

Optimal constant picewise vaccination policies for COVID-19

SDIV, FPA, DGS, GASV

Abstract

BACKGROUND.
FINDINGS.
IMPLICATIONS.

1. Introduction

Main contribution and its relevance.

Background.

Vaccine development.

Problem setup.

Litterature review.

Papaer structure.

2. Covid-19 spread dynamics

Uncontrolled dynamics. We split the the constant population N in a base SEIR structure with segregation infected classes according with manifestation of symptoms. We also postulate the extra state Y_{IS} to fit commulative symptomatic cases reported in the databases from Mexico city during the exponential grow phase. Our dynamics reads

$$\begin{aligned} L' &= \theta\mu N^* - \epsilon\lambda L - \delta_L L - \mu L, \\ S' &= (1 - \theta)\mu N^* + \delta_L L + \delta_R R - (\lambda + \mu)S, \\ E' &= \lambda(\epsilon L + S) - (\kappa + \mu)E, \\ I_S' &= p\kappa E - (\gamma_S + \delta_H + \mu_{IS}^{\text{SDIV}} + \mu)I_S, \\ I_A' &= (1 - p)\kappa E - (\gamma_A + \mu)I_A, \\ H' &= \delta_H I_S - (\gamma_H + \mu_H + \mu)H, \\ R' &= \gamma_S I_S + \gamma_A I_A + \gamma_H H - (\delta_R + \mu)R, \\ D' &= \mu_{IS}^{\text{SDIV}} I_S + \mu_H H, \\ \frac{dY_{IS}}{dt} &= p\kappa E, \\ \lambda &:= \frac{\beta_A I_A + \beta_S I_S}{N^*}, \\ N^*(t) &= L + S + E + I_S + I_A + H + R. \end{aligned} \tag{1}$$

See table[*] for notation and values.

Hypothesis. We consider that susceptible individuals become infected when they are in contact with asymptomatic individuals and individuals with symptoms, we will propose that a proportion of asymptomatic individuals have a way to get relief and not die. A proportion of individuals infected with symptoms may die from the disease or may be relieved.

We callibrate parameters of our base dynamics in (1) via Multichain Montecarlo (MCMC). To this end, we assume that the comulative incidence of new infected symptomatic cases CI_S follows a Poisson distribution with mean $\lambda_t = IC_s(t)$. Further, following [] we postulate priors for p and κ

$$\begin{aligned} Y_t &\sim \text{Poisson}(\lambda_t), \\ \lambda_t &= \int_0^t p\delta_e E, \\ p &\sim \text{Uniform}(0.3, 0.8), \\ \kappa &\sim \text{Gamma}(10, 50). \end{aligned} \tag{2}$$

Using the reproductive number definition of VanDenDrishe, we obtain

$$R_0 := \frac{N^*(\beta_S p \kappa + \beta_A \kappa (1 - p))}{(\mu - \kappa)(\gamma_S + \mu_{IS} + \gamma_A + \mu)N^* \mu}$$

Figure [...] displays data of coumulative confirmed cases of COVID-19 of Mexico city, and the fitt of our model in eqs. (1) and (2).

3. Imperfect-preventive Covid-19 vaccination

Preventive vaccines.

Efficacy and vaccine-induced immunity.

Actual vaccine stage development.

Vaccination reproductive number.

Vaccination rate λ_V estimate.

Parameter	Description
μ	Death rate
β_S	Infection rate between susceptible and symptomatic infected
β_A	Infection rate between susceptible and asymptomatic infected
λ_V	Vaccination rate
δ_V^{-1}	Vaccine-induced immunity
ε	Vaccine efficacy
κ^{-1}	Average incubation time
p	New asymptomatic generation proportion
θ	Proportion of individuals under lockdown
γ_S^{-1}	Average time of symptomatic recovery
γ_A^{-1}	Recovery average time of asymptomatic individuals
γ_H^{-1}	Recovery average time by hospitalization
δ_R^{-1}	Natural immunity
δ_H	Infected symptomatic hospitalization rate

Table 1: Parameters definition of model in eq. (1).

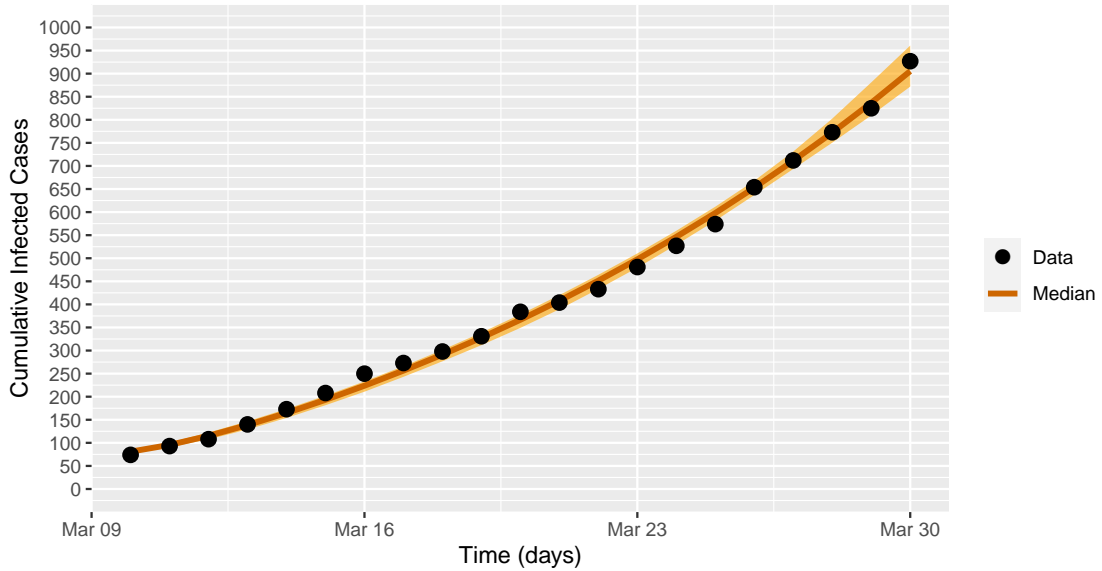


Figure 1: Fit of diary new cases of Mexico city during exponential growth.

Feasibility regions according to efficacy and vaccination rate.

$$\begin{aligned}
L' &= \theta \mu N^* - (\epsilon \lambda + \delta_L + \mu) L \\
S' &= (1 - \theta) \mu N^* + \delta_L L + \delta_V V + \delta_R R \\
&\quad - (\lambda + \lambda_V + \mu) S \\
E' &= \lambda (\epsilon L + (1 - \varepsilon) V + S) - (\kappa + \mu) E \\
I_S' &= p \kappa E - (\delta_H + \gamma_S + \mu_{I_S} + \mu) I_S \\
I_A' &= (1 - p) \kappa E - (\gamma_A + \mu) I_A \\
H' &= \delta_H I_S - (\gamma_H + \mu_H + \mu) H \\
R' &= \gamma_S I_S + \gamma_A I_A + \gamma_H H - (\delta_R + \mu) R \\
D' &= \mu_{I_S} I_S + \mu_H H \\
V' &= \lambda_V S - [(1 - \varepsilon) \lambda + \delta_V + \mu] V
\end{aligned}$$

4. Optimal controlled version

Controlled Model. Now we model vaccination, treatment and lockdown as an optimal control problem. According to dynamics in eq. (1), we modulate the vaccination rate by a time-dependent control signal $u_V(t)$. We add compartment X_{vac} to count all the vaccine applications of susceptible, exposed, asymptomatic and recovered individuals. This process is modeled by

$$X'(t) = (\lambda_V + u_V(t))(S + E + I_A + R) \quad (4)$$

and describes the number of applied vaccines at time t . Consider

$$x(t) := (L, S, E, I_S, I_A, H, R, D, V, X_{vac})^\top(t)$$

$$\begin{aligned}
\frac{dX_{vac}}{dt} &= (u_V(t) + \lambda_V) [L + S + E + I_A + R] \\
\frac{dY_{I_S}}{dt} &= p \kappa E \\
\lambda &:= \frac{\beta_A I_A + \beta_S I_S}{N^*}
\end{aligned}$$

and control signal $u_v(\cdot)$. We quantify the cost and reward of a vaccine strategy policy via the penalization functional

$$J(u_V) := \int_0^T a_S I_S + a_D D + \frac{1}{2} c_V u_v^2 ds. \quad (5)$$

In other words, we assume in functional J that pandemic cost is proportional to the symptomatic and death reported cases and that a vaccination policy implies quadratic consumption of resources.

Further, since we aim to simulate vaccination policies at different coverage scenarios, we impose the vaccination counter state's final time condition $X(T)$

$$\begin{aligned} x(T) &= (\cdot, \cdot, \cdot, \cdot, \cdot, X_{vac}(T))^\top, \in \Omega \\ X_{vac}(T) &= x_{coverage}, \\ x_{coverage} &\in \{\text{Low}(0.2), \text{Mid}(0.5), \text{High}(0.8)\}. \end{aligned} \quad (6)$$

Thus, given the time horizon T , we impose that the last fraction of vaccinated populations corresponds to 20%, 50% or 80%, and the rest of final states as free. We also impose the path constraint

$$\Phi(x, t) := \kappa I_S(t) \leq B, \quad \forall t \in [0, T], \quad (7)$$

to ensure that healthcare services will not be overloaded. Here κ denotes hospitalization rate, and B is the load capacity of a health system.

Given a fixed time horizon and vaccine efficiency, we estimate the constant vaccination rate as the solution of

$$x_{coverage} = 1 - \exp(-\lambda_V T). \quad (8)$$

That is, λ_v denotes the constant rate to cover a fraction $x_{coverage}$ in time horizon T . Thus, according to this vaccination rate, we postulate a policy u_v that modulates vaccination rate according to λ_V as a baseline. That is, optimal vaccination amplifies or attenuates the estimated baseline λ_V in a interval $[\lambda_v^{\min}, \lambda_v^{\max}]$ to optimize functional $J(\cdot)$ —minimizing symptomatic, death reported cases and optimizing resources.

Our objective is minimize the cost functional (5)—over an appropriated functional space—subject to the dynamics in eqs. (1) and (4), boundary conditions, and the path constrain in (7). That is, we search for vaccination policies $u_V(\cdot)$, which solve the following optimal control problem

(OCP).

$$\begin{aligned} \min_{u \in \mathcal{U}} J(u) &:= \int_0^T [(a_D \mu_s + a_H \delta_H) I_S(r) + a_{I_S} p \kappa E(r)] dr \\ \text{s. t.} \\ L' &= \theta \mu N^* - \epsilon \lambda L - u_L(t) L - \mu L \\ S' &= (1 - \theta) \mu N^* + u_L(t) L + \delta_v V + \delta_R R \\ &\quad - [\lambda + (\lambda_V + u_V(t)) + \mu] S \\ E' &= \lambda(\epsilon L + (1 - \epsilon) V + S) - (\kappa + \mu) E \\ I_S' &= p \kappa E - (\gamma_S + \mu_{I_S} + \delta_H + \mu) I_S \\ I_A' &= (1 - p) \kappa E - (\gamma_A + \mu) I_A \\ H' &= \delta_H I_S - (\gamma_H + \mu_H + \mu) H \\ R' &= \gamma_S I_S + \gamma_A I_A + \gamma_H H - (\delta_R + \mu) R \\ D' &= \mu_{I_S} I_S + \mu_H H \\ V' &= (\lambda_V + u_V(t)) S - [(1 - \epsilon) \lambda + \delta_V + \mu] V \end{aligned}$$

$$\begin{aligned} \frac{dX_{vac}}{dt} &= (u_V(t) + \lambda_V) [L + S + E + I_A + R] \\ \frac{dY_{I_S}}{dt} &= p \kappa E \\ \lambda &:= \frac{\beta_A I_A + \beta_S I_S}{N^*} \end{aligned}$$

$$\begin{aligned} L(0) &= L_0, \quad S(0) = S_0, \quad E(0) = E_0, \quad I_S(0) = I_{S_0}, \\ I_A(0) &= I_{A_0}, \quad H(0) = H_0, \quad R(0) = R_0, \quad D(0) = D_0, \\ V(0) &= 0, \quad X_{vac}(0) = 0, \quad u_V(\cdot) \in [u_{\min}, u_{\max}], \\ X_{vac}(T) &= x_{coverage}, \quad \kappa I_S(t) \leq B, \quad \forall t \in [0, T], \\ N^*(t) &= L + S + E + I_S + I_A + H + R + V \end{aligned} \quad (9)$$

Bayesian estimation.

5. Parameters

6. Optimal control problem

7. Numerical Results

Bayesian.

8. Appendix

Consider the following cost functional that we want to minimize

$$\int_0^T C(t, X(t), u(t)) dt \quad (10)$$

subject to the dynamics

$$\dot{X}(t) = f(t, X(t), u(t)), \quad 0 \leq t \leq T, \quad (11)$$

and the initial state $X(0) = x_0$. Let $t_0 < t_1 < \dots < t_n$, with $t_0 = 0$ and $t_n = T$, be a partition of the interval $[0, T]$. We consider *piecewise constant controls* \tilde{u} of the form

$$\tilde{u}(t) = a_j \quad t_j \leq t < t_{j+1} \quad (12)$$

Parameter	Median	Reference
q_r, ϵ	0.4, 0.3, 0.1	this study
β_S	$q_r \times 8.690\,483 \times 10^{-1}$	this study
β_A	$q_r \times 7.738\,431 \times 10^{-1}$	this study
κ	0.196\,078\,43	*
p	0.1213	*
θ	0.2,	this study
δ_L	0.04	postulated
δ_H	0.2	*
δ_V	0.002\,739\,726\,027\,397\,260\,3	$\delta_V^{-1} = 2$ years CanSinoBIO
δ_R	0.005\,555\,56	$\delta_R^{-1} \approx 180$ days
μ	$3.913\,894 \times 10^{-5}$	**
μ_{I_S}	0.0	
μ_H	0.016\,32	[FENG]
γ_S	0.092\,506\,94	*
γ_A	0.167\,504\,19	*
γ_H	$5.079\,869 \times 10^{-1}$	*
λ_V	0.000\,611\,35	
ε	0.7, 0.80, 0.9, 0.95	[PRESS RELESASES]
N	26\,446\,435	**
L_0	0.266\,260\,097\,021\,127\,96	
S_0	0.463\,606\,046\,009\,872	
E_0	0.000\,670\,33	*
I_{S_0}	9.283×10^{-5}	* * *
I_{A_0}	0.001\,209\,86	*
H_0	$1.341\,579\,69 \times 10^{-4}$	**
R_0	$2.661\,259\,39 \times 10^{-1}$	
D_0	0.001\,900\,74	**
X_{vac}^0	0.0	
V_0	0.0	
$Y_{I_S}^0$	0.122\,581\,64	
B	0.000\,359\,216\,658\,124\,242\,5	9500 beds/ N
a_{I_S}	0.002\,012\,775\,543\,825\,648\,6	DALY def
a_H	0.001\,411\,888\,738\,103\,725, or $a_H(x) := 0.001\,411\,888\,738\,103\,725 \log(\frac{1}{B-\kappa I_S})$	DALY def [Jo 2020]
a_D	7.25	DALY def

for $j = 0, \dots, n-1$.

ASSUMPTION 1.

ASSUMPTION 2.

By Assumption 1, the system

$$\dot{X}(t) = f(t, X(t), a_0), \quad X(0) = x_0, \quad 0 \leq t \leq t_1,$$

has a unique solution $\tilde{X}_0(t; x_0, a_0)$ which is continuous in (x_0, a_0) . Next, put $x_1 := \tilde{X}_0(t_1; x_0, a_0)$ and consider the system

$$\dot{X}(t) = f(t, X(t), a_1), \quad X(t_1) = x_1, \quad t_1 \leq t \leq t_2,$$

which, again by Assumption 1, has a unique solution $\tilde{X}_1(t; x_1, a_1)$ continuous in (x_1, a_1) . By following this procedure, we end up having a recursive solution

$$\tilde{X}_{n-1}(t; x_{n-1}, a_{n-1}), \quad x_{n-1} := \tilde{X}_{n-2}(t_{n-1}; x_{n-2}, a_{n-1}),$$

Thus, for a control \tilde{u} of the form (12) and the corresponding solution path \tilde{X} , we have

$$\int_0^T C(t, \tilde{X}(t), \tilde{u}(t)) dt = \sum_{j=0}^{n-1} \int_{t_j}^{t_{j+1}} C(t, \tilde{X}_j(t), a_j) dt.$$

Notice that each \tilde{X}_j is a continuous function of (a_0, \dots, a_j) and x_0 . Therefore, by Assumption 2, the mapping

$$(a_0, \dots, a_{n-1}) \mapsto \sum_{j=0}^{n-1} \int_{t_j}^{t_{j+1}} C(t, \tilde{X}_j(t), a_j) dt$$

is continuous.

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