#### Population infection estimation from wastewater surveillance for SARS-CoV-2 in Nagpur, India during the second pandemic wave

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### Abstract

Wastewater-based epidemiology (WBE) has emerged as an effective environmental surveillance tool for predicting severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) disease outbreaks in high-income countries (HICs) with centralized sewage infrastructure. However, few studies have applied WBE alongside epidemic disease modelling to estimate the prevalence of SARS-CoV-2 in low-resource settings. This study aimed to explore the feasibility of collecting untreated wastewater samples from rural and urban catchment areas of Nagpur district, to detect and quantify SARS-CoV-2 using real-time qPCR, to compare geographic differences in viral loads, and to integrate the wastewater data into a modified

Susceptible-Exposed-Infectious-Confirmed Positives-Recovered (SEIPR) model. Of the 983 wastewater samples analyzed for SARS-CoV-2 RNA, we detected significantly

higher sample positivity rates,  $43.7\%$  (95% confidence interval (CI) 40.1, 47.4) and 30.4% (95% CI 24.66, 36.66), and higher viral loads for the urban compared with rural samples, respectively. The Basic reproductive number,  $R_0$ , positively correlated with population density and negatively correlated with humidity, a proxy for rainfall and dilution of waste in the sewers. The SEIPR model estimated the rate of unreported coronavirus disease 2019 (COVID-19) cases at the start of the wave as 13.97 [95% CI (10.17, 17.0)] times that of confirmed cases, representing a material difference in cases and healthcare resource burden. Wastewater surveillance might prove to be a more reliable way to prepare for surges in COVID-19 cases during future waves for authorities.

# **Introduction**

Wastewater-based epidemiology (WBE) has emerged as a valuable and cost-effective strategy for monitoring the prevalence of severe acute respiratory syndrome coronavirus  $2$  (SARS-CoV-2) within communities and predicting disease outbreaks  $[1, 2]$  $[1, 2]$ . This approach capitalizes on the detection of SARS-CoV-2 RNA in wastewater samples and <sup>5</sup> has been widely employed using samples obtained from wastewater treatment plants (WWTPs) in nations with centralized sewage networks [\[1,](#page-14-0) [2\]](#page-14-1). While initially applied in <sup>7</sup> countries with centralized sewage networks, predominantly through wastewater treatment plant (WWTP) samples  $[3, 4]$  $[3, 4]$ , the applicability of WBE has transcended geographical constraints, encompassing a diverse range of sources such as river water, <sup>10</sup> airport wastewater, hospital effluents, marketplaces, and municipal drains [\[5–](#page-14-4)[8\]](#page-14-5). <sup>11</sup>

This research endeavours to explore the feasibility of a cross-sectional 12 wastewater-based sampling strategy aimed at detecting and quantifying SARS-CoV-2 13 viral loads in untreated wastewater within the Nagpur district, located in Maharashtra, <sup>14</sup> Central India. Notably, the sampling done for this study coincides with the second wave <sup>15</sup> of the COVID-19 pandemic in India in 2021, marked by an unprecedented surge in <sup>16</sup> transmission and heightened disease impact. However, the comprehensive integration of  $\frac{1}{17}$ WBE into disease surveillance systems, particularly in low- and middle-income countries 18 (LMICs), is limited due to inadequate centralized sanitation facilities [\[3,](#page-14-2) [4\]](#page-14-3). One of the <sup>19</sup> major reasons for this underutilization of WBE in LMICs, despite its huge potential, is  $\infty$ that in such countries centralized sanitation facilities are often lacking [\[7\]](#page-14-6). <sup>21</sup>

In the course of carrying out this research, though there are several related 22 works  $[9-13]$  $[9-13]$ , one seminal study in the field of WBE that this research utilised was conducted by McMahon *et al.* [\[9\]](#page-14-7). Their study investigates the use of wastewater  $\frac{24}{4}$ samples to monitor community-level transmission of  $SARS-CoV-2$ , the virus responsible  $25$ for COVID-19. The authors employ a Susceptible-Exposed-Infectious-Recovered (SEIR)  $_{26}$ model to estimate the number of infected individuals based on SARS-CoV-2 RNA  $_{27}$ concentrations detected in wastewater. Via their rigorous analysis, McMahon *et al.* [\[14\]](#page-15-1)  $_{28}$ demonstrate the utility of the SEIR model in predicting infections by considering <sup>29</sup> various parameters such as transmission rates and viral shedding dynamics. In addition, so their work introduces a simplified equation that aids in estimating infections from  $\frac{31}{21}$ wastewater data, enhancing the accessibility of the model's application. The study's use  $\frac{32}{2}$ of Monte Carlo simulations further strengthens the accuracy of predictions, revealing a 33 notable discrepancy between estimated infections and confirmed cases, thus highlighting  $\frac{34}{4}$ the potential value of the SEIR model in informing public health strategies  $[14]$ .

To this end, the integration of wastewater-based estimates complements traditional  $\frac{36}{10}$ clinical testing and bolsters the accuracy of surveillance efforts, especially in  $\frac{37}{20}$ resource-constrained settings where extensive clinical testing might be challenging [\[15\]](#page-15-2). 38 Thus, by integrating data from wastewater samples with demographic information and <sup>39</sup> clinical data, a model is proposed which generates robust estimates of the number of <sup>40</sup> COVID-19 infections within a given population. Crucially, this approach provides a <sup>41</sup> comprehensive perspective on viral transmission dynamics, assisting public health <sup>42</sup> officials in understanding the disease's impact on a broader scale. The imperative of <sup>43</sup> this study is to develop wastewater-based surveillance systems in LMICs, particularly <sup>44</sup> those with resource limitations and complex infrastructural challenges and underscores  $\frac{45}{100}$ the necessity of adapting WBE to a broader global context  $[16, 17]$  $[16, 17]$ . In these settings,  $\frac{46}{10}$ the translation of wastewater surveillance data into effective public health tools requires  $\frac{47}{47}$ the integration of mathematical models and simulations.

To address these challenges, the adaptation of mathematical models is crucial. The <sup>49</sup> use of a modified version of SEIR modelling and Monte Carlo simulation  $(MC)$  in this  $\sim$ study is motivated by the ability to effectively capture and analyze the transmission  $\frac{51}{100}$ dynamics of infectious diseases, such as SARS-CoV-2. The SEIR model and MC <sup>52</sup> simulation have established themselves as valuable tools in epidemiological research  $\frac{53}{2}$ because of their ability to provide insights into the complex systems involved in <sup>54</sup> infection transmission, population dynamics, and uncertainty analysis. The SEIR 55 compartment model forms the foundation for understanding disease transmission  $\frac{56}{100}$ dynamics [\[18](#page-15-5)[–22\]](#page-15-6). The SEIR model categorizes individuals into different compartments  $\frac{57}{2}$ based on their disease status, encompassing susceptible, exposed, infectious, and  $\frac{58}{100}$ recovered individuals. This model enables the estimation of disease prevalence over time, aiding in the interpretation of wastewater surveillance data and its linkage to  $\frac{60}{60}$ community infection dynamics. MC simulations, on the other hand, are a robust  $61$ computational technique used to account for uncertainties and variations in parameters.  $\epsilon_2$ By generating multiple simulations with randomly sampled inputs,  $MC$  simulations  $\frac{63}{1000}$ enable the exploration of a range of possible outcomes. This is particularly valuable in  $\epsilon$ epidemiological studies where factors such as contact rates, transmission probabilities, <sub>65</sub> and intervention effects can vary or are uncertain. MC simulations provide a way to  $\frac{66}{66}$ quantify the uncertainty associated with model predictions, helping researchers 67 understand the potential variability in their results  $[14, 23-27]$  $[14, 23-27]$  $[14, 23-27]$ .

This research initiative represents a pioneering effort in the Indian context, <sup>69</sup> harnessing the SEIPR model and MC simulations to illuminate the transmission  $\frac{70}{20}$ patterns of SARS-CoV-2 through wastewater. By addressing critical knowledge gaps  $\frac{71}{2}$ within LMICs and regions confronting infrastructural limitations, this study contributes  $\frac{72}{2}$ not only to scientific advancement but also furnishes actionable insights for policy  $\frac{73}{2}$ formulation and disease mitigation. Amidst the complex landscape of the COVID-19 <sup>74</sup> pandemic, this endeavour augments the global repository of knowledge, empowering  $\frac{75}{5}$ communities and authorities alike to respond effectively to this ongoing public health  $\tau$  $\Box$ challenge.  $\Box$ 77

In this study, we explored the feasibility of conducting a cross-sectional  $\frac{78}{78}$ wastewater-based sampling study for the detection, determination, and comparison of  $\frac{79}{20}$  $SARS-CoV-2$  viral loads from untreated wastewater in urban and rural areas of Nagpur  $\approx$ district, Maharashtra, Central India. We selected our sampling period during the second <sup>81</sup> wave of COVID-19 in India in 2021. We next developed a modified version of the SEIR  $\frac{1}{82}$ compartment mathematical model that has been frequently used to model COVID-19 83 dynamics in different populations  $[18, 19, 22]$  $[18, 19, 22]$  $[18, 19, 22]$ , herein termed the "SEIPR model" to  $\frac{84}{9}$ predict the number of infected individuals within specific Nagpur district partitioned  $\frac{1}{1000}$ zones and the total urban population under study. After predicting the number of  $\frac{86}{100}$ infected individuals, the estimates were used to perform Monte-Carlo simulations to  $\frac{87}{87}$ model the variations in the concentration of SARS-CoV-2 RNA in wastewater over time. <sup>88</sup> These modelled changes were then compared to the actual measurements recorded to  $\frac{89}{89}$ evaluate the accuracy of our SEIPR model. The urban incident COVID-19 cases were  $\Box$ also used to calculate the basic reproduction number  $R_0$  based on the SEIPR model.  $\Box$ This data was correlated to air temperature, relative humidity (a loose proxy for rainfall  $_{92}$ as we did not have the precise precipitation data), and population density to enhance  $\frac{93}{2}$  epidemiological understanding of environmental and human factors that may impact <sup>94</sup>  $SARS-CoV-2$  transmission dynamics in Central India. To the best of our understanding,  $\frac{95}{2}$ this is the first Indian report that has employed the SEIPR model to measure the  $\frac{96}{96}$ transmission patterns of SARS-CoV-2 through wastewater. This study could prove <sup>97</sup> valuable for local authorities and government officials as it provides important insights  $\frac{98}{96}$ to make well-informed policy decisions.

# $\mathbf M$ aterials and methods  $\begin{array}{ccc} \text{100} & \text{100} & \text{100} \end{array}$

## Wastewater sampling,  $SARS-CoV-2$  detection, and quantification  $_{101}$

Untreated (raw) wastewater samples were collected prospectively from the drainage  $102$ systems in the Nagpur district of Maharashtra, India, during the second wave of the 103 COVID-19 pandemic from January 31st to July 9th, 2021. Nagpur district is divided  $_{104}$ into 13 rural talukas and the Nagpur urban region, governed under Nagpur Municipal 105 Corporation (NMC). The Nagpur urban region is further divided into ten municipality 106 zones with each further divided into municipal wards. Individual grab samples were  $_{107}$ collected from sewers within each urban municipality zone as well as open  $108$ drains/groundwater sources of rural talukas representing the complete Nagpur district, <sup>109</sup> as illustrated in Fig [1](#page-3-0) right panel (urban taluka) and left panel (rural talukas in relation <sup>110</sup> to urban taluka). Each sample  $(1000 \text{ mL})$  was collected in sterile wide-mouth  $111$ autoclaved plastic bottles sealed in plastic bags and transported under a cold chain at  $_{112}$ 4 ◦C within 18-24 hours. All sampling was conducted during the morning hours between <sup>113</sup> 07:30 to midday using appropriate COVID-19 precautions. Samples were transported to  $_{114}$ Dr B. Lal Institute of Biotechnology, Jaipur, for pre-processing, RNA extraction and 115 SARS-CoV-2 detection by RT-qPCR, as previously described [\[28\]](#page-16-1). No specific permits  $_{116}$ were required for this study for field site access. We have only informed NMC regarding  $_{117}$ this study. Detailed sample processing methodology is presented in [S1 Appendix](#page-13-0) of 118 Supplementary information. 119

<span id="page-3-0"></span>Fig 1. Map of Nagpur district (study area) showing sampling locations for wastewater study. Each dot represents a location of wastewater collection in Nagpur urban and rural talukas. The map was created using the ArcGIS 10.4 version from a GIS student. Source of map used "ESRI, Maxar, Earthstar, Geographics and the GIS user Community".

#### Data collection for COVID-19 cases and environmental 120 characteristics in the characteristics

Demographics, and climatic factors including the presence of rainfall, air temperature, 122 and relative humidity, along with GPS coordinates, were also recorded by field workers 123 based at the Central Indian Institute of Medical Sciences (CIIMS) and assisted by the <sup>124</sup> NMC. Daily laboratory-confirmed COVID-19 positive cases and deaths between 1st 125 February and 30th July 2021 within the ten different municipality zones in urban 126 Nagpur were obtained from the health department of the NMC.

## Epidemiological modelling and estimation of infected individuals  $\frac{1}{128}$

We based our study of the transmission of  $SARS-CoV-2$  infections on a deterministic  $129$ ordinary differential equation (ODE) disease model in which the individuals in an entire <sup>130</sup> population can present in five mutually exclusive compartments according to their <sup>131</sup>

disease status and other measures. These compartments are susceptible, exposed,  $_{132}$ infectious, confirmed positive and recovered, abbreviated as the SEIPR model, which is <sup>133</sup> a modification of the SEIR model and that described by Acheampong *et al.* [\[20\]](#page-15-9), where  $_{134}$ additional compartments were given to reflect the Ghanaian environment [\[20\]](#page-15-9). We 135 denote the proportion of susceptible individuals by  $S(t)$ , the proportion of  $136$ asymptomatic infected individuals by  $E(t)$ , the proportion of symptomatic infectious  $137$ individuals by  $I(t)$ , the proportion of confirmed positive infectious individuals by  $P(t)$  138 and the proportion of recovered individuals by  $R(t)$ . It must be noted here that 139 individuals in the confirmed-positive class are carriers of the SARS-CoV-2 virus who <sup>140</sup> have had clinical confirmation of this status. However, individuals in an infectious class  $_{141}$ show clear symptoms and have high infectivity but have not yet been clinically  $_{142}$ confirmed positive. Notably, as highlighted by Acheampong et al. [\[20\]](#page-15-9), individuals <sup>143</sup> classified within the infectious class  $I(t)$  represent an abstract concept that is often  $144$ unmeasurable. This underscores the significance of introducing a compartment like the <sup>145</sup> confirmed-positive class  $P(t)$ , enabling comparison with the actual reported cases  $146$ within the population. The SEIPR model was applied to study COVID-19 dynamics in  $_{147}$ ten zones within Nagpur's urban area. Each zone operates independently. Disease <sup>148</sup> transmission is driven by a force of infection  $(\lambda)$ , determined by the effective contact rate per day  $(\beta_1)$  and reductions in transmissibility for exposed  $(\beta_2)$  and confirmed 150 positive  $(\beta_3)$  individuals. Disease-induced deaths are assumed to only occur within the 151 infectious  $(I)$  and confirmed positive  $(P)$  compartments. The model describes how 152 individuals transition between these compartments based on rates of entry and exit, such as exposure to infection ( $\lambda$ ), testing ( $\omega$ ), recovery ( $\rho$ ), and disease-induced death 154 (d). Recovery of individuals  $(R)$  depends on recovery rates from the confirmed positive 155 (P) and symptomatic infectious (I) compartments ( $\rho P(t)$  and  $\rho \gamma P(t)$ , respectively). Additionally, no natural birth and death are considered, and their exclusion may be justified given the assumed short-term focus on COVID-19 dynamics and the neglect of  $\frac{158}{158}$ population-level demographic changes, simplifying the model for this specific <sup>159</sup> epidemiological context. These underlying assumptions guide the model's representation <sup>160</sup> of COVID-19 transmission and progression in the Nagpur urban area. The transmission <sup>161</sup> dynamics of the SARS-CoV-2 infections are described by the five nonlinear systems of <sup>162</sup>  $ODEs$  shown in Eq  $(1)$ : 163

<span id="page-4-0"></span>
$$
\frac{\mathrm{d}}{\mathrm{d}t}X_t = f(X_t, t, \theta),\tag{1}
$$

with  $X_0 = [S_0, E_0, I_0, P_0, R_0]^T$  is initial number of individuals, t denotes time,  $X_t = [S, E, I, P, R]^T$  denotes the number of individuals in these compartments at time t, 165 T denotes matrix transposition, denotes the parameter vector and  $f(\cdot)$  denotes the 166 nonlinear relationship describing the state variable (see [S2 Appendix](#page-14-8) in the 167 Supplementary Information for detailed mathematical derivation of SEIPR model). The 168 force of infection used in this model is  $\lambda = \beta_1(\beta_2 E(t) + \beta_3 P(t) + I(t))$ , with  $\beta_1$  denoting 169 the effective contact rate per day, and  $\beta_2$  and  $\beta_3$  respectively accounts for the reduction  $\frac{170}{20}$ in disease transmissibility of exposed and confirmed positive individuals. A value of  $\frac{1}{171}$ epidemiological importance in infectious disease modelling is the basic reproductive  $\frac{172}{172}$ number, which in this study is referred to as the number of secondary SARS-CoV-2  $_{173}$ infections generated by a single active SARS-CoV-2 infected individual during the entire <sup>174</sup> infectious period [\[29\]](#page-16-2). It is given by the Eq  $(2)$ : 175

<span id="page-4-1"></span>
$$
R_0 = \beta_1 S^0 \left( \frac{\beta_2}{\epsilon} + \frac{\beta_3 (1 - \omega)}{i_T} + \frac{\delta (1 - \omega) + i_T \omega}{i_T p_T} \right),\tag{2}
$$

where  $i_T = \delta + \gamma \rho + d$  and  $p_T = \rho + d$ . The effective reproductive number  $(R_0)$  is made

up of contributions from secondary infections from the exposed class generated by  $\frac{177}{20}$ asymptomatic individuals (first term), confirmed positive individuals' class (second <sup>178</sup> term), and the infected (symptomatic) class (third term).  $S^0$  is the proportion of the 179 population that is initially susceptible. Other parameters in Eq  $(2)$  are defined as  $180$ follows:  $\epsilon$  denotes the incubation period,  $\sigma$  denotes the progression rate of susceptible 181 individuals to the confirmed positive class via testing per day,  $\delta$  denotes the progression  $_{182}$ rate of infectious individuals to the confirmed positive class via testing per day,  $d_{183}$ denotes the disease-induced death rate per day,  $\omega$  denotes the fraction of exposed 184 individuals that transient to confirmed positive class,  $\gamma$  denotes the fraction of  $\frac{185}{185}$ infectious individual that transient to recovery class and  $\rho$  denotes the recovery rate of  $_{186}$ confirmed positive individuals per day. In this study, the nonlinear least squares scheme 187 is used to estimate the parameters involved in the calculation of  $R_0$ . The model fitting 188 was first carried out for each zone to obtain zone-specific parameter estimates and 189 secondly for all zones put together as a single unit. Further details about model 190 [d](#page-14-8)erivation and parameter estimation can be found in the Supplementary (see [S2](#page-14-8) 191 [Appendix](#page-14-8) for a full description of model parameters and variables). For this study, the <sup>192</sup> number of SARS-CoV-2 infected individuals within urban Nagpur was estimated using <sup>193</sup> the modelling approach proposed by McMahon *et al.* [\[14\]](#page-15-1), which combines our disease  $_{194}$ model (SEIPR) to the viral concentration estimations [\[14\]](#page-15-1). As already mentioned, there 195 are ten zones within urban Nagpur and each zone is modelled independently. Based on <sup>196</sup> McMahon *et al.* [\[14\]](#page-15-1), using our SEIPR disease model, the number of newly detected 197 infections on the jth day  $I_j^n$  is modelled as a Poisson process with rate parameter 198  $N\beta_1[\beta_2E(j) + \beta_3P(j) + I(j)]$ , which is expressed as Eq [\(3\)](#page-5-0):

<span id="page-5-0"></span>
$$
\mathcal{I}_j^n \sim \text{Poisson}\{N\beta_1[\beta_2 E(j) + \beta_3 P(j) + I(j)]\}, \text{ for } j = 1, 2, \dots, J,
$$
 (3)

where N is the total number of individuals that reside in the zone of the drainage  $_{200}$ systems. The viral load being introduced into the drainage system at time  $t$  is 201

<span id="page-5-3"></span>
$$
V_0(t) = \sum_{j:j \le t} \sum_{i=1}^{T_j^n} V_{ij}(t),
$$
\n(4)

where  $V_{ij}(t)$  is the number of copies of SARS-CoV-2 RNA entering the drainage  $_{202}$ systems via faeces of the *i*<sup>th</sup> individual of out the  $\mathcal{I}_j^n$  who became infected on day j is 203 modelled according to the Eq  $(5)$  204

<span id="page-5-1"></span>
$$
V_{ij}(t) = \vartheta_{ij} \left\{ 10^{\frac{\phi_{ij}(t-j)}{5}} I(j < t \le 5+j) + 10^{\psi_{ij}^{\frac{(\phi_{ij} - \psi_{ij})(t-5-j)}{5}} I(t > 5+j) \right\},\tag{5}
$$

for  $i = 1, 2, \ldots, \mathcal{I}_j^n$  (infected individuals) and  $j = 1, 2, \ldots, J$  (days). In Eq [\(5\)](#page-5-1),  $\vartheta_{ij}$  205 denotes the  $log_{10}$  g of faeces per *i*th individual who gets infected on the *j*th day, modelled as a normal distribution with mean of 2.41 and standard deviation of 0.25 per  $_{207}$ data from lower-middle-income countries [\[15\]](#page-15-2),  $\phi_{ij}$  denotes the log<sub>10</sub> maximum RNA  $\qquad$ <sub>208</sub> copies per g being of faeces shed 5 days after being infected, modelled as a normal <sup>209</sup> distribution with mean of 7.6 and standard deviation of 0.8 [\[14\]](#page-15-1) and  $\psi_{ij}$  denotes the 210  $log_{10}$  RNA copies per g being of faeces shed 25 days after being infected, modelled as a  $_{211}$ normal distribution with mean of 3.5 and standard deviation of 0.4. To correlate the 212 viral load being introduced into the drainage system to that being measured, McMahon <sup>213</sup> *et al.* [\[14\]](#page-15-1) proposed the Eq [\(6\)](#page-5-2) called the downstream RNA copies measured,  $V(t, \tau)$  to 214 account for the time-dependent degradation in the drainage system, <sup>215</sup>

<span id="page-5-2"></span>
$$
V(t,\tau) = V_0(t) \left(\frac{1}{2}\right)^{\tau/\tau^*},\tag{6}
$$

where  $\tau$  is the time elapsed between waste excretion and arrival at the drainage systems  $\tau$ modelled as a uniform distribution from  $\tau = 1$ h to  $\tau = 1.5$ h,  $V_0(t)$  is the viral load introduced into the drainage system modelled by Eq  $(4)$ ,  $\tau^*$  is the 218 temperature-dependent half-life modelled according to Eq [\(7\)](#page-6-0) <sup>219</sup>

<span id="page-6-0"></span>
$$
\tau^* = \tau_0^* Q_0^{(T - T_0)/10^{\circ}C},\tag{7}
$$

where  $T$  is the current temperature of the drainage system, modelled as a uniform  $_{220}$ distribution from  $T = 19$ °C to  $T = 31$ °C,  $\tau_0^*$  is the half-life (h) at an ambient 221 temperature of  $T_0$ , modelled as a normal distribution with a means of 3 h and 30 h  $_{222}$ respectively, with standard deviations of 0.7 and 1.5,  $Q_0$  is the temperature-dependent  $\frac{223}{2}$ rate of change, modelled as a normal distribution with a mean of 5.5 and standard 224 deviation of 0.5. The choice of distributions and parameter ranges were informed by previous research as well as actual measurements or observations of SARS-CoV-2 in <sup>226</sup> wastewater to inform their selection of parameter ranges for the Monte Carlo simulation. <sub>227</sub> All the above information was used to simulate the viral load of infected individuals  $_{228}$ generated by our proposed disease model via 500 Monte Carlo simulations, since beyond <sup>229</sup> this number of Monte-Carlo simulations, the value of the simulated RNA copies does 230 not significantly change. Importantly, the number of Monte Carlo samples depends on  $_{231}$ various factors including the complexity of the model, which is the case here.

Finally, McMahon *et al.* [\[14\]](#page-15-1) proposed a model for estimating the number of infected <sub>233</sub> individuals in each day given the measured RNA copies quantified from samples <sup>234</sup> collected from the drainage systems and is given by the Eq  $(8)$  235

<span id="page-6-1"></span>
$$
\mathcal{J}_t = \frac{Q \times V}{A \times B},\tag{8}
$$

where  $Q$  denotes the average flow rate at the drainage system in L per day,  $V$  denotes 236 the virus copies per L,  $\tilde{A}$  is the rate of faeces production per person in g per day with  $_{237}$  $A = 2 \times 128$  for developing countries [\[15\]](#page-15-2), and B denotes the maximum rate at which  $\frac{238}{2}$ the virus is shed in RNA copies for g of faeces per day with  $B = 10^{7.6} \times 128$  [\[14\]](#page-15-1). In this 239 study, Q was calculated as a point estimate using the product of the at-home <sup>240</sup> population in the catchment of each zone, and the observed average per capita <sup>241</sup> wastewater rate, which we assumed to be either 120 or 135 L/person/day (based on the  $_{242}$ Ministry of Housing and Urban affairs suggested benchmark for urban water supply). <sub>243</sub>

#### Statistical analyses <sup>244</sup>

Due to the lack of COVID-19 incidence data for the rural areas in Nagpur, we explored <sub>245</sub> catchment areas within urban Nagpur by zones to gain insight into the concentration of <sup>246</sup> SARS-CoV-2 viral load in the collected wastewater samples. Based on the model 247 parameter estimates, the distribution of the RNA copies per day existing in the <sup>248</sup> drainage systems by zones was estimated, where we used the 2011 population census <sup>249</sup> data as an estimate for each population zone. Of note, the use of the Monte Carlo 250 simulation approach can help estimate uncertainties and account for variability in the 251 data, which provides some indication of potential uncertainty and variability in 252 prevalence estimates despite the limitations of using this census data, making the <sup>253</sup> margin of error not a major problem. Data on continuous variables are presented as  $_{254}$ median with interquartile ranges  $(IQR)$ . Categorical variables are shown as counts and  $_{255}$ percentages in parentheses. The normality of data was assessed using the Shapiro-Wilk <sup>256</sup> test. Student's t-test was used for comparing variables which were normally distributed. <sup>257</sup> Mann-Whitney test was used when the normality assumption was violated. The 258 Fisher's exact test and Proportion tests were applied to compare categorical variables. <sup>259</sup> All p– values and confidence intervals (CIs) are two-sided and a p–value of  $< 0.05$  is 260

<span id="page-7-0"></span>

Table 1. Summary of climatic characteristics and RT-PCR results of wastewater samples collected within urban and rural Nagpur catchment.

Data are presented n  $(\%)$  or median (IQR). NA = not applicable. a: RT-PCR results for wastewater samples; ct: cycle threshold; n.s.: not significant

> Of the 10 sampled urban catchment zones, two zones (7 and 9) yielded no SARS-CoV-2 <sup>281</sup> RNA detection but did record the highest humidity levels (Table [2\)](#page-8-0). Only 3 zones  $_{282}$ experienced rainfall; zones 1 and 8, where rainfall was recorded 1 day prior to sample 283 collection, and zone 7, where sample collection took place during heavy rainfall. It is <sup>284</sup> likely that these rainfall events would also contribute to diluting the sewage prior to  $\frac{285}{265}$ sampling. Moreover, rainfall events would also contribute to more rapid and effective 286 flushing out within the sewers. In zone 9, wastewater sampling followed the conclusion <sub>287</sub>

<span id="page-8-0"></span>of the main COVID-19 infection wave, and therefore, the cases of COVID-19 at the time <sup>288</sup> of sampling were expected to be very low, as illustrated in Fig S1A –S1E [\(S3 Appendix\)](#page-14-9). <sup>289</sup> The distributions of the continuous data and their normality plots by individual zones  $_{290}$ are shown in Figures S3A-D [\(S3 Appendix\)](#page-14-9). The respective significance  $p-$  values 291 shown on the plots are all less than 0.05, indicating the data is not normally distributed.  $_{292}$ Table S1 [\(S3 Appendix\)](#page-14-9) summarises the demographic characteristics of the catchment 293 zones where wastewater samples were collected. The demographic and environmental <sup>294</sup> characteristics by zones are presented in Tables S2 and S3 [\(S3 Appendix\)](#page-14-9). <sup>295</sup>

Table 2. SARS-CoV-2 RT-PCR results detected per unit of time and detected viral load results of the wastewater samples with climatic and population census information for each Nagpur catchment zone.

Catchment	Population <sup><math>a</math></sup>	Temperature	Humidity	RT-PCR Result <sup>b</sup>	Genome Copy
		$(^\circ\mathrm{C})$	$(\%)$	(Positive)	$(10^5$ Copies per L)
Zone $1$	239171	$24(22-26)$	$52(40-65)$	24(24.5)	$\overline{1.135}$ $(0.875 - 1.359)$
Zone 2	159458	$24(22-26)$	$33(25-39)$	47(39.5)	17.003 (3.463 - 298.375
Zone 3	232247	$32(30-34)$	$23(18-34)$	57(87.7)	$2.390(0.776 - 3.753)$
Zone 4	208426	$30(27.5 - 33.5)$	$(16.5 - 41)$ 21	39(83.0)	$\overline{0.883}$ $(0.560 - 2.552)$
Zone 5	243953	$31(30-32)$	$39(27-45)$	46(73.0)	$2.168$ $(1.533 - 2.864)$
Zone 6	204438	$31(29-32)$	$36(33 - 44.2)$	36(60.0)	$2.509$ $(1.309 - 3.439)$
Zone 7	187044	$27(26-29)$	$92(88-94)$	0(0.0)	NA
Zone 8	346287	$33(30-34)$	$40(28.5 - 52.5)$	5(6.7)	$0.066$ $(0.036 - 0.124)$
Zone 9	317321	$29(27-32)$	$83(74-94)$	0(0.0)	NA
Zone $10$	267320	$27(25-29)$	$27(24-31)$	71(71.7)	$0.705(0.376 - 1.185)$

Data are presented n  $(\%)$  or median (IQR); a: 2011 census data; b: RT-PCR results for wastewater samples per unit of time for each zone, NA: not available.

#### Estimation of infected individuals 296

We fitted our proposed SEIPR model to the reported confirmed COVID-19 positive  $_{297}$ cases and deaths in urban Nagpur via the nonlinear least squares method. Fig [2](#page-9-0) (a and <sup>298</sup> b) shows the representative model fit for the SEIPR model to data for all 10 Nagpur <sup>299</sup> catchment zones combined as a single unit for the period of March to July 2021. <sup>300</sup>

Both plots show an increase in confirmed positive cases and deaths up to the first 50  $\frac{301}{201}$ days and then a decrease over the last  $100$  days. Thus, the SEIPR model predicts a  $\frac{302}{2}$ decrease in the susceptible population as individuals become exposed, infected,  $\frac{303}{200}$ confirmed positive, and then either recover or are confirmed dead. The remaining model  $_{304}$ fittings for the urban zones are presented in Figure S5 [\(S3 Appendix\)](#page-14-9). The <sup>305</sup> corresponding model parameter estimates for the respective catchment zones and  $R_0$  as  $\frac{306}{200}$ calculated using clinical incident data only, are presented in Table [3.](#page-9-1) Each urban <sup>307</sup> catchment zone exhibited different effective contact rates,  $\beta_1$ , signifying different  $\frac{308}{208}$ contact patterns. In addition, the basic reproduction number,  $R_0$  is different for each  $\sim$ catchment zone with the highest  $R_0$  observed in zone 9 and lowest in zone 2. All the  $\sim$  310 zones have an  $R_0$  greater than 1 except for zone 2. All the zones, when combined as a  $\frac{311}{2}$ single unit, gave an  $R_0$  of 1.11. Linear regression analysis to investigate the variation in  $\frac{312}{21}$  $R_0$  and  $\beta_1$  between the zones revealed a statistically significant positive correlation  $\beta_3$ between  $R_0$  and population density  $[R^2 = 0.40, p-value= 0.05]$  whilst for effective  $\frac{314}{2}$ contact rate  $(\beta_1)$  and  $R_0$ , there was a negative correlation with humidity  $[R^2 = 0.49, \ldots]$ p-value= 0.02]. No significant relationship was seen between temperature and  $R_0$  or  $\beta_1$  316 (Figure S8 in [\(S3 Appendix\)](#page-14-9)).

Taking all zones combined, Fig [2](#page-9-0) (c) depicts the distribution of the RNA copies per  $\frac{318}{2}$ day, similar to the dynamics observed by McMahon *et al.* [\[14\]](#page-15-1).  $\qquad \qquad \qquad \ldots$ 

Catchment		Parameter									
	$\beta_1$	$\beta_2$	$\beta_3$	$\epsilon^*$	$\sigma$	$\omega$	$\delta$	$\gamma$		d	$R_0$
	per	$(10^{-4})$	$(10^{-4})$	(per	$(10^{-4})$ per	$(10^{-4} \text{ per})$	$(10^{-4}$ per		$10^{-}$ per	$10^{-}$ $J^{-4}$ per	
	$\rm day)$			$\bf day)$	$\rm day)$	$\rm day)$	$\rm day)$		$\rm day)$	$\rm day)$	
Zone 1	0.91	0.42	143.79	0.2	2.48	19.16	21.81	1.00	1773.70	1.51	1.02
Zone 2	0.80	8.20	170.81	0.2	2.70	5.74	21.192	1.00	1627.00	1.58	0.98
Zone 3	0.92	0.01	8.22	0.2	2.64	10.13	25.811	1.00	1714.30	1.81	1.06
Zone 4	0.93	306.13	139.39	0.2	1.49	8.95	10.09	1.00	1344.40	1.03	1.52
Zone 5	0.99	7.12	1969.00	0.2	2.04	2.66	23.67	1.00	1557.30	1.88	1.26
Zone 6	0.76	1194.60	0.37	0.2	1.08	22.97	0.67	0.99	1333.90	0.56	1.59
Zone 7	0.75	0.02	11.05	0.2	0.522	2.70	3.51	1.00	1376.80	0.06	1.09
Zone 8	0.82	20.31	9990.80	0.2	0.69	1.66	8.46	0.98	1639.90	0.62	1.02
Zone 9	0.49	0.03	4998.80	0.2	0.88	27.16	0.01	0.59	1001.50	0.42	1.66
Zone $10$	0.92	1.27	2.07	0.2	2.07	5.51	11.77	1.00	1583.00	0.87	1.15
All Zones	0.80	209.22	242.52	0.2	1.53	0.32	12.65	0.98	1570.60	0.98	1.11

<span id="page-9-1"></span>Table 3. Model parameter estimates and basic reproduction number  $(R_0)$  for each catchment zone of Nagpur district.

\*: fixed parameter estimate adapted from Zhang *et al.*, \*\*: Computation of  $R_0$  and all model parameters are based on clinical incidence data and not wastewater samples. Note:  $\beta_1$  denotes the effective contact rate per day,  $\beta_2$  and  $\beta_3$  respectively account for the reduction in disease transmissibility of exposed and confirmed positive individuals.  $\epsilon$  denotes the incubation period,  $\sigma$  denotes the progression rate of susceptible individuals to confirmed positive class via testing per day,  $\delta$  denotes the progression rate of infectious individuals to confirmed positive class via testing per day, d denotes the disease-induced death rate per day,  $\omega$  denotes the fraction of exposed individuals that transient to confirmed positive class,  $\gamma$  denotes the fraction of infectious individual that transient to recovery class and  $\rho$  denotes the recovery rate of confirmed positive individuals per day.

> <span id="page-9-0"></span>Fig 2. Considering a half-life of 30 h. Model fit to the proportion of the population. (a) (left) confirmed positive COVID-19 infections and (b) (right) confirmed deaths from COVID-19 infections for all zones as a single unit. (c) SEIPR model [\(1\)](#page-4-0) prediction for the mass rate of SARS-CoV-2 RNA in wastewater over time via Monte-Carlo simulation represented by black points. (d) Zoomed-in plot of predicted number of active COVID-19 cases versus SARS-CoV-2 RNA mass rate with individual Monte-Carlo simulations represented by grey points, where 75% CI and 95% CI are denoted by the green and red solid lines, respectively. Coloured datapoints denote the measured RNA mass rates and estimated infectious individuals based on Eq [\(8\)](#page-6-1) as presented in Table [4,](#page-11-0) respectively for an assumed average per capita wastewater rates of 120 L per person per day (red solid points) and 135 L per person per day (blue solid points) for all zones as a single unit.

There is a positive correlation between the concentration of SARS-CoV-2 RNA in the  $\frac{320}{200}$ wastewater and the number of confirmed positive individuals as well as recovering  $\frac{321}{20}$ individuals and shedding rates. There was a different association between the measured <sub>322</sub> viral RNA concentration and the confirmed positive cases during the earlier stages 323 (January and February 2021) of the wastewater sampling, with high wastewater viral  $_{324}$ concentrations but low numbers of confirmed positive individuals. Therefore, zone 1 and 325 zone 2 were not considered for the viral RNA load SEIPR modelling (see Figure S2 in  $\frac{326}{2}$ the [S3 Appendix\)](#page-14-9). Fig [2](#page-9-0) (d) depicts a zoomed-in plot of the predicted number of active  $\frac{327}{20}$ COVID-19 cases versus SARS-CoV-2 RNA mass rate with individual Monte-Carlo <sup>328</sup> simulations represented by grey points. The measured RNA mass rates and estimated  $\frac{329}{2}$ number of infectious individuals based on Eq  $(8)$  are denoted by the coloured datapoints  $\frac{330}{2}$ and fall within the  $95\%$  CI denoted by the red solid lines. In this particular study, the  $\frac{331}{2}$ sensitivity of the model regarding the viral half-life at an ambient temperature of the  $\frac{332}{20}$ 

drainage is explored. It is observed that for a viral half-life of 3 h, the association  $\frac{333}{333}$ between the mass rate of gene copies detected in wastewater and the confirmed positive <sup>334</sup> cases is affected (see Supplementary figures S7A in the [S3 Appendix\)](#page-14-9). Data for all other <sup>335</sup> catchment zones are given in the Supplementary figures S6-S8 [\(S3 Appendix\)](#page-14-9)) except  $\frac{336}{10}$ for zone 7 and zone 9, where wastewater samples from these zones tested negative  $\frac{337}{337}$ (Table [2\)](#page-8-0). Furthermore, all plots depicting the entire Monte-Carlo simulations of the <sup>338</sup> predicted number of active COVID-19 cases versus SARS-CoV-2 RNA mass rate are  $\frac{339}{2}$ presented in Figures S9-S10 [\(S3 Appendix\)](#page-14-9).

Table [4](#page-11-0) presents the SARS-CoV-2 RNA wastewater concentrations in samples taken  $_{341}$ from all the catchment zones considered as a single unit between 1st March and 27th of <sup>342</sup> May, 2021. Results of the other catchment zones are presented in the Supplementary. <sub>343</sub> Each row corresponds to a specific date on which the wastewater samples were taken. <sub>344</sub> The "RNA (copy per L)" column provides the concentration of SARS-CoV-2 genetic 345 material in wastewater, providing insights into the prevalence of the virus in the  $\frac{346}{2}$ population. The following columns, titled "Option 1" and "Option 2", present two <sup>347</sup> separate scenarios based on different wastewater rates per capita (120 L/person/day for  $\frac{348}{2}$ Alternative 1). and 135 L/person/day for 'Option 2). These scenarios are important for  $\frac{349}{2}$ estimating the number of infected individuals using RNA concentrations as an indicator <sup>350</sup> of viral activity. Calculated RNA levels are provided for each scenario, showing the rate <sup>351</sup> of change in viral RNA levels per day. In addition, the "Estimated number of infected <sup>352</sup> individuals" column quantifies the number of potential COVID-19 cases inferred from  $\frac{353}{2}$ RNA levels, providing a way to assess community spread of the virus.

Direct comparison with clinically observed cases is presented in the column 355 "Clinically observed COVID-19 positive cases", showing actual confirmed positive cases <sup>356</sup> reported by clinical diagnoses. This actual data is used as a benchmark to evaluate the  $\frac{357}{200}$ validity of the estimates obtained through wastewater analysis. Side-by-side estimating <sub>358</sub> infected individuals with observed clinical cases helps assess the reliability of using  $\frac{359}{2}$ wastewater RNA concentrations as a predictor to monitor trends in COVID-19. Overall,  $_{360}$ this table highlights the importance of leveraging wastewater-based epidemiology to  $_{361}$ better understand viral prevalence. The ratio of unreported to reported cases under  $\frac{362}{362}$ options 1 and 2 are respectively computed to be  $12.42$  (95% CI 9.04, 15.15) and 13.97  $\frac{363}{2}$  $(95\% \text{ CI } (10.17, 17.0).$ 

## $\sum$  iscussion  $365$

WBE has been used as a tool for surveillance of COVID-19 infections at the  $_{366}$ community-level and complements clinical-based surveillance and screening, which is  $\frac{367}{267}$ limited by cost, turnaround time, and the bias associated with uncharacterized  $\frac{368}{368}$ asymptomatic infections and their contribution to infection spread. WBE captures the <sup>369</sup> totality of symptomatic, pre-symptomatic and asymptomatic carriers within a specific  $\frac{370}{200}$ community [\[16,](#page-15-3) [17\]](#page-15-4) This study is the first to successfully pilot and assess WBE as a  $\frac{371}{371}$ methodology for the detection and quantification of SARS-CoV-2 viral RNA in  $\frac{372}{272}$ community sewers in Nagpur district of Central India during the second wave of the  $\frac{373}{2}$ pandemic in 2021. Whilst several epidemiological models have been described and <sup>374</sup> compared for transmission of SARS-CoV-2 [\[2,](#page-14-1) [21,](#page-15-10) [30\]](#page-16-3), this study employed a new SEIPR  $\frac{375}{275}$ model, which adds the extra compartment of "confirmed positive" to estimate the <sup>376</sup> number of infected individuals and was further used to estimate the mass rate of RNA  $_{377}$ in the wastewater. We observed a low number of clinical cases early in the COVID-19  $\frac{378}{276}$ wave that was out of proportion to the observed high SARS-CoV-2 concentration in the  $\frac{379}{2}$ wastewater. If we use our modelling results from later in the study and apply them to  $\frac{380}{100}$ this earlier period, it reveals that the clinical surveillance data underestimated the level  $\frac{381}{381}$ of COVID-19 transmission in the Nagpur district. The model predicts the unreported <sup>382</sup>

<span id="page-11-0"></span>

Date	$\overline{\textbf{RNA}}$ $(10^5$	Option 1 Option 2			<b>Clinically observed</b>	
	copies per $L$ <sup>*</sup>					number of Covid-19
		<b>RNA</b> rate	<b>Estimated</b>	<b>RNA</b> rate	Estimated	positive cases
		$(10^{14}$ copies	number of	$(10^{14}$ copies	number of	
		per day) <sup><i>a</i></sup>	infected	per day) <sup><i>a</i></sup>	infected	
			individuals		individuals	
			$(10^{14})$		$(10^{14})$	
01/03/2021	6.10	1.76	17.28	1.98	19.40	777
02/03/2021	4.01	1.16	11.37	$\overline{1.30}$	12.79	897
03/03/2021	3.50	1.01	9.91	1.14	11.15	845
04/03/2021	3.74	1.08	10.59	1.21	11.92	1172
08/03/2021	41.31	1.92	116.98	13.41	131.61	1049
09/03/2021	34.1	9.84	96.59	11.07	108.66	1433
10/03/2021	6.26	1.81	17.73	2.03	19.95	1604
05/04/2021	15.1	4.36	42.77	4.90	48.12	2652
06/04/2021	14.10	4.07	3.94	4.58	44.93	3283
07/04/2021	97.40	28.12	275.89	31.63	310.38	2881
08/04/2021	65.10	18.80	184.46	21.15	207.52	4016
13/04/2021	2.73	0.79	7.73	0.89	$\overline{8.70}$	3613
15/04/2021	9.34	2.70	$\overline{26.46}$	$\overline{3.03}$	29.77	3779
19/04/2021	11.30	3.27	32.06	3.68	36.06	4878
20/04/2021	708.00	204.39	2005.44	229.93	2256.12	4787
21/04/2021	24.70	7.13	69.96	8.02	78.71	4619
23/04/2021	$\overline{20.10}$	5.80	56.95	6.53	64.06	4936
24/04/2021	23.10	6.67	65.43	7.50	73.61	4720
27/04/2021	19.90	5.74	56.37	6.46	63.41	4803
28/04/2021	6.81	1.97	19.29	$\overline{2.21}$	21.70	4422
29/04/2021	19.00	5.48	53.82	6.17	60.55	3649
30/04/2021	9.79	02.83	27.73	3.18	31.20	4085
03/05/2021	14.90	04.30	42.20	4.84	47.48	2498
04/05/2021	12.70	03.67	35.97	4.12	40.47	2534
06/05/2021	10.90	03.15	30.87	3.54	34.73	2255
07/05/2021	2.63	0.76	7.44	0.85	8.37	2016
25/05/2021	$\overline{0.07}$	$\overline{0.02}$	0.20	$\overline{0.02}$	0.23	339
27/05/2021	$\overline{0.18}$	$\overline{0.05}$	0.52	$\overline{0.06}$	0.58	$\overline{216}$

Table 4. SARS-CoV-2 RNA concentrations, estimated RNA rate and number of infected individuals from all the catchment zones as a single unit.

Option 1 assumes an average per capita wastewater rate of 120 L/person/day; Option 2 assumes an average per capita wastewater rate of 135 L/person/day; a: based on the numerator of Eq [\(8\)](#page-6-1); b: based on Eq (8); \* aggregate SARS-CoV-2 RNA concentration if samples are taken from different locations measured on the day.

> number of cases under the per capita wastewater rates of  $120L/person/day$  and 383 135L/person/day to be 12.42 (95% CI 9.04, 15.15) and 13.97 (95% CI (10.17, 17.0) <sup>384</sup> times higher than the reported number of cases, respectively. Hence, SARS-CoV-2 RNA 385 detected in community wastewaters may have come from pre-symptomatic,  $_{386}$ symptomatic, or asymptomatic cases who did not self-report to their local health <sup>387</sup> monitoring unit due to fear of social stigma, isolation, or quarantine, or simply because  $\frac{1}{388}$ they did not know they were infected  $[31, 32]$  $[31, 32]$ . Under-reporting bias in the clinical  $\frac{389}{2}$ incident data is also likely to have arisen due to the limitation of testing resources  $\frac{390}{2}$ (analytical kits, personnel), coverage, and accessibility of testing sites [\[33\]](#page-16-6). We observed <sup>391</sup>

that SARS-CoV-2 seemed to be suppressed in samples collected from catchment zones <sub>392</sub> recording higher relative humidity, a loose proxy for rainfall. This was further 393 substantiated by observing a statistically significant negative correlation between  $R_0$   $\qquad$   $\frac{394}{994}$ (effective reproductive number) and humidity, and  $\beta_1$  (effective contact rate per day)  $\qquad$  395 and humidity, but not temperature. These results partly agree with those reported  $\frac{396}{2}$ elsewhere in which temperature and humidity were inversely correlated with daily new  $\frac{397}{2}$ cases and deaths of COVID-19 with several studies reporting that  $SARS-CoV-2$  is  $\qquad \qquad \text{398}$ sensitive to high temperatures and humidity  $[34, 35]$  $[34, 35]$ . It is likely that rainfall events prior  $\frac{399}{2}$ to or during the sampling phase may have contributed to the lack of detection of  $\frac{400}{400}$  $SARS-CoV-2 RNA due to the dilutional effect. The substantial variation in parameter  $401$$ values across the ten geographic zones, as detailed in Table 3, is a consequence of the  $\frac{402}{402}$ inherent complexity and diversity of real-world conditions being modelled. These  $403$ variations are influenced by factors such as population density, healthcare infrastructure, <sup>404</sup> interventions, and social behaviours specific to each zone. While these differences may <sup>405</sup> appear significant, they are expected in epidemiological modelling and reflect the diverse nature of disease spread in different settings. Rather than indicating issues with  $_{407}$ the model, these variations underscore the need for tailored, context-specific modelling  $\frac{408}{408}$ to capture the nuanced dynamics within each zone accurately. This diversity in parameters enhances the model's ability to represent the unique characteristics of each <sup>410</sup> zone. Moreover, the calculation of  $R_0$  considers the complex interplay of these  $411$ parameters, and the model offers valuable insights into the dynamics of COVID-19 <sup>412</sup> within a geographically diverse urban area like Nagpur. The variation in  $R_0$  estimates  $\frac{413}{413}$  $(0.98-1.66)$  between the different zones in Nagpur urban district may be due to  $414$ additional factors such as variation in socio-behavioural habits (personal hygiene, <sup>415</sup> wearing masks, handwashing, social distancing, vaccine uptake, social gatherings), <sup>416</sup> sociodemographic, educational levels and dietary factors. Factors such as high levels of  $\frac{417}{417}$ youth, income inequality, high population density and social media usage are associated  $\frac{418}{418}$ with high  $R_0$  and may be important influences shaping zonal-wise variation in  $R_0$  in  $\phantom{R_0}$ Nagpur as reported across countries [\[36\]](#page-16-9). Overall, these  $R_0$  estimates for the second  $\frac{420}{420}$ wave of COVID-19 in India are consistent with a baseline  $R_0$  of 1.450 recorded for  $\frac{421}{221}$ Maharashtra and 1.379 for India by Marimuthu *et al.* [\[37\]](#page-16-10) but fall below earlier  $422$ estimates calculated by Shil et al who reported  $R_0$  in the range of 2-3 during the initial  $\frac{423}{423}$ wave of infection for the majority of Indian districts (March-June 2020) [\[38\]](#page-17-1). This  $424$ depicts that the use of 2011 population census data as a proxy for the modelling process  $\frac{425}{425}$ in this study was not out of place as the estimated  $R_0$  in this study is consistent with  $\frac{426}{4}$ what other studies have found. This feasibility study identified a unique set of  $\frac{427}{427}$ challenges in the implementation of WBE in Central India which mirrors those observed  $_{428}$ in other LMIC settings such as Bangladesh  $[4]$ . These include establishing a sampling  $429$ plan and schedule that is representative of the different urban and rural catchment <sup>430</sup> populations, underdeveloped sewage systems in rural areas necessitating onsite <sup>431</sup> sanitation epidemiology/sampling; development and validation of standardized protocols  $\frac{432}{4}$ for lab analysis; complex collaborative efforts from government agencies, public health  $\frac{433}{433}$ units and academia and resource limitations (e.g., autosamplers not suitable for large  $_{434}$ rapid monitoring where passive sampling techniques are more easily implemented) [\[39\]](#page-17-2). <sup>435</sup> Supply chain issues for essential goods such as PPE and PCR diagnostic reagents, and <sup>436</sup> logistical constraints such as inaccessibility and poor transport systems made it difficult  $_{437}$ to reach rural communities in remote areas. In recognition of these challenges, we acknowledge several study limitations. Although we did assess and compare the  $\frac{439}{439}$ abundance of SARS-CoV-2 viral concentration in untreated wastewater samples <sup>440</sup> between urban and rural areas, in line with other wastewater research studies in <sup>441</sup> India [\[40,](#page-17-3) [41\]](#page-17-4) most of our sampling sites were from urban zones of Nagpur, introducing  $\frac{442}{4}$ sampling bias. Due to the lack of COVID-19 clinical incident data for the rural areas  $\frac{443}{4}$  sampled, we were not able to apply our SEIPR model to model infectious burden in rural Nagpur. We also had to base our model assumptions on historical rather than <sup>445</sup> current census data which is not available from Nagpur district. Due to the limitation <sup>446</sup> of resources and skilled personnel, we were not able to undertake 24-hour composite and <sup>447</sup> longitudinal sampling which we recognize would have made our data more representative, to assess the impact of seasonality or to obtain detailed information on <sup>449</sup> spatiotemporal trends. Moreover, we were unable to record physiochemical, hydrologic,  $\frac{450}{200}$ and anthropogenic parameters of the wastewater samples which would have affected  $_{451}$ RNA concentrations, and consequently, SARS-CoV-2 RNA detection [\[31\]](#page-16-4). Although we  $\frac{452}{452}$ did not collect daily rainfall measurements and instead used relative humidity as a  $\frac{453}{453}$ proxy for rainfall, the majority of the sampling period was conducted during periods of <sup>454</sup> no rain. We acknowledge the use of air temperature as a surrogate for wastewater  $\frac{455}{455}$ temperature in the absence of direct wastewater temperature data, particularly in open <sup>456</sup> drainage systems. While this substitution is a common practice in environmental <sup>457</sup> modelling due to data limitations, it's essential to recognize its potential limitations and  $\frac{458}{458}$ the possible impact on the results. Wastewater temperature can be influenced by  $\frac{459}{459}$ various factors beyond just air temperature in open drainage systems, such as ground  $460$ temperature, flow rates, and interactions with other environmental factors. This  $\frac{461}{461}$ assumption may introduce some level of uncertainty into the model, and future studies <sup>462</sup> should aim to collect specific wastewater temperature data to improve the accuracy of  $\frac{463}{463}$ the modelling. However, given the data constraints, the use of air temperature can provide a reasonable estimation of wastewater temperature and is a common approach <sup>465</sup> in the field. We recognize that with any modelling efforts, it will be important to <sup>466</sup> explore the sensitivity of the model to different assumptions in future research. Future  $_{467}$ studies should also adopt the use of rapid in-field testing of SARS-CoV-2 or any  $\frac{468}{468}$ pathogenic target as opposed to bringing samples back to a central lab with <sup>469</sup> appropriately trained personnel. This technology is already in proof-of-concept stages  $470$ and could be easily operationalized ahead of future outbreaks or pandemics.

# Conclusion  $\frac{472}{472}$

We have established a quantitative framework to estimate COVID-19 prevalence and  $473$ predict SARS-CoV-2 transmission through integrating wastewater-based surveillance <sup>474</sup> data into a SEIPR model. The constructed model may be used to provide accurate and <sup>475</sup> robust estimates of future waves of the COVID-19 pandemic and could usefully be  $476$ applied to study other infectious diseases or expanded to consider reinfected <sup>477</sup> populations. Our findings showcase the translational value of utilizing WBE to study <sup>478</sup> the health of a population for epidemiological inference and in informing public health  $_{479}$ actions, particularly where comprehensive individual testing is severely constrained by a <sup>480</sup> shortage of resources and logistical challenges. However, to realize the true value of this  $_{481}$ tool in India and other LMICs, it will be important for governmental and other funding  $_{482}$ agencies to invest heavily in building laboratory capacity and sample collection teams.  $\frac{483}{1000}$ Such efforts should also help re-emphasize the criticality of clean water, sanitation, and  $484$ waste management as potential control points in the fight against COVID-19 and future  $\frac{485}{485}$  $\mathbf{p}$ andemics.  $\begin{array}{ccc} & & & \text{if } &$ 

# Supporting information

<span id="page-13-0"></span>S1 Appendix. Pre-processing of wastewater samples, nucleic acid extraction and  $SARS-CoV-2$  qualitative and quantitative detection.  $(PDF)$  490

<span id="page-14-9"></span><span id="page-14-8"></span>

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