

Clinical Utility of Elosulfase Alfa in the Treatment of Morquio A Syndrome

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Abstract: Mucopolysaccharidosis type IVA (MPS IVA or Morquio A) is an autosomal recessive disorder and is one of the lysosomal storage diseases. Patients with MPS IVA have a striking skeletal phenotype but normal intellect. The phenotypic continuum of MPS IVA ranges from severe and rapid progress to mild and slow progress. The diagnosis of MPS IVA is usually suspected based on abnormal bone findings and dysplasia on physical examination and radiographic investigation in the preschool years. In the past, only supportive care was available. Due to the early and severe skeletal abnormalities, the orthopedic specialist was usually the main care provider. However, patients need aggressive monitoring and management of their systemic disease. Therefore, they need an interdisciplinary team for their care, comprising medical geneticists, cardiologists, pulmonary specialists, gastroenterologists, otolaryngologists, audiologists, and ophthalmologists. After the US Food and Drug Administration approved elosulfase alfa in 2014, patients older than 5 years could benefit from this treatment. Clinical trials showed clinically meaningful improvements with once-a-week intravenous dosing (2.0 mg/kg per week), significantly improving the 6min walk test, the 3min stair climb test, and respiratory function when compared with placebo. Elosulfase alfa is well-tolerated, and there is a good response indicated by decreasing urine glycosaminoglycans.

Keywords: elosulfasealfa, enzyme replacement therapy, GALNS, lysosomal storage disorder, Morquio A, MPS IVA, mucopolysaccharidosis

Introduction

Mucopolysaccharidosis IVA (MPS IVA, Morquio A syndrome, Morquio–Brailsford syndrome, OMIM 253000) is a lysosomal storage disease (LSD). It is an inherited autosomal recessive disorder characterized by a deficiency in the enzyme N-acetylgalactosamine-6-sulfatase (GALNS; EC 3.1.6.4) due to mutations of the *GALNS* gene. The result is an accumulation of the glycosaminoglycans (GAGs), keratan sulfate (KS), and chondroitin-6-sulfate (C6S) in the tissues, bones, and major organs.¹⁻⁵

Morquio A syndrome was first described by a pediatric physician, Dr. Luis Morquio, from Uruguay, and an English radiologist, Dr. James Brailsford, from Birmingham, UK, in 1929.^{6,7} Therefore, the disease is sometimes called the Morquio–Brailsford syndrome. Dr. Morquio described 4 siblings from a Swedish family with corneal clouding, aortic valve disease, and high levels of urinary KS (uKS).⁶

The prevalence of MPS IVA is 1/323,000 in Denmark, 1/599,000 in the UK, 1/926,000 in Australia, and 1/1,872,000 in Malaysia. The birth prevalence rate of MPS

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IVA ranges from 1/71,000 in the United Arab Emirates to 1/500,000 in Japan.^{8,9} Patients with Morquio A syndrome appear healthy at birth and, subsequently, develop multiorgan signs and symptoms. They may be diagnosed as early as 6 months old.

The *GALNS* gene is located on chromosome 16q24.¹⁰ It contains 14 exons with 2339 base pairs and is approximately 50 kb. It encodes a 522-amino acid enzyme. There are numerous heterogeneous mutations of the *GALNS* gene which can occur throughout the coding sequence. According to ClinVar Database, as of August 2021, 738 known variants of the *GALNS* gene were found, of which 276 are pathogenic or likely pathogenic, 327 are of unknown clinical significance, and 135 are benign or likely benign.

The signs and symptoms of patients with MPS IVA include short stature for their age with a short neck, abnormal skeletal development and spinal deformities, joint hypermobility and laxity, genu valgum (knock knees), large elbows and wrists, a waddling gait, and a bell-shaped chest. They may have deficient tooth enamel, abnormal heart development, respiratory compromise, corneal clouding, hearing impairment, and mild hepatosplenomegaly^{11–16}. Patients also have greater anesthetic risks because they have macroglossia, temporomandibular joint stiffness, difficult or failed intubations, abnormal laryngeal anatomy, trachea deformity, and subglottic narrowing.¹⁷ Figure 1 showed computed tomography angiogram in a 16-year-old boy with MPS IVA.¹⁸ The deficient enzyme activity and resulting abnormal GAG excretion can be detected in the first trimester.^{19,20}

Compared with other types of MPS, Morquio A does not involve the brain or cause significant impairment of cognitive development.^{21,22} The range of clinical manifestations is diverse in Morquio A. Due to the genetic heterogeneity, a spectrum of disease progression is present (ie, severe/classical, intermediate, and mild/attenuated). Patients with milder “nonclassical/attenuated” forms can have a normal life expectancy.²³ According to a study by Lavery et al,²⁴ the main cause of death in nearly two-thirds of patients (63%) with Morquio A is respiratory failure. Other causes are cardiac failure (11%), posttraumatic organ failure (11%), complications of surgery (11%), and myocardial infarction (4%).

Diagnosis of Morquio A

Morquio A syndrome is initially suspected because of the appearance of the symptoms described above. However,

differentiating it from other disorders is difficult because of its heterogeneous expressions, which may delay the diagnosis.^{23,25} At ages 2 to 3 years, the symptoms progress due to a significant GAG accumulation in the tissues, bones, and organs. Because there is a natural decline in uKS levels with age, age-appropriate reference ranges in screening tests are needed to avoid false negatives, especially in older patients.^{26,27} To confirm the diagnosis of Morquio A syndrome, we must detect low *GALNS* activity in leukocytes and fibroblasts and perform molecular testing.^{23,26}

Management of Morquio A

A multidisciplinary approach is necessary for managing Morquio A. Having orthopedic intervention in MPS IVA patients is important.²⁸ They need an experienced anesthesia care team because of the possible airway complications secondary to GAGs accumulation in the upper and lower airways. There may be odontoid hypoplasia, impaired respiratory function, and cardiac problems.^{29,30} Appropriate growth charts should be used to avoid severe growth abnormalities in children.⁴ Because of the abnormalities of the spine and malalignment of both upper and lower extremities, MPS IVA patients usually have disabilities. They should have a close evaluation by orthopedics specialists and neurosurgeons at least annually. Because of the risk of spinal cord compression, they should have a neurological examination at least every 6 months, including annual radiography of the spine and lower extremities and whole-spine MRI or when clinically indicated.

The involvement of a pulmonary specialist is needed to manage respiratory compromise and sleep apnea.^{28,31,32} A cardiology team should also closely follow patients to monitor and treat valvular heart disease.^{33,34} For hearing impairment, an audiology team should treat them aggressively as necessary, and eye examinations and intervention for corneal clouding will help to optimize visual outcomes.³ If patients have conductive hearing loss due to middle ear fluid, they should be treated with ventilation tubes. Patients should have slit lamp biomicroscope of the cornea. If vision is reduced due to corneal clouding, keratoplasty is considered. Although hepatomegaly is not acutely dangerous, other organs may be affected by the size of the liver because MPS IVA patients have short stature. Furthermore, pain and cosmetic difficulties may develop due to dental abnormalities if patients do not have appropriate dental care early.³ To reduce the total number of hospital visits and facilitate care



Figure 1 Computed tomography angiogram in a 16-year-old boy with mucopolysaccharidosis type IVA. **(A)** Three-dimensional reconstruction of the trachea in a sagittal oblique projection shows a severe narrowing of the trachea at the thoracic inlet (arrow). **(B)** Sagittal image shows that the brachiocephalic artery (dashed arrow) contributes to crowding at the thoracic inlet but does not directly indent the trachea (solid arrow). **(C)** Three-dimensional coronal reconstruction posteriorly in the chest shows tortuosity of the descending thoracic aorta, crossing the midline (arrow). Reprinted by permission from Springer Nature, Averill LV, Kecskemethy HH, Theroux MC, et al. Tracheal narrowing in children and adults with mucopolysaccharidosis type IVA: evaluation with computed tomography angiography. *Pediatr Radiol.* 2021;51(7):1202–1213. Figure 1 is copyright protected and excluded from the open access licence.¹⁸

coordination, multidisciplinary clinics can improve patient satisfaction and quality of life.

The care team needs to improve the learning and social environment for MPS IVA patients due to their numerous systemic complications. MPS IVA patients have normal intelligence; thus, optimizing the environment and neurologic protection can prevent damage and facilitate good outcomes for patients.²⁸ The “International guidelines for

the management and treatment of MPS IVA syndrome” was released in 2014. The guidelines summarized the extensive experience in managing MPS IVA based on two meetings of international experts. It includes an extensive baseline evaluation and reevaluation at intervals to identify systemic changes early to allow for monitoring and intervention. Table 1 shows the monitoring recommendations for MPS IVA patients.^{22,28,35}

Table I Monitoring Recommendations for Mucopolysaccharidosis Type IVA Patients

At Diagnosis	Every 6 Mo	Annually	Every 1–3 Yr
Medical history Physical examination Growth evaluation Neurological examination Eye examination Hearing test Pulmonary function Sleep study Electrocardiogram Echocardiogram Dental evaluation Endurance test Quality of life and pain assessment Functional/ADL assessment Hip/pelvis imaging Whole Spine imaging Spine MRI	Medical history Physical examination Growth evaluation Neurological examination Dental evaluation	Eye examination Hearing test Pulmonary function Electrocardiogram Echocardiogram Endurance test Quality of life and pain assessment Cervical spine imaging	Electrocardiogram Echocardiogram Spine MRI Sleep study

Notes: Reproduced with permission from Dove Medical Press, Regier DS, Tanpaiboon P. Role of elosulfase alfa in mucopolysaccharidosis IVA. *Appl Clin Genet*. 2016;9:67–74.³⁵

Abbreviations: mo, months; yr, years; ADL, activities of daily living.

Until recently, MPS IVA patients only had supportive care for their symptoms like adenotonsillectomy for obstructive sleep apnea, hearing aids for sensorineural hearing loss, and keratoplasty for corneal clouding. Patients also had aggressive supportive and symptomatic surgical and nonsurgical interventions to treat skeletal abnormalities. There are benefits from hematopoietic stem cell transplantation (HSCT) in MPS. According to Yabe et al in 2016,³⁶ there were four MPS IVA patients with HSCT being followed at least 10 years. All of them had normal level of GALNS activity in lymphocytes after HSCT. Only one patient had osteotomy in both legs and others had no orthopedic surgical intervention. Their activity of daily living (ADL) was better than untreated patients. However, due to few MPS IVA patients with HSCT, there are no clear mortalities about HSCT for MPS IVA having been reported.³⁷

For many forms of LSD, enzyme replacement therapy (ERT) has become the standard treatment. In 1996, the first ERT was approved for Gaucher disease. ERT has many limitations, including high cost, low penetration of the central nervous system, and infusion reactions. However, with more clinical experience, infusion reactions could be prevented. The low penetration of enzymes through the blood–brain barrier reduces central nervous system penetration and limits the efficacy of all ERTs.

For instance, type 1 Gaucher disease patients have improvements in systemic symptoms after ERT. However, peripheral administration of ERT cannot improve neurological regression in MPS type II and III patients who often die from neurological complications. Because MPS IVA patients do not have intellectual disabilities, ERT can provide beneficial outcomes, including preventing disease progression to a certain extent. Nevertheless, it cannot reverse most pathology that had developed before treatment. There are five types of MPSs for which ERT has currently been approved: MPS I (larosulfase; Aldurazyme®, Genzyme, Boston, MA, USA), II (idursulfase; Elaprase®, Shire, Dublin, Republic of Ireland), IVA (elosulfasealfa; Vimizim®, Biomarin, Novato, CA, USA), VI (galsulfase; Naglazyme®, Biomarin), and VII (vestronidase alfa; Mepsevii®, Ultragenyx, Novato, CA, USA).

Elosulfase Alfa Mechanism of Action

ERT can only be administered by intravenous infusion. The mechanism of action of elosulfase alfa is to provide enzymatic activity to reduce the enzyme deficiency caused by MPS IVA. Elosulfase alfa is a purified human enzyme; recombinant DNA technology (recombinant human enzyme or rhGALNS) in the Chinese hamster ovary cell line is used to produce it. It is identical in sequence and enzymatic

activity with the human form of GALNS. Elosulfase alfa and human GALNS have the same amino acid sequence and N-linked glycosylation.^{38,39} Because it contains bis-mannose 6 phosphate, one of the oligosaccharide chains can bind to a cell surface receptor (cation-independent mannose-6-phosphate receptor). Elosulfase alfa metabolizes GAGs, C6S, and KS.^{40,41} Elosulfase alfa is active in a low pH environment; thus, it has a low activity outside the lysosome.

Pharmacokinetics and Pharmacodynamics

Qi et al reported the pharmacokinetics of elosulfase alfa⁴² based on a Phase III trial for 24 weeks in 2014.² They analyzed the pharmacokinetics during and after the initial infusion of elosulfase alfa. Figure 2 shows the pharmacokinetics during and after week 22 of infusion.^{42,43} This figure reflects the mean plasma concentration profiles of elosulfase alfa during and after infusion. They collected blood samples before and 15, 60, and 120 min after beginning the infusion and 5, 15, 30, 60, 120, and 180 min after completing the infusion. Noncompartmental analysis of pharmacokinetic parameters was done with WinNonlin® (Certara, L.P., Princeton, NJ, USA) software. According to this figure, it shows that after 22 weeks treatment, the dose of 2 mg/kg/QW could keep higher level of elosulfase alfa than the dose of 2 mg/kg/QOW. Table 2 shows that after repeated administration over 22 weeks, the plasma half-life of elosulfase alfa increased from approximately 7 min in the first infusion to approximately 36 min.^{42,43} The intracellular half-life of elosulfase alfa is estimated to be 5–7 days. This was determined in vitro in human

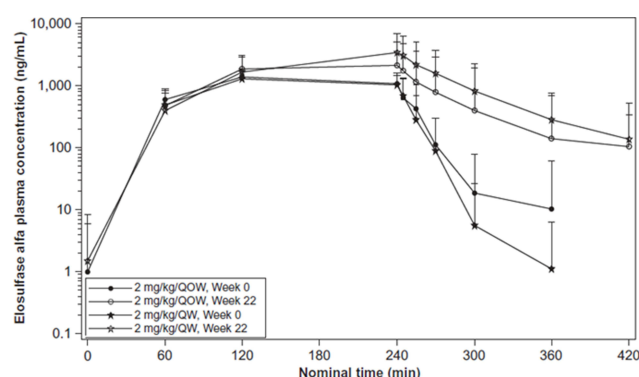


Figure 2 Mean plasma concentration profiles of elosulfase alfa during and after infusion. QOW: every other week; QW: once per week. Reprinted by permission from Springer Nature, Qi Y, Musson DG, Schweighardt B, et al. Pharmacokinetic and pharmacodynamic evaluation of elosulfase alfa, an enzyme replacement therapy in patients with Morquio A syndrome. *Clin Pharmacokinet*. 2014;53(12):1137–1147. Figure 2 is copyright protected and excluded from the open access licence.⁴²

Morquio A fibroblasts.⁴⁴ The appearance of neutralizing anti-elosulfase alfa antibodies in patients is thought to be associated with the increase in the plasma half-life.⁴² The total exposure assessed by AUC_{0-last} increased 181% or 192% after 22 weeks of administration. The pharmacodynamics, efficacy, or safety outcomes in patients are not influenced by this apparent increase in exposure.

Elosulfase alfa normalizes the levels of the deficient endogenous enzyme in lysosomes via a mannose-6-phosphate-mediated mechanism.⁴⁵ According to pre-clinical and clinical studies, the major pharmacodynamic effect of elosulfase alfa is to reduce the uKS by increasing the lysosomal degradation of this metabolite^{45–47}. Although we could use uKS to differentiate MPS IVA from other forms of MPS and to demonstrate pharmacodynamic effects of ERT treatment, we could not predict the skeletal improvement during ERT for MPS IVA by uKS.⁴⁸ Biodistribution throughout the heart layers, the entire growth plate thickness, and in macrophages and hepatocytes was observed with the intravenous administration of elosulfase alfa to wild-type mice in preclinical studies.⁴⁵

Preclinical Data

In a study to determine the safety, efficacy, and tolerability of elosulfase alfa, using MPS IVA mice, after 12 weeks of treatment, there was an apparent reduction of GAGs storage in the visceral organs, bone marrow, heart valves, connective tissues, and ligaments.⁴⁹ There was further evaluation in newborn MPS IVA mice at birth. According to the bone pathology study, although the chondrocytes were vacuolated, the columnar structure remained organized. This result showed that before the cartilage cell layer became mature bone in MPS IVA, the enzyme entered the cartilage and prevented the disorganization of the cartilage structure. It also showed that prompt treatment with ERT could prevent the bone pathology of MPS IVA.⁵⁰ According to Tomatsu et al in 2010,⁵¹ the treatment effect is better when we treat mice for a longer term or from birth. GALNS with hexaglutamate sequence (E6) tagged (E6-GALNS) could clear the storage materials in bone, bone marrow, and heart valves substantially, especially after 24 weekly infusions.

Clinical Trial Data

In MOR-002 trial (Patients with MPS IVA, who enrolled in a prior BioMarin sponsored clinical study of BMN 110; NCT00884949; Study Identification Number MOR-002;

Table 2 Pharmacokinetic Parameters for Elosulfase Alfa in Patients with Morquio a Syndrome

Parameter	Elosulfase alfa 2.0 mg/kg/QOW	Elosulfase alfa 2.0 mg/kg/QW	Ratio of Elosulfase alfa QOW/QW ^a (%)
Week 0			
n	24	22	
AUC _{0-∞} , ng min/mL	287,597 (96,432.1), 14	231,074 (103,207.4), 15	124.5
AUC _{last} , ng min/mL	248,720 (97,063.7), 24	237,884 (100,328.6), 22	104.6
C _{max} , ng/mL	1,438 (435.3), 24	1,494 (534.1), 22	96.2
CL, mL/min/kg	7.54 (2.002), 14	10.04 (3.733), 15	75.1
V _{ss} , mL/kg	219.42 (95.483), 12	395.74 (315.636), 14	55.4
V _d , mL/kg	68.79 (34.008), 14	123.66 (144.115), 15	55.6
t _{1/2} , min	6.57 (3.110), 14	7.52 (5.484), 15	87.4
t _{max} , min	150 (58.1), 24	172 (75.3), 22	87.2
Week 22			
n	23	22	
AUC _{0-∞} , ng min/mL	463,460 (491,418.9), 19	619,080 (422,048.3), 20	74.9
AUC _{last} , ng min/mL	411,687 (420,279.7), 23	577,371 (416,316.6), 22	71.3
C _{max} , ng/mL	2,616 (2,702.1), 23	4,036 (3,237.1), 22	64.8
CL, mL/min/kg	6.50 (2.942), 19	7.08 (12.997), 20	91.8
V _{ss} , mL/kg	245.19 (273.145), 17	649.67 (1,841.703), 20	37.7
V _d , mL/kg	120.11 (71.076), 19	299.52 (543.309), 20	40.1
t _{1/2} , min	19.25 (19.217), 19	35.86 (21.485), 20	53.7
t _{max} , min	159 (60.6), 23	202 (90.8), 22	78.5
Week 22/Week 0^b (%)			
n	23	21	
AUC _{0-∞} , ng min/mL	179.2	328.6	
AUC _{last} , ng min/mL	176.3	280.6	
C _{max} , ng/mL	183.6	291.6	
CL, mL/min/kg	87	46.4	
V _{ss} , mL/kg	127	188.9	
V _d , mL/kg	147	246	
t _{1/2} , min	280	696	
t _{max} , min	119.8	145.7	

Notes: Values are expressed as mean (SD), n unless otherwise indicated.

For patients who have missing values of AUC_{0-∞}, t_{1/2}, CL, V_d and V_{ss}, the parameters could not be estimated due to insufficient data in the terminal phase of the plasma profile. For patients who have missing values of V_{ss} only, their V_{ss} was not reported due to a negative value. Adjusting for infusion caused a negative MRTinf value. The V_{ss} value was also negative because of the relationship: V_{ss} = MRTinf-CL.

AUC_{0-∞} area under the plasma concentration–time curve from time zero to infinity, AUC_{last} area under the plasma concentration–time curve from time zero to the time of last measurable concentration, CL total clearance of drug after intravenous administration, C_{max} observed maximum plasma concentration, MRTinf mean residence time extrapolated to infinity, SD standard deviation, QOW every other week, QW weekly, V_{ss} apparent volume of distribution at steady-state, V_d apparent volume of distribution based upon the terminal phase, t_{1/2} elimination half-life.

^aRatio is ratio of means

^bOnly patients with pharmacokinetic data available for both visits are included. Reprinted by permission from Springer Nature, Qi Y, Musson DG, Schweighardt B, et al. Pharmacokinetic and pharmacodynamic evaluation of elosulfase alfa, an enzyme replacement therapy in patients with Morquio A syndrome. Clin Pharmacokinet. 2014;53 (12):1137–1147. Table 2 is copyright protected and excluded from the open access licence.⁴²

Abbreviations: AUC_{0-∞}, area under the plasma concentration–time curve from the start of infusion to the last measurable observation; CL, total clearance of drug after intravenous administration; C_{max}, maximum observed concentration; t_{max}, time of observed C_{max}; t_{1/2}, plasma elimination half-life; V_{ss}, apparent volume of distribution at steady state; V_d, apparent volume of distribution based upon the terminal phase t_{1/2}; QOW, every other week; QW, once per week.

Baseline and every 12 weeks for up to 72 weeks), elosulfase alfa was generally well tolerated at all doses. Over 5 years of treatment, no new safety concerns emerged during sustained treatment with elosulfase alfa 2.0 mg/kg per week. Adverse events occurred in patients in all studies; however, most were mild or moderate in severity. Hypersensitivity reactions were frequent; however, they could be managed by additional medications and altering

the infusion rate. There was one patient with a type I hypersensitivity reaction and another with recurrent infusion reactions. These patients discontinued the study. Nevertheless, in the subsequent Phase 3 trial, no hypersensitivity adverse events were seen.²

In the MOR-002 trial (72 weeks) and the MOR-100 long-term extension (192 weeks), patients could maintain their performance in the 6min walk test (6MWT) and 3min

stair climb test (3MSCT). One patient even had slightly better results in both tests. However, there was no exclusion of surgical procedures during the studies. The surgical procedures could affect the 6MWT and 3MSCT results. Due to the small sample size and absence of a control group, we could not evaluate efficacy. In the Morquio A Clinical Assessment Program (MorCAP) natural history study, which followed the progression of untreated Morquio A syndrome over 2 years, 6MWT distances in untreated patients decreased over this period.⁵² These results contrast the MOR-002 trial and the MOR-100 long-term extension. However, we should note that the mean age of patients at baseline was higher in the MorCAP observational study than in the MOR-002 study (14.4 ± 11.97 years vs 8.4 ± 2.90 years).

MPS IVA patients risk declining pulmonary function as the condition progresses; however, in the MOR-002 and MOR-100 studies, there were increases in forced vital capacity (FVC), forced expiratory volume in 1s (FEV1), and maximum voluntary ventilation (MVV). However, since there was no comparison with an age-matched group of untreated patients, these results could be partially attributed to growth.⁵³ Therefore, the true impact of treatment on pulmonary function is yet to be determined. Furthermore, there were no negative effects in the quality-of-life measures in children with Morquio A syndrome,⁵⁴ even though the quality-of-life measures would be expected to decline as the disease progresses. In the MOR-002 and MOR-100 studies, patients maintained a consistent level of function in the domains of self-care, mobility, and caregiver assistance over the 5-year study according to the MPS health assessment questionnaire (MPS-HAQ).

The 2.0 mg/kg per week elosulfase alfa dosing demonstrated the greatest reduction of uKS levels. Additionally, there was superior plasma availability with the 2.0 mg/kg per week dose. Concerning pharmacokinetic parameters, the maximum observed concentration in plasma (C_{max}) increased. The areas under the curve from time 0 to last measurable concentration (AUC_{0-}) were nonlinear and far exceeded the dose increases from 0.1 to 1.0 to 2.0 mg/kg per week. We also observed that the decreases of clearance, the volume of distribution based on the terminal rate constant (V_{dz}), and the volume of distribution at steady state (V_{dss}) were also nonlinear. This indicated the possible saturation of clearance mechanisms such as protease degradation and/or cation-independent mannose

6-phosphate receptor (CI-M6PR) receptor-mediated cellular uptake.

The impact of neutralizing antibodies on the pharmacokinetics, seen in MOR-002 and MOR-100, is consistent with the placebo-controlled MOR-004 study.^{55,56} In the MOR-004 study, we were also aware that there were no relationships between efficacy outcomes and neutralizing antibody positivity or titer according to the alternative cell-based flow cytometry neutralizing antibodies assay.⁵⁷ This result indicated that neither total antibodies nor neutralizing antibodies are related to the incidence of adverse events, loss of efficacy, or altered pharmacodynamics.

Treatment

Based on infusion volume, 2.0 mg/kg per week intravenous infusion over 3.5–4.5 h is currently the recommended dose. For patients <25 kg and >25 kg, the final volume is 100 mL and 250 mL respectively. The ERT is diluted with 0.9% sodium chloride injection. For patients <25 kg and ≥ 25 kg, the infusion rate should be 3 mL/h and 6 mL/h for the first 15 min, increasing to 6 mL/h and 12 mL/h for the next 15 min. If this rate is tolerated, the rate can be increased by 6 mL/h for patients <25 kg and 12 mL/h for patients ≥ 25 kg every 15 min. The maximum rates are 36 mL/h for patients <25 kg and 72 mL/h for patients ≥ 25 kg. An antihistamine with or without antipyretics should be administered 30–60 minutes before infusion. Due to the risk of sleep apnea and airways difficulties, a nonsedating antihistamine is preferred.

Adverse Events and Reactions

Clinical studies investigating adverse events enrolled 235 patients in all.^{39,58} Symptoms and signs of anaphylaxis (cough, erythema, throat tightness, urticaria, hypotension, dyspnea, gastrointestinal symptoms) occurred in 16 (6.8%) patients. Anaphylaxis occurred as early as 30 minutes after beginning the infusion to 3 hours after completing it. Subsequent elosulfase alfa infusion with infusion rate adjustments and/or medical intervention were completed in all patients except two. The rate of hypersensitivity reactions was 18.7%, and the reactions were observed as early as 30 minutes after starting the infusion to as late as 6 days afterward. According to the MOR-004 clinical trial (Time Frame: Baseline to Week 24), the most common adverse events were headache (20.3%), pyrexia (18.6%), vomiting (15.3%), nausea (13.6%), diarrhea (11.9%), fatigue (11.9%), upper abdominal pain (8.5%), cough (8.5%), oropharyngeal pain (5.1%), abdominal pain (3.4%), and

chills (1.7%).² Based on the study of children under 5 years of age who received 2.0 mg/kg per week for 52 weeks,⁵⁹ the rate of adverse events requiring infusion interruption and medical intervention was 0.8%. After treatment, growth velocity and uKS levels were improved.

Prevention and Management of Adverse Reactions

The adverse reaction risk in phase I/II clinical trials could be reduced by nonsedating antihistamines with or without antipyretics.² If patients have a history of adverse reactions or allergies, we should consider premedication like sedating antihistamines, antipyretics, H₂ blockers, leukotriene receptor antagonists, oral cromolyn sodium (personal communication), and/or steroids. Additionally, the infusion rate should be decreased by 25%–50%, especially during the first 1–2 hours.³⁶ Patients should have an airway patency evaluation before beginning ERT if they have airway management or positive pressure during sleep to manage sleep apnea. These patients also need oxygen during infusion therapy.

Because of the increased risk of hypersensitivity reactions, ERT should be avoided if patients have an acute infection, a fever, or respiratory illness.⁴⁰ Patients should have immediate medical care if they develop an acute hypersensitivity reaction. For mild-to-moderate reactions, the infusion could be slowed or stopped and antihistamines, antipyretics, and/or steroids administered. For

severe anaphylaxis reactions, the infusion should be stopped immediately and breathing, and circulation evaluated. One should stabilize the airway, give oxygen supplementation, and administer intramuscular epinephrine and intravenous fluids for hypotension.

Expert Commentary

Pharmacokinetic studies outside of clinical trials and clinical settings are needed. Immunogenicity is not correlated with the efficacy of elosulfase alfa. Although only a small proportion of patients are hypersensitive,⁶⁰ some clinicians remain concerned about the drug's immunological risk. They believe the risk of infusion-associated reactions and infusion-site reactions may increase due to the presence of anti-elosulfase alfa antibodies. However, clinical trials have shown no correlation between the immune response and infusion-associated or infusion-site reactions.³⁹ Only a small minority of patients have an elevated immune response.

We cannot currently predict which Morquio A patients will respond to ERT.⁶¹ About 10% of patients have an excellent response that exceeds expectations. There is also a small group of patients who do not respond to ERT at all. Therefore, Morquio A patients receiving elosulfase alfa should have at least 12 months of treatment before assessing their response. The current quality-of-life assessment tools do not adequately capture the treatment benefits for patients.⁶² We must develop improved and more innovative daily lifestyle measures to adequately assess clinical response. Lifestyle tracker apps could provide an all-day record of activity and sleep habits in

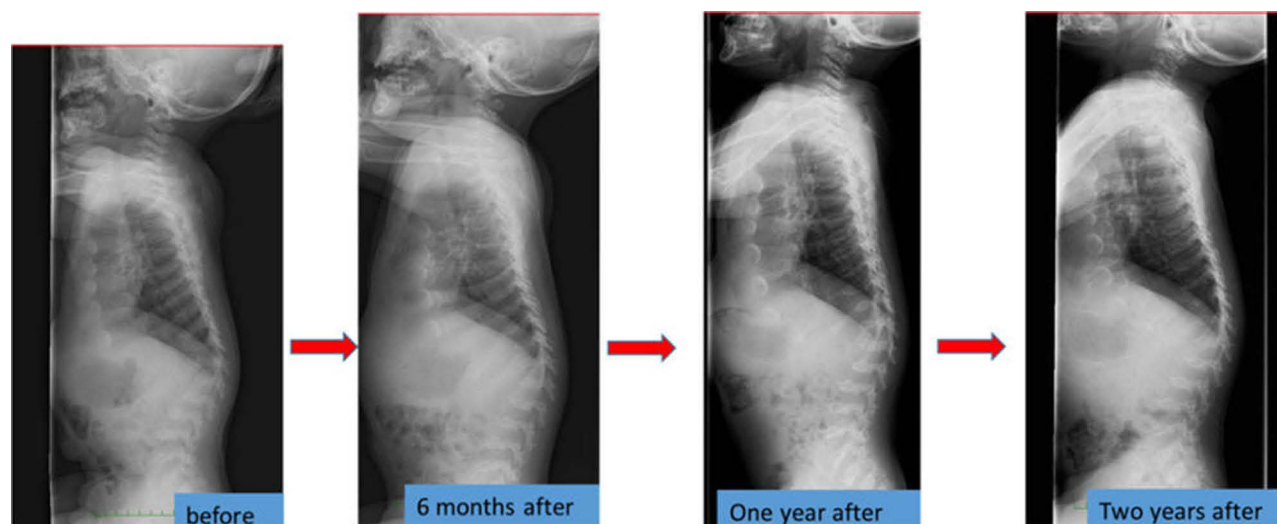


Figure 3 The spine X-ray findings before and after the enzyme replacement therapy in a Japanese boy with MPS IVA. Reproduced from Nakamura-Utsunomiya A, Nakamae T, Kagawa R, et al. A case report of a Japanese boy with MorquioA syndrome: effects of enzyme replacement therapy initiated at the age of 24 months. *Int J Mol Sci.* 2020;21(3):989.⁷²

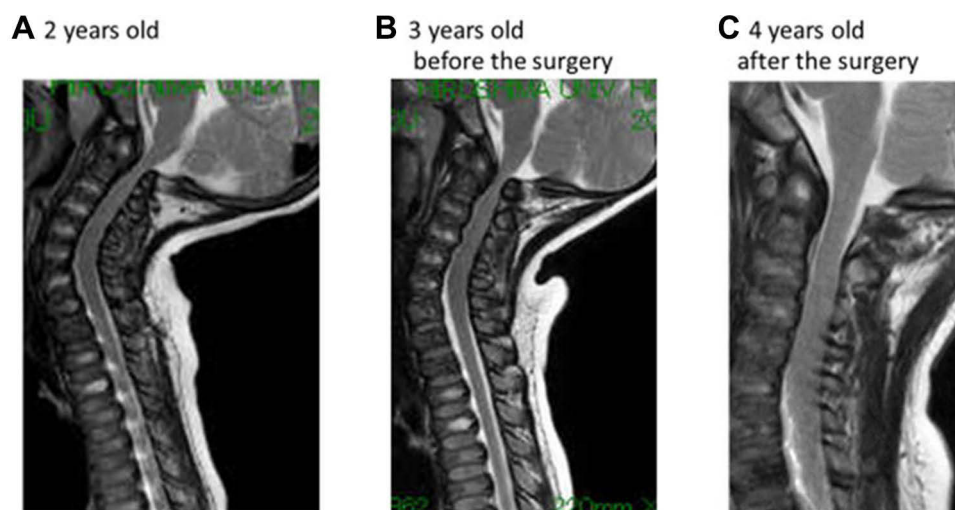


Figure 4 The changes in MRI findings. **(A)** The spinal cord compression at C1 level was already present at the first examination. **(B)** The lesion was slowly progressive after the ERT started, **(C)** which was relieved by resecting the posterior arch of the atlas vertebra. Reproduced from Nakamura-Utsunomiya A, Nakamae T, Kagawa R, et al. A case report of a Japanese boy with Morquio A syndrome: effects of enzyme replacement therapy initiated at the age of 24 months. *Int J Mol Sci.* 2020;21(3):989.⁷²

the future^{63–69}. Blood KS directly derived from the bone is an important biomarker of improvement in true clinical endpoints including bone pathology or any other skeletal signs and symptoms.⁷⁰

Orthopedic surgeries are necessary because ERT have provided only modest benefit for skeletal deformities.⁷¹ X-ray and MRI of spine showed that ERT had limited effects on bones⁷² (Figures 3 and 4). Patients with MPS IVA usually have odontoid hypoplasia with subluxation, gibbus deformity, hip dysplasia and osteonecrosis, and lower extremity valgus. Due to these symptoms, patients should necessitate monitoring and often surgical intervention like realignment osteotomy, corrective knee surgery and hip replacement.

Most patients in the UK and Canada are infused at home. Conversely, patients in the USA are infused in infusion centers which increases treatment costs. Nevertheless, infusion centers offer additional benefits like emotional support from other patients, families, and healthcare workers. Experienced clinicians and healthcare providers, including infusion nurses, are important. They can provide the proper and effective rate of ERT infusion and drug dose and act to prevent or manage adverse reactions.

Elosulfase alfa has improved the walking distance of MPS IVA patients and their respiratory function. However, compared with the growth charts for untreated patients, patients treated with ERT did not show any significant increase in growth in any age group, even if they had early ERT intervention before 5 years of age.⁷³ We still

have much to learn to improve our management of Morquio A syndrome. There was no effective drug for Morquio A before elosulfase alfa; however, now, there is hope for the future in this field.

Five-Year View

ERT could only slow skeletal disease progression even when started in infancy before 2 months of age.⁷⁴ There are studies aimed at increasing the effectiveness of ERT with targeted therapies.^{69,75} In preclinical research, gene-targeted therapy has shown some efficacy in treating Morquio A^{75–77}. It may be more effective to use small, specific, targeted molecules in addition to ERT to treat Morquio A, which will solve the problems of immunogenicity and bone uptake in the future.^{75,76} In the next 5 years, we should be able to predict which mutations of the *GALNS* gene are responsive to ERT.

According to a study by Donida et al in a cohort of MPS IVA patients receiving ERT therapy,⁷⁸ interleukin-6 and decreased glutathione levels were elevated. This suggests a proinflammatory and prooxidant state. However, there were no MPS IVA patients who had not received ERT for comparison. Further investigation of combined antioxidant and anti-inflammatory agent therapy with ERT is needed to optimize patient outcomes.

Conclusions

Morquio A is not just a skeletal disorder; there is a multiorgan involvement. Elosulfase alfa could be effective in improving

patient endurance, pulmonary function, and quality of life. It is important to have long-term outcome studies to determine the effect of elosulfase alfa on the bone outcomes of patients. Moreover, we must evaluate the efficacy of ERT treatment in moderating the long-term progression of MPS IVA.

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Disclosure

The authors declare that they have no conflicts of interest.

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