Un modello Agent-Based per studiare la diffusione del virus SARS-CoV-2

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IRFS Piemonte - 23 Febbraio 2021

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Objectives of the model

- We propose an agent-based model to simulate the Covid-19 epidemic diffusion, with Susceptible, Infected, symptomatic, asymptomatic, and Recovered people: hence the name S.I.s.a.R. The scheme comes from S.I.R. models, with (i) infected agents categorized as symptomatic and asymptomatic and (ii) the places of contagion specified in a detailed way, thanks to agent-based modeling capabilities.
- The infection transmission is related to three factors: the infected person's characteristics and those of the susceptible one, plus those of the space in which contact occurs.
- The model includes the structural data of Piedmont, but it can be readily calibrated for other areas. The model manages a realistic calendar (e.g., national or local government decisions), via a script interpreter.

Tool and links

- We use NetLogo, at https://ccl.northwestern.edu/netlogo/.
- S.I.s.a.R. is at https://terna.to.it/simul/SIsaR.html with information on model construction, the draft of a paper also reporting results, and an online executable version of the simulation program, built using NetLogo.
- A short paper related to this presentation is at https://rofasss.org/2020/10/20/sisar/
 G. Pescarmona, P. Terna, A. Acquadro, P. Pescarmona, G. Russo, and S. Terna. How Can ABM Models Become Part of the Policy-Making Process in Times of Emergencies—The SISAR Epidemic Model. *RofASSS*, 2020.

The scale and the items

- 1:1000.
- Houses.
- Schools.
- Hospitals.
- Nursing homes,
- Factories.

The interface and the information sheet

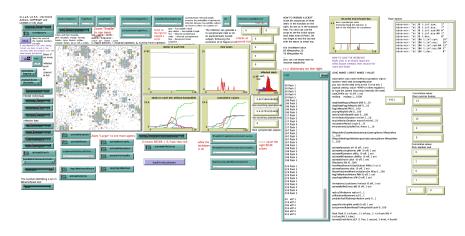


Figure 1: The interface

The interface and the information sheet

WHY THE NAME?	VIEWING THE MODEL		
S - Susceptible I - Infected s - symptomatic a - asymptomatic R - Recovered	In the deak version, we can use both the continuous view, observing all the agents' movements and the "on tick" one, leaser but updating the screen only at the end of each tick. To set the view, we use a chooser in the top part of the sort. In the web version we have uniquely the on fick feature.		
with capital letters referring the classic <u>S.I.R. model</u> , also in <u>Wikipadia</u> .	The desk version also allows 3D view: right click on the map of the world and choose Switch to 3D View.		
Website	HOW IT WORKS		
The model has a website.	Scripting capability		
PRELIMINARY NOTE			
This is a simulation with random events, please do not take it as a sure forecasting machine; it is a reasoning machine, a cort of very complex "what!" mental experience. The New Yor Time Shore sure an explayer to the Time Cond.15 Bibliot. Why Doos the Virus Mallins Sures. Exists and Shore Chemical Shore	The code can manage a script to set the parameter modifications occurring while the simulation is running. Explanations in the right side of the interface. The simulation starts at list 1, but we can already set the initial values at list 0. A brick () via setting set all the distall values for the experiments at tick 0 and then (ii) modify those that we want change before filting go.		
to be some combination of the above factors, as well as one other mentioned by researchers: sheer luck. In the same way, in the simulations run with this model we can have very different outcomes as we change the	Special item "flash"		
in the same way, in the simulations run with this model we can have very distinct outcomes as we change the initial seed of the random numbers. Those values are determining mainly the movements at a tiny scale and so the interactions-infections chains.	With the "flash" item, followed by 1 or 2 or 3, we obtain a flash output in the window to the far right, with data the beginning of the tack for (1) total infected symptomatic people; (2) total infected symptomatic people in NH; (4) total infected symptomatic people in NH; (4) total infected symptomatic people in NH; (5) total infected symptomatic people in NH; (6) total infected symptomatic people in NH; (7) total infected symptomatic people in NH; (8) total infected symptomatic people in NH; (9) total infected symptomatic people in NH; (1) total infecte		
We can use the model in a comparative way, observing different range of results with different initial conditions (parameters).			
Finally, to have a reference at an actual situation, the model is related to the Piedmont scale, with 4,350 agents vs. 4.35 millions of inhabitants. The scale 1 to 1000 is over-represented in the case of schools, with their	Special item "activate check point"		
vs. 4.5 millions on imbolants. The scale it is 1000 is over-represented in the case of schools, with their classrooms with a realistic number of students, apartments with a realistic quantity of inhabitants, and likewise workspaces, hospitals, nursing homes.	With the "aCP" (activate checkpoint) item, followed by n with n in [1,6] as closed interval, we collect data at the		

Figure 2: The information sheet, about 20 pages

We look also to the time series of the total infected people in Piedmont.

tick of the aCP command. In BehaviorSpace, we can send (or not) the checkpoint data to the table of the results; we have six possible checkpoints, each reporting stocks of infected symptomatic people, infected

asymptomatic people, deceased people. The data are at the beginning of the tick,

The world

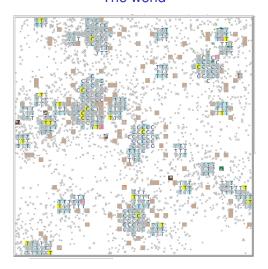


Figure 3: The world



The world 3D

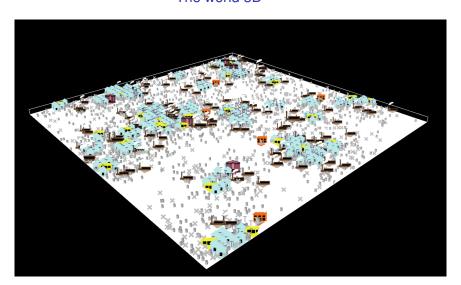


Figure 4: The world 3D



The agents





Figure 5: Probes to different agents

Contagion representation

- The model allows analyzing the sequences of contagions in simulated epidemics, reporting the places where the contagion occur.
- We represent each infecting agent as a horizontal segment with a vertical connections to another agent receiving the infection. We represent the infected agents via further segments at an upper layer.
- With colors, line thickness, and styles, we display multiple information.
- This enables understanding at a glance how an epidemic episode is developing.
 In this way, it is easier to reason about countermeasures and, thus, to develop intervention policies.

An example (1/2)

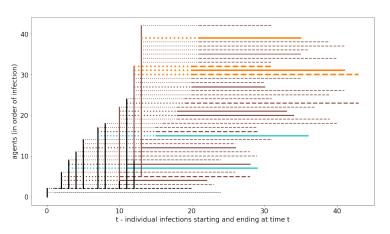


Figure 6: A case with containment measures, first 40 infections: workplaces (brown) and nursing homes (orange) strictly interweaving

An example (2/2), more contagions

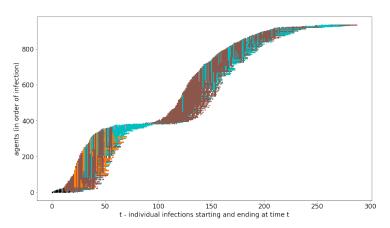


Figure 7: A Case with containment measures, the whole epidemics: workplaces (brown) and nursing homes (orange) and then houses (cyan), with a bridge connecting two waves

Other examples (i) on the left, an epidemic without containment measures; (ii) on the right, an epidemic with basic non-pharmaceutical containment measures

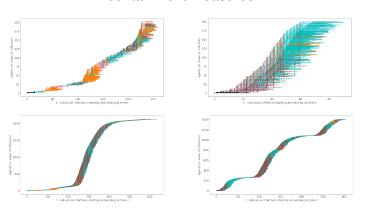


Figure 8: Two cases with initial and full periods

A contagion sequence suggesting policies: in Fig. 9 we can look both at the places where contagions occur and at the dynamics emerging with different levels of intervention.

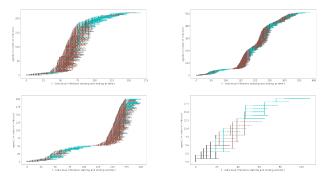


Figure 9: (top left) an epidemic with regular containment measures, showing a highly significant effect of workplaces (brown); (top right) the effects of stopping fragile workers at day 20, with a positive result, but home contagions (cyan) keep alive the pandemic, exploding again in workplaces (brown); (bottom left) the same analyzing the first 200 infections with evidence of the event around day 110 with the new phase due to a unique asymptomatic worker, and (bottom right) stopping fragile workers and any case of fragility at day 15, also isolating nursing homes

Simulation batches

- We explore systematically the introduction of factual, counterfactual, and prospective interventions to control the spread of the contagions.
- Each simulation run—whose length coincides with the disappearance of symptomatic or asymptomatic contagion cases—is a datum in a wide scenario of variability in time and effects.
- Consequently, we need to represent compactly the results emerging from batches
 of simulation repetitions, to compare the consequences of the basic assumptions
 adopted for each specific batch.
- We use blocs of ten thousand repetitions. Besides summarizing the results with the usual statistical indicators, we adopt the technique of the heatmaps.
- Each heatmap reports the duration of each simulated epidemic in the x axis and the number of the symptomatic, asymptomatic, and deceased agents in the y axis. The z axis is represented by the colors, in logarithmic scale.
- In our batches we have 10,000 runs.

Ten thousand epidemics without control in Piedmont

	symptomatic	totalInfected&Deceased	duration
count	10000.00	10000.00	10000.00
mean	969.46	2500.45	303.10
std	308.80	802.88	93.50

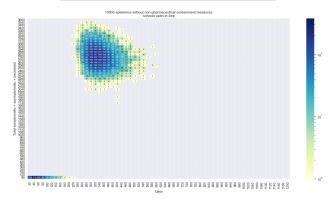


Figure 10: Without non-pharmaceutical containment measures

Ten thousand epidemics with basic control in Piedmont

	symptomatic	totalInfected&Deceased	duration
count	10000.00	10000.00	10000.00
mean	344.22	851.64	277.93
std	368.49	916.41	213.48

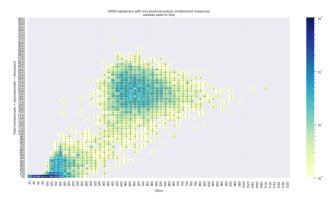


Figure 11: First wave with non-pharmaceutical containment measures

Actual data

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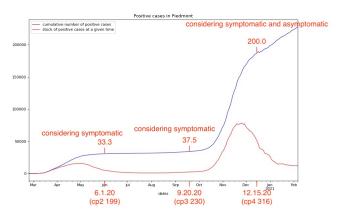


Figure 12: Data in Piedmont

Selecting realistic spontaneous second waves

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	symptomatic Dec20	totalInfected&Deceased Dec20	duration
count	140.00	140.00	170.00
mean	605.79	1528.31	535.19
std	264.29	644.34	167.42

170 epidemics out of 10,000 stable in Summer 2020, with: at 6.1.20 select sym. (10, 70] actual v. 33.3 & at 9.20.20 select sym. (20, 90] actual value 37.5. 140 residual epidemics at 12.15.20 (actual symptomatic + asymptomatic people: 200.0).



Figure 13: First wave with non-pharmaceutical containment measures, spontaneous second wave, without specific measures

Selecting realistic second waves, with new infections from outside

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	symptomatic Dec20	totalInfected&Deceased Dec20	duration
count	1044.00	1044.00	1407.00
mean	588.67	1474.10	527.85
std	251.96	618.87	184.76

1,407 epidemics out of 10,000 stable in Summer 2020, with: at 6.1.20 select sym. (10, 70] actual v. 33.3 & at 9.20.20 select sym. (20, 90] actual value 37.5. 1,044 residual epidemics at 12.15.20 (actual symptomatic + asymptomatic people: 200.0).

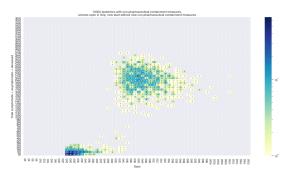


Figure 14: First wave with non-pharmaceutical containment measures, forcing the second wave, without specific measures

Selecting realistic second waves, with new infections from outside

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	symptomatic Dec20	totalInfected&Deceased Dec20	duration
count	874.00	874.00	1407.00
mean	223.61	594.97	404.36
std	138.52	372.63	137.51

1,407 epidemics out of 10,000 stable in Summer 2020, with: at 6.1.20 select sym. (10, 70] actual v. 33.3 & at 9.20.20 select sym. (20, 90] actual value 37.5. 874 residual epidemics at 12.15.20 (actual symptomatic + asymptomatic people: 200.0).



Figure 15: First wave with non-pharmaceutical containment measures, forcing the second wave, with new specific non-pharmaceutical containment measures

Selecting realistic second waves, with new infections from outside

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	symptomatic Dec20	totalInfected&Deceased Dec20	duration
count	987.00	987.00	1407.00
mean	286.09	753.06	443.05
std	164.33	424.90	151.25

1,407 epidemics out of 10,000 stable in Summer 2020, with: at 6.1.20 select sym. (10, 70] actual v. 33.3 & at 9.20.20 select sym. (20, 90] actual value 37.5. 987 residual epidemics at 12.15.20 (actual symptomatic + asymptomatic people: 200.0).



Figure 16: First wave with non-pharmaceutical containment measures, forcing the second wave; in second wave, uniquely stop to fragile people of any kind (including workers)

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To recap

Scenarios		total sym.	total sym., asympt., deceased	duration
no control in first wave				
	count	10000.00	10000.00	10000.00
	mean	969.46	2500.45	303.10
	std	308.80	802.88	93.50
basic controls in first wave				
	count	10000.00	10000.00	10000.00
	mean	344.22	851.64	277.93
	std	368.49	916.41	213.48
basic controls in first wave				
forcing realistic second	count	1044.00	1044.00	1407.00
wave, without new controls	mean	588.67	1474.10	527.85
,	std	251.96	618.87	184.76
basic controls in first wave				
forcing realistic second	count	874.00	874.00	1407.00
wave, with new controls	mean	223.61	594.97	404.36
,	std	138.52	372.63	137.51
basic controls in first wave				
forcing realistic second	count	987.00	987.00	1407.00
wave, with stop to frag.	mean	286.09	753.06	443.05
	std		424.90	151.25
people (incl. workers)	std	164.33	424.90	151.

Table 1: Report of the key results, with count, mean, and std

Exploring vaccinations

Exploring vaccination sequences (using genetic algorithms or reinforcement learning).

A new model: the map

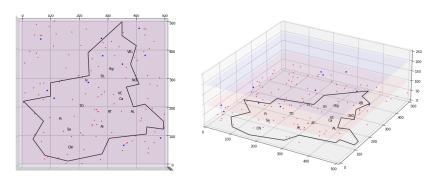


Figure 17: 3D Piedmont

A new model: the scale and the items

- 1:100.
- Infection engine, https://terna.to.it/simul/InfectionEngine.pdf.
- Houses.
- Schools.
- Hospitals.
- · Nursing homes,
- · Factories.
- Transportations.
- Aggregation places: happy hours, night life, sport stadiums, discotheques, ...

The tool: S.L.A.P.P.

Scientific advertising: https://terna.github.io/SLAPP/

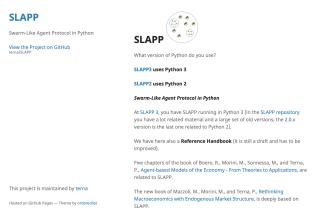


Figure 18: Swarm-Like Agent Protocol in Python

A few considerations:

- The importance of High Performance Computing.
- From S.I.s.a.R. to a model of the society and of the economy of Piedmont.
- The S.I.s.a.R. model is a tool for comparative analyses, not for forecasting (the enormous standard deviation values are intrinsic to the problem).
- The model is highly parametric and more it will be.
- New crisis calling for immediate simulation could take a substantial advantage from the parametric structure of the model.

The slides are at https://terna.to.it/simul/TernaIRES20210223.pdf.

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The image, elaborating Fig. 17, is dedicated to Pietro Greco (Barano d'Ischia, April 20th, 1955—Ischia, December 18th, 2020)