the cell growth and hence copper accumulation is negatively correlated to the gestational age.

Chorionic villus sampling
First trimester prenatal diagnosis is carried out by determining the total copper content in chorionic villi by neutron activation analysis and was initiated in 1983. As the normal copper content in this tissue is very low (1 mg copper per mg tissue), the test is highly susceptible to exogenous copper contamination, which may give false positive results. The whole procedure of chorionic villus sampling must therefore be absolutely free of copper contamination and, as this condition is not always fulfilled, DNA analysis become a valuable supplement. False positive or negative results in biochemical analysis may also occur when the separation of the maternal tissue from the chorionic villi is not optimal. Verification of the prenatal diagnosis can be carried out by direct measurement of copper accumulation in the placenta and this method has also proven to be valuable for neonatal diagnosis of males at risk.

DNA analysis
DNA based prenatal diagnosis of MD is now possible and in some cases may be superior to biochemical diagnosis (fig 4). However, when the mutation in the family has not yet been identified, biochemical diagnosis will still be preferred as mutation detection in MD is quite challenging for the reasons mentioned above. Even in the cases where the genetic defect is known, the mutation detection method used has to be optimised before performing the prenatal diagnosis, whereas biochemical diagnosis is already well established. Diagnostic possibilities and risk factors should therefore be evaluated critically for each case.

CARRIER DIAGNOSIS
Carrier determination by measuring radioactive copper accumulation in cultured fibroblasts is possible. However, owing to random inactivation of one of the X chromosomes, negative results are not reliable and mutation analyses will therefore provide the ultimate proof of carriership (figs 4 and 5). Identification of carriers will also have an impact on the number of the prenatal diagnoses referred to our institute, as about 80% of the male fetuses tested are not affected, indicating that a substantial number of the females are not carriers. Using mutation analyses we have performed carrier diagnosis in about 15 families and in 11 of these families the mother was heterozygous for the mutation found in the index patient (unpublished data).

In families where the mutation is unknown, intragenic polymorphic markers may also enable carrier diagnosis. A BglI polymorphism has been identified within the coding region of MNK and it is localised to exon 10 (unpublished data). Furthermore, two polymorphic CA repeats within the gene (introns 2 and 5) were identified by us and others and these are currently being used for carrier diag-

nosis in the families where the mutation is as yet unknown (unpublished data).

Treatment of Menkes disease
In MD copper uptake is normal, but a defect in MNK disturbs the intracellular copper homeostasis and copper requiring enzymes cannot receive the copper necessary for their normal function. The objective of a treatment is thus to provide copper to the intracellular compartments where the copper enzymes are synthesised. However, parenteral administration of various copper preparations (as copper sulphate or copper-EDTA) did not produce substantial clinical improvement in MD patients. On the other hand, copper-histidine, the physiological copper complex found in human serum, had a positive effect in four unrelated MD patients, who are the oldest surviving patients receiving this therapy. They are now alive between the ages of 7 and 19 years with a milder clinical course resembling OHS. In two of these patients we have identified the genetic defect and they both have severe mutations (premature stop codons in exon 4 and exon 12) which would have resulted in a progressive clinical course if untreated. However, patients receiving copper-histidine after the first few months of age do not benefit in the same way (though survival may be prolonged), suggesting that the therapy should be initiated very early, before the occurrence of irreversible neurodegeneration. Studies with the mouse model of MD imply that there is a critical stage in brain development at which copper is essential, suggesting that induced premature delivery of affected infants for early treatment might be beneficial.

In the four patients mentioned above, though the neurological symptoms improved, the connective tissue abnormalities persisted. As lysyl oxidase is the enzyme involved in cross linking of collagen and elastin in connective tissues, these results suggest that the function of this enzyme does not appear to be corrected with copper-histidine administration. However, as indicated by the improvement of the neurological symptoms, copper seems to be delivered to some of the copper requiring enzymes using copper-histidine.

Wilson disease
Wilson disease (WD) is an autosomal recessive disorder of copper metabolism resulting from the toxic effects of copper, in contrast to MD mimicking copper deficiency (table 2). Though these two diseases have different clinical progression, the serum copper and ceruloplasmin are low in both cases. WD is mainly characterised by different degrees of liver disease, neurological or psychiatric symptoms, and clinical variability. The clinical features of WD are attributable to the toxic accumulation of copper in the liver and other tissues, such as kidney, brain, and cornea (Kaysen-Fleischer rings).

The Wilson disease locus (WND) has been assigned to chromosome 13q and was localised within an approximately 1.6 cM region at 13q14-21 by extensive linkage studies. In
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contrast to MD, visible cytogenetic rearrangements, which could lead to straightforward isolation of the gene, were not reported for WD. Identification of MNK encoding a putative copper transporting ATPase, and its hardly detectable expression in liver, led to the suggestion that WD might be caused by a defective liver specific copper transporter. Soon after, WND was cloned using sequences specific to MNK and to a heavy metal binding site within the amyloid β protein precursor. The 7.5 kb mRNA transcript is expressed predominantly in liver, kidney, and placenta, while the message is low in heart, brain, lung, muscle, and pancreas. The putative protein product is also a copper binding P type ATPase (designated ATP7B), highly homologous to ATP7A (57%). The WND gene (or ATP7B) has 21 exons in liver transcripts and 22 exons in kidney transcripts and its genomic structure shows a remarkable similarity to MNK, organised in 23 exons.

The Long-Evans Cinnamon (LEC) rat showing liver disturbances is suggested to be an animal model for Wilson disease. The rat homologue of WND has recently been isolated using human specific sequences and it has been shown to be defective in the LEC rat, confirming its status as a murine model for WD. The predicted protein product of the rat gene is also a copper binding P type ATPase (atp7b) showing 82% sequence similarity to ATP7B.

Isolation of WND now enables DNA based diagnosis of Wilson disease. Differing from Menkes disease, only point mutations or very small rearrangements are detected in Wilson disease. Mutation detection in Wilson disease is also challenging because of the large number of mutations in a 4.1 kb coding region. However, the presence of haplotypes specific to WND chromosomes may facilitate the mutation analyses as each haplotype is shown to be generally associated with a specific mutation. Mutation analysis in WD would be of great importance especially in diagnosing potential patients with no family history of Wilson disease and in early diagnosis of the sibs of affected patients.

New insights into normal and defective copper metabolism

Copper homeostasis depends on a balance between intestinal absorption and biliary excretion. Copper absorbed from the intestine is attached mainly to albumin and transported to the liver, the central organ of copper homeostasis, where storage and biliary excretion of copper and ceruloplasmin synthesis take place. Copper is secreted from the hepatocytes to the plasma bound to ceruloplasmin, but the mechanism by which it is delivered to different tissues is not fully understood. Regulation of intracellular copper homeostasis, from uptake to transport of the metal to its functional destination and export from the cell, is not well known either.

In Menkes disease, as in Wilson disease, serum copper and ceruloplasmin levels are low. In MD intestinal absorption of copper is grossly diminished. Copper accumulates in intestinal mucosa, kidney, spleen, lung, pancreas, muscle, and skin, while in the liver and brain the copper levels are below normal.

The finding of potential CpG islands has been the first molecular indication that MNK might be a housekeeping gene, required by all or most tissues, in line with the multisystemic character of the disorder. Consistent with the tissue distribution of copper in Menkes patients, the MNK transcript is also widely expressed. However, two organs, brain and liver, where the copper levels are low, need special attention. The MNK transcript is expressed in brain, supporting the suggestion that in MD low copper levels in this organ might be a secondary effect, owing to diminished transport across the blood-brain barrier. In liver the MNK transcript is hardly detectable and a straightforward explanation for the involvement of liver in MD awaits further studies.

In MD cellular copper uptake is normal and a defect in MNK causes intracellular accumulation of the metal. Copper enzymes are

<table>
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<th>Table 2: Comparison of Menkes disease with Wilson disease</th>
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<td>Incidence</td>
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<td>Serum Cu ‡</td>
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<td>Serum ceruloplasmin ‡</td>
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<tr>
<td>Genomic organisation</td>
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<tr>
<td>1500 AA copper binding ATPase</td>
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<tr>
<td>All tissues, hardly detectable in liver</td>
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<tr>
<td>23 exons spanning 150 kb genomic region</td>
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<td>1% cyogenetic abnormalities</td>
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<td>20% gross rearrangements</td>
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<td>80% small base pair changes</td>
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</table>
deprived of copper necessary for their normal function, while the metal is bound to metallothionein, a cysteine rich heavy metal binding protein, which protects the cell from toxic effects of the free ion. It is thus conceivable that ATP7A delivers copper to the enzymes and is also involved in copper efflux. However, copper accumulation is likely to result from a defect in copper translocation across an intracellular compartment, rather than a defective copper export across the plasma membrane. ATP7A is thus likely to be located in one or more intracellular compartments or organelles, possibly endoplasmic reticulum or golgi appara- ratus or both, and is involved in the regulation of the intracellular copper pool by translocating the metal through membranes.

In WD it is likely that ATP7B plays a direct role in incorporation of copper into ceruloplas- min and in biliary copper excretion. A defect in WD results in deficient production of ceruloplasmin and copper starts accumulating in the hepatocytes bound to metallothionein. However, in later stages copper accumulation reaches a toxic level and subsequent overflow from liver to other tissues, like brain, cornea, and kidney, leads to a variety of damage to these organs also. Expression of the WD transcript predominantly in liver is in line with the clinical course and the underlying biochemical defect.

MD is characterised by an overall copper deficiency, while WD is associated with copper toxicity. However, the functions of ATP7A and ATP7B appear to be quite similar. They are both predicted to be copper translocating membrane proteins likely to be located in the intracellular compartments or organelles. ATP7A has a role in the delivery of copper to different enzymes, while ATP7B is involved in supplying copper to ceruloplasmin. Furthermore, both proteins are likely to be involved in the excretion of copper from the cell via intracellular compartments or organelles.

Though an important step has been taken by the cloning of the genes defective in Wilson and Menkes diseases, and their murine homologues, the enigma of intracellular copper metabolism is not solved yet. Identification of other potential proteins involved in the copper uptake and intracellular transport will undoubtedly increase our understanding of cellular copper homeostasis.

Copper translocating P type ATPases in other species

ATP7A shows a remarkable sequence similarity to P type ATPases involved in copper translocation in *Enterococcus hirae,* leading to the prediction of the first intracellular copper binding P type ATPase in eukaryotes. Since then, a number of related genes have been described in eukaryotes and prokaryotes. These are all predicted to encode P type ATPases with one or more copper binding domains at the amino-terminus. The human members of this copper associated subfamily of the P type ATPases, ATP7A and ATP7B, show high sequence homology. They both have six predicted copper binding domains, including the consensus GMXXCXCG motif, and copper binding of one of these domains has recently been reported for ATP7B.

The two other eukaryotic members of this family are the mouse protein ATP7a with six copper binding domains and the rat protein ATP7b with five. Both proteins show remarkable sequence homology to their human counterparts.

In *Saccharomyces cerevisiae* two genes mapping to different chromosomes were identified and they were predicted to encode for proteins (Pca1 and Ccc2) belonging to this family. Ccc2 contains two copper binding motifs and may provide copper from the cytosol into an extracytosolic compartment. This is indeed analogous to the mechanism suggested for ATP7A and ATP7B. Pca1 has a single copper binding motif and may have a role in copper extrusion from the cell. Among the bacterial members of this family, there are three copper ATPases, Cop A and Cop B described in *Enterococcus hirae* and CtaA described in *Synechococcus* 7942. All these proteins are involved in copper homeostasis, where CopA and CtaA are serving in the uptake and CopB in the extrusion of copper, in the respective bacteria. It is likely that not only the eukaryotic cells but most of the prokaryotic cells as well have copper enzymes involved in electron transfer and therefore require copper transport systems. These systems in bacteria and yeast may thus provide an experimentally accessible model for understanding the overall intracellular copper homeostasis in higher eukaryotes. Recently, a role of Ccc2 in iron metabolism has been described in yeast and it was shown to be providing copper to a ceruloplasmin-like oxidase required for iron uptake. Identification of other potential copper transport proteins in bacteria or yeast, and understanding their function, may thus provide insight also into the connection of copper and iron homeostasis in man.

Despite recent advances, copper homeostasis is still not fully understood. However, to solve this enigma we now have the most necessary tools, the animal models, and simple biological systems such as bacteria and yeast.

Menkes disease: recent advances and new aspects


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Menkes disease: recent advances and new aspects.

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