



Research Article

Received: date: 05.09.2023 Accepted: date: 28.10.2023 Published: date: 31.12.2023

Mathematical Modeling of the Spread of Sars-Cov-2 at the Onset of Vaccination Against Covid-19 with CoronaVac in Türkiye

Ersin Sener 1* Ummu Sahin Sener 2

¹Department of Mathematics, Faculty of Science and Arts, Kırklareli University, Kırklareli, Türkiye; ersinsener@klu.edu.tr ²Department of Mathematics, Faculty of Science and Arts, Kırklareli University, Kırklareli, Türkiye; ummusahin@klu.edu.tr Orcid: 0000-0002-5934-3652¹ Orcid: 0000-0001-9055-8734²

*Correspondence: ersinsener@klu.edu.tr

Abstract: The Sars-CoV-2 virus, first detected in Wuhan, China, became a global crisis that affected the entire world and was declared a pandemic by the World Health Organization (WHO) in March 2020. The most basic protective measure in the fight against pandemics facing humanity is vaccination. From this point of view, data is collected between January 13 and February 11, 2021 by taking the number of daily cases, deaths and recovered patients in Türkiye. During this period, vaccination against Covid-19 with Sinovac's CoronaVac vaccine is started in Türkiye. Mathematical predictive models of the observed values are constructed and compared using polynomial regression (up to the 3rd degree) and nonlinear regression, i.e., curve fitting methods, and **SIR** (Susceptible-Infected-**R**emoved), which is a system of ordinary differential equations (ODEs). The efficiencies of these prediction models are tested, validated, and the most effective mathematical prediction models are proposed. The values of root mean square error (**RMSE**) and mean absolute percentage error (**MAPE**) are used as performance measures to compare the methods. The proposed prediction models are also used for forecasting. The number of new cases occurring each day is predicted using the time-dependent equations of the SIR method, which are solved using the Euler method. It is found that the SIR method is quite successful in predicting the observed values compared to the other methods, but the QR method are given more successful results in predicting the total number of deaths. **Keywords:** Curve fitting, mathematical modeling, polynomial regression, SARS-CoV-2, SIR

1. Introduction

In early December 2019, a case of pneumonia of unknown etiology is discovered in Wuhan city, Hubei province, China, and the disease is reported to the World Health Organization (WHO) in late December 2019 [1]. In January 2020, a novel coronavirus, severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), is isolated from patients with infected pneumonia, and the disease caused by this virus is designated as coronavirus disease 2019 (COVID -19) in February 2020 [2, 3]. Since this virus spreads very rapidly within a few months and infects patients around the world, it is recognized as a pandemic by WHO in March 2020 [4]. Pandemic is the general term for epidemic diseases that spread rapidly over a large area in more than one country or continent in the world and cause deaths.

Building mathematical models using computational methods in pandemics is important to determine the rate of spread of the disease and the actions that need to be taken to prevent the spread of the disease. Mathematical modeling of the spread of epidemics has a long history and is initiated in the 1700s by Daniel Bernoulli, who developed a mathematical model to analyze the mortality caused by smallpox [5, 6]. Not much work had been done on this topic until the publication of Ross, which is developed in the

Citation: E. Sener and Ü.S. Sener, "Mathematical Modeling of the Spread of Sars-Cov-2 at the Onset of Vaccination Against Covid-19 with CoronaVac in Türkiye", Journal of Statistics and Applied Sciences, no. 8, pp. 1-14, Dec. 2023, doi:10.52693/ jsas.1355520

early 1900s and used a mechanistic a priori modeling approach with a set of equations to approximate discrete time dynamics that can be considered the foundation of mathematical epidemiology.

Mathematical and computational methods used in epidemiology can make important contributions to the spread, incidence, analysis, and control of disease [7]. The use of mathematical models in epidemiology has made it possible to define complex data, determine general rules for epidemic dynamics, predict parameters that cannot be directly measured, identify problems that might threaten public health, and select an optimal experimental design [8, 9]. The models are generally used to predict and explain trends in disease recurrence, spread, morbidity, or mortality [10]. With the correct interpretation of systems of dynamic equations, the development of analytical solutions, and the advancement of numerical methods, the methods used in epidemic modeling have evolved considerably [11]. By implementing infection prevention and control measures, it may be possible to reduce the spread of infection in the community and thus reduce the number of people who become infected in the early stages of the pandemic.

In the last 2 years, modeling the spread of the virus has been the main problem for researchers in the Covid-19 trial, which has negatively affected our lives and even stalled. A number of studies have used mathematical epidemiological models to analyze the transmission dynamics of COVID -19. SIR is one of the most commonly used mathematical epidemiological models [12]. Covid-19 data from seven countries-China, South Korea, Italy, Spain, Brazil, Germany, and France during the period from February to July 2020 are modeled using machine learning methods, SIR, and a time window SIR (TW-SIR) prediction model [13]. For Türkiye, Covid-19 data between March 11, 2020 and February 22, 2021 are used to examine monthly case counts with time series. In this study, a hybrid model of seasonal autoregressive integrated moving average (SARIMA) and neural network nonlinear autoregressive (NNAR) hybrid model is implemented [14]. The spread of Covid-19, many methods of time series analysis, and mathematical modeling methods of epidemiology are discussed very extensively [15]. In accordance with the information obtained from the literature review, it appears that mathematical epidemiological models are used for the early periods of Covid-19, but curve fitting methods are not used in these studies. The studies conducted to date have attempted to predict when the virus will peak.

epidemiological models are used for the early periods of Covid-19, but curve fitting methods are not used in these studies. The studies conducted to date have attempted to predict when the virus will peak. This study focuses on the period between January 13 and February 11, 2021, when vaccination against COVID -19 started and the first vaccine dose is administered. Our primary objective is to create mathematical prediction models for the spread of SARS-CoV-2 virus during the first dose of vaccination against SARS-CoV-2 coronavirus with Sinovac's CoronaVac vaccine, which is licensed for emergency use in Türkiye. The secondary objective is to propose an optimal model by comparing the success of the SIR model [13–16], which is a system of ordinary differential equations used in the construction of predictive models, linear regression (LR), polynomial regression (quadratic regression (QR) and cubic regression (CR)), and nonlinear regression (NLR) in estimating the general parameters of the spread of the pandemic [17]. In general, one of the objectives of this study is to determine which mathematical models can predict and create a priori the spread of the disease in the event of a possible pandemic in our globalized world. In addition, predictions of the daily number of cases and the daily number of deaths for the period February 12-15, 2021, are made using optimal mathematical predictive models.

2. Materials and Methods

2.1. Data of SARS-Cov-2

Our dataset consists of daily values of infected individuals (column 1), removed individuals (column 2), and deceased individuals (column 3) between January 13 and February 11, 2021 for Türkiye [18]. The dataset is split into two terms for modeling, the training term, and the test term. The first term is the training term between January 13 and February 5, 2021, and the training term is the percentage of 80 % of the total data. The second term is the test term between February 6 and February 11, 2021. The test term is used for model validation. The mathematical modeling processes of the data in the study are given in Figure 1 as a flow-chart.

In addition to the data set, the initial values needed to solve the model SIR using Euler's method are determined as follows: At t = 0 (January 13, 2021), there are susceptible individuals ($S_0 = 81992782$), infected individuals ($I_0 = 104669$), removed individuals ($R_0 = 2241616$), and the population is (N = 84339067).



Figure 1. Mathematical modeling flow-chart

2.2. Methodology

2.2.1. SIR Model

....

In mathematical modeling of epidemics, the ordinary differential equation model, the so-called SIR (Susceptible-Infected-Removed (recovered and dead)) model, is one of the basic models. Individuals in a population of N are assumed to belong to one of three groups at time t [10].

S(t): The class of individuals that are not infected now but will be infected later (Susceptible-S).

I(t): The class of individuals who have contracted the disease and are now ill, i.e., individuals who infect others or are associated with an infection and infect others.

R(t): The class of individuals who are removed (recovered and died).

The number of individuals in each of these groups changes with time, i.e. S(t), I(t), and R(t) are functions of time t. The sum of the individuals in these three groups gives the total population size N in other words $N(t) = S(t) + I(t) + R(t) = constant, t \ge 0$. As long as the pandemic continues, people from the "S" group can progress to the "I" group, and people from the "I" group can progress to the "R" group.

$$\frac{dS}{dt} = -\beta S(t)I(t), S(0) = S_0 \tag{1}$$

$$\frac{dI}{dt} = \beta S(t)I(t) - \gamma I(t), I(0) = I_0$$
⁽²⁾

$$\frac{dR}{dt} = \gamma I(t), R(0) = R_0 \tag{3}$$

 $\beta > 0$ is the contact rate or infection rate of the disease, $\gamma > 0$ is the recovery rate from infected persons to recovered persons. In the Eq. (1), the infection rate of healthy people at a given time is proportional to the ratio of healthy to infected people. That is, it is proportional to the product of S(t), and I(t). The rate of change is negative because the healthy population is infected, and the number of healthy people always decreases. In Eq. (2) the rate of change of infected people is given by the difference between the rate at which healthy people become infected and the rate at which infected people move into the group of those who removed according to Eq. (3), the rate at which people who recover or die leave the infected group is directly proportional to the number of infected people. The relationship between S(t), I(t), and R(t) can be seen in the flowchart of the model SIR in Figure 2.



Figure 2. Flowchart of the SIR pandemic model

Assuming that the time value *t* goes to infinity $\lim_{t\to\infty} S(t) = S_{\infty}$ and $\lim_{t\to\infty} R(t) = R_{\infty}$. The number of infected individuals can fall to zero or behave nonmonotonically by first increasing to a maximum and then falling to zero. When $I'(0) = \beta S(0)I(0) - \gamma I(0) > 0$, prevalence begins to increase. The necessary

and sufficient condition for the first increase in the number of infected is $\beta S(0) - \gamma > 0$ or $\frac{\beta S(0)}{\gamma} > 1$.

Dividing Eq. (1) by Eq. (3);

$$\frac{dS}{dR} = \frac{-\beta S(t)I(t)}{\gamma I(t)} = -\frac{\beta S(t)}{\gamma}$$
(4)

Solving this equation for t, Eq. (5) can be obtained as follows:

$$S = S(0)e^{-\beta R/\gamma} \ge S(0)e^{-\beta N/\gamma} > 0$$
(5)

From this it follows that $S_{\infty} > 0$ and S_{∞} is the final size of the pandemic, the pandemic is extinguished when $\lim_{t \to \infty} I(t) = I_{\infty} = 0$ and R_{∞} is bounded by *N*. The equations given in Eq. (1-3) for the model SIR are ODEs that can be solved using Euler's formula.

$$S_{n+1}(t) = S_n(t) - \beta S_n(t) I_n(t) \Delta t = S_n(t) [1 - \beta I_n(t) \Delta t]$$
(6)

$$I_{n+1}(t) = I_n(t) + (\beta S_n(t) I_n(t)) \Delta t = I_n(t) [1 + (\beta S_n(t) - \gamma) \Delta t]$$
(7)

$$R_{n+1}(t) = R_n(t) + (\gamma I_n(t))\Delta t$$
(8)

where $\Delta t = t_{n+1} - t_n$ is a small time change, $S_{n+1}(t)$, $I_{n+1}(t)$, and $R_{n+1}(t)$ are susceptible, infected and recovered individuals, respectively, calculated from the previous step.

2.2.2. Polynomial Regression

Regression analysis is one of the most common statistical methods to study and model the relationship between variables. It identifies the relationship between a dependent variable and one or more independent variables. Assuming a model of the relationship between variables and estimates of parameter values, a predictive regression equation is developed.

The linear regression model, the simplest regression model given by the equation $\hat{y} = X\beta + \epsilon$, is a general model used to construct any linear relationship in the unknown parameters β . $\hat{y} = \beta_0 + \beta_1 x + \epsilon$

is called the linear regression model [19]. x is the independent variable, that is predictor or regressor variable, and y is called the dependent variable or, in other words, the response variable. Since this equation involves only one regressor variable, this model is called a simple linear regression model. The linear regression model is a general model that can be used in cases where the relationship between explanatory variables and response variable is linear.

Even in complex nonlinear relationships, polynomials can be used extensively in situations where the response is curvilinear, since modeling can be done by fitting polynomials over small intervals of x. Polynomial regression is a special case of the general linear regression model and includes the quadratic and higher order values of the independent variable(s) to make the regression function curvilinear [20]. In general, a k^{th} order polynomial model for one variable is given in Eq. (9).

$$\hat{y} = \beta_0 + \beta_1 x + \beta_2 x^2 + \dots + \beta_k x^k + \epsilon$$
(9)

The model given in Eq. (9) can be solved by the least squares method. Taking k = 1, one obtains a linear function, taking k = 2, a quadratic function, and taking k = 3, one obtains a cubic function.

Unless required by the nature of the data or for other reasons, the degree of the polynomial used should be kept as low as possible. In our study, by using polynomials up to the 3^{th} degree, an attempt is made to find out at which polynomial of what degree the best result is obtained, and it is found that the most suitable curve was in the third-degree polynomial regression.

2.2.3. Non-linear Regression

Note that the shape and parameters of the curve can be determined by a nonlinear regression approach as in Eq. (10) if the data set can be represented by a nonlinear regression curve when the observed values are plotted [21].

$$\hat{y} = ae^{bx} + \epsilon \tag{10}$$

2.2.4. Model Goodness of Fit

The evaluation of the fitted prediction models can be done with the Root Mean Squared Error (*RMSE*) and the Mean Absolute Percentage Error (*MAPE*). *RMSE* is the square root of the variance of the residuals and is given in Eq. (11).

$$RMSE = \sqrt{\frac{1}{n} \sum_{i=1}^{n} (y_i - \hat{y}_i)}$$
(11)

where *n* is the total number of observations, y_i is the *i*th observed values, and y_i is the *i*th predicted values. The fact that the *RMSE* value is very close to 0 indicates the absolute fit of the model to our data set (the lower *RMSE*, the better model fit). The second criterion in the examination of model fit, *MAPE*, is a measure of the estimation accuracy of an estimation method in curve fitting [22]. Usually, accuracy is expressed as a ratio defined by the following formula Eq. (12).

$$MAPE = \frac{100\%}{n} \sum_{t=1}^{n} \left| \frac{y_t - \hat{y}_t}{y_t} \right|$$
(12)

where y_t is the observed value, and \hat{y}_t is the predicted value at time t. The absolute value of the ratio in Eq. (12) is summed for each predicted time point and divided by the n number of observed values. The *MAPE* value is indeed very close to 1, indicating a relatively good fit to our data set. To investigate the optimal model fit for the datasets we have mathematically modeled, these two values are calculated. The proposed optimal model is constructed considering these criteria.

3. Results

This section is focused to the development of the models and the comparison of their performances. For this purpose, the data for the period February 6-11, 2021, corresponding to 20% of our dataset, are used as a test dataset to check the prediction performance of all the proposed models. The analyzes of the prediction models according to the explanatory variables are presented in the following subheadings.

3.1. SIR Model Parameters Estimation

When modeling the spread of Sars-Cov-2 virus using SIR, it is important to determine the model parameters. For this reason, the observed parameters γ -recovery rate and \Re_0 -reproduction number are calculated from the data sets are given in Figure 3a-3b.



Figure 3. (a)- Daily Recovery rate, (b)-Daily Reproduction number

The daily recovery rate, which indicates that those infected with the virus are less likely to transmit the disease, i.e., that they have recovered, is calculated daily using Eq. (8) and shown in Figure 3a. The Ministry of Health of the Republic of Türkiye has set the average time for an individual infected with Sars-CoV-2 virus to recover at 14 days [23]. Therefore, the recovery rate for an infected individual is $\gamma = 1/14 = 0.071$, which is shown in Figure 3a. The daily recovery rate we calculated shows that the recovery period of infected individuals lasts more than 14 days [24]. However, considering the statements of the official authorities, the daily recovery rate was set as $\gamma = 0.071$ in the analyzes.

The average infectivity of transmission of the virus to another individual by an individual infected with Sars CoV-2 virus can be defined as reproduction.

The reproduction number \Re_0 , which is an important value for determining actions to be taken depending on the course of the pandemic, is calculated daily [7]. \Re_0 is the average of the daily reproduction number $\Re_0 = 0.886$, as shown in Figure 3b.

At $\Re_0 > 1$ the risk of infection continues, $\Re_0 < 1$ the risk of infection decreases and may end, and

 $\Re_0 = 1$ the risk of disease remains constant [7]. The \Re_0 value calculated for the period in question is expected to end in the following days if current conditions are maintained.

3.2. Proposed Prediction Models of Susceptible (S)

The Susceptible (S) class of individuals that are not yet infected but may become infected later. Five different methods are used to model S in population N, namely SIR, linear regression (LR), quadratic regression (QR), cubic regression (CR), and non-linear regression (NLR), and the results of the predictive models are shown in Table 1.

The metrics *RMSE* and *MAPE* are used to determine the fitting success of curve fitting methods used to determine the curve that best predicts the available data. The $RMSE_s$ and $MAPE_s$ values calculated by each prediction method for the train and test terms are given in Table 1. The five different methods used to predict the *S* values are compared for the train and test terms into which we divided the data set to establish a model.

Model	Model Parameters	$RMSE_S$		MAPE _S	
Widdei		Train	Test	Train	Test
SIR	$\beta = 7.76e - 10$ $\gamma = 0.071$	1604.66	2712.66	1.62E-03	3.23E-03
LR	$\beta_0 = 81988284$ $\beta_1 = -6777.42$	2217.73	7857.92	2.36E-03	9.22E-03
QR*	$\beta_0 = 81984576$ $\beta_1 = -5953.33$ $\beta_2 = -31.696$	1659.99	2185.15	1.74E-03	2.44E-03
CR	$\beta_0 = 81989358$ $\beta_1 = -7966.15$ $\beta_2 = 158.1$ $\beta_3 = -4.87$	858.21	8675.56	8.40E-04	9.84E-03
NLR	a = 81988315 b = -8.27e - 5	2246.92	7355.40	2.42E-03	8.59E-03

Table 1. Results of proposed prediction models of *S*

When we examine the $RMSE_s$ and $MAPE_s$ values for the train term, the CR method yields the lowest $RMSE_s = 858.21$ and $MAPE_s = 8.40E - 04$ values. However, the CR method has the largest error values with $RMSE_s = 8675.56$ and $MAPE_s = 9.84E - 03$ in the test term. When estimating *S*, the $RMSE_s$ values are expected to be approximately the same for the train and test terms. The method QR, which has similar error values $RMSE_s = 1659.99, 2185.15$, for the model and test terms, respectively. Instead of evaluating the metrics $RMSE_s$ and $MAPE_s$ separately for the train and test terms, the curve fitting method that provides the best fit is determined by evaluating both values together. From this point of view, it is decided that the QR, which gives the smallest change in both metrics for both periods, would be a useful curve fitting method for estimating the susceptible individuals. The model QR given in Eq. (13) is proposed as the optimal model for predicting *S* in the population. The QR-quadratic regression prediction model of *S* is given in Eq. (13).

$$\hat{y} = 81984576 - 5953.33 \times day - 31.696 \times day^2 + \epsilon \tag{13}$$

See Figure 4 for a tangible representation of the predictions of the methods and the observed values. The predictions of the models are given below in Figure 4 to see the extent to which the prediction methods we used to fit the curve predict the observed values.



Figure 4. The graph of proposed prediction models of Susceptible (*S*)

The train and test term data are the observed values. Our goal is to build the predictive models over the train term and validate the built predictive models over the test term. To this end, in Figure 4, we see the predictions of S obtained using the method QR, and the predicted values of S are overlapped with the observed values. This overlap is quite successful. In addition, our forecasting is shown with the QR model between February 12-15, 2021, in Figure 4. In these projections, S is forecasted that will be approximately 81745522 via QR model by using Eq. (13) on February 15, 2021.

3.3. Proposed Prediction Models of Infected Individuals (I)

The class (I) of individuals who have the disease and are now infected, i.e., people who infect others or are associated with the infection and infect others. Five different methods are used to model I in population N SIR, LR, QR, CR, and NLR, respectively, and the results of the predictive models are shown in Table 2.

Model	Model Parameters	RN	ISE _I	MAPE _I	
		Train	Test	Train	Test
SIR*	$\beta = 7.76e - 10$ $\gamma = 0.071$	681.19	902.40	0.579	0.927
LR	$\beta_0 = 105127$ $\beta_1 = -815.603$	954.55	2166.10	0.817	1.97
QR	$\beta_0 = 105414$ $\beta_1 = -879.480$ $\beta_2 = 2.57$	947.72	1781.86	0.796	1.62
CR	$\beta_0 = 105378$ $\beta_1 = -864.11$ $\beta_2 = 1.008$ $\beta_3 = 0.037$	947.66	1707.44	0.794	1.57
NLR	a = 105543.64 b = -0.009	949.01	1639.94	0.792	1.54

Table 2. Results of proposed prediction models of I

The $RMSE_{I}$ and $MAPE_{I}$ values calculated for I from each prediction method for the train and test terms are given in Table 2. The five different methods used to predict the I values are compared for the train and test terms into which we divided the dataset to build a predictive model.

When we examine the $RMSE_i$ values for the train and the test terms, the method SIR, which is a system of ODEs, yields the lowest value with $RMSE_i = 681.19, 902.40$, respectively. When examining the $MAPE_i$ values for the train and the test terms, the method SIR provides the lowest value with $MAPE_i = 0.579, 0.927$. In predicting I, the values of $RMSE_i$, and $MAPE_i$ are expected to be approximately equal for the train and test terms. Thus, the model SIR is the most effective model for predicting I. The SIR prediction model of I is given in Eq. 14.

$$\hat{I} = 7.76e - 10 \times SI - 0.071 \times I \tag{14}$$

where \hat{I} is the predictor of I. A projection of all the prediction methods and the observed values can be found in Figure 5. The predictions of the models are given below in Figure 4 to see the extent to which the prediction methods we used to fit the curve predict the observed values of I.

Predictions of I obtained with the SIR method overlap with the observed values of I, as can be seen in Figure 5. This overlap is quite successful. In addition, our forecasting with the SIR model during February 12-15, 2021, are shown in Figure 5. In these projections, I is forecasted that will be about 82693 for February 15, 2021, using the SIR model with Eq. 14.



Figure 5. The graph of proposed prediction models of Infected (*I*)

3.4. Proposed Prediction Models of Removed Individuals (R)

The class of removed individuals (R) that are removed from the timely infected individuals (recovered and dead). Five different methods are used to model R in population N, namely SIR, LR, QR, CR, and NLR, and the results of the predictive models are shown in Table 3.

The $RMSE_R$ and $MAPE_R$ values calculated for R from each prediction method for the train and test terms are given in Table 3. In addition, Table 3 compares the five different methods used to predict R values for the train and test terms.

Model	Model Parameters	$RMSE_R$		MAPE _R	
		Train	Test	Train	Test
SIR*	$\beta = 7.76e - 10$ $\gamma = 0.071$	1646.72	902.40	0.0628	0.0908
LR	$\beta_0 = 2245656$ $\beta_1 = 7593.027$	2499.15	5943.90	0.0912	0.0239
QR	$\beta_0 = 2249077$ $\beta_1 = -6832.811$ $\beta_2 = 29.239$	2098.88	909.61	0.0702	0.0327
CR	$\beta_0 = 2244331$ $\beta_1 = 8830.256$ $\beta_2 = -159.108$ $\beta_3 = 4.829$	1554.66	10223.73	0.0561	0.373
NLR	a = 2247053 b = 0.003	2238.77	3698.38	0.0806	0.149

Table 3. Results of proposed prediction models of R.

When examine the $RMSE_R$ values for the model term, the CR method yields the lowest $RMSE_R = 1554.66$ value. For the test term, the CR method has the largest error value with $RMSE_R = 10223.73$. In predicting R, the values of $RMSE_R$ and $MAPE_R$ are expected to be approximately the same for the training and test terms. The method SIR, which has similar error values with $RMSE_R = 1646.72, 902.40$, $MAPE_R = 9.08E - 02$ for the train and test terms, respectively, is the most effective method in predicting R. The SIR prediction model of R is given in Eq. 15.

$$\hat{R} = 0.071 \times I$$

(15)

where \hat{R} is the predictor of R. For a projection of all prediction methods and observed values, see Figure 6 for R. Figure 6 is shown below to visually illustrate the extent to which the prediction methods predict the observed values of R and to see the forecasted values for the 4-day period February 12-15, 2021.



Figure 6. The graph of proposed prediction models of Removed individuals (R)

Predicting changes in the number of daily new cases (NC_{daily}) is of paramount importance to decision makers in the spread of the virus. Precisely for this purpose, we sought an answer to the question: **How to estimate the number of daily new cases?** By simple mathematical operations with the values *S*, *I*, and *R* that we calculated before this section, it is possible to predict the number of (NC_{daily}) by the equation in Eq. 16 to predict. The number of the NC_{daily} can be calculated in time *t* via SIR method in Eq. 16.

$$I_t^{SIR} + R_t^{SIR} = TC_t^{SIR}$$
(16)

where, I_{t}^{SIR} is the predicted value of I, R_{t}^{SIR} is the predicted value of R, and TC_{t}^{SIR} is the predicted value of Total Case (*TC*). The number of NC_{daily} can be calculated by subtracting the *TC* at time t = 1 from t = 0 (*time* as *day*), in other words, the increase in *TC C* gives the number of NC_{daily} in Eq. 17.

$$NC_{daily} = TC_{t+1}^{SIR} - TC_t^{SIR}$$
(17)

The observed and predicted values of NC_{daily} are shown in Figure 7. As can be seen in Figure 7, the predictions of NC_{daily} obtained by the method SIR via Eq. 17 obtained, overlap with the observed value. This overlap is quite successful. In addition to this overlap, our forecasting with the SIR during February 12-15, 2021 are shown in Figure 7. In these projections, the NC_{daily} is forecasted to add approximately 7865 patients to the infected individuals on February 15, 2021.



Figure 7. The graph of prediction models of daily new cases (NC_{daib}) via SIR method

3.5. Prediction Models of Total Death (D_{total})

Three different methods are used to model total mortality (D_{total}) in population N, namely LR, QR, and NLR, and the results of the predictive models are shown in Table 4.

The *RMSE* and *MAPE* values calculated for D_{total} from each prediction method for the train and test terms are given in Table 4. In addition, three different methods for predicting D_{total} values for the train and test terms are compared in Table 4.

Model	Model Parameters	$RMSE_R$		MAPE _R	
		Train	Test	Train	Test
LR	$\beta_{0} = 23319$	67.04	243.85	0.231	0.874
	$\beta_1 = 139.716$				
QR*	$\beta_{_{0}} = 23151$	6.5	15.35	0.0218	0.0567
	$\beta_{1} = 177.104$				
	$\beta_{2} = 1.438$				
NLR	a = 23369 b = 0.006	84.93	309.08	0.293	1.11

Table 4. Results of proposed prediction models of D_{total}

When examining the $RMSE_{D_{nucl}}$ values for train and test terms, the QR method yields the lowest value with $RMSE_{D_{nucl}} = 6.5$, 15.35, respectively. When examining the $MAPE_{D_{nucl}}$ values for the train and test terms, the QR method provides the lowest value with $MAPE_{D_{nucl}} = 0.0218$, 0.0567.

In the predictor of D_{total} , it is expected that there will be approximately similar $RMSE_{D_{total}}$, and $MAPE_{D_{total}}$ values closest to 0 for train and test terms. Thus, the QR model is the most effective model for predicting D_{total} . The QR model of D_{total} is given below in Eq. 18.

$$\hat{y} = 23151 - 177.104 \times day - 1.438 \times day^2 + \epsilon \tag{17}$$

The extent to which prediction methods predict the observed values of D_{total} and the forecasting values over the 4-day period between February 12-15, 2021 is given in Figure 8.



Figure 8. The graph of prediction models of daily new cases (D_{total})

The predictions of D_{total} obtained by the method QR are overlapped with the observations. This overlap is quite successful. In addition to this overlap, our forecasting is shown with the QR method between

February 12 and February 15, 2021 in Figure 8. In these results, it is forecasted that the number of D_{total} will be about 27510 individuals on February 15, 2021. The sum of those who recovered and those who died is the number of people who recovered from the Sars-CoV-2 virus. In this case, we need to look at what percentage of people who died from Sars-CoV-2 make up the total number of people infected. Considering this situation, the following Figure 9 shows what percentage of infected persons die every day in Türkiye.

In the period we studied, the average mortality rate of infected persons is 1.04 % percent. In other words, about one patient in 100 infected persons is dead.



Figure 9. The graph of Death data per case (%)

4. Conclusions

In this study, which we conducted to investigate the situation of Covid-19 pandemic in Turkey, modeling, and further estimations for the daily number of recovered and deceased cases are made. The method SIR, the commonly used polynomial regression method (up to 3rd order) and nonlinear regression methods that are commonly used in modeling the spread of a disease are applied and these methods are compared to determine the most successful method.

In our analysis, the QR method best succeeded in predicting susceptible individuals in the population, whereas the SIR method was best able to predict infected and removed individuals. Modeling of the estimate of the number of new cases transmitted each day, which has not previously occurred, was demonstrated in this study using the method SIR. The powerful overlap of the observed values of the daily number of cases and the values estimated by the method SIR shows that the method with the value $RMSE(NC_{daily}) = 61.718$ is a very suitable method for our sample. The number of individuals who died due to Sars-CoV-2 shows that the QR method is the best predictor of the dataset. In this study, where we made a forecasting, on February 15, 2021, in the population S = 81745522, I = 82693, R = 2508720, $NC_{daily} = 7865$, $D_{total} = 25710$, and the mortality rate of individuals exposed to the virus is calculated as 1.04%.

Evaluating the research and forecasting regarding the pandemic's course together, it is expected that the pandemic will persist, as it is calculated to be R = 69531 on February 15, 2021. It seems possible to get rid of the Sars-CoV-2 virus by taking precautions to reduce human contact with each other and by continuing vaccination effectively and rapidly.

Author Contributions: Conceptualization, E.S. and U.S.S.; methodology, E.S. and U.S.S; software, E.S.; validation, E.S. and U.S.S; formal analysis, E.S.; investigation, E.S. and U.S.S; resources, E.S. and U.S.S; data curation, E.S.;

writing—original draft preparation, E.S. and U.S.S; writing—review and editing, E.S. and U.S.S; visualization, E.S.; supervision, E.S.; project administration, E.S.; funding acquisition, E.S. and U.S.S.

Funding: This research received no external funding. Data is available in a public-open access repository, the Health Ministry of Türkiye Republic data archive in [21].

Acknowledgments: Some of the results of this study were presented at the 10th International Eurasian Conference on Mathematical Sciences and Applications (IECMSA-2021).

Conflicts of Interest: The authors declare no conflict of interest. The funders had no role in the design of the study; in the collection, analyses, or interpretation of data; in the writing of the manuscript, or in the decision to publish the results.

References

- F. Zhou *et al.*, "Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: a retrospective cohort study," *The Lancet*, vol. 395, no. 10229, pp. 1054–1062, Mar. 2020, doi: 10.1016/S0140-6736(20)30566-3.
- [2] A. L. Phelan, R. Katz, and L. O. Gostin, "The Novel Coronavirus Originating in Wuhan, China: Challenges for Global Health Governance," JAMA, vol. 323, no. 8, pp. 709–710, Feb. 2020, doi: 10.1001/jama.2020.1097.
- [3] Coronaviridae Study Group of the International Committee on Taxonomy of Viruses *et al.*, "The species Severe acute respiratory syndrome-related coronavirus: classifying 2019-nCoV and naming it SARS-CoV-2," *Nat. Microbiol.*, vol. 5, no. 4, pp. 536–544, Mar. 2020, doi: 10.1038/s41564-020-0695-z.
- [4] J. W. M. Chan *et al.*, "Short term outcome and risk factors for adverse clinical outcomes in adults with severe acute respiratory syndrome (SARS)," *Thorax*, vol. 58, no. 8, pp. 686–689, Aug. 2003, doi: 10.1136/thorax.58.8.686.
- [5] M. J. Keeling and P. Rohani, Modeling Infectious Diseases in Humans and Animals. Princeton University Press, 2011. doi: 10.1515/9781400841035.
- [6] O. Diekmann and J. A. P. Heesterbeek, Mathematical Epidemiology of Infectious Diseases: Model Building, Analysis and Interpretation. John Wiley & Sons, 2000.
- [7] F. Arroyo-Marioli, F. Bullano, S. Kucinskas, and C. Rondón-Moreno, "Tracking R of COVID-19: A new real-time estimation using the Kalman filter," PLOS ONE, vol. 16, no. 1, p. e0244474, Jan. 2021, doi: 10.1371/journal.pone.0244474.
- [8] A.-J. Valleron, "Roles of mathematical modelling in epidemiology," COMPTES RENDUS-Acad. Sci. PARIS Ser. 3, vol. 323, no. 5, pp. 429–434, 2000.
- [9] F. Brauer, "Lecture Notes in Mathematical Epidemiology".
- [10] M. Martcheva, An Introduction to Mathematical Epidemiology, vol. 61. in Texts in Applied Mathematics, vol. 61. Boston, MA: Springer US, 2015. doi: 10.1007/978-1-4899-7612-3.
- [11] N. M. Ferguson, "Mathematical prediction in infection," Medicine (Baltimore), vol. 37, no. 10, pp. 507–509, Oct. 2009, doi: 10.1016/j.mpmed.2009.07.004.
- [12] M. Abotaleb et al., "Modeling Covid-19 Infection Cases and Vaccine in 5 Countries Highly Vaccinations," Turk. J. Math. Comput. Sci., vol. 13, no. 2, pp. 403–417, Dec. 2021, doi: 10.47000/tjmcs.905508.
- [13] Z. Liao, P. Lan, Z. Liao, Y. Zhang, and S. Liu, "TW-SIR: time-window based SIR for COVID-19 forecasts," *Sci. Rep.*, vol. 10, no. 1, Art. no. 1, Dec. 2020, doi: 10.1038/s41598-020-80007-8.
- [14] İ. Demir and M. Kirisci, "Forecasting COVID-19 Disease Cases Using the SARIMA-NNAR Hybrid Model," Univers. J. Math. Appl., vol. 5, no. 1, Art. no. 1, Mar. 2022, doi: 10.32323/ujma.1010490.
- [15] A. Adiga, D. Dubhashi, B. Lewis, M. Marathe, S. Venkatramanan, and A. Vullikanti, "Mathematical Models for COVID-19 Pandemic: A Comparative Analysis," J. Indian Inst. Sci., vol. 100, no. 4, pp. 793–807, Oct. 2020, doi: 10.1007/s41745-020-00200-6.
- [16] E. Eroglu, A. A. Esenpinar, E. Bozkurt, and S. Tek, "Mathematical Modeling of Covid-19 Phenomenon; The Cases: Germany, Israel and Canada," *Fresenius Environ. Bull.*, vol. 29, no. 10, pp. 9063–9074.
- [17] B. M. Ndiaye, L. Tendeng, and D. Seck, "Analysis of the COVID-19 pandemic by SIR model and machine learning technics for forecasting." arXiv, Apr. 03, 2020. doi: 10.48550/arXiv.2004.01574.
- [18] A. A. Toda, "Susceptible-Infected-Recovered (SIR) Dynamics of COVID-19 and Economic Impact." arXiv, Mar. 26, 2020. doi: 10.48550/arXiv.2003.11221.
- [19] H. (Howie) Weiss, "The SIR model and the Foundations of Public Health," Mater. Matemàtics, pp. 1–17, 2013.
- [20] B. Barrett, "Regression Analysis: Concepts and Applications," *Technometrics*, vol. 37, no. 2, pp. 229–229, 1995, doi: 10.1080/00401706.1995.10484308.
- [21] "Genel Koronavirüs Tablosu." Accessed: Sep. 05, 2023. [Online]. Available: https://covid19.saglik.gov.tr/TR-66935/genelkoronavirus-tablosu.html
- [22] H. L. Seal, "Studies in the History of Probability and Statistics. XV: The Historical Development of the Gauss Linear Model," *Biometrika*, vol. 54, no. 1/2, pp. 1–24, 1967, doi: 10.2307/2333849.
- [23] J. D. Gergonne, "The application of the method of least squares to the interpolation of sequences," *Hist. Math.*, vol. 1, no. 4, pp. 439–447, Nov. 1974, doi: 10.1016/0315-0860(74)90034-2.
- [24] S. Bakanlığı, "Temaslı takibi, salgın yönetimi, evde hasta izlemi ve filyasyon," Bilimsel Danışma Kurulu Ank. Halk Sağlığı Genel Müdürlüğü, 2020.