# Stability and Bifurcation Analysis of a Mathematical Modeling of Measles Incorporating Vitamin A Supplement 

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#### Abstract

Measles is transmissible disease that is common among children. The death caused by measles among children of five years and below is alarming in spite of the safe and effective vaccine. It has been discovered that Vitamin A Deficiency (VAD) in children increases their chances of measles infection. In this paper, the mathematical model of measles incorporating Vitamin A supplement as treatment was formulated and analyzed. The equilibrium points are obtained and analyzed for stability. Bifurcation and sensitivity analyses were carried out to gain further insight into the spread and control of measles. The stability analysis revealed that Disease Free Equilibrium $(D F E)$ is stable if $R_{0}<1$. The bifurcation analysis revealed forward bifurcation while the sensitivity analysis shows the most sensitive parameters of the model that are responsible for the spread and control of the diseases. The effect of sensitive parameters on Basic Reproduction Number, $R_{0}$ were presented graphically. Vaccination, recovery and Vitamin A supplement rates have been shown from the graphical presentation as the important parameter that will eradicate the measles from the population while contact and loss of immunity rates have shown that measles will persist in the population. People should be sensitized on the danger of living with infected persons. Government should do more in routine immunization and administration of Vitamin A Supplement.


Keywords: Stability, equilibrium, measles, trace-determinant, Lyapunov function

## 1. Introduction

Measles has become a serious public health challenge to the little children globally. It is transmitted through the fluid from the nose, mouth or throat of infected persons (Atkinson, 2011). Research shows that $90 \%$ of susceptible individuals who have contact with infected person are likely to be infected (Leuridan et al., 2012). High fever, a runny nose and red eyes are the signs and symptoms of measles. Blindness, severe diarrhoea and pneumonia are many at times are measles complications. In 2018 over 140,000 people died from measles (WHO, 2019). The Nigerian Centre for Disease Control (NCDC) in the beginning of 2019 recorded over

22,000 suspected cases of measles. Over $50 \%$ happen in the terrorist- affected Borno State as well as in 37 internally displaced persons (IDP) camps, this resulted to 13,753 suspected cases and 72 deaths (NCDC, 2019). Deficiency of Vitamin A has been identified as the major factor responsible for the death and other health complications resulting from measles. Almost 200 million children have been affected by its adverse health challenges, including death. Hence, WHO recommends that Vitamin A supplement be given to children from 6 months to 5 years of age (Mayo-Wilson et al., 2011). They also recommend that, Vitamin A supplement be given two times in 24 hours to the children with measles (Imdad et al., 2017).

Abubakar, et al. (2012), formulate the model of Measles dynamics using SIR model and obtained the equilibrium points. They used Belman and cook theory to carried out the stability analysis of endemic equilibrium. Abubakar, et al. (2013), analyzed the stability of endemic equilibrium using Hopf;s bifurcation theory. In Somma et al., (2015), they modified the existing Maternally-Immune Susceptible Infected Recovered (MSIR) model by incorporating vaccination and disease-induce death rates. They obtained the equilibrium points and analyzed same for stabilities. Somma et al. (2017) used Lyapunov functions to carry out the stability analysis. Fred et al. (2014) studied mathematical modeling of measles with vaccination as control measure but did not include Maternally-derived -immunity in their model. Okyere-Siabouh and Adetunde (2013) formulated a mathematical model of measles using Susceptible-Exposed-Infected-Recovered (SEIR) epidemiological model. In their study they assumed that everybody is born into the susceptible class. Peter et al. (2018), Sinha et al. (2019) and Garba et al. (2020) also proposed a mathematical model of measles disease dynamics with emphasis on vaccination as means of control. Their models did not consider Vitamin A supplement.

In this paper, we obtain the Disease Free Equilibrium (DFE) and Endemic Equilibrium (EE) in terms of force of infection. The $R_{0}$ was computed and used in the stability, bifurcation and sensitivity analyses. We use trace-determinant method and Lyapunov function for the stability analyses. The sensitive parameters and $R_{0}$ are presented graphically.

## 2 Materials and Methods

### 2.1 Model Formulation

The model equation is divided into four compartments based on the epidemiological status of individuals: with $N(t)$ total population, Maternally-Derive-Immunity $M(t)$, Susceptible $S(t)$, Infected $I(t)$ and Recovered/Immune $R(t)$, where $t$ is time. The new babies are born into $M$ class and susceptible class at constant rate $\Lambda . \Lambda \beta$ is the proportion of children born with passive immunity while $\Lambda(1-\beta)$ is the proportion of children born without passive immunity. The new babies loss their immunity after some time at a rate $\theta$ and move to susceptible class. The susceptible individuals become infected with measles at a contact rate $\alpha$. The susceptible class is vaccinated at a rate $v$ and move to recovered/immuned class. The infected individuals are treated and given Vitamin A supplement at recovery rate $\gamma$ and Vitamin A supplement rate $\tau$, respectively and moved to recovered/immune class. The death rate due to disease is $\delta$ and the natural death rate of the entire population is $\mu$. The schematic diagram and model equations for the measles transmission as discuss in this paper is presented in Fig 2.1:

### 2.2 Model Equations

$$
\begin{align*}
& M^{\prime}=\Lambda \beta-(\theta+\mu) M  \tag{2.1}\\
& S^{\prime}=\Lambda(1-\beta)-\frac{\alpha S I}{N}+\theta M-(\mu+v) S \tag{2.2}
\end{align*}
$$

$$
\begin{align*}
& I^{\prime}=\frac{\alpha S I}{N}-(\gamma+\tau+\mu+\delta) I  \tag{2.3}\\
& R^{\prime}=(\gamma+\tau) I-\mu R+v S \tag{2.4}
\end{align*}
$$

Where,

$$
\begin{equation*}
N=M+S+I+R \tag{2.5}
\end{equation*}
$$



Figure 2.1: Schematic Diagram of the Model

### 2.3 Existence of Equilibrium Points

At equilibrium $M^{\prime}=S^{\prime}=I^{\prime}=R^{\prime}=0$
Let
$E^{*}=\left(M^{*}, S^{*}, I^{*}, R^{*}\right)=(M, S, I, R)$
$\Lambda \beta-A_{1} M^{*}=0$
$\Lambda(1-\beta)-\frac{\alpha S^{*} I^{*}}{N^{*}}+\theta M^{*}-A_{2} S^{*}=0$
$\frac{\alpha S^{*} I^{*}}{N^{*}}-A_{3} I^{*}=0$
$A_{4} I^{*}+v S^{*}-\mu R^{*}=0$
Where, $A_{1}=(\theta+\mu), A_{2}=(\mu+v), A_{3}=(\gamma+\tau+\mu+\delta), A_{4}=(\gamma+\tau)$

Table 1: Description of Variables and Parameters of the Model

| Variables/Parameter | Description |
| :---: | :--- |
| $\mathbf{N}$ | Total Population |
| $M$ | Maternally-Derived -Immunity |
| $S$ | Susceptible |
| $I$ | Infected |
| $R$ | Recovered/Immune |
| $\Lambda$ | Recruitment rate |
| $\beta$ | Birth with Temporary Passive immunity |
| $\theta$ | Loss of Immunity Rate |
| $\alpha$ | Contact Rate |
| $\delta$ | Death Rate due to Disease |
| $\gamma$ | Recovery Rate |
| $\tau$ | Vitamin A supplement Rate |
| $\mu$ | Natural Death Rate |
| $\nu$ | Vaccination Rate |

From (2.10)
$I^{*}=0$
or
$\left(\frac{\alpha S^{*}}{N^{*}}-A_{3}\right)=0$
It is shown from (2.12) and (2.13) that there exist two equilibria; (2.12) is the Disease- Free Equilibrium (DFE) while (2.13) is the Endemic Equilibrium (EE).
Let,
$E^{0}=\left(M^{0}, S^{0}, I^{0}, R^{0}\right)=(M, S, I, R)$
be the DFE points
Substituting (2.12) into (2.8) to (2.11) and solve simultaneously gives the DFE:
$\left(M^{0}, S^{0}, I^{0}, R^{0}\right)=\left(\frac{\Lambda \beta}{A_{1}}, \frac{A_{6}}{A_{1} A_{2}}, 0, \frac{v A_{6}}{A_{1} A_{2} \mu}\right)$
Where, $A_{5}=\Lambda(1-\beta), A_{6}=A_{1} A_{5}+\Lambda \beta \theta$
At DFE
$\left.\begin{array}{l}N^{0}=M^{0}+S^{0}+I^{0}+R^{0} \\ N^{0}=\frac{A_{7}}{A_{1} \mu}\end{array}\right\}$
Where, $A_{7}=\Lambda \beta \mu+A_{6}$

### 2.4 The Basic Reproduction Number $R_{0}$.

Basic Reproduction Number, is the largest eigenvalue or spectral radius of $F V^{-1}$ is the basic reproduction number of the model. The approach of (Driessche and Watmough, 2002) will be used to compute the $R_{0}$.
$F=\frac{\alpha \mu A_{6}}{A_{2} A_{7}}$
$V=A_{3}$
$V^{-1}=\frac{1}{A_{3}}$
$F V^{-1}=\frac{\alpha \mu A_{6}}{A_{2} A_{3} A_{7}}$
Hence,

$$
\begin{equation*}
R_{0}=\frac{\alpha \mu A_{6}}{A_{2} A_{3} A_{7}} \tag{2.21}
\end{equation*}
$$

### 2.5 Stability Analysis of the Model

Theorem 2. 1: The DFE point $E^{0}$ of the model is Locally Stable if $R_{0}<1$.
Proof: The trace-determinant approach of will be implored to prove the above theorem. The $E^{0}$ is locally stable if trace of the Jacobian Matrix is negative and determinant of the same matrix is positive.
Therefore the Jacobian of the model at $E^{0}$ is given as
$J\left(E^{0}\right)=\left[\begin{array}{cccc}-A_{1} & 0 & 0 & 0 \\ \theta & -A_{2} & -A_{8} & 0 \\ 0 & 0 & A_{8}-A_{3} & 0 \\ 0 & v & A_{4} & -\mu\end{array}\right]$
Where, $A_{8}=\frac{\alpha \mu A_{6}}{A_{2} A_{7}}$
$\operatorname{tr} J\left(E^{0}\right)=\left(-A_{1}\right)+\left(-A_{2}\right)+\left(A_{8}-A_{3}\right)+(-\mu)$
It is shown from (2.23) that the trace of $J\left(E^{0}\right)<0$ if $\left(A_{8}-A_{3}\right)<0$
This implies that
$\left(A_{8}-A_{3}\right)<0$
Further simplification of (2.24) gives

$$
\frac{\alpha \mu A_{6}}{A_{2} A_{3} A_{7}}<1
$$

The Left Hand Side (LHS) of (2.25) is equivalent to (2.21). Hence,
$R_{0}<1$
$\operatorname{det} J\left(E^{0}\right)=\mu\left(A_{1} A_{2} A_{3}-A_{1} A_{2} A_{8}\right)$
$\operatorname{det} J\left(E^{0}\right)>0$ implies
$A_{3}>A_{8}$
Equation (2.2) also implies
$A_{8}<A_{3}$
Further simplification of (2.29) gives

$$
\begin{equation*}
R_{0}<1 \tag{2.30}
\end{equation*}
$$

Hence, equation (2.26) and (2.30) proves the Theorem 2.1 that is the DFE is Locally Stable.

Theorem 2: The DFE point $E^{0}$ of the model is Globally Stable if $R_{0} \leq 1$.

## Proof:

Consider the Lyapunov candidate
$V=A_{3} I$
Taking the time derivative of (2.31) we have
$\frac{d V}{d t}=A_{3}\left[\frac{\alpha S}{N}-A_{3}\right] I$
Since $S \leq S^{0}$ and $N \leq N^{0}$
$\frac{d V}{d t} \leq A_{3}\left[\frac{\alpha S^{0}}{N^{0}}-A_{3}\right] I$
Further simplification of (2.33) gives
$\frac{d V}{d t} \leq A_{3}{ }^{2}\left(R_{0}-1\right) I$
Hence, from equation (2.34) $\frac{d V}{d t} \leq 0$ if, $R_{0} \leq 1$. We conclude that $V(M, S, I, R)$ is negative definite and this proves that the DFE, $E^{0}$ is Global Stability.

### 2.6 Endemic Equilibrium (EE) Point in Terms of Force of Infection

Let,

$$
\begin{equation*}
E^{* *}=\left(M^{* *}, S^{* *}, I^{* *}, R^{* *}\right)=(M, S, I, R) \tag{2.35}
\end{equation*}
$$

be the Endemic Equilibrium point
Therefore, equation (2.8) to (2.11) becomes,

$$
\left.\begin{array}{l}
\Lambda \beta-A_{1} M^{* *}=0 \\
\Lambda(1-\beta)-\lambda^{* *} S^{* *}+\theta M^{* *}-A_{2} S^{* *}=0  \tag{2.36}\\
\lambda^{* *} S^{* *}-A_{3} I^{* *}=0 \\
A_{4} I^{* *}+v S^{* *}-\mu R^{* *}=0
\end{array}\right\}
$$

Where,

$$
\begin{equation*}
\lambda^{* *}=\frac{\alpha I^{* *}}{N^{* *}} \tag{2.37}
\end{equation*}
$$

Solving (2.36) simultaneously gives the Endemic Equilibrium points Infection in terms of force of infection
$\left(\begin{array}{c}M^{* *} \\ S^{* * *} \\ I^{* * *} \\ R^{* * *}\end{array}\right)=\left(\begin{array}{c}\frac{\Lambda \beta}{A_{1}} \\ \frac{A_{6}}{A_{1}\left(\lambda^{* *}+A_{2}\right)} \\ \frac{\lambda^{* *} A_{6}}{A_{1} A_{3}\left(\lambda^{* *}+A_{2}\right)} \\ \frac{\lambda^{* *} A_{4} A_{6}+v A_{3} A_{6}}{\mu A_{1} A_{3}\left(\lambda^{* *}+A_{2}\right)}\end{array}\right)$
At Endemic Equilibrium
$\left.\begin{array}{l}N^{* *}=M^{* *}+S^{* *}+I^{* *}+R^{* *} \\ N^{* *}=\frac{A_{9} \lambda^{* *}+A_{2} A_{3} A_{7}}{\mu A_{1} A_{3}\left(\lambda^{* *}+A_{2}\right)}\end{array}\right\}$
Where, $A_{9}=\left(\Lambda \beta \mu A_{3}+\mu A_{6}+A_{4} A_{6}\right)$
Substituting $I^{* *}$ and $N^{* *}$ into (2.37) and simplify gives
$\lambda^{* *}=\frac{A_{2} A_{3} A_{7}}{A_{9}}\left(R_{0}-1\right)$
Equation (2.40) is significant in this work because that is what we are going to use to carry out the bifurcation analysis.

## 3. Results and Discussion

Table 3.1 is the table of estimated values of variables and parameters of the model that are used to carry out the bifurcation analysis, sensitivity analysis and the graphical presentation of sensitive parameters against $R_{0}$. The details of the estimation of values are in Appendices.

Table 3.1: Estimated Values of Variables and Parameters of the Model

| Variables | Values per year | Source |
| :---: | :---: | :---: |
| $M(0)$ | $4,445,373$ | B 9 |
| $S(0)$ | $191,497,210$ | B 10 |
| $I(0)$ | 61,931 | B 3 |
| $R(0)$ | $4,959,085$ | B 4 |
| $N$ | $200,963,599$ | B 1 |
| $\Lambda$ | $7,534,530$ | B 2 |
| $\beta$ | 0.75 | B 13 |
| $\alpha$ | 0.9 | B 12 |
| $\delta$ | 0.005 | B 6 |
| $\gamma$ | 0.99 | B 5 |
| $\tau$ | 0.83 | B 14 |
| $\mu$ | 0.12 | B 7 |
| $\theta$ | 0.39 | B 11 |
| $v$ | 0.65 | B 8 |

### 3.1 Bifurcation Analysis

The stability of DFE at $R_{0}<1$ is not sufficient to conclude that the measles can be eradicated of in the population. However, we carried out the bifurcation analysis using equation (2.40) to see whether the model exhibit Backward or Forward bifurcation.


Figure 3.1: Forward Bifurcation Diagram for the Model
In figure 3.1, it was revealed that the stabilities of DFE and EE points depending on the value of $R_{0}$. A forward bifurcation in the equilibrium points occur at $R_{0}=1$. The DFE is stable if, $R_{0}<1$ and unstable if $R_{0}>1$. However, $R_{0}>1$, implies the stable EE. Thus, the model exhibit forward bifurcation because the stable endemic equilibrium does not exist when $R_{0}<1$. The force of infection, $\lambda^{* *}$ is an increasing function of $R_{0}$.

### 3.2 Sensitivity Analysis of the $R_{0}$ with Some Parameter of the Model

The approach of Abdulrahman et al. (2013) will be followed to carry out the sensitivity analysis. The normalized forward sensitivity indices with respect to a parameter values, $Q$ is defined as

$$
\begin{equation*}
S_{Q}^{R_{0}}=\frac{\partial R_{0}}{\partial Q} \times \frac{Q}{R_{0}} \tag{4.1}
\end{equation*}
$$

Where,

$$
\begin{equation*}
Q=\{\alpha, \gamma, v, \theta, \tau\} \tag{4.2}
\end{equation*}
$$

Maple 13 software is used to calculate the sensitivity indices using the values in Table 3.1.

Table 3.2: Sensitivity Indices of the Parameters of the Model

| Parameter | Low transmission <br> Sensitivity Index | High transmission <br> Sensitivity Index |
| :---: | :---: | :---: |
| $\alpha$ | 1.000000000 | 1.00000000 |
| $\gamma$ | -0.0861244019 | -0.5089974293 |
| $\nu$ | -0.6756756756 | -0.8441558441 |
| $\theta$ | 0.2507331378 | 0.1638655463 |
| $\tau$ | -0.171003717 | -0.4267352185 |

It is shown in Table 3.2 that the parameters have either positive or negative effects on the $R_{0}$. The positive parameters will increase the $R_{0}$ while the negative parameters will decrease the $R_{0}$. The contact rate $\alpha$ has the highest sensitivity index follow by vaccination rate, $v$ and other parameters follow.

### 3.3 Graphical Presentation of $R_{0}$ and Some Parameters of the Model

Figures 3.2 to 3.6 are the graphical presentations of sensitive parameters and $R_{0}$


Figure 3.2: The Effect of Contacts Rate Graph on $R_{0}$
Figure 3.2 shows that as contact rate increases with time the $R_{0}$ increases. This implies that vulnerable people should avoid contact with infected persons. It is observed from figure 3.3 that as recovery rate increases with time the $R_{0}$, decreases to almost zero. This shows the vital role the treatment of the
symptoms of measles can play in eradicating it from the population. The more people are treated the less the $R_{0}$ and it will result to total eradication from the population.


Figure 3.3: The Effect of Recovery Rates on $R_{0}$


Figure 3.4: The Effect of Vaccination Rates on $R_{0}$
In Figure 3.4 it is revealed that as vaccination rate increases with time the $R_{0}$ decreases. Vaccination of children between nine months to one year of age is crucial to eradicate measles. It is shown from Figure
3.5 that as loss of immunity increase with time the $R_{0}$, increases. Strong immunity is needed in the fight to eradicate measles. In Figure 3.6 as the Vitamin A supplement rate increases with time the basic reproduction number $R_{0}$, decreases. Vitamin A supplement is recommended to boost the immune system of the children infected with measles.


Figure 3.5: The Effect of Loss of Immunity Rate on $R_{0}$


Figure 3.6: The Effect of Vitamin A Supplement Rates on $R_{0}$

## 4. Conclusion

In this paper, we obtained the DFE and EE in terms of force of infection and analyzed the Local and Global stabilities of DFE. Trace -determinant method was used to analyze the local stability and

Lyapunov function to analyzed the global stability. The DFE is locally and globally stable. Measles will be put under control if $R_{0}<1$. Bifurcation analysis reveals that the model exhibit forward bifurcation (i.e. measles can be eradicated when $R_{0}<1$ ). It is evident from the sensitivity analysis that the contact rate $\alpha$, is the most sensitive parameter to increase the $R_{0}$ and vaccination rate $v$ is the most sensitive parameter to reduce the $R_{0}$. Graphical presentation shows that, vaccination rate and recovery rate are important parameters in eradicating the measles from the population. It was also discovered from Figure 3.2 that increase in contact with people infected with measles increases the basic reproduction number. Hence, the infected people should be separated from the susceptible individuals in order to reduce the spread of epidemics. It is also recommended that government should do more in routine immunization of measles and administration of Vitamin A supplement to the infected people.

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## APPENDICES

Appendix A: Reported Measles Cases in Nigeria by WHO Region 2006 to 2020 as of April, 2020
Table A1a: Reported Measles Cases in Nigeria by WHO Region 2006 to 2020 as of April, 2020

Measles cases: Nigeria
World Health
Organization




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Table A1b: Reported Measles Cases in Nigeria by WHO Region 2006 to 2020 as of April, 2020

| Hemar | Comifitinimed Canger |
| :---: | :---: |
| 200E | -1015 |
| 2007 | 3123 |
| 200us | -10453 |
| 2009 | -4583 |
| 2010 | 5072 |
| 2017 | -506z |
| 2012 | B423 |
| 2013 | E53G2 |
| 2014 | Exsod |
| 2015 | -2351 |
| 207E | $77^{\text {THEA }}$ |
| 2077 | - - - - - |
| 20- | 7018 |
| 2015 | 28302 |
| 2020 | 2376 |

Source: WHO (2020)
Table A2: Distribution of Measles Cases in Nigeria, Year and Month

| Year | Jan | Feb | Mar | Apr | May | Jun | Jul | Aug | Sep | Oct | Nov | Dec | TOTAL |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2011 | 4514 | 3970 | 4499 | 2518 | 1054 | 610 | 431 | 434 | 232 | 250 | 341 | 209 | 19062 |
| 2012 | 374 | 845 | 1186 | 796 | 355 | 220 | 137 | 190 | 229 | 189 | 807 | 1095 | 6423 |
| 2013 | 6604 | 12228 | 17785 | 9417 | 4385 | 1502 | 966 | 685 | 543 | 551 | 507 | 219 | 55392 |
| 2014 | 1557 | 1881 | 1285 | 685 | 449 | 287 | 68 | 80 | 55 | 115 | 243 | 145 | 6850 |
| 2015 | 641 | 1558 | 1927 | 1686 | 1390 | 932 | 691 | 599 | 664 | 969 | 924 | 410 | 12391 |
| 2016 | 2638 | 3947 | 3052 | 1823 | 1547 | 972 | 699 | 976 | 781 | 581 | 316 | 249 | 17581 |
| 2017 | 1142 | 1142 | 1913 | 1548 | 1231 | 1041 | 696 | 805 | 696 | 578 | 242 | 154 | 11188 |
| 2018 | 1223 | 1457 | 1699 | 906 | 425 | 238 | 228 | 155 | 183 | 190 | 166 | 148 | 7018 |
| 2019 | 3004 | 4935 | 7344 | 5808 | 3570 | 1325 | 825 | 318 | 305 | 340 | 292 | 236 | 28302 |
| 2020 | 927 | 779 | 607 | 63 | 0 |  |  |  |  |  |  |  | 2376 |

Source: WHO (2020)

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Table 3: Summary of Reported Suspected Measles Cases by State in 2019

| States | Dectembar | Deathe in Deatamber | Wheele 01- 202 | Tetal Peatis weele 1-52 (Clirses) |
| :---: | :---: | :---: | :---: | :---: |
| Abia | 17 | 0 | sog | 0 |
| Adamawa | 79 | 0 | 1,864 | $46(2596)$ |
| Akwa Ibom | 8 | 0 | 429 | 0 |
| Anamibra | 48 | 0 | 861 | $2(0.259)$ |
| Bauchi | 24 | 0 | 1,244 | 0 |
| Bayelsa | 7 | 0 | 368 | 0 |
| Berme | 0 | 0 | 325 | 0 |
| Bramo | 118 | 0 | 22,234 | 118 (0.5-5) |
| Cross River | 23 | 0 | 432 | 0 |
| Delta | 29 | 0 | 682 | 0 |
| Eboanyi | 8 | 0 | 346 | 0 |
| Edio | 4 | 0 | 409 | 0 |
| Elsiti | 28 | 0 | 911 | 0 |
| Enuga | 22 | 0 | 546 | 2 (0.4-6) |
| FCI | 0 | 0 | 216 | 0 |
| Grombe | 17 | 0 | 493 | 4 (0.8-59) |
| Imo | 11 | 0 | 712 | 0 |
| Jigawa | 0 | 0 | 1,160 | 0 |
| Faduma | 75 | 0 | 1,829 | 14 (0.8-9) |
| Famo | 3 | 0 | 4,141 | 27 (0.7-9) |
| Fatsina | 252 | 0 | 9,353 | 53 (0.6.9) |
| Kebbi | 64 | 3 | 801 | 4 (0.5-6) |
| Kogi | 12 | 0 | 277 | 0 |
| Kwara | 2 | 0 | 228 | $11.0 .596)$ |
| Lagas | 0 | 0 | 763 | 0 |
| Nasarawa | 5 | 0 | 279 | 0 |
| Nizger | 5 | 0 | 223 | 0 |
| Ogm | 22 | 0 | 735 | 0 |
| Ondo | 9 | 0 | 467 | 0 |
| Osm | 37 | 0 | 865 | 0 |
| Oyo | 5 | 0 | 1,062 | 0 |
| Platean | 3 | 0 | 4 Sl | 2 (0.4-5) |
| Rivers | 20 | 0 | 435 | 0 |
| Solkoto | 91 | 0 | 1,503 | $2(0.159)$ |
| Taraba | 11 | 0 | 143 | 0 |
| Yobe | 47 | 0 | 3,897 | 16 (0.4-5) |
| Zamffara | 3 | 0 | 405 | 0 |
| Total Cases | 1,109 | 3 | 61,931 | $291.00596)$ |

Source: (NCDC)

## Appendix B: Estimation of Variables and Parameter Values

The values of variables and parameters were estimated based on the available data from the World Health Organization (WHO), Population Reference Bureau and reliable related literature.

## B1: The Total Population, $N$

According to Worldometer, the Nigeria total population for 2019, is 200,963,599.

$$
N=200,963,599
$$

## B2: Recruitment Number, $\Lambda$

According to Knoema the number of new birth in Nigeria in 2019 is 7,534,530.
Therefore,
$\Lambda=7,534,530$

## B3: Number of Infected, $I$

The Nigeria Centre for Disease Control (NCDC) estimate that, there are 61,931 reported cases of measles in Nigeria in 2019, resulting in 291 deaths. (See Table A2)

$$
I=61,931
$$

## B4: Number of Recovered/Immune, $R$

$R=$ recovered + immune
From B3 the number of cases is 61,931 and number of death is 291.
Recovered $=61,931-291=61,640$
According to UNICEF Data: Monitoring the situation of children and women
the number of infants in Nigeria in 2019 is $7,534,530$ and the percentage of vaccination is $65 \%$.
Therefore,
Vaccinated $=65 \%$ of $7,534,530=4,897,445$.
Hence,
Recovered/Immune Human population, $R=61,640+4,897,445$
$R=4,959,085$
B5: Recovery Rate, $\gamma$

From B3 and B4

$$
\begin{aligned}
& \gamma=\frac{\text { Recovered }}{\text { Number of cases }} \\
& \gamma=\frac{61,640}{61,931}=0.99
\end{aligned}
$$

## B6: Disease Induce death rate, $\delta$

From B3 the number of cases of measles is 61,931 and the number of death from measles is 291

$$
\begin{aligned}
& \delta=\frac{\text { Number of Death from measles }}{\text { Number of cases }} \\
& \delta=\frac{291}{61,931}=0.005
\end{aligned}
$$

## B7: Natural Death Rate, $\mu$

According to UNICEF, the death rate is 120 deaths per 1,000 live births. Therefore,

$$
\mu=\frac{120}{1000}=0.12
$$

## B8: Vaccination rate, $v$

According to, WHO in 2019, about $65 \%$ of the Nigeria's children received one dose of measles vaccine. Therefore,

$$
v=0.65
$$

## B9: Maternally-Derived-Immunity, $M$

According to Millennium Development Goal (MDG4), close to $41 \%$ of deaths among newborn infants, occurred in their first 28 days of life or the neonatal period.
$M=59 \%$ of $7,534,530$
$M=4,445,373$
B10: Number of Susceptible, $S$
Recall $N=M+S+I+R$ therefore,

$$
S=N-(M+I+R)
$$

$S=200,963,599-(4,445,373+61,931+4,959,085)$
$S=191,497,210$

B11: Loss of immunity, $\theta$
According to WHO Immunization coverage fact sheet, national immunization schedule reported that, only $65 \%$ of children received 1 doses of measles. Therefore,
$\theta=35 \%=0.35$

## B12: Contact Rate, $\alpha$

$$
\alpha=\frac{9}{10}=0.9
$$

## B13: Birth with Temporary Passive immunity

$\beta=0.75$
B14: Vitamin A, Supplementation Coverage Rate:
According to Trading Economics the Vitamin A, supplementation coverage in 2018 is 83\% $\tau=0.83$

