

Mathematical Modeling of Infectious Disease Spread: Analyzing the SIR Model and Its Treatment Strategies

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Communicated by Nasreen Kausar

MSC 2010 Classifications: Primary 33C20; Secondary 33C65.

Keywords and phrases: Mathematical modeling, Ordinary differential equations, SIR model, Public health, Disease control strategies.

Sreelatha Devi was very thankful to the NBHM (project 02011/12/ 2020NBHM(R.P)/R&D II/7867) for their necessary support and facility.

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Abstract In this paper, we introduce spreads of infectious diseases, along with especially mathematical models based on ODEs that represent the most important areas within epidemiology in understanding a disease's dynamics and epidemics patterns predictions. One of the very basic frames developed is the SIR model. There are three classes of individuals in this model: the susceptible ones, infected with active transmission, and recovered, either immunized or not transmitting. The ODE system describes how, with time-variable illness transmission and recovery rates, the number of people in each compartment changes. Real epidemics from such models include measles, COVID-19, and influenza. Modifications of these models include the SEIR model with an exposed, non-infectious stage and the SIRV model that includes vaccination. For example, the SIR model was applied to enable the public health officers to forecast the course of the virus, peak infection rates, and effectiveness of social distancing and vaccination programs during the COVID-19 outbreak. Methods based on ODEs are important tools for managing infectious disease spread between populations and can inform policy by incorporating factors related to diseases into the model.

1 Introduction

One of the more important topics in epidemiology is that of infectious disease and transmission among humans. The context of today's global health crises, for example with COVID-19, creates a great deal of interest and urgency in the use of these models. One of the very basic, widely used, and the most rudimentary frameworks divides people into three groups, which includes susceptibility to infection, people are ill, and thus are still infectious, and then the recovered individuals who are no longer infectious [1]. The SIR model gives a mathematical description of outbreaks of diseases through its interpretation of how people flow among different compartments based on certain rates of transmission and rates of recovery. Due to its flexibility, the SIR model was extended to include the SIRV model (vaccine) and the SEIR model (exposed compartment for infected but non-infectious individuals). These models have proven to be extremely important in the fight against real-world diseases such as measles, influenza, and more recently, COVID-19 [2] [3]. The ODE-based models have really helped in forecasting the course of the COVID-19 epidemic, appraising the effectiveness of the public health interventions, and have served information for decision support. To study the infectious disease dynamics and inform design for the public health management strategies, this paper shall survey the SIR model along with its variants. All these models give valuable insights into the pathways of transmission, the role of immunity, and the possible effects of vaccination and other preventive measures [4] [5] .

The rapid progress in technology and theory has dramatically enhanced our arsenal in

the fight against epidemics, and we are getting better on it. The global surveillance network is growing under an intensive effort at the worldwide level. We can now produce effective vaccines and antiviral drugs and knowledge goes deep in details such as the molecular structure of a variety of viruses [6]. It is developing a tremendous and intense research for the design of better drugs and vaccines. But research warns that a new pandemic the worst fear is of the influenza-type is sooner or later on the horizon. The key question(s) is not if but when it will arise, how it is going to spread, how fatal it will be, who should receive the vaccine when not everyone can, how probable are several waves of re-emergence and what kind of intervention may be used in order to prevent the spread. Unfortunately, we still don't have robust answers with all the progress [7]. Since epidemics and pandemics are concerns to everyone around; researchers from all over the globe are putting their efforts to combat the disease. Mathematicians also contributed to medical sciences and provided numerous mathematical models of epidemic diseases. These models provide information related to the forthcoming behavior of the disease. They also use these models to predict when a particular disease is at its peak and guess when the disease will die off. The SIR epidemic model has been preferred over other compartmental epidemic models because the available data are limited [8].

This paper is an investigation into the beautiful legacy, merging mathematical rigor and epidemiological insight into navigating the realm of infectious disease dynamics through differential equations toward untangling complex threads in transmission, immunity, and intervention [9]. The population density, the vaccination rate, and the disease parameters make the model rich, bringing it nearer to the intricacies of reality. This study holds a great significance both theoretically and practically for measures of public health. It fills the gap between mathematical abstractions and real challenges; through it, one can seek a finer view of the course of disease outbreaks and how interventions work [10]

2 Mathematical Modeling of SIR

A very simple yet powerful mathematical method to describe the dynamics of infectious diseases in a population is the SIR model. The population is split up into three groups, that is,

- (i) S(t): People susceptible to being infected with the disease.
- (ii) I(t): Infected persons who can transmit the infection to susceptible persons.
- (iii) R(t): Recovered subjects, which have successfully immunized and cannot become infected.

A system of ordinary differential equations (ODEs) describing how the number of people in each group changes over time controls transitions between these compartments. The two important factors determining such changes are: say the transmission rate, β determines how often susceptible individuals contract the infection, and the rate at which infected individuals recover and transition to the recovered compartment is determined by the recovery rate, γ . Mathematical modeling and simulation enable fast assessment. A problem with the cost of data collection being prohibitively expensive or a large number of experimental conditions to test is often approached through simulation. Quite a massive number of approaches have been proposed in looking at the problem from different angles over the years. SIR-type models have been further extended to include demographics like age distributions, mortality and spatial dependence of the spread in account of diffusion and migration effects as well as genetic mutations in the interacting populations, thereby enhancing their realism [11] [12]. Many scientists have investigated the nature and diagnosis of this disease; these investigations are either experimental or theoretical. The language of mathematics is widely used for describing a number of naturally occurring phenomena. This is because mathematical modeling, together with theoretical analysis enables to provide a qualitative description or understanding of systems as they are. It can also be used to make inferences about the model parameters from which one anticipates dynamical behavior from a given system, or it may facilitate the comprehension of which parameters are most important for the behavior. No rigorous analytical solutions have been reported, to our knowledge, for non-steady-state glucose, insulin and β -cell mass concentration [13].

2.1 ODE System for the SIR Framework

The SIR model is expressed mathematically by the following set of equations:

1. Susceptible Population (S):

In a sufficiently large population, the number of infected individuals at the initial stages of the infection is well below the population size. Under certain conditions, it may stay small in comparison to the number of susceptible individuals remaining [14].

$$\frac{dS}{dt} = -\beta SI$$

The rate of change in the susceptible population is represented by this equation. $S(t)$ decreases as vulnerable individuals migrate into the infected compartment after becoming infected. The number of susceptible people and the number of infected people are directly correlated with the rate of new infections.

2. Infected Population (I):

$$\frac{dI}{dt} = \beta SI - \gamma I$$

As vulnerable people contract the infection, the number of sick people rises, and as infected people recover, the infected population falls. The infection rate is represented by the first component

$$\beta SI$$

, while the recovery rate is represented by the second term, γSI . [15].

3. Recovered Population (R):

$$\frac{dR}{dt} = \gamma I$$

When infected individuals recover at a rate γ , the recovered population increases. People are thought to be immune and incapable of contracting the infection again once they enter the recovered compartment [16].

2.2 Parameters and Assumptions

The probability of a disease spreading each interaction between a susceptible person and an infected person is represented by the transmission rate β . The recovery rate γ , indicates how quickly an infected person recovers and develops immunity. One important measure that comes from the model is the basic reproduction number R_0 ,

$$R_0 = \frac{\beta}{\gamma},$$

. In a population that is totally susceptible, R_0 is the average number of new infections that one infected person causes. The sickness spreads if $R_0 > 1$, and it dies out if

$$R_0 < 1$$

[17].

2.3 Dynamics of the SIR Model

This sickness takes toll readily at the beginning thus a ramp up of infected persons population, after the large group of susceptibles coupled by small recoveries ones. Although there

are fewer of this group, but these who continue to rise with large infectee who recovered, consequently decreasing overall disease rates, the infection rate slowly decreases and the patients uptake is taken on an increase in scale; subsequently, it tends to stop absolutely at the falling point of graph [18].

2.4 Extensions of the SIR Model

1. SIRV Model (Vaccination):

$V(t)$, can be added in order to include vaccination on the model. Vaccinated individuals bypass the infected category and move directly from their susceptible category to their respective recovered category [19]. The corresponding equations for the SIRV model can be presented as:

$$\frac{dS}{dt} = -\beta SI - \nu S$$

$$\frac{dI}{dt} = \beta SI - \gamma I$$

$$\frac{dR}{dt} = \gamma I + \nu S$$

Where ν is the vaccination rate.

2. SEIR Model (Latency Period):

An exposed compartment $E(t)$ is added to the SEIR model to include those infected but not yet infectious, accounting for the latent period. The model becomes the number of susceptible individuals decreases as more people become infected and then recover, reducing the overall transmission of the disease [20]. As the population acquires immunity through natural infection or recovery, eventually the infection rate declines, the number of infected increases and then declines.

$$\frac{dS}{dt} = -\beta SI$$

$$\frac{dS}{dt} = \beta SI - \sigma E$$

$$\frac{dI}{dt} = \sigma E - \gamma I$$

$$\frac{dR}{dt} = \gamma I$$

Where σ is the rate at which exposed individuals become infectious.

2.5 Application to Epidemic Prediction and Control

The SIR, among its extended models, proves to be a useful tool in predicting infections' spreading and evaluating intervention strategies, such as social distance, vaccination programs, and quarantines. Using SIR and other such models, researchers and public health professionals can:

- (i) Obtain the total number of infected people across time and the peak of infection.
- (ii) Estimate the duration of the epidemic and the eventual proportion of the population that will either be immune or infected.
- (iii) The efficacy of the control measures may be comprehended by including a vaccine or modifying other parameters, such as the transmission rate (η).

Several modifications of the SIR model were used in the time of the COVID-19 outbreak

to forecast the trends of the outbreak, estimate the success of the taken measures, and allocate the healthcare resources properly. Such models could be fitted appropriately to the actual data that were available at that time so that informed decisions could be made based on wise choices [21].

2.6 Solving The Mathematical Modeling

We use the following system of equations to solve the system of ordinary differential equations for the SIR model:

$$\begin{aligned}\frac{dS}{dt} &= -\beta SI \\ \frac{dI}{dt} &= \beta SI - \gamma I \\ \frac{dR}{dt} &= \gamma I\end{aligned}$$

Since this system of equations is nonlinear, it can be quite often difficult to find a solution in the analytical way. However, we can easily solve this problem numerically using very simple methods like Runge-Kutta or Euler.

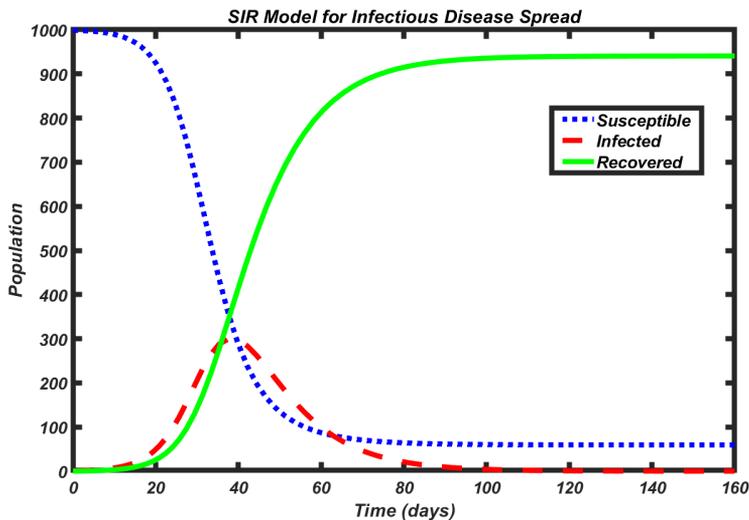


Figure 1. SIR model for infectious Disease Spread

Steps for Numerical Solution:

Step 1. We consider initial values for $S(0)$, $I(0)$, and $R(0)$.

Step 2. We consider the rate for recovery as γ and for transmission as β .

Step 3. Use a numerical method, such as Runge-Kutta 4th order, to solve a system of differential equations.

We set up and solve an example.

Parameters to be assumed for simulation are $S(0) = 0.99$, $R(0) = 0$, $I(0) = 0.01$, $\beta = 0.3$, $\gamma = 0.1$

By putting these parameters in the numerical solution we get the graph as shown in Figure 1 :

The graph in Figure 1 represents the numerical solution of the SIR model over 160 days of the given compartments:

Susceptible (S): Basically, everyone is susceptible at the beginning. As the disease spreads, this number continues to decline because people keep moving into the infected compartment.

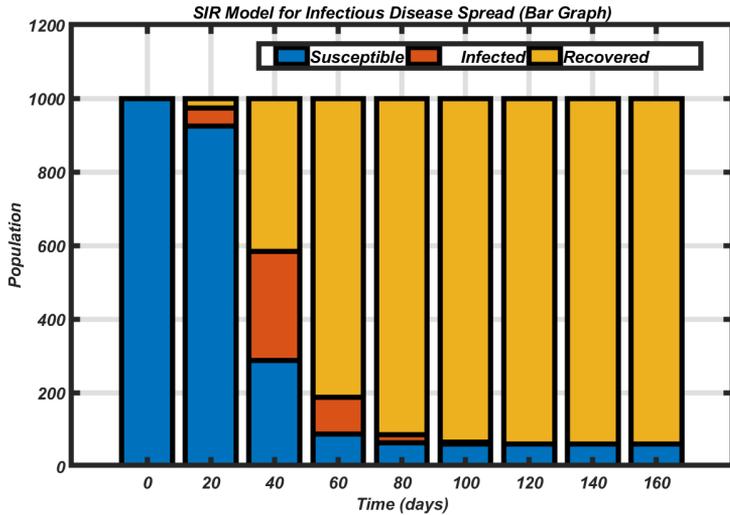


Figure 2. Bar graph of SIR model for infectious Disease Spread

Infected (I): In the beginning, the infected population grows because more and more susceptible individuals become infected. Then, at the top of the curve, the infected number starts to fall as recovery increases and fewer individuals remain susceptible.

Recovered (R): More and more infected individuals continue recovering, getting into this category at any point in time. This simulation illustrates typical dynamics of an infectious disease in a population. The number of affected people reaches its highest during the peak of the infection, after which there is a fall as the disease eventually goes extinct and a sizable section of the population recovers and becomes immune [22] [23] [24]. For better understanding, SIR model for infectious disease spread is shown in Bar graph in Figure 2.

2.7 Model Analysis of the SIR Model

The SIR model is the set of ordinary differential equations representing the dynamics that the spread of infectious disease follows in a population. Such insights obtained from the study of the solution of the system are long overdue in revealing critical points in the disease’s behavior, such as infection peaks, recovery rates, and overall epidemic control. Using the numerical solution we had obtained, let us outline the analysis.

1. Primary Phase: Disease Acquisition

At the start of the outbreak, nearly all population members are in the susceptible compartment as represented by 99% of the example. This heavy proportion of susceptibles within the population leads to fast development of infected population out of a small initial population, such as 1% in this case. Infection rate, depends on the transmission rate along with the interaction between a susceptible and an infected member. In our model, when $\beta = 0.3$, this means an individual who is sick could infect 30% of the contacts daily. Consequently, with a high-probability occurrence of such interactions between the susceptible people and the infected ones within the initial days of any outbreak, infections occurs [25].

2. Growth of the Epidemic: Infection Peaks

The number of infections rises quickly at initially, but gradually peaks and the number of new infections begins to fall. This peak denotes the maximum number of infected individuals at any one moment. The peak happens because fewer new infections result from an increasing number of infected individuals decreasing the vulnerable population. The pandemic begins to slow down when there are fewer susceptible people available to spread the disease. The growth

of the epidemic can be explained by the basic reproduction number,

$$R_0 = \frac{\beta}{\gamma} = \frac{0.3}{0.1} = 3$$

In a fully susceptible population, this means that each infected individual is predicted to infect three more people. The sickness spreads when $R_0 > 1$, and the pandemic peaks and starts to decline as R_0 approaches 1. When the proportion of vulnerable people falls below a threshold that prevents the disease from spreading as quickly, the infection peaks. When more people develop immunity, the illness begins to decline [26].

3.Recovery and Immunity

Those who are infected relocate into the recovered compartment when they become better. The diseased population declines over time, but the restored population increases dramatically. 10% of the sick population recovers on average per day, according to the recovery rate $\gamma = 0.1$. Over time, the 'herd immunity' increases since it is believed that the recovered persons are immune to the disease or are no longer able to spread it. Because fewer vulnerable people are available for transmission due to the rise in the recovered population, the pandemic is coming to an end. The pandemic eventually comes to an end as the recovered population grows and the infected population decreases. One important factor in stopping the disease's spread is herd immunity [27].

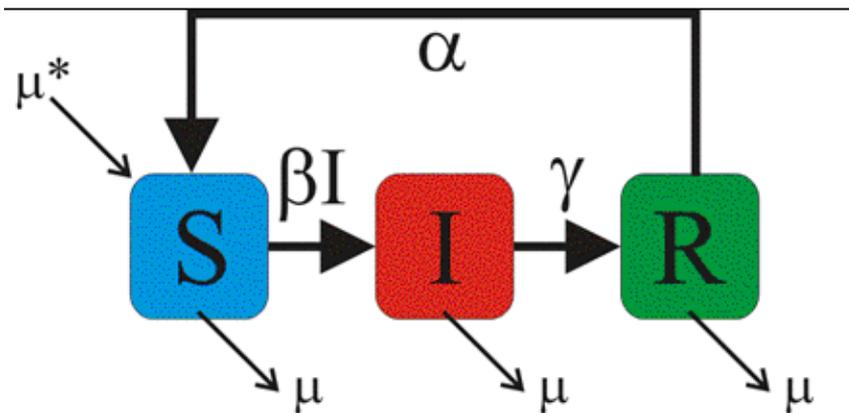


Figure 3. Compartment model of transmission rate (β) and recovery rate (η)

4.Epidemic Resolution

The susceptible population eventually declines to a level where the epidemic cannot continue. As a result, the number of people sick drops to zero because the disease can no longer spread. An estimate of the total population proportion infected by the disease during the epidemic can be obtained from the final size of the recovered population. After the disease has had its full course and the infected population has completely disappeared, there is a sizable recovered population and a much smaller susceptible group remaining. When social separation or vaccinations are not used, an epidemic will naturally come to this end [28][29]. Comparison of transmission rate (β) and recovery rate (η) is shown in Figure 3.

3 Model Analysis's Key Findings

1.Disease Control

The fundamental reproduction number,

$$R_0 = \frac{\beta}{\gamma}$$

, is essential for determining the infectiousness of a disease [4]. The sickness spreads throughout the population if $R_0 > 1$. Reducing R_0 below 1 is the goal of interventions in order to contain an

outbreak. This can be achieved by decreasing the rate of transmission β by implementing strategies such as mask-wearing, social distancing, or quarantine, and increasing the rate of recovery γ by implementing efficient medical interventions or shortening the period of infectiousness.

2. Peak Infection and Healthcare Capacity

The model illustrates how crucial the peak infection is. Overcrowding of infectious individuals might overload healthcare services. Lowering the peak and spreading infections across a longer time span can be achieved by flattening the curve by lowering the transmission rate through public health initiatives.

3. Herd immunity

As more people get better and develop immunity, the disease’s spread diminishes. The model emphasizes the function of herd immunity, which is the moment at which a sizable proportion of the population possesses immunity, essentially stopping the illness from spreading even in the absence of universal vaccination.

4. Role of vaccination

By transferring people straight to the recovered (immune) compartment, vaccination reduces the susceptible population, according to an extension of the SIR model. This can stop major epidemics and quicken the process of herd immunity. Immunization efficiently reduces the number that is susceptible, resulting in an earlier control of the epidemic [30] [31] [32].

3.1 Equilibrium Points of the SIR model

Equilibrium points in mathematical modeling are those conditions in which the system remains constant across time. Equilibrium for the SIR model is reached when all three compartments’ rates of change $S(t)$, $I(t)$, and $R(t)$ are zero.

$\frac{dS}{dt} = 0, \frac{dI}{dt} = 0, \frac{dR}{dt} = 0$ Let’s analyse the equilibrium points by examining the SIR model’s differential equations:[33] [34]

$$\begin{aligned} \frac{dS}{dt} &= -\beta SI \\ \frac{dI}{dt} &= \beta SI - \gamma I \\ \frac{dR}{dt} &= \gamma I \end{aligned}$$

Step 1: Equilibrium Conditions

There must be zero rates of change for equilibrium to exist.

(i). $\gamma I = 0$ from $\frac{dR}{dt} = \gamma I$. This suggests that $I = 0$ or $\gamma = 0$, which is implausible given that γ denotes a non-zero recovery rate. Therefore, $I = 0$ implies that there cannot be any infected persons for equilibrium to exist.

(ii). Utilizing $\frac{dS}{dt} = -\beta SI$ Since we know that $I = 0$ at equilibrium, this equation becomes:
 $\frac{dS}{dt} = -\beta S \cdot 0 = 0$

This equation is always satisfied when $I = 0$. Thus, the value of S is not directly constrained by this equation.

(iii). From $\frac{dI}{dt} = \beta SI - \gamma I$, Substituting $I = 0$ into this equation, we get:

$\frac{dI}{dt} = \beta S \cdot 0 - \gamma \cdot 0 = 0$ This equation is also automatically satisfied when $I = 0$.

Step 2: Identifying Equilibrium Points

When the infected population (I) equals zero, the equilibrium sites are discovered. There are two main scenarios, or states of equilibrium:

1. Equilibrium without disease

When the population reaches the disease-free equilibrium, no one is infected and everyone is either susceptible or recovered. In this case $I = 0$ (there is no infection), If there has been no recovery, $S = N$ (the entire population is still vulnerable), and $R = 0$ denotes the absence of infections, meaning that no one has recovered. The following yields the disease-free equilibrium:

$(S, I, R) = (N, 0, 0)$

This indicates that there isn’t an epidemic outbreak and that the whole population is still vulnerable. If $R_0 = \frac{\beta}{\gamma} < 1$, the basic reproduction number, this equilibrium is stable. In this instance, the disease cannot spread since a small number of infections will disappear before they have a

chance to trigger a significant outbreak.

2. Endemic Equilibrium

An endemic equilibrium occurs when the disease is present in the population for an extended period, but the number of cases is low or stable. A portion of the population has recovered, and a portion is still vulnerable. In an endemic equilibrium, $I = 0$ indicates that there are no new infections, $S = S^*$ indicates that a portion of the population is still vulnerable, as indicated by R_0 , and $R = N - S^*$ indicates that the remainder of the population has recovered.

We use the requirement that the virus not spread at equilibrium to get the value of $\frac{dI}{dt} = 0 = \beta S^* I - \gamma I$

We rewrite the equation to determine the equilibrium value for S^* since $I = 0$,

$S^* = \frac{\gamma}{\beta} = \frac{1}{R_0}$ This indicates that the fundamental reproduction number R_0 and the number of susceptible people at equilibrium are inversely related. The endemic equilibrium is therefore provided by:

$(S^*, I^*, R^*) = (1/R_0, 0, N - 1/R_0)$ If $R_0 > 1$, there is an endemic equilibrium that is stable. This suggests that the disease will continue in the population, albeit at an endemic level that is controllable, if it is sufficiently contagious (i.e., R_0 exceeds 1). $R_0 < 1$ indicates the presence of Disease-Free Equilibrium ($S = N, I = 0, R = 0$). Infections swiftly disappear, and the illness does not spread. When $R_0 > 1$, endemic equilibrium ($S^* = 1/R_0, I = 0, R = N - S^*$) occurs. There are still some susceptible people in the population as the disease stabilizes.

3.2 Observation

The SIR model's equilibrium point analysis offers important insights into how infectious illnesses behave in a community. When the fundamental reproduction number $R_0 < 1$, the infection cannot spread and eventually dies out naturally, a state known as the disease-free equilibrium. In the context of disease control operations, this is a good result because it suggests that an outbreak will not spread throughout the community. On the other hand, when $R_0 > 1$, the endemic equilibrium occurs, indicating that the disease continues to exist in the community and that a consistent percentage of people are still susceptible and have recovered. The situation of an endemic disease, when the infection is present but under control, is reflected in this equilibrium. These balance factors are important when planning for public health. The goal of measures to lower R_0 is to move the system closer to the equilibrium when there is no sickness, which can be achieved by therapy, social distancing, or immunization. These equilibria's stability emphasizes how crucial it is to keep R_0 below 1 in order to stop the disease from spreading and to encourage its eventual eradication.

4 Results and Discussion of the SIR Model

Important insights into the dynamics of disease transmission, such as the infection peak, recovery trends, and overall epidemic control, are offered by the numerical simulation of the SIR model. Here, we go over the outcomes of solving the differential equation system and consider how these discoveries might be applied to actual pandemic situations.

Time-series data for the number of susceptible $S(t)$, infected $I(t)$, and recovered $R(t)$ persons over time are produced by the SIR model. The outcomes can be categorized into multiple crucial stages:

a. Early Infection Growth

At the beginning of the simulation, almost all of the population $S(0) \approx N$ is susceptible, only a small portion is infected $I(0)$, and there are no recoveries $R(0) = 0$. Because there are many susceptible people and the transmission rate (β) is large, the number of infected persons $I(t)$ increases quickly. The disease spreads rapidly as infected persons contact the vulnerable populace. Because most people are susceptible, there is an initial high rate of infection, which causes the diseased population to grow exponentially.

b. Maximum Infection

The susceptible population $S(t)$ starts to decrease as the number of sick people rises. The disease's transmission rate slows down with time as fewer people become susceptible

to it. The number of infected people at any particular time is the maximum number that the infected population eventually achieves a peak. This represents the greatest strain on medical systems, making it an important factor to consider while managing healthcare resources. It is observed that when the susceptible population drops below a crucial threshold, the incidence of new infections slows and the number of active infections starts to reduce, marking the peak of the infection.

c. Decline in Infections and Rise in Recoveries

More people shift to the recovered compartment as the number of new infections starts to decline after the infection peak. As more people heal, the number of afflicted persons decreases. When afflicted people recover and develop immunity to the illness, the recovered population $R(t)$ increases gradually. This slows the disease’s spread even more and strengthens the herd immunity. It is observed that as the pandemic progresses, fewer people become infected and the majority of people either recover or maintain their immunity.

d. End of the Epidemic

As the majority of the population either recovers or stays susceptible but is no longer exposed to diseased individuals, the number of infected individuals eventually approaches zero. The population’s final state reveals a high percentage of recovered people and a comparatively low number of susceptible people. It is observed that when there are no more cases of infection, the epidemic ends. The population achieves a stable equilibrium when there are no longer any new infections, putting the existing susceptible individuals at danger. Population distribution across various compartments are shown in Table 1.

Time (days)	Susceptible	Infected	Recovered
0.0	999.0	1.0	1.0
20.0	895.2	76.5	28.3
40.0	765.4	150.8	83.8
60.0	650.1	200.3	149.6
80.0	560.3	215.7	224.0
100.0	499.5	202.5	298.0
120.0	458.8	173.4	367.8
140.0	429.9	138.2	431.9
160.0	407.3	103.6	489.1

Table 1. shows the population distribution across the compartments at each specified time point.

5 Strategies and Treatments for the SIR Model

A number of intervention and treatment techniques can be used in conjunction with the SIR (Susceptible-Infected-Recovered) paradigm to stop the spread of infectious diseases. In order to move the epidemic toward a controllable or eliminated state, these tactics seek to lower the susceptibility population, raise the recovery rate, or decrease the transmission rate β . The primary methods of therapy and control are listed below:

(I).Vaccination Effect:

By transferring susceptible individuals from the susceptible compartment (S) to the recovered compartment (R) without requiring them to go through the infected condition, vaccination directly lowers the number of susceptible individuals. The objective is to decrease the fundamental reproduction number R_0 to less than 1 and to attain herd immunity.

(II). Social Distancing and Quarantine Effect:

These strategies lessen the frequency of interaction between susceptible and sick people, which lowers the rate of transmission β . Reducing interactions slows down the disease’s spread and postpones the infection peak. Impact on Mathematics is Reducing β shortens the term β , hence slowing the pace of new infections. By doing so, the epidemic curve is flattened, the peak is postponed, and the strain on healthcare systems is distributed across a longer time frame. The objective is to lower the infection rate and avoid an unexpected spike in cases that would over-

burden the healthcare system.

(III). Antiviral Treatments and Medical Care Impact: Antiviral medications or therapies that hasten recovery shorten the duration of an individual's infection, hence accelerating the rate of recovery γ . Improved medical care can also shorten the length of an illness and lower mortality. Impact of Mathematics is by increasing γ , the number of active infections is decreased overall by increasing the flow of persons from the infected compartment to the recovered compartment. Moreover, this reduces the fundamental reproduction number, or $R_0 = \frac{\beta}{\gamma}$. The objective is to decrease the number of reproductions, shorten the infectiousness period, and speed up healing.

(IV). Isolation of Infected persons Effect:

By keeping infected persons far from the susceptible population, isolation considerably lowers the risk of transmission. During the infectious phase, infected individuals are isolated from the general community. Mathematical Impact shows Isolation lowers β because it decreases the effective contact rate between susceptible and infected individuals. It safeguards individuals who are still vulnerable and slows the disease's spread.

(V). Contact Tracing Effect:

The disease can be prevented from spreading by identifying and separating those who have come into touch with an affected person. Before they may infect others, prospective new cases can be promptly isolated with the aid of contact tracing. Mathematical Impact predicts by reducing the number of encounters between susceptible and infected individuals, contact tracing effectively lowers β . For people who are more susceptible to illness, it serves as a focused type of quarantine.

(VI). Reducing Susceptibility with Public Health Interventions Effect:

Generally speaking, public health initiatives like mask use, better cleanliness (hand washing, sanitization), and encouraging healthy lifestyles can lessen people's vulnerability to illness. Mathematical Impact shows by reducing the likelihood that a vulnerable person would contract the infection from an infected person, these interventions indirectly diminish β . Reduce the overall rate of transmission and stop the spread of new illnesses.

6 Conclusion

A fundamental framework for comprehending the transmission of infectious diseases and the efficacy of different intervention tactics is offered by the SIR model. The model illustrates the main dynamics of disease transmission, including the significance of regulating the basic reproduction number R_0 and the effects of various public health initiatives, through the examination of susceptible, infected, and recovered populations. To manage infection peaks, prevent transmission, and develop herd immunity, vaccination, social distancing, isolation, and improved medical care are essential. Through the modification of parameters like the recovery rate γ and transmission rate β , the SIR model illustrates how prompt and efficient actions can lessen healthcare costs, avert major outbreaks, and eventually bring an epidemic under control or eradicate it. The model's simplicity makes it an effective guiding tool. Because of its simplicity, the model can be used to inform public health policies and enhance readiness for future outbreaks. Although the SIR model oversimplifies real-world complexity, it provides insightful information on managing epidemics, assisting policymakers in putting into practice efficient public health strategies. Based on the SIR model, the mathematical modeling described in this work provides important insights into the dynamics of illness transmission and control measures. Through the process of solving the system of ordinary differential equations, we were able to comprehend the spread of infection and the significance of important variables like rates of transmission and recovery.

NOMENCLATURE

Symbol	Description
$S(t)$	Number of susceptible individuals at time t
$I(t)$	Number of infected individuals at time t
$R(t)$	Number of recovered (or removed) individuals at time t
β	Transmission rate (rate of disease spread from infected to susceptible)
γ	Recovery rate (rate at which infected individuals recover)
N	Total population, $N = S(t) + I(t) + R(t)$
$\frac{dS}{dt}$	Rate of change of the susceptible population with respect to time
$\frac{dI}{dt}$	Rate of change of the infected population with respect to time
$\frac{dR}{dt}$	Rate of change of the recovered population with respect to time
R_0	Basic reproduction number, $R_0 = \frac{\beta}{\gamma}$
t	Time variable

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Received: 2025-01-01.

Accepted: 2025-05-29.