A 1 year and 4 month old child with Mucopolysaccharidoses type-II: a clinical case report from Ethiopia

Solomie Jebessa Deribessa MD, MPH,1* Mekdes Endale Bisrat, MD2, Zewdu Terefework, PHD3 Shane C. Quinonez, MD4

Key Clinical message: Mucopolysaccharidosis-type-II/Hunter syndrome is an X-linked recessive lysosomal storage disorder. We report a 1 year and 4 month old boy with coarse facial appearance, macrocephaly, dermal melanocytosis, widened wrists, kyphotic deformity and left sided inguinal hernia, consistent with MPSII, subsequently confirmed with genetic tests. There is no family history, parents are counselled.

Abstract:

1 Infectious Disease consultant, department of pediatrics and child health St.Paul’s Hospital Millennium Medical College, Email: solomejebessa@gmail.com; *Dr. Solomie Jebessa is first and corresponding author
2 Pediatrics Resident, department of pediatrics and child health St.Paul’s Hospital Millennium Medical College, Email: mekdesendale21@gmail.com
3 MRC-ET, Advanced Laboratory, Bole suncity, Gerj, Addis Ababa Ethiopia. Email: zterefework@gmail.com
4 Department of pediatrics and communicable disease, Division of genetics, Metabolism and Genomic Medicine, University of Michigan, USA, Email: squinon@med.umich.edu

This is the author manuscript accepted for publication and has undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi: 10.1002/CCR3.5122

This article is protected by copyright. All rights reserved
Background: Mucopolysaccharidoses (MPSs) are a class of lysosomal storage disorders resulting in progressive disease manifestations and are caused by pathogenic variants in genes coding for enzymes needed to degrade glycosaminoglycans. While most of the seven MPSs are autosomal recessive disorders, MPS-II, also known as Hunter syndrome, is inherited in an X-linked recessive manner and is the most common MPS.

Case: Here we report a 1 year and 4 month old boy who presented with delayed developmental milestones, back deformity and left scrotal swelling noticed by parents at one year of age. He has coarse facial appearance with macrocephaly, widened wrists, congenital dermal melanocytosis on his back, kyphotic deformity in the thoracolumbar area and left sided inguinal hernia all consistent with a suspected MPS-II diagnosis. The MPS-II diagnosis was subsequently confirmed with genetic testing of the IDS gene.

Conclusion: To our knowledge this is the first case of MPS-II reported from Ethiopia. This case shows the importance of early clinical recognition of genetic conditions and the utility of genetic testing for confirmation. The diagnosis provided important surveillance and natural history information for the patient’s providers and family.

Keywords: Mucopolysaccharidosis, Glycosaminoglycans (GAG), iduronate-2-sulfatase (IDS), Hunter syndrome

Introduction

Mucopolysaccharidoses (MPS) are a group of lysosomal storage disorders resulting in progressive disease manifestations and are caused by pathogenic variants of genes coding for enzymes needed to degrade glycosaminoglycans. Glycosaminoglycans (GAG) are long-chain complex carbohydrates composed of uronic acids, aminosugars and neutral sugars. The major GAGs are chondroitin-4-sulfate, chondroitin-6-sulfate, heparin sulfate, dermatan sulfatan, keratin sulfate and hyaluronan. These substances are synthesized, with the exception of hyaluronan, linked to protein to form proteoglycans and are major constituents of the ground substance of connective tissue and of nuclear and cell membranes. Degradation of proteoglycans starts with proteolytic removal of the protein core, followed by the stepwise degradation of the GAG moiety. 1
Each MPS is caused by the deficiency of distinct lysosomal enzymes required for the stepwise degradation of GAGs. Failure of this degradation process, due to an absence or dysfunctional lysosomal enzyme, results in the intralysosomal accumulation of GAG fragments. Distended lysosomes accumulate in the cell, interfere with cell function and lead to characteristic patterns of clinical, radiologic and biochemical abnormalities.  

Seven types of MPS are classified based on deficiency of one of the eleven specific lysosomal enzymes and are numbered MPS-I through MPS-IX (excluding MPSV and VIII which are no longer used). The difference in clinical finding of these seven MPSs are indicated in the table below (Table 1). Most mucopolysaccharidoses (MPS-I-H/Hurler syndrome, MPS-I-S/Scheie syndrome, MPS-III/Sanilippo syndrome, MPS-IV/Morquio syndrome, MPS-VI/Maroteaux-Lamy syndrome, MPS-VII/Sly syndrome, MPS-IX, MPS-Plus Syndrome/MPS-PS) are autosomal recessive disorders with the exception being MPS-II (Hunter syndrome) which is X-linked recessive. Given this, Hunter syndrome typically affects males however it has been rarely seen in females secondary to skewed X-inactivation of the normal X-chromosomes and expression of the maternally inherited mutated IDS allele.  

Table 1: I-H, Hurler syndrome; I-S, Scheie syndrome; II, Hunter syndrome; III, Sanfilippo syndrome; IV, Morquio syndrome; VI, Maroteaux-Lamy syndrome; VII, Sly syndrome.

| Manifestations          | MUCOPOLYSACCHARIDOSIS (MPS) TYPE |
|-------------------------|----------------------------------|
|                         | I-H | I-S | II | III | IV | VI | VII |
| Intellectual disability | +   | -   | ±  | +   | -  | -  | ±   |
| Coarse facial features  | +   | (+) | +  | +   | -  | +  | ±   |
| Corneal clouding        | +   | +   | -  | -   | (+)| +  | ±   |
| Visceromegaly           | +   | (+) | +  | (+)| -  | +  | +   |
| Short stature           | +   | (+) | +  | -   | +  | +  | +   |
| Joint contractures      | +   | +   | +  | -   | -  | +  | +   |
| Dysostosis multiplex    | +   | (+) | +  | (+)| +  | +  | +   |
| Leucocyte inclusions    | +   | (+) | +  | +   | -  | +  | +   |
| Mucopolysacchariduria   | +   | +   | +  | +   | +  | +  | +   |

+, Presence of manifestation, −, absence of manifestation; ±, possible presence of manifestation; (+), mild manifestation.

This article is protected by copyright. All rights reserved.
Hunter syndrome (Mucopolysaccharidosis type II, OMIM# 309900) was first described by Charles Hunter in 1917, a Canadian Professor of Medicine, when he reported medical histories of two affected brothers. Hunter syndrome is one of the most common MPSs with an estimated prevalence of 1 in 100,000 to 170,000 male births. It is a life-limiting multi-systemic disorder caused by deficiency of iduronate 2-sulfatase (IDS). The IDS gene is mapped to Xq28 with pathogenic sequence variants of IDS detected in about 80% of patients with MPS-II with major deletion or rearrangements of IDS found in the remaining 20% with these usually associated with a more severe clinical phenotype. The deficiency of IDS leads to an accumulation of heparan sulfate and chondroitin sulfate B (dermatan sulfate) in cellular lysosomes, interfering with their function.

The clinical presentation of patients with Hunter syndrome (MPS-II) vary from severe to milder forms with signs and symptoms apparent usually by two to four years of age. Findings of affected patients include macrocephaly, short neck, broad chest, delayed tooth eruption, hearing loss, coarse facial features with thick lips, thick nostrils and macroglossia. Patients have also impaired growth, short stature, joint stiffness with restriction of movements and dysostosis multiplex noted on radiographs. Chronic diarrhea may also occur as a result of GAG accumulation in the gastrointestinal system with other features including hepatosplenomegaly, inguinal and umbilical hernias. Neurologic system involvement includes intellectual disability, delayed developmental milestones, communicating hydrocephalus due to thickened meninges and spastic paraplegia. Cutaneous manifestations include congenital dermal melanocytosis observed in African and Asian patients with grouped skin papules found in some patients. A distinguishing feature of MPS-II is an absence of corneal clouding which is often present in patients with MPS-I. In patients with MPS II total urinary GAGs are elevated, particularly Dermatan and Heparan sulfate, with a definitive diagnosis made by identifying deficient iduronate 2-sulfatase activity or a pathogenic IDS variant.

Individuals with milder forms of MPS-II will have slow progression of somatic symptoms with minimal neurologic involvement and have been reported to live 65 and 87 years. Severly affected patients with MPS-II will show progressive symptoms including significant neurologic deficits which may be present for 10-15 years preceding death.

Case Presentation
The proband is an Ethiopian boy seen initially at 1 year and 4 months of age whose parents brought him to St. Paul’s Hospital in Addis Ababa, due to left side scrotal swelling, lower back swelling, and delayed sitting, standing and walking initially noted at one year of age. The family identified the back swelling as a possible cause for the delayed sitting, standing and walking. The mother claimed that she saw the scrotal swelling especially during periods of the patient crying. He had no history of swallowing difficulty, vomiting or diarrhea with no difficulty breathing or shortness of breath or abnormal body movements. The patient was exclusively breastfed for the first 6 months of his life then complemented with porridge made of mixed cereals and cow’s milk. Gradually his parents noted decreased oral intake of uncertain etiology with a resultant decrease in growth. He was exposed to sunlight at least 3 times per week since the age of 3 months without the application of any ointment. Currently, he is fully vaccinated as per the Ethiopian vaccination program schedule.

The proband’s mother has no history of previous abortions or stillbirths with normal antenatal follow-up and negative routine tests. She delivered the proband via spontaneous vaginal delivery, with the baby noted to cry immediately and not requiring a neonatal intensive care unit (NICU) admission. He is the first child from a non-consanguineous marriage of an orthodox Christian family. The mother is 25 years old and the father is 32 years old with no similar family history present though the two maternal aunts have no children yet.

On the initial presentation to our hospital, the diagnosis of rickets was made with wrist x-ray and was treated with megadose of Vitamin D and Calcium supplementation. On examination (Figures 1, 2 and 3) he had coarse facial features with a broad and large forehead and a flattened nasal bridge. At his initial evaluation his pulse rate was 122 beats per minute, respiratory rate was 18/minute and axillary body temperature was 36.8 ° C. His anthropometries showed a body weight of 10.3 kg (between 0 and -1 SD for his age), length of 73 centimeters (below -3SD for his age); weight/length ratio between +1 and +2 SD and a head circumference of 52 centimeters which is over+3SD for his age. He had pale conjunctiva with no icterus or corneal clouding. His tongue was not enlarged. His neck was short with no lymphadenopathy. His chest was broad and symmetric, without deformity and clear lung sounds with good air entry. Cardiovascular examination revealed a well heard and normal S1 and S2 without a gallop or murmurs. His abdomen was protuberant, soft and moved with respiration with no

5 * SD: Standard Deviation: all SD measurements according to the World Health Organization growth charts for children under five years of age.

This article is protected by copyright. All rights reserved
tenderness or mass and normoactive bowel sounds. There was a 4 X 5 centimeter left reducible mass extending from the inguinal area to the scrotum. On genitourinary examination, there was a normal external male genitalia with bilaterally descended testicles. On musculoskeletal examination, he had broad hands and widened wrists with kyphosis of the thoraco-lumbar area which was non tender. On cutaneous examination, he multiple hyper-pigmented macules and patches were noted on his back, the dorsum of his left hand and on the left side of his face. Neurologically he was alert, both pupils were midsized and reactive to light, with normal tone and 5/5 strength with deep tender reflexes of 2/4 in all extremities.

**Ophthalmologic evaluation:** Normal visual acuity in both eyes for his age, normal intraocular pressure in both eyes, pupils were symmetrically round, reactive and regular in both eyes. Normal eyelids and well oriented lashes, normal conjunctiva, some dark-brown pigmenitary changes on the sclera (benign melanocytic nevi), clear and normal sized cornea, deep anterior chamber, brown and normal iris, clear lens, clear vitreous, pink oval and normal optic disc, shiny macula, no pigmentary retinal changes and normal vasculature.

**ENT evaluation:** Bilaterally normal hearing tests.

**Laboratory Workup:** His white blood count was 8300/mm³ (normal for age: 4000 – 12000/mm³) with Neutrophils of 38.7%, Lymphocytes of 54%. Hemoglobin was 6.6 g/dl (normal for age 10.5-14 g/dl) hematocrit of 23.8% (normal for age 32-42%), platelet count of 235,000/mm³ (normal for age: 150,000 – 400,000/mm³), serum Phosphorous of 0.88mmol/L (normal for age: 1.25-2.10mmol/L), Alkaline phosphatase 1158 U/L (Normal for age 145-420 U/L), ionized calcium 1.18 mmol/L (normal for age 1.2-1.38 mmol/L) and serum vitamin D,25-hydroxy of 40.18 (normal value 75-250).

**Imaging results:**

Radiologic evaluation included thoracolumbar xrays that showed normal bone mineral density though with 17 degree dextroscoliosis from T4 to T11 and kyphosis of the lumbar vertebrae at L1-3. There was also hypoplasia of the L2 vertebral body with grade one retrospondylolisthesis, cortical discontinuity of the pedicle of L2 vertebra noted as well as irregularity of the L2 vertebra. (Figures 4 and 5) MR imaging of the same area showed anterior irregular beaking of the L2 vertebra, narrow L1-L2 intervertebral disc space with no other abnormalities detected. There was though noted decreased bone mineralization with cupping and fraying over the distal ulnar and radial metaphysis of both right and left wrist x-rays.

This article is protected by copyright. All rights reserved
(Figure 6) Skull x-ray was reported as normal. (Figure 7) An echocardiogram revealed concentric hypertrophy of the left atrium and left ventricles with no other abnormalities detected.

**Molecular Analysis:**

The Multiplex ligation-dependent probe amplification (MLPA) method (Schouten et al. 2002) kits P125, P164 and P309 (MRC-Holland, The Netherlands) were used to screen the DNA which was obtained from peripheral blood sample of the Proband and his mother for copy number changes, deletions and duplication on the IDS gene located on the Xq28 region of the X chromosome.

The MLPA investigation also included searching for other possible causes of metabolic disorders as a result of genetic alteration on the chromosomal and mitochondrial DNA. The extent of rearrangement or copy number changes on the Xq28 region was investigated using MLPA probes targeting genes (FMR1 and AFF2) located upstream and (IDSP1, MTM1, MTMR1, FLNA, DKC1) located downstream of the IDS gene. Testing showed deletions of exons 4, 5, 6 and 7 of the IDS gene (Figure 8); further sequencing could not be done since it is not available in our setup/country.

The MLPA analysis was also performed on a DNA obtained from the mother. The result showed heterozygous deletions of exons 4, 5, 6 and 7 of the IDS gene.(Figure 9)

In addition, deletion of the RNR1 and RNR2 genes, which code for mitochondrial ribosomal RNA, were incidentally noted (Figure 10). The significance of the incidental finding, a “deletion” of the mitochondrial 16s rRNA gene, to the clinical scenario cannot be determined from this test and requires a more thorough investigation as this was identified by deletion of four MLPA probes flanking the gene. Further enzyme testing and sequencing of both the IDS and rRNA genes are required to identify the molecular etiology, which we don’t have in our setting. The deletions observed on the RNR1 and RNR2 genes were also present in the mother just as in the proband.(Figure 11)
Discussion:

Hunter syndrome is an X-linked recessive, multisystemic and progressive disease. Here we present the first case of Hunter Syndrome reported from Ethiopia. Patients with Hunter syndrome typically present between 2 to 4 years of age though severe forms may present between 6 and 24 months of age. Based on the age of presentation, phenotype consistent with MPS-II and the multi-exonic IDS deletion, this patient’s presentation is consistent with a severe Hunter Syndrome presentation. This is especially important to note as a previously reported patient with a similar IDS variant presented with an attenuated form of the disease. \(^9\)

The proband had coarse facial features with macrocephaly, short neck, broad chest, inguinal hernia with impaired growth and short stature similar to individuals with Hunter syndrome. The patient did though lack other features of the condition including thick lips, thick nostrils delayed tooth eruption, large tongue, hearing loss and chronic diarrhea though with time these features may develop.\(^3\)

Individuals with Hunter syndrome often have joint stiffness with restriction of movements and dysostosis multiplex with thickened ribs and ovoid vertebrae. The proband at the time of evaluation was 1 year and 4 months old and was unable to sit, stand and walk, parents ascribed this to the back deformity, his imaging also showed dextroscoliosis of 17° from T4 to T11, kyphosis of lumbar vertebrae at L1-3, with hypoplasia of the L2 vertebral body with grade one retrospontylolisthesis, cortical discontinuity of pedicle of L2 vertebra with irregularity of the right superior end plate and ovoid lumbar (1-4) vertebra.\(^3\)

Neurologic system involvement of MPS-II typically includes intellectual disability, delayed developmental milestones for age, communicating hydrocephalus and spastic paraplegia. Our patient had noted delayed developmental milestone but no other neurologic deficits. Cutaneous manifestations include congenital dermal melanocytosis observed in African and Asian patients and grouped skin papules which was seen in our patient.\(^3,7\)

Though we were unable to measure urinary GAGs or iduronate-2-sulfatase activity the clinical findings and IDS molecular analysis support the MPS-II diagnosis. We identified deletions on exons 4, 5, 6 and 7 of the IDS gene on both the proband and maternal DNA. Similar variant has been described previously in a patient with an attenuated form of MPS II.\(^9\) Among the previously described

This article is protected by copyright. All rights reserved
alternations, 28.2 % are large alterations that include complete and partial deletions and rearrangements and 71.8 % are small deletions and single nucleotide polymorphisms resulting in various mutations.\textsuperscript{10, 11} Additionally Complex rearrangements due to illegitimate recombinations of these regions resulting in Hunter syndrome are reported.\textsuperscript{12, 13, 14}

The incidental finding of the $RNR1$ and $RNR2$ deletions is of uncertain significance at this point. The $IDS$ exon deletions were felt to be causative as similar deletions have been seen in previously reported Hunter syndrome patients.\textsuperscript{9, 15} Further enzyme testing and sequencing are required to identify the significance further (which we don’t have in our setting); though pathogenic variants in these 2 genes have not yet been identified to cause human disease. The maternal screening confirms the inheritance of the deleted exons on the $IDS$ gene as well as the deletion of the maternally inherited mitochondrial rRNA gene deletion.

In patients with Hunter syndrome early enzyme replacement therapy is indicated before irreversible organ damage occurs, however this management is not available in Ethiopia. Follow-up of this patient should involve a multidisciplinary team including pediatricians, orthopedists, neurosurgeons, ophthalmologists, otorhinolaryngologists and cardiologists; which is feasible in our setting for the proband.

\textbf{Aknowledgemnt}

Authors aknowledgement extends to \textit{MRC-ET, Advanced Laboratory}, in Addis Ababa for doing the genetics test for free; to pediatrics department; to the radiology department; to the ophthalmology department (particularly Dr. Melessew Merkebie) and to the ENT department at St.Paul’s Hospital Millennium Medical college, Addis Ababa.

\textbf{Authors' Contribution}

1st Author: Solomie Jebessa Deribessa, who is also the corresponding author :Have made substantial contributions to conception and design, or acquisition of data, or analysis and interpretation of data; and has been involved in drafting the manuscript or revising it critically for important intellectual content.

This article is protected by copyright. All rights reserved
Hunter syndrome a case report from Ethiopia

2nd Author: Mekdes Endale Bisrat: Have made substantial contributions to conception and design, or acquisition of data, and has been involved in drafting the manuscript.

3rd Author: Zewdu Terefework: Have made substantial contributions to design, or acquisition of data, or analysis and interpretation of data; and has been involved in drafting the manuscript or revising it critically for important intellectual content.

4th Author: Shane C. Quinonez, Have made substantial contributions to design, interpretation of data; and has been involved in drafting the manuscript or revising it critically for important intellectual content.

Authors do not have any conflict of interest.

Written consent has been obtained from the patient parent.

References

1 Spranger, J.W. (2016). Mucopolysaccharidoses. In: Nelson textbook of pediatrics. Elsevier, pp.1551–1552.

2 Guillén-Navarro, E., Domingo-Jiménez, M., Alcalde-Martín, C., Cancho-Candela, R., Couce, M., Galán-Gómez, E. and Alonso-Luengo, O. (2013). Clinical manifestations in female carriers of mucopolysaccharidosis type II: a spanish cross-sectional study. Orphanet Journal of Rare Diseases, 8(1), p.92.

3 Khan, S.A., Hira Peracha, Ballhausen, D. and Shunji Tomatsu (2017). Epidemiology of mucopolysaccharidoses. [online] ResearchGate. Available at: https://www.researchgate.net/publication/317242102_Epidemiology_of_mucopolysaccharidoses [Accessed 21 Jun. 2021].

4 Germaine, L Defendi. “Hunter Syndrome (Mucopolysaccharidosis Type II): Background, Pathophysiology, Epidemiology.” EMedicine, 4 2018, emedicine.medscape.com/article/944723-overview#:~:text=In%201917%20at%20The%20Royal.

This article is protected by copyright. All rights reserved
5 Burton, B.K., Jego, V., Mikl, J. and Jones, S.A. (2017). Survival in idursulfase-treated and untreated patients with mucopolysaccharidosis type II: data from the Hunter Outcome Survey (HOS). Journal of Inherited Metabolic Disease, 40(6), pp.867–874.

6 Richter, T., Nestler-Parr, S., Babela, R., Khan, Z.M., Tesoro, T., Molsen, E., Hughes, D.A. and International Society for Pharmacoeconomics and Outcomes Research Rare Disease Special Interest Group (2015). Rare Disease Terminology and Definitions—A Systematic Global Review: Report of the ISPOR Rare Disease Special Interest Group. Value in health : the journal of the International Society for Pharmacoeconomics and Outcomes Research, [online] 18(6), pp.906–14. Available at: https://www.ncbi.nlm.nih.gov/pubmed/26409619 [Accessed 31 Oct. 2019].

7 Alkhalil, M., Alabsi, M. and Alabsi, N. (2016). Hunter Syndrome. British Journal of Medicine and Medical Research, 15(12), pp.1–7.

8 Jan P. Schouten JP et al. (2002). Relative quantification of 40 nucleic acid sequences by multiplex ligation-dependent probe amplification. Nucleic Acids Res. 2002 June 15; 30(12):e57. Doi:10.1093/nar/gnf056

9 Birot, A.M.; Bouton, O.; Froissart, R.; Maire, I.; Bozon, D. IDS Gene-pseudogene exchange responsible for an intragenic deletion in a Hunter patient. Hum. Mutat. 1996, 8, 44–50.

10 Froissart, R., Da Silva, I. M. & Maire, I. Mucopolysaccharidosis type II: an update on mutation spectrum. Acta. Paediatr. Suppl. 96, 71–77 (2007)

11 HY Lin · 2020. Identification and Functional Characterization of IDS Gene Mutations Underlying Taiwanese Hunter Syndrome (Mucopolysaccharidosis Type II) Int. J. Mol. Sci. 2020, 21(1), 114

12 Karsten S, et al. Novel type of genetic rearrangement in the iduronate-2-sulfatase (IDS) gene involving deletion, duplications, and inversions. Hum Mutat. 1999;14(6):471-6. doi: 10.1002/(SICI)1098-1004(199912)14:6<471::AID-HUMU5>3.0.CO;2-5.PMID: 10571944

13 Galvis, J., Gonzáles, J., Uribe, A., Velasco, H. (2015). Deep Genotyping of the IDS Gene in Colombian Patients with Hunter Syndrome. JIMD Rep. February, 19, 101–109.

14 Bunge, S., et al. Homologous nonallelic recombinations between the iduronate-sulfatase gene and pseudogene cause various intragenic deletions and inversions in patients with mucopolysaccharidosis type II. Eur. J. Hum. Genet. 6, 492–500 (1998).

15 Peter D. Stenson et al, Human Gene Mutaion Database (HGMD) 2003update; Human Mutaion 21:577-581(2003)
This article is protected by copyright. All rights reserved