Linear mixed model analysis to evaluate correlations between remdesivir adverse effects with age and gender of patients with mild Covid-19 pneumonia

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Abstract
We aimed to assess longitudinal changes in clinical indexes of corona disease 2019 (Covid-19) patients with mild pulmonary infection during 5 days of remdesivir therapy and determine the effect of age and gender on remdesivir adverse effects (AE). Patients’ clinical data including inflammatory markers, liver and renal function tests, and heart rate (HR) were extracted from medical records. Linear mixed model (LMM) was used to analyze longitudinal changes in patients’ clinical indexes. Gender and age were inserted in LMM as covariates to find their correlation with AE and clinical indexes. Of 84 patients, 35 patients met our criteria for the study. There were significant increases in mean levels of white blood cell (WBC; \( p = 0.005 \)), alanine aminotransferase (ALT; \( p = 0.001 \)), aspartate aminotransferase (\( p = 0.001 \)), blood urea nitrogen (BUN; \( p = 0.001 \)), and creatinine (\( p = 0.006 \)), whereas mean levels of erythrocyte sedimentation rate (\( p = 0.005 \)), C-reactive protein (\( p = 0.001 \)), alkaline phosphatase (\( p = 0.001 \)), and potassium (\( p = 0.003 \)) decreased significantly. Estimated glomerular filtration rate (\( p = 0.001 \)) and HR (\( p = 0.001 \)) showed a notable decline over the course of treatment. LMM analysis showed that mean changes in WBC (\( \beta = 0.94, p = 0.029 \)), creatinine (\( \beta = 0.12, p = 0.020 \)), and HR (\( \beta = 6.47, p = 0.008 \)) were greater in males than in females. Also, age of patients had a significant effect on the mean changes of WBC (\( \beta = -0.02, p = 0.023 \)), sodium (\( \beta = -0.06, p = 0.010 \)), BUN (\( \beta = 0.23, p = 0.001 \)), and HR (\( \beta = -0.29, p = 0.001 \)). Despite no renal and liver dysfunction, Covid-19 patients with mild pulmonary infection may develop some remdesivir AE and attributed side effects might be affected by gender and age of patients.

KEYWORDS
adverse effects, Covid-19, linear mixed model, remdesivir

1 | INTRODUCTION

Corona disease 2019 (Covid-19), caused by a novel severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), is a viral infection of the respiratory tract typified with fever, dry cough, fatigue, and worsening dyspnea.1 During the current pandemic, several treatment protocols for Covid-19 including chloroquine/hydroxychloroquine, favipiravir, lopinavir/ritonavir, and azithromycin have been tested and used.2 Remdesivir, an adenosine triphosphate (ATP) analog that inhibits viral RNA polymerase, has been approved by the US Food and Drug Administration (FDA) for hospitalized patients with Covid-19.3,4 Despite the beneficial effects of remdesivir in Covid-19 treatment, hepatotoxicity and nephrotoxicity have been reported as adverse effects (AEs) in Covid-19 patients receiving
intravenous remdesivir because of accumulation of sulfobutylether-β-cyclodextrin (SBECD) carrier.5

Despite the emergent and widespread use of remdesivir for the Covid-19 treatment, some AE including cardiac, renal, and liver complications have been reported in patients receiving remdesivir.6 Among the remdesivir AEs, anemia, infusion site reactions, increased liver enzymes, cutaneous rash, kidney injuries, and hypotension are the most common. According to the FDA Emergency Use Authorization, a loading dose of 200 mg (5 mg/kg) once in a day in patients ≥40 kg and 100 mg from Day 2–5 (2.5 mg/kg) has been proposed. Clinical investigations on patients with normal kidney function indicated that remdesivir and its active metabolite, remdesivir triphosphate, are predominantly (74%) eliminated by the kidneys.7,8 Moreover, transaminase elevations have been observed in Covid-19 patients receiving remdesivir and, therefore, liver function tests (LFTs) must be monitored daily, and remdesivir is discontinued in individuals with alanine aminotransferase (ALT) more than five times the upper limit of normal healthy individuals.7 Regarding cardiovascular complications, sinus bradycardia, hypotension, T-wave abnormalities, atrial fibrillation (AF), and a prolonged QT interval are the most common AEs.9

It has been shown that remdesivir is adequately tolerated but does not provide significant therapeutic effects in seriously COVID-19 patients. Furthermore, there is a lack of clinical evidence to evaluate the appearance of remdesivir AE and outcomes in men and women and different age groups.10 The goal of the current study is to investigate longitudinal changes in clinical indexes of hospitalized Covid-19 patients with mild pulmonary infection during a 5-day course for remdesivir therapy.

2 | PATIENTS AND METHODS

2.1 | Study design and subjects

This retrospective longitudinal study was conducted on Covid-19 hospitalized patients receiving intravenous remdesivir in the regular ward between June and September 2021. The protocol of the study was approved by the ethics and research review committee of the Iran University of Medical Sciences (IR. IUMS. REC.1400.072). All participants gave written informed consent, and the study was carried out following...
the guidelines of the Declaration of Helsinki. Inclusion criteria to enter in the study were patients who had no history of kidney and liver dysfunction, estimated glomerular filtration rate (eGFR) cutoffs of 30 or 50 ml/min per 1.73 m², elevated LFT less than five times the upper limit of normal, oxygen saturation above 90%, and pulmonary infection of <20% in lung high-resolution computed tomography. The estimated sample size was 36 calculated based on Type I error (5%), power (80%), an effect size of 2.9, and change SD 6 for ALT before and after remdesivir reported by Aiswarya et al.12 All patients were treated according to protocol with dexamethasone 6 mg daily, remdesivir (a loading dose of 200 mg on day 1 followed by 100 mg for 4 days), and enoxaparin 60 mg daily. Demographic data and medical records of patients were extracted from Hospital Information System and analyzed.

2.2 | Clinical indexes

To follow Covid-19 disease inflammatory state during 5 days remdesivir therapy, white blood cell (WBC) count, C-reactive protein (CRP), and erythrocyte sedimentation rate (ESR) were assessed in the patients. Also,

| TABLE 1 | Characteristics of Covid-19 patients |
|---|---|
| Age (year) | 56.22 ± 17.86 |
| Gender (n, %) |  |
| Male | 15 (42.9) |
| Female | 20 (57.1) |
| Fever (n, %) | 14 (40) |
| Dyspnea (n, %) | 27 (77.1) |
| Dry cough (n, %) | 31 (88.6) |
| HTN (n, %) | 11 (31.4) |
| DM (n, %) | 6 (17.1) |
| Favipiravir consumption (n, %) | 22 (62.9) |
| Spo2 (%) | 92.22 ± 2.30 |

Abbreviations: Covid-19, corona disease 2019; DM, diabetes mellitus; HTN, hypertension.

aContinues data are presented as mean ± SD.
bCategorical data are presented as frequency (percentage).

| TABLE 2 | Clinical indexes of Covid-19 patients over the course of remdesivir therapy |
|---|---|
| Clinical index | Reference range | Gender | Days of remdesivir therapy | Days of remdesivir therapy | Days of remdesivir therapy | Days of remdesivir therapy | Days of remdesivir therapy | p |
|---|---|---|---|---|---|---|---|---|---|
| WBC | 4.5–11 (×10^3/micL) | M | 5.30 ± 1.84 | 7.41 ± 2.25 | 6.94 ± 2.28 | 0.006 |
| F | 4.63 ± 1.53 | 6.23 ± 1.84 | 6.05 ± 1.06 | 0.001 |
| ESR | Up to 15 (mm/h) | M | 41.27 ± 26.44 | 41.01 ± 29.78 | 22.0 ± 12.66 | 0.003 |
| F | 44.50 ± 20.58 | 34.81 ± 17.48 | 30.60 ± 14.85 | 0.001 |
| CRP | Up to 6 (mg/L) | M | 27.11 ± 26.99 | 16.57 ± 15.53 | 10.88 ± 11.91 | 0.001 |
| F | 22.34 ± 18.54 | 14.58 ± 15.04 | 10.49 ± 10.66 | 0.002 |
| ALT | Up to 40 (U/L) | M | 32.73 ± 19.60 | 41.01 ± 29.78 | 22.0 ± 12.66 | 0.003 |
| F | 43.10 ± 23.68 | 34.81 ± 17.48 | 30.60 ± 14.85 | 0.001 |
| AST | Up to 37 (U/L) | M | 37.86 ± 8.77 | 47.86 ± 11.35 | 51.93 ± 15.76 | 0.001 |
| F | 43.51 ± 15.54 | 55.50 ± 18.59 | 60.95 ± 18.98 | 0.001 |
| ALP | 65–305 (U/L) | M | 198.94 ± 65.25 | 134.93 ± 43.47 | 126.40 ± 27.75 | 0.001 |
| F | 234.60 ± 115.08 | 152.10 ± 58.75 | 128.20 ± 58.41 | 0.001 |
| Sodium | 135–145 (mEq/L) | M | 135.67 ± 3.31 | 138.47 ± 3.74 | 137.87 ± 3.81 | 0.026 |
| F | 137.80 ± 4.12 | 137.30 ± 2.72 | 138.00 ± 1.98 | 0.195 |
| Potassium | 3.5–5.5 (mEq/L) | M | 4.07 ± 0.19 | 3.91 ± 0.17 | 3.70 ± 0.16 | 0.017 |
| F | 3.90 ± 0.37 | 3.69 ± 0.29 | 3.63 ± 0.21 | 0.004 |
| BUN | 14–45 (mg/dl) | M | 29.67 ± 12.52 | 40.33 ± 14.48 | 41.01 ± 6.47 | 0.001 |
| F | 27.99 ± 9.78 | 36.60 ± 9.12 | 38.31 ± 5.98 | 0.001 |
| Creatinine | 0.7–1.4 (mg/dl) | M | 1.06 ± 0.25 | 1.05 ± 0.21 | 1.04 ± 0.20 | 0.848 |
| F | 0.86 ± 0.15 | 0.83 ± 0.17 | 1.00 ± 0.16 | 0.001 |
| eGFR | ≥60 ml/min/1.73 m² | M | 81.07 ± 22.18 | 79.33 ± 19.74 | 78.73 ± 17.19 | 0.007 |
| F | 79.31 ± 18.88 | 75.90 ± 17.30 | 62.05 ± 14.37 | 0.001 |
| Hear rate | 60–100 beats/min | M | 81.13 ± 12.19 | 62.33 ± 12.14 | 58.13 ± 14.06 | 0.001 |
| F | 77.30 ± 10.30 | 60.85 ± 6.23 | 51.15 ± 4.34 | 0.001 |

Abbreviations: ALT, alanine transaminase; ALP, alkaline phosphatase; AST, aspartate aminotransferase; BUN, blood urea nitrogen; Covid-19, corona disease 2019; CRP, C-reactive protein; eGFR, estimated glomerular filtration rate; ESR, erythrocyte sedimentation rate; F, female; M, male; WBC, white blood cell.

aContinues data are presented as mean ± SD.
remdesivir AEs on the function of the liver, kidney, and heart were monitored through checking clinical indexes including aspartate transaminase (AST), ALT, alkaline phosphatase (ALP) for liver function, blood urea nitrogen (BUN), creatinine, eGFR, sodium, and potassium for kidney function, and heart rate (HR) on the first, third, and fifth days of remdesivir therapy.

### 2.3 Statistical analysis

Linear mixed model (LMM) is one of the longitudinal data analysis methods that describes the relationship between a response variable and other explanatory variables that have been obtained along with the response. We used LMM to assess mean changes of clinical indexes in patients over the time of remdesivir therapy. Age and gender were inserted in the model as covariates to analyze their effects on the mean changes in clinical indexes. Restricted maximum likelihood with slope and intercept random effect was used to estimate all coefficients in the model. Continuous variables are presented as mean ± SD and categorical variables are presented as frequency (%). All statistical analyses were done using IBM SPSS version 24.0 (IBM, Inc.). $p < 0.05$ were considered significant.

**FIGURE 2** Changes in mean white blood cell (WBC), C-reactive protein (CRP), and erythrocyte sedimentation rate (ESR) index estimated by linear mixed model (LMM) analysis on the first, third, and fifth day of remdesivir therapy over a 5-day course of treatment in corona disease 2019 (Covid-19) patients (reference: first day).

**FIGURE 3** Changes in mean alanine transaminase (ALT), aspartate transaminase (AST), and alkaline phosphatase (ALP) index estimated by linear mixed model (LMM) analysis on the first, third, and fifth day of remdesivir therapy over a 5-day course of treatment in corona disease 2019 (Covid-19) patients (reference: first day).
3 | RESULT

3.1 | Characteristics of Covid-19 patients

As depicted in Figure 1, among 84 Covid-19 patients who were admitted for remdesivir therapy, 35 patients met our inclusion criteria and enter into the study. Demographic and clinical characteristics of Covid-19 patients are summarized in Table 1. The mean age of patients was 56.22 ± 17.86 years and 57.1% of them were women. Dyspnea (77%) and dry cough (88.6%) were the most common symptoms of the disease and 62.9% of patients received oral Favipiravir before starting remdesivir therapy.

3.2 | Remdesivir AE in Covid-19 patients during treatment

As represented in Table 2, mean WBC increased significantly (male: \( p = 0.006 \), female: \( p = 0.001 \)), whereas mean ESR (male: \( p = 0.003 \), female: \( p = 0.001 \)) and CRP (male: \( p = 0.001 \), female: \( p = 0.002 \)) showed a notable decline through remdesivir therapy. Figure 2 shows changes in mean WBC, CRP, and ESR index for male and female Covid-19 patients in more detail during 5 days of remdesivir injection.

Regarding remdesivir AE, we found a significant rise in mean levels of ALT (male: \( p = 0.001 \), female: \( p = 0.001 \)) and AST (male: \( p = 0.001 \), female: \( p = 0.001 \)), and a considerable decline in mean ALP (male: \( p = 0.001 \), female: \( p = 0.001 \)) during treatment. Figure 3 illustrates changes in mean ALT, AST, and ALP in further detail over the course of remdesivir therapy. Mean sodium in males showed a notable increase (\( p = 0.026 \)), whereas mean potassium in both males (\( p = 0.017 \)) and females (\( p = 0.004 \)) decreased significantly. However, sodium and potassium values remained within the normal range during treatment. Also, mean BUN in both genders (\( p = 0.001 \)) increased notably, while just females experienced a significant rise in creatinine levels (\( p = 0.001 \)). Figure 4 provides more information about changes in mean levels of BUN, creatinine, sodium, and potassium on the first, third, and fifth day of remdesivir injection.

Furthermore, females experienced a notable decrease in eGFR (\( p = 0.001 \)) and mean HR of patients decreased significantly (male: \( p = 0.001 \), female: \( p = 0.001 \)) during remdesivir therapy but HR gradually restored to normal after drug discontinuation. Figure 5 presents changes in mean eGFR and HR in more detail over the course of treatment.
The clinical syndrome caused by Covid-19 infection primarily affects respiratory tracts, but other organs including the heart, gastrointestinal system, and liver may also be affected. Recent reports have shown that some patients with the severe form of the Covid-19 infection experience an intensified immune response similar to cytokine release syndrome and multisystem inflammatory syndrome. Consistent with us, a prior study by Stoeckle et al. have shown that CRP levels decrease significantly after remdesivir treatment in none intubated Covid-19 patients, indicating that remdesivir mitigates inflammatory response in a certain group of patients but not in all. Moreover, women patients in the study had lower CRP levels than men that may be due to the lower viral load and higher number of CD4+ T cells, and higher levels of antibodies in women. Interestingly, producing interleukin-6, which actively

3.3 Relationship of remdesivir AE with age and gender of patients

As described in the Section 2.3, age and gender were inserted in the LMM as covariates to find their effects on changes in clinical indexes in patients treated with remdesivir. Our results are summarized in Table 3 and indicate that gender as a covariate was significantly related to the changes in patients' indexes. Mean changes in WBC ($\beta = 1.03, p = 0.029$), creatinine ($\beta = 0.15, p = 0.009$), and HR ($\beta = 6.22, p = 0.015$) attributed to the remdesivir were greater for men than women. In addition, age of patients had a significant effect on the mean changes of WBC ($\beta = -0.02, p = 0.023$), sodium ($\beta = -0.06, p = 0.026$), BUN ($\beta = 0.23, p = 0.001$), and HR ($\beta = -0.29, p = 0.001$) over the time of remdesivir therapy.

4 DISCUSSION

In this single-center study, clinical outcomes of 35 Covid-19 patients with mild pulmonary infection treated with remdesivir were evaluated. We did not find serious AEs attributed to the remdesivir therapy but bradycardia with renal and liver dysfunction were seen in the Covid-19 patients. In addition, age and gender of patients had significant effects on some AEs of remdesivir.

### Table 3: Effects of gender and age on remdesivir AEs assessed by LMM analysis

| Variables | Mean ± SE | Gender | Age |
|-----------|-----------|--------|-----|
| WBC       | Male = 6.57 ± 0.33, $\beta = 1.03, p = 0.029$; Female = 5.63 ± 0.30, $\beta = 0.023$ |
| ESR       | Male = 29.83 ± 5.52, $\beta = -2.69, p = 0.13$; Female = 40.30 ± 4.81, $\beta = 0.404$ |
| CRP       | Male = 17.80 ± 2.84, $\beta = 1.54, p = 0.062$; Female = 16.26 ± 2.65, $\beta = 0.770$ |
| ALT       | Male = 48.09 ± 6.81, $\beta = 1.42, p = 0.09$; Female = 49.51 ± 6.08, $\beta = 0.735$ |
| AST       | Male = 46.29 ± 3.49, $\beta = -7.60, p = 0.05$; Female = 50.13 ± 3.16, $\beta = 0.708$ |
| ALP       | Male = 151.47 ± 15.73, $\beta = -21.62, p = 0.43$; Female = 173.09 ± 13.03, $\beta = 0.478$ |
| Sodium    | Male = 137.78 ± 0.56, $\beta = 0.15, p = 0.06$; Female = 137.64 ± 0.49, $\beta = 0.026$ |
| Potassium | Male = 3.90 ± 0.04, $\beta = 0.12, p = 0.001$; Female = 3.78 ± 0.03, $\beta = 0.597$ |
| BUN       | Male = 35.52 ± 1.54, $\beta = 0.73, p = 0.23$; Female = 34.78 ± 1.46, $\beta = 0.001$ |
| Creatinine| Male = 1.04 ± 0.04, $\beta = 0.15, p = 0.02$; Female = 0.91 ± 0.03, $\beta = 0.460$ |
| eGFR      | Male = 84.04 ± 4.08, $\beta = 10.79, p = 0.16$; Female = 71.15 ± 3.66, $\beta = 0.294$ |
| Heart rate| Male = 68.55 ± 1.76, $\beta = 6.22, p = 0.001$; Female = 62.10 ± 1.53, $\beta = 0.015$ |

Abbreviations: AE, adverse effect; ALT, alanine transaminase; ALP, alkaline phosphatase; AST, aspartate aminotransferase; BUN, blood urea nitrogen; CRP, C-reactive protein; eGFR, estimated glomerular filtration rate; ESR, erythrocyte sedimentation rate; LMM, linear mixed model; WBC, white blood cell.
fat, different gastric motility, and lower GFR in comparison to men. These anatomical and physiological differences can affect the pharmacokinetics and pharmacodynamics of the drugs in the body including drug absorption, distribution, metabolism, and elimination.\textsuperscript{31}

The present study has some potential limitations. Our investigation was a single-center study and the main limitations of our investigation were short-term follow-up and small sample size. Also, the study involved hospitalized Covid-19 patients treated with remdesivir and the lack of Covid-19 control patients who did not treat with remdesivir is another limitation. Thus, we cannot exactly conclude that AEs observed in the patients are due to remdesivir or SARS-CoV-2. Hence, more extensive studies with a control group are necessary to investigate remdesivir AEs and predict sex and age differences in remdesivir treatment outcomes.

5 | CONCLUSION

We found that patients with mild Covid-19 pneumonia may develop AE attributed to the remdesivir, despite having no liver and renal dysfunction. Moreover, remdesivir AE might be affected by gender and age of patients and clinicians should be more cautious about additive AE when other medicines are used in addition to remdesivir therapy.

AUTHOR CONTRIBUTIONS

Mohsen Sedighi contributed to conceptualization, project administration, methodology, and writing the original draft. Alireza Amanollahi contributed to data analysis, methodology, and writing the original draft. Nader Tavakoli, Omid M. Moghaddam, Hamed B. Ghafoori, and Seyede E. Hoseini contributed to the data collection and investigation. Nader Tavakoli contributed to funding acquisition, supervision, writing, and review, and editing.

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CONFLICTS OF INTEREST

The authors declare no conflicts of interest.

DATA AVAILABILITY STATEMENT

Data are available from the corresponding author upon reasonable request.

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