Clinical Study

Effect of Hydrocortisone on Intradialytic Hypotension: A Preliminary Investigational Study

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Introduction. Approximately 15 to 33% of all dialysis treatments are complicated by intradialytic hypotension (IDH). In this study, we tested the hypothesis that the intravenous administration of hydrocortisone prior to HD treatment could prevent IDH or at least decrease the drop in the blood pressure resulting from IDH. Methods. This study was approved by our local ethics committee/IRB (2017/87) and by the Jordan Food and Drug Administration (7/clinical/18). Additionally, it is registered on ClinicalTrials.gov (NCT03465007). In this preliminary investigational study, we screened all chronic hemodialysis patients at our clinic who were 18 years of age or older (n = 82) for IDH. There were 14 patients included in the interventional part of this study; patients were given IV hydrocortisone for 3 consecutive HD sessions, followed or preceded by 3 intervention-free sessions where they were given 5 ml of saline as a placebo. Results. The initial total sample size was 82 patients. The frequency of IDH at our clinic was 24.4%. Fourteen out of the 20 patients who were diagnosed with IDH agreed to enroll in the interventional part of our study. The mean age of the patients in the interventional part of our study was 53.5 years (±10.3). These patients included 5 (35.7%) men and 9 (64.3%) women. Upon comparing the number of hypotensive attacks with and without the hydrocortisone administration, we found a significant difference (p = 0.003) between the hydrocortisone and placebo treatments in which 12 (85.7%) patients had fewer IDH episodes with the hydrocortisone treatment than with placebo. Conclusion. This preliminary investigational study found that the administration of a stress dose of hydrocortisone prior to hemodialysis could be an effective measure for preventing or minimizing the risk of IDH episodes. Additional prospective studies on this subject are needed. Ruling out adrenal insufficiency in patients diagnosed with IDH is also crucial.

1. Introduction

Approximately 15 to 33% of all dialysis treatments are complicated by intradialytic hypotension (IDH), which is associated with possible serious consequences [1–4]. The Kidney Disease Outcomes Quality Initiative (KDOQI) and European Best Practice Guidelines (EBPG) define intradialytic hypotension as a decrease in systolic blood pressure by ≥20 mmHg or a decrease in mean arterial pressure by 10 mmHg, and this is associated with clinical events and the need for nursing interventions [5].

IDH is associated with increased all-cause mortality, cardiovascular morbidity and mortality, and other diseases including myocardial infarction, fluid overload/heart failure, and stroke [6]. Hemodialysis is considered a stressful event that requires a significant increase in ACTH and cortisol levels [7]. Serum cortisol starts to rise 3-4 hours after starting hemodialysis, it
peaks at 6 hours, and it returns to the predialysis level 24 hours after dialysis treatment [8]. Hemodialysis could remove some of the already present serum cortisol and along with the supposed delay in adrenocortical response; this will put our HD patients at risk for adrenal crisis [8].

Based on above, we believe that hemodialysis patients will not have enough time to physiologically raise their cortisol level to cover the stress imposed by the HD treatment, and this could be one of the main contributing factors to the development of IDH. Therefore, in this study, we tested the hypothesis that the intravenous administration of 100 mg of hydrocortisone given 30 minutes prior to HD treatment could prevent IDH or at least decrease the drop in the blood pressure during HD treatment.

2. Methods

This study was approved by our local ethics committee/IRB (2017/87) and by the Jordan Food and Drug Administration (7/clinical/18). Additionally, it is registered on ClinicalTrials.gov (NCT03465007). All participants signed an informed consent form, first to participate in the screening for intradialytic hypotension, then to participate in the interventional part of the study. This study adheres to CONSORT 2010 guidelines [9].

2.1. Design. This preliminary investigational study was designed as a randomized, placebo-controlled, double-blind, crossover trial. All patients included in the interventional part of this study were given an IV of hydrocortisone for 3 consecutive sessions, followed by or preceded by 3 intervention-free sessions where we used saline via an IV as a placebo.

The patients were allocated to hydrocortisone or placebo as the first intervention randomly using Randomizer.org. Patients and the providing nurses were blinded to the intervention, and the doses were given by the nurses (either hydrocortisone or placebo) to the patient.

2.2. Assessments. This comparative study was conducted from March 2018 through September 2018 at Jordan University Hospital (JUH), a tertiary medical center in Amman, Jordan.

We screened all chronic hemodialysis patients who are 18 years of age or older (n = 82) at the HD clinic at our university hospital for intradialytic hypotension. We defined intradialytic hypotension based on the KDOQI and European Best Practice Guidelines definitions as a decrease in systolic blood pressure of ≥20 mmHg or a decrease in mean arterial pressure by 10 mmHg, providing that this is associated with clinical events and the need for nursing interventions [5].

The BP was measured 30 minutes before HD initiation, at the beginning of HD, and every thirty minutes thereafter. The occurrence of IDH was defined as any drop in BP during HD with respect to the lower BP reading between the BP thirty minutes prior to HD initiation and the BP at the beginning of HD. Two of the authors (A.M and K.O) supervised the whole process and were present during the whole time of hemodialysis sessions, and they were the ones who documented the clinical events associated with the drop in blood pressure and also checked the blood pressure with any associated symptoms in addition to the every thirty-minute blood pressure measurement. Symptoms included mainly dizziness and fatigue, and the clinical interventions included stopping the ultrafiltration, Trendelenburg positioning, and intravenous fluid boluses. The annotation of these episodes was blinded with respect to the intervention.

Twenty of the 82 patients were diagnosed with intradialytic hypotension based on the three HD sessions. Fourteen of the 20 patients agreed to enroll in our study, and 6 declined. We screened all 14 of the patients for adrenal insufficiency by first taking random early morning serum cortisol level measurements. Serum cortisol was determined by the ADVIA Centaur cortisol assay, a competitive immunoassay using direct chemiluminescent technology (Bayer Diagnostics, UCSF Clinical Labs-Chemistry, San Francisco, CA 94143, USA). The normal reference range for morning cortisol was determined to be 4.3–22.4 mcg/dl. Seven of the 14 patients had random morning cortisol level > 10 mcg/dl (>276 nmol/l) and were without clinical symptoms or signs of adrenal insufficiency other than intradialytic hypotension, so we did not further investigate these patients for adrenal insufficiency [10]. The other 7 patients had a random early morning cortisol level < 10 mcg/dl (<276 nmol/l), so we proceeded with performing the adrenocorticotropic hormone (ACTH) stimulation test to rule out adrenal insufficiency. Four of the 7 patients had a normal ACTH stimulation test result with the cortisol level rising to >18 mcg/dl (>497 nmol/l). The other 3 were diagnosed with adrenal insufficiency, given that no rise in cortisol > 18 mcg/dl (>497 nmol/l) was shown. One of these three patients was already taking chronic 5 mg/day oral prednisone, which explains his findings. All three were referred to the endocrinology clinic. ACTH level measurements and pituitary MRI were done, and the results were consistent with those of idiopathic central adrenal insufficiency. ACTH was determined by the Elecsys ACTH test system using a quantitative electrochemiluminescence immunoassay (ECLIA) (Roche Diagnostics, Indianapolis, IN, USA, 2010). The normal reference range for ACTH was determined to be 7.2–63.3 pg/ml. Two patients declined receiving oral steroid replacement therapy but agreed to proceed with our clinical trial, and the third patient was kept on his same dose of prednisone and agreed to proceed with our clinical trial. Figure 1 details the flowchart of the patient inclusion protocol used in this study.

2.3. Intervention and Measurements. Each eligible participant was instructed to not change his medication schedule or his diet. We did not implement any other new intervention to help with the intradialytic hypotension during the study period.

After obtaining proper consents from patients, we proceeded with administering 100 mg of intravenous hydrocortisone or saline, which was given 30 minutes prior to the initiation of HD [11]. We measured the BP thirty minutes prior to starting HD, at the beginning of HD, and every thirty minutes thereafter. We used our own upper arm automated
blood pressure device (Fresenius 4008S) for the blood pressure measurement, which is the normal routine at our HD clinic. Blood pressure measurements were performed by our regular HD nurses (blinded to the intervention). We obtained the age, weight, height, TSH level, and early morning serum cortisol level from each patient. We also performed an echocardiogram for patients who complained of any cardiac symptoms during their daily activities such as exertional shortness of breath.

2.4. Statistical Analysis. We used SPSS version 24.0 (Chicago, USA) to perform the statistical analysis in our study. We used the mean (±standard deviation) to describe continuous variables (e.g., age). We used the count (frequency) to describe

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**Figure 1**: The flowchart of the patient inclusion protocol used in this study.
other nominal variables (e.g., sex). $p < 0.05$ was assigned as the $\alpha$. Data were assessed for normality using the Shapiro-Wilk test, histograms, and Q-Q plots.

We used the Wilcoxon signed-rank test to analyze the difference between the frequency of hypotensive attacks with and without the hydrocortisone administration, and we reported the results as the number of negative ranks (cases in which the frequency of hypotensive attacks with the intervention was less than the frequency of hypotensive attacks without the intervention), positive ranks (cases in which the frequency of hypotensive attacks with the intervention was more than the frequency of hypotensive attacks without the intervention), and ties (the frequency of hypotensive attacks with the intervention equals the frequency of hypotensive attacks without the intervention).

We used a paired sample $t$-test to analyze the mean blood pressure between each day with the intervention and the days without the intervention.

We used the Pearson test to analyze the correlation between the mean systolic blood pressure measurements and age, weight, and TSH and morning cortisol levels.

### 3. Results

The total sample size was 82 HD patients. The frequency of IDH in our clinic was 20/82 (24.4%). The frequency of adrenal insufficiency was 3/14 (21.4%) in the patients who were diagnosed with IDH and agreed to enroll in the interventional part of our study.

We included 14 patients with IDH in the second part (interventional) of our study. The statistical analysis of our study showed that the patients had a mean age of 53.5 years ($\pm$10.3 years). There were 5 (35.7%) men and 9 (64.3%) women included in this study. Eight (57.1%) of the included patients were known to have hypertension. Details of the patient sample included in the interventional part of the study are shown in Table 1.

Upon comparing the numbers of hypotensive attacks with and without the hydrocortisone administration, we found a significant difference ($p = 0.003$) as follows:

(i) Negative ranks (i.e., cases in which the frequency of hypotensive attacks with the intervention was less than the frequency of hypotensive attacks without the intervention): 12

(ii) Positive ranks (i.e., cases in which the frequency of hypotensive attacks with the intervention was more than the frequency of hypotensive attacks without the intervention): 1

(iii) Ties: 1

The frequencies of IDH episodes with and without the intervention for each patient and the characteristics of each patient are shown in Tables 2(a) and 2(b).

A comparison between mean SBP for the included sample at days 1, 2, and 3 with and without hydrocortisone showed no significant differences, as shown in Table 3.

We did not find any significant correlation between the mean blood pressure measurements and the age, weight, and TSH and morning cortisol levels of the patients.

Echocardiogram was done for nine out of the 14 patients who complained of any cardiac symptoms during their daily activities such as exertional shortness of breath. It showed ejection fraction ranging around 50-60% and normal to grade 1 diastolic dysfunction.

No side effects were reported during or after the administration of intravenous hydrocortisone.

### 4. Discussion

This randomized controlled, double-blind, crossover study showed that the intravenous administration of 100 mg of hydrocortisone given thirty minutes prior to starting...
treatment for HD could significantly decrease the number of IDH episodes in HD patients \((p = 0.003)\).

The prevalence of IDH at our HD clinic was 24.3\%, which is consistent with the prevalence of IDH in other studies \([1–4]\).

IDH is associated with increased all-cause mortality, cardiovascular morbidity and mortality, and other diseases, including myocardial infarction, fluid overload and heart failure, and stroke \([6]\).

Adrenal insufficiency is not uncommon in chronic HD patients, and IDH can be one of the nonspecific signs of adrenal insufficiency in chronic HD patients \([12]\), so it is crucial to rule out adrenal insufficiency in these patients, in addition to the fact that these patients will also need a daily maintenance dose of glucocorticoids beside the glucocorticoid stress dosing prior to stressful events like hemodialysis \([7, 13]\).

This IDH etiology could be attributed to different possible causes, such as an imbalance between ultrafiltration and intravascular volume refilling rate, abnormal adaptive responses to ultrafiltration, cardiovascular diseases, age, autonomic dysfunction, and diabetes \([1, 14]\).

Common interventions to treat IDH could include reducing the ultrafiltration rate, UF modeling, cooling the dialysate, midodrine, intravenous normal saline, adjusting blood pressure medications, adjusting the dry weight, sodium modeling in the dialysate, and others \([1, 15]\).

|   | Day 1 | Day 2 | Day 3 | Day 1+HC | Day 2+HC | Day 3+HC |
|---|-------|-------|-------|----------|----------|----------|
| Patient 1 | Y     | Y     | Y     | N        | N        | N        |
| Patient 2 | Y     | N     | N     | Y        | Y        | N        |
| Patient 3 | Y     | Y     | Y     | N        | N        | N        |
| Patient 4 | Y     | Y     | N     | N        | N        | N        |
| Patient 5 | Y     | N     | Y     | N        | Y        | Y        |
| Patient 6 | Y     | N     | Y     | N        | N        | N        |
| Patient 7 | Y     | Y     | Y     | Y        | N        | Y        |
| Patient 8 | Y     | Y     | Y     | N        | Y        | Y        |
| Patient 9 | Y     | Y     | Y     | Y        | N        | Y        |
| Patient 10 | N    | Y     | N     | N        | N        | N        |
| Patient 11 | Y    | Y     | N     | N        | N        | N        |
| Patient 12 | N    | Y     | Y     | N        | N        | N        |
| Patient 13 | Y    | N     | Y     | N        | N        | N        |
| Patient 14 | Y    | Y     | N     | N        | N        | N        |
| Total     | 12   | 10   | 9     | 3        | 3        | 4        |

IDH: intradialytic hypotension; HC: hydrocortisone; Y: yes; N: no.
Midodrine could be an effective treatment for IDH, but it could be associated with higher long-term mortality [16]. A high dialysate sodium level could have some hemodynamic benefits in HD patients, but it will lead to sodium loading and more weight gain between HD treatments, which could eventually lead to more IDH episodes [17]. Cooling the dialysate could be an effective measure for lowering the frequency of IDH by promoting peripheral vasoconstriction, but there was a higher rate of feeling cold among all patients [18]. One case report was published when this study was running showing that fludrocortisone could be helpful for the treatment of IDH [19], but in this study, we studied hydrocortisone instead which has both glucocorticoid and mineralocorticoid activity.

A hydrocortisone dosage of 300-450 mg/day was given to critically ill patients requiring chronic renal replacement therapy, and this resulted in the normalization of serum sodium and potassium in 4 of the clinical cases [20]. The serum cortisol level starts to rise 3-4 hours after starting hemodialysis, it peaks at 6 hours, and it returns to the predialysis level 24 hours after dialysis treatment [8]. Hemodialysis could remove some of the already present serum cortisol and along with the supposed delay in adrenocortical response; this will put our HD patients at risk for adrenal crisis [8].

Long-term use of supraphysiologic doses of glucocorticoids, even if just intermittent, may have adverse effects on many major organ systems; however, the risk-benefit ratio of its use can be improved by careful monitoring and following of preventive strategies to minimize its potential side effects [21]. This would include providing appropriate immunizations prior to the institution of therapy, also assessment for the presence of any preexisting conditions whose control may be affected by glucocorticoid use such as diabetes mellitus and hypertension [22]. For example, in patients who develop steroid-induced hyperglycemia, adding insulin treatment or adjusting its dose is a common and effective treatment and the same applies to patients with hypertension by adjusting BP medications if needed.

Based on our literature review (PubMed and Google), there is no clinical trial that has studied the possible therapeutic effects of hydrocortisone in the prevention of IDH. We strongly believe that using intravenous hydrocortisone as mentioned above could be a breakthrough, as it is an inexpensive, safe, and available medication for the management of IDH.

Our study has some limitations. Our study has a small sample size, but despite this, we found statistically and clinically significant results. Longitudinal multicenter studies are needed to assess the morbidity and mortality benefits of this treatment for patients with IDH.

5. Conclusion

This preliminary investigational study showed that the administration of a stress dose of hydrocortisone prior to hemodialysis could be an effective measure for preventing or minimizing the risk of intradialytic hypotensive episodes. Additional prospective studies on this subject are needed to further evaluate this topic. Ruling out adrenal insufficiency in patients diagnosed with IDH is also crucial.

Abbreviations

ACTH: Adrenocorticotropic hormone  
IDH: Intradialytic hypotension  
HD: Hemodialysis  
KDOQI: Kidney Disease Outcomes Quality Initiative  
BP: Blood Pressure  
IV: Intravenous.

Data Availability

Data are available from the corresponding author upon request.

Conflicts of Interest

All authors declare that there are no conflicts of interest.

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Table 3: A comparison between mean SBP for the included sample at days 1, 2, and 3 with and without hydrocortisone.

| Day  | Mean SBP (mmHg) | SD  | Mean difference | SD  | p value |
|------|----------------|-----|----------------|-----|---------|
|      | Without HC     |     |                |     |         |
| Day 1| 118            | 22.7| 2.0            | 10.3 | 0.47    |
|      | 116            | 23.1|                |     |         |
| Day 2| 112            | 23.2| 2.5            | 9.7  | 0.36    |
|      | 109            | 21.9|                |     |         |
| Day 3| 105            | 19.4| -6.3           | 12.2 | 0.09    |
|      | 111            | 22.9|                |     |         |

HC: hydrocortisone; SBP: systolic blood pressure.
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