

## Research Article

# Intestinal Dialysis Research Progress and the Early Treatment of a Non-Diabetic Patient with Symptomatic Uremia and Fatty Liver with Intestinal Dialysis: The Practice of Evidence-Based Medicine

Aamir Jalal Al-Mosawi\*

Advisor in Pediatrics and Pediatric Psychiatry the National Training and Development Center and Baghdad Medical City Dubai UAE.

**Corresponding Author:** Aamir Jalal Al-Mosawi. Advisor in Pediatrics and Pediatric Psychiatry the National Training and Development Center and Baghdad Medical City Dubai UAE.

Received: 📅 2024 Feb 09

Accepted: 📅 2024 Feb 29

Published: 📅 2024 Mar 07

## Abstract

**Background:** Chronic renal failure results from a variety of pathophysiological mechanisms and etiologies and is associated with progressive and irreversible damage and loss of the kidneys' tissue leading to failure of the kidneys to excrete waste products, and also failure to perform some other functions. Many patients with chronic renal failure in a country like Iraq has been reported to be reluctant to accept dialysis therapies because of the wide spread notion of its association with high mortality. The lack of effective, convenient, and affordable therapy for chronic renal failure in many regions of the world should not mean should that the patients with advanced chronic renal failure are left without other suitable, convenient and acceptable care. The aim of this paper is to describe the early treatment of a patient with symptomatic uremia with intestinal dialysis.

**Patients and Methods:** A 60-year non-diabetic male patient was experiencing progressive symptomatic uremia. On the 18th of November 2023, blood urea was elevated at 217 mg/dL, serum creatine was 5.2 mg/dL, and he had symptomatic uremia with nausea, vomiting, fatigue, pruritus and anemia. The patient was treated with intestinal dialysis (Acacia gum supplementation plus conservative dietary and pharmacological management of chronic renal failure) which was prescribed according to the latest published guidelines. It was necessary during the first week of treatment to eliminate almost all dietary protein, and his diet was consisting mainly of high calorie juices, grapes, and water melon.

**Results:** When the patient was seen on the 2nd of December 2023, treatment was associated with marked symptomatic and laboratory improvements. Blood urea was 115 mg /dL, serum creatine was 3.6 mg /dL However, and serum calcium was 5.3 mg/dL (Normal ranges: 8-10.5 mg/dL). Therefore, oral alphacalcidol was added in a dose of 1 microgram daily.

**Conclusion:** Intestinal dialysis will continue to be used to improve the management of chronic renal failure and symptomatic uremia as long as there is no convenient and affordable therapy for chronic renal failure in many regions of the world.

**Keywords:** Symptomatic Uremia, Intestinal Dialysis, Fatty Liver, Educational Article and Expert Opinion.

## 1. Introduction

Chronic renal failure results from a variety of pathophysiological mechanisms and etiologies and is associated with progressive and irreversible damage and loss of the kidneys' tissue leading to failure of the kidneys to excrete waste products, and also failure to perform some other functions. The incidence of chronic renal failure has been disturbingly increasing during the previous decades, and it has been increasingly considered as a global health problem. The availability of renal replacement therapy and its quality are much

less in developing countries than in the advanced countries. That was generally attributed to the associated high cost and the complexity of its technology. Economically disadvantaged courtiers have priorities to provide the basic health services and to improve them rather than to offer expensive therapeutic technologies that are considered by a significant number of their populations as inconvenient.

Many patients with chronic renal failure in a country like Iraq has been reported to be reluctant to accept dialysis therapies

because of the wide spread notion of its association with high mortality. The lack of convenient and affordable therapy for chronic renal failure in many regions of the world should not mean should that the patients with advanced chronic renal failure are left without other suitable, convenient and acceptable care. Therefore, the need for a more robust and more convenient therapy for chronic renal failure has been increasingly emphasized during the previous two decades. A novel urea lowering dietary therapy that could provide a novel paradigm for the management of chronic renal failure has been described. This new dietary therapy was used in combination with the traditional known therapies of chronic renal failure and has been increasingly known as intestinal dialysis [1-3].

The aim of this paper is to describe the initial treatment of a patient with symptomatic uremia with intestinal dialysis.

## 2. Patients and Methods

A 60-year non-diabetic male patient was experiencing progressive symptomatic uremia with anorexia, fatigue, pruritus and anemia over the previous weeks. On the 12th of November 2023, blood urea was elevated at 220 mg/dL, serum creatine was 5.2 mg/dL, and he was anemic with hemoglobin at 9.7 g/dL (Normal ranges: 11.5-16.5 g/dL). Urinalysis didn't show important findings. He was seen by more than two doctors, and all referred him for dialysis treatment which he and his family consistently rejected.

On the 18th of November 2023, blood urea was elevated at 217 mg/dL, serum creatine was 5.2 mg/dL, and he had symptomatic uremia with nausea, vomiting, fatigue, pruritus and anemia. He was also experiencing reduction of the urine output and anemic with hemoglobin at 9.7 g/dL (Normal ranges: 11.5-16.5 g/dL). The patient was treated with intestinal dialysis (Acacia gum supplementation plus conservative dietary and pharmacological management of chronic renal failure) which was prescribed according to the latest published guidelines [4-13].

### He received

- Bumetanide 1 mg twice daily to improve urine output.
- Oral ferrous sulfate 200 mg three times daily, to be continued and adjusted according to hemoglobin level.
- Oral calcium carbonate 1000 mg daily, to be continued indefinitely.
- Intramuscular pyridoxine 100 mg daily for 7 days.
- Intramuscular Vitamine B complex daily for 7 days.
- Oral alphacalcidol 1 mcg daily, to be continued with dose adjustment as necessary according to the serum calcium level.
- Oral cetirizine hydrochloride 10 mg once at night and topical crotamiton were used to control pruritus as long as they were needed.

He also received acacia gum powder 25g dissolved in 250 ml Diet 7Up and given three times daily before meals. It was necessary to give him oral domperidone (Molium) 10 mg three times daily to prevent and reduce the occurrence of abdominal distention and discomfort of the intake. It was necessary during the first week of treatment to eliminate al-

most all dietary protein, and his diet was consisting mainly of high calorie juices, grapes, and water melon. Therefore, nutritional support with oral royal jelly capsules was added to prevent any unexpected nutritional deficiency.

## 3. Results

When the patient was seen on the 2nd of December 2023, treatment was associated with marked symptomatic and laboratory improvements. Blood urea was 115 mg /dL, serum creatine was 3.6 mg /dL, but he was still anemic with hemoglobin at 9.4 g/dL (Normal ranges: 11.5-16.5 g/dL). Therefore, a one-week course of daily intramuscular iron dextran (100 mg daily) was prescribed. Serum calcium was 5.3 mg/dL (Normal ranges: 8-10.5 mg/dL). Therefore, oral alphacalcidol was added in a dose of 1 microgram daily (The dose to be adjusted as necessary according to serum calcium level).

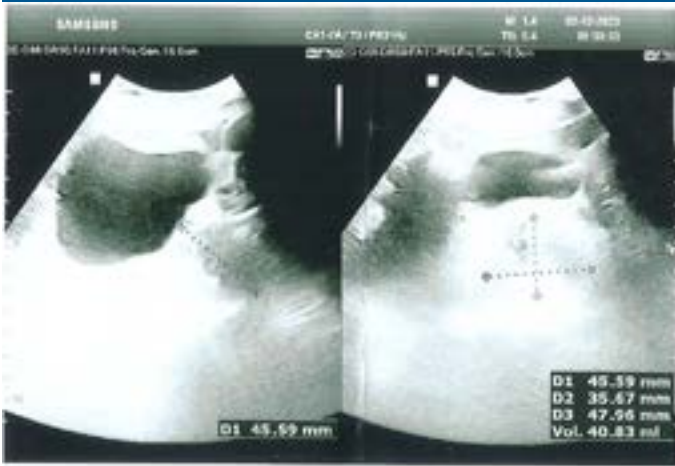
Ultrasound imaging (Figure-1) showed reduction in renal size (RK: 9.6 x 4.5 cm, LK: 9.5 x 4.5 cm) [Adults male normal renal size is 10 x 9-14x13 cm] with increased parenchymal echotexture suggestive of bilateral parenchymal renal disease). Medullary pyramid thickness of both kidneys was 13 mm. There was no pelvi-calyceal dilatation, but there were coarse crystals with the collecting system of both kidneys. Therefore, the patient received oral essential oil terpenes (Urinex) based on the evidence presented by Al-Mosawi AJ [14-17].

Ultrasound imaging also showed gaseous bowel loop, the spleen, pancreas, and gall bladder were normal in appearance, the common bile duct was not dilated (4 mm in diameter), and the main portal vein was not dilated. The liver was of normal size but was showing mild fatty changes. Therefore, oral silymarin (Legalon) 75 mg once daily was prescribed based on the evidence provided by Buturova and colleagues (2010), Luis (2015) [18-19].

The urinary bladder was normal and had normal bladder thickness (Pre-voiding volume was 150 ml, and the re-voiding volume was 4 ml, but the prostate was enlarged with smooth surface and a volume of 40.8 ml (Normal: 25 ml). Therefore, Oral finasteride was added.



Figure-1A: Renal ultrasound



**Figure-1B:** Abdominal ultrasound

#### 4. Discussion

Obesity and diabetes are generally considered the most important risk factors for developing fatty changes in the liver [2]. Although, the patient in this report was not diabetic or obese, he had an ultrasound evidence of fatty changes in the liver. However, in 2015, Ludovico Abenavoli from Italy and his research team considered fatty liver that is not related to alcohol to be the most common liver disorder throughout the world [20, 21].

Zobair Younossi from the United States and his international research group emphasized that although obesity is the major risk factor for the development of non-alcoholic liver disease, a large number of patients with the condition are not obese [22].

The lack of convenient and affordable therapy for chronic renal failure in many regions of the world should not mean should that the patients with advanced chronic renal failure are left without other suitable, convenient and acceptable care.

Therefore, the need for a more robust and more convenient therapy for chronic renal failure has been increasingly emphasized during the previous two decades. A novel urea lowering dietary therapy that could provide a novel paradigm for the management of chronic renal failure has been described. This new dietary therapy was used in combination with the traditional known therapies of chronic renal failure and has been increasingly known as intestinal dialysis [1-3].

The use of a dietary material to increase extra-renal excretion and shift the urinary excretion of urea to the intestinal excretion has been increasingly called "Intestinal dialysis technology [23, 24].

The clinical use of intestinal dialysis has been increasing reported as early as the 2000s and was first endorsed the Ira Greifer (Figure-2), a pioneer of pediatric nephrology [1-3, 23-32].



**Figure-2:** Ira Greifer, a Pioneer of Pediatric Nephrology

Introducing acacia gum which is a safe dietary fiber as a medicine for its urea lowering effect was first suggested in 2006, and its role in the management of chronic renal failure was emphasized [33-36].

#### 5. Conclusion

Intestinal dialysis will continue to be used to improve the management of chronic renal failure and symptomatic uremia as long as there is no convenient and affordable therapy for chronic renal failure in many regions of the world.

#### Acknowledgment

The author has the copyright of the sketch in this paper.

#### References

1. Al Mosawi, A. J. (2019). Pervasive developmental disorders in Iraqi children. *Journal of Psychiatry Research Reviews & Reports. SRC/JPSRR-102*. DOI: doi.org/10.47363/JPSRR/2019 (1), 102, 2-8.
2. Al Mosawi, A. J. (2019). Pervasive developmental disorders in Iraqi children. *Journal of Psychiatry Research Reviews & Reports. SRC/JPSRR-102*. DOI: doi.org/10.47363/JPSRR/2019 (1), 102, 2-8.
3. Al Mosawi, A. (2018). *Pediatric psychiatry: An accredited training course*. LAP LAMBERT Academic Publishing.
4. Al-Mosawi, A. J. (2020). *Case studies in pediatric psychiatry: An approach to deep learning*.
5. Al-Mosawi, A. J. (2020). Clinical uses of cerebrolysin in pediatric neuropsychiatry. *Science World Journal of Pharmaceutical Sciences*, 1(1), 1-4.
6. Al Mosawi, A. (2018). *Asperger syndrome and regressive autism*. LAP LAMBERT Academic Publishing.
7. Al-Mosawi, A. J. (2019). New therapies for Rett syndrome. *J Bio Innov*, 8(3), 301-307.
8. Al-Mosawi, A. J. (2019). *Childhood dementia: Heller syndrome*. (2019) Baghdad. Iraq Headquarter of Copernicus Scientists International Panel Publishing.
9. Al Mosawi, A. J. (2019). Heller syndrome in two Iraqi children. *Clinical Research and Trials*, 5, 1-3.
10. Al-Mosawi, A. J. (2019). The use of cerebrolysin and citicoline in autism and Asperger syndrome. *J Bio Inn*

- ov, 8(1), 99-108.
11. Al-Mosawi, A. J. (2020). A Unique experience with mental and developmental retardation: Innovative Medical therapies for idiopathic mental retardation. *EC Clinical and Medical Case Reports*, 3(5), 42-54.
  12. Al-Mosawi, A. J. (2020). Clinical uses of cerebrolysin in pediatric neuropsychiatry. *Science World Journal of Pharmaceutical Sciences*, 1(1), 1-4.
  13. Al-Mosawi, A. J. (2019). The etiology of mental retardation in Iraqi children. *autism*, 1, 4-7.
  14. Al Mosawi, A. (2018). A novel therapeutic approach for idiopathic mental retardation. LAP LAMBERT Academic Publishing.
  15. Al-Mosawi, A. J. (2019). The pattern of mental retardation in Iraqi children. Saarbrücken LAP Lambert Academic Publishing.
  16. Al-Mosawi, A. J. (2019). The pattern of cerebral palsy in Iraqi children. *Med Life Clin*, 1(1), 1001.
  17. Al-Mosawi, A. J. (2019). The pattern of cerebral palsy in Iraqi children. *Med Life Clin*, 1(1), 1001.
  18. Al-Mosawi, A. J. (2020). Cure of autistic disorders: Mission impossible is possible in an illustrated pioneering experience. *Archives of Health Science*, 4(1), 1-26.
  19. Al Mosawi, A. J. (2021). A Typical Autism Associated with Elevated Gonadotrophin and Precocious Puberty: A Very Rare Association or a New Clinical Syndrome? *development*, 33(2).
  20. Al-Mosawi, A. J. (2022). Catatonia: A Rare Manifestation of Autism. *Medp Psychiatry Behav Sci*, 1(1).
  21. Al-Mosawi, A. J. (2023). The association of autism with self-injurious behaviors: An educational article. *Journal of Innovations in Medical Research*, 2(1), 5-10.
  22. Al-Mosawi, A. J. Autism with Severe Mental Retardation: a Therapeutic Challenge and Expert Opinion.
  23. Al-Mosawi, A. J. (2021). Autosomal Recessive Autism: Cure of the Major Autistic Features. *Scholars International Journal of Anatomy and Physiology*, 4(8), 120-126.
  24. Al-Mosawi, A. J. (2022). Atypical Genetic Autism: Cure of the Major Autistic Features and the Need for Cognitive Improvement and Rehabilitation. *Medp Psychiatry Behav Sci*, 1(1).
  25. Al-Mosawi AJ. Books of Aamir Jalal Al-Mosawi included in Book authority's list of Best Books of All Time on December 15, 2021.
  26. Al-Mosawi, A. J. (2022). Cerebral Palsy and Autism Associated with Periventricular White Matter Hyperintensity on Brain Magnetic Resonance Imaging: A New Disorder and Its Treatment. *Medp Psychiatry Behav Sci*, 1(1).
  27. Al-Mosawi, A. J. (2022). Cerebral Palsy and Autism Associated with Periventricular White Matter Hyperintensity on Brain Magnetic Resonance Imaging: A New Disorder and Its Treatment. *Medp Psychiatry Behav Sci*, 1(1).
  28. Al-Mosawi, A. J. (2022). Treatment of a girl from Tunisia with typical autism: Evidence-based medicine and expert opinion. *Biomedical and Biotechnological Sciences*, 1(2), 1-5.
  29. Al-Mosawi, A. J. (2022). A girl from Pakistan with atypical autism: Expert opinion and a therapeutic recommendation. *World Journal of Radiology and Imaging*, 1(1), 38-41.
  30. Al-Mosawi, A. (2023). A Case of Atypical Autism with Mental Retardation in an Adult from Canada: An Educational Article and Expert Opinion. *J Brain Neurol Dis*, 6(4), 1-5.
  31. Al-Mosawi, A. J. (2020). Clinical uses of cerebrolysin in pediatric neuropsychiatry. *Science World Journal of Pharmaceutical Sciences*, 1(1), 1-4.
  32. Al-Mosawi, A. J. (2019). New medical therapies for the treatment of myelomeningocele. *Surgical Medicine Open Access Journal*, 2(4), 1-4.
  33. Al-Mosawi, A. J. (2018). A novel therapy for pediatric juvenile spinal muscular atrophy. Saarbrücken.
  34. Al-Mosawi, A. J. (2020). The use of cerebrolysin in pediatric Wohlfart Kugelberg Welander syndrome. *MOJ Clinical & Medical Case Reports (e-ISSN: 2381-179X)*, 10(1), 20-23.
  35. Al Mosawi, A. (2018). A novel therapy for pediatric Charcot Marie Tooth disease. LAP LAMBERT Academic Publishing.
  36. Al Mosawi, A. J. *Journal of Neurological Research and Therapy*