# **Epidemics** logistics

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natural causes (e.g. influenza outbreak)



deliberate terrorist actions (bioterrorism)





natural disasters

man-made disasters

**Epidemics Logistics** 

# Introduction



"Nothing for you to worry about. We're cleaning students' lockers."

Containment of an outbreak:

- Non-pharmaceutical interventions
- Pharmaceutical interventions
- Harmonized approaches.



The containment effort consists of four **phases**: Preparedness - Outbreak investigation – Response - Evaluation



# **Epidemics control and logistics operations: Preparedness**

- Maintaining a certain level of available resources so as to reduce morbidity and mortality when an outbreak occurs
- Pharmaceuticals and supplies in large quantities in order to assist a prompt response, if necessary
- Procurement of vaccines and medical supplies and their exact storing location play a crucial role.





# **Epidemics control and logistics operations:**

# **Outbreak investigation**

- Detection of any suspected outbreak and its confirmation through laboratory testing.
- Surveillance systems must be placed to provide the essential information regarding any unexplained infection increases seen over a period of time.
- Determine the type and magnitude of the containment effort once epidemic thresholds have been reached.



A global view of HIV infection 39.5 million people [34.1-47.1] living with HIV in 2006



The boundaries and names shown and the designations used on this map 4 not imply the copression of any opinion whatsever on the part of the Woold Health Cognation conventing the legal status of any country, twintry, city or area or of at authorises, or concerning the definitation of at forefires or boundaries. Dotted lines on maps sepresent approximate border lines for which there may not get be that agreement. Data Source: WHO / UNAIDS Map Production: Public Health Mapping and GIS Communicable Diseases (CDS) World Health Organization





#### **Epidemics Logistics**

# **Epidemics control and logistics operations: Response**

- Once an epidemic outbreak is confirmed, measures and control strategies must be implemented as soon as possible at a regional and/or national level.
- Treatment centers should be established and available resources such as medical supplies and personnel should be deployed rapidly in order to contain the epidemic.





# **Epidemics control and logistics operations: Evaluation**

- The evaluation phase is very useful as it provides strong insights towards a series of modifications that need to be made in order to increase the resilience of the control mechanisms in future epidemic outbreaks.
- Despite the fact that the evaluation phase entails limited physical movement of medical supplies and complementary commodities, it remains important from a logistical point of view.





# **Epidemics control and logistics operations:**

### Actions



# Establish an emergency supply chain

- Manufacturers should produce vaccines, antiretroviral drugs and complementary medical supplies
- Governments and public health institutions should purchase and stockpile well in advance
- Transportation and distribution of these supplies from central warehouses to regional store sites and then to local Points of Dispensing (PODs).



# **Epidemics control and logistics operations: Managerial issues**



- Affected people proceeding to treatment centers - patient flow operations - dispensing activities of the medical supplies
  - Reverse logistics activities (dangerous wastes must be treated carefully or disposed)
  - Coordination among manufacturers, governments, primary health care institutes, possibly military agencies, NGOs, etc.
  - Managing the information regarding the demand for medical supplies as well as the flow of funds.



# **Epidemics control and logistics operations: Materials Flow (Dasaklis et al., 2012)**



Materials flow of the epidemics control supply chain (End-to-End approach).



# Literature review

Classification based on a three-level assessment framework:

- First level: the various logistical features incorporated in each reference are classified according to the time framework in which they take place (pre-event or post-event)
- Second level: a context-specific classification is made where the logistics features are correlated to the nature of the outbreak
- Third level: the logistical features and the methodologies applied (qualitative or quantitative) for solving the problem tackled are classified.



# Literature review



A framework for epidemics logistics literature classification.



# Literature review

### Facts:

- Epidemic control measures may vary depending on the nature of the outbreak and their implementation time horizon
- Communicable disease outbreaks may also occur in the aftermath of natural disasters where certain sub-populations are dislocated or subject to a humanitarian crisis
- Disease outbreaks may also occur during big events and mass gatherings such as the Olympic Games
- The time frame in which control measures apply may also be different.



# Literature review: Essential question

How and to what extent the availability of certain quantities of essential medical supplies (vaccines) could affect the progression of a disease outbreak?

Need to define:

- the amount of commodities to be utilized during the control effort
- the selection of modes of transportation/distribution and relevant capacities
- the inventory level of medical supplies and commodities held in each node (facility) of the network
- the capacity of stockpile centers (National, Regional and Local)



# Literature review: Epidemics control network



**Epidemics control logistics network configuration.** 



### **Epidemics Logistics**

**OR/MS** contribution to resources' allocation

# and scheduling for epidemics control

### Problems:

- Limited resources' allocation among subpopulations
- Scheduling resources for epidemics control

•Etc.

### Commonly used objectives:

• Maximize the number of Quality Adjusted Life Years gained

• Maximize the number of infections averted (equivalently minimize the cumulative number of new infections).

### Techniques:

- Linear and integer programming models
- Numerical analysis procedures
- Cost-effectiveness analysis
- Simulation
- Non-linear optimization
- Control theory techniques
- Heuristic algorithms.



# **Problems addressed**

- 1. Controlling infectious disease outbreaks: A deterministic allocation-scheduling model with multiple discrete resources.
- 2. Emergency supply chain management for controlling a smallpox outbreak: The case for regional mass vaccination.



# Multiple resources problem (Rachaniotis et al., 2017)

Find the optimal schedule-allocation of limited discrete resources (mobile medical teams) employed in parallel in a time horizon to implement a vaccination campaign for infected subpopulations unable to proceed to vaccination centres either because they are house bound (elderly, incapacitated etc.) or they are in institutions in several distinct areas in order to minimize the total number of new infections (or, equivalently, to maximize the total number of infections averted).



# Assumptions

- The mobile medical teams can be considered as parallel (identical or non-identical) unrelated resources with constant service rates
- More than one medical team may be allocated to a specific regional population
- Pre-emption is not allowed
- Control actions rely on vaccination of specific groups of the population (house bound, institutionalized etc.)
- All the available medical teams at any time are employed for controlling the epidemic
- The resources' traveling times are assumed to be negligible.



# **Notation and objective**

Let



 $P = \{P_1, P_2, ..., P_n\}$  be the set of n populations in different regions, and let  $N_i$  be the size of  $P_i$ , i=1,...,n.  $t_0 > 0$  be the common for all populations time required for the resources to commence vaccination. t be the discrete time units (days).

 $t_{end}$  be the end of the vaccination campaign in all regions. This time is not known in advance, since it depends on whether additional (resources) medical teams become available and when (time and resource-dependent problem).  $m_t$  be the resources (medical teams) available at time t.

 $(r_1(t), r_2(t), ..., r_n(t))$  be the vector of the number of medical teams assigned for vaccination in every regional population at time *t*, where  $r_i(t) \in \{0, 1, ..., m_t\}$ , i=1, ..., n. This is the problem's decision vector variable.

 $R_{Ei}$  be the effective reproduction number in  $P_i$ .

 $I_i(r_i(t))$  be the number of new infections therefore infective in  $P_i$  at time t.

 $C(r_i(t))$  be the completion time of the vaccination campaign for controlling the epidemic in  $P_i$  at time t (i.e.  $R_{Ei} \le l$ ), having  $r_i(t)$  medical teams assigned to region *i*.

The objective is to minimize the total number of new infections, given the available number of mobile medical teams:

$$\min \sum_{t=t_0}^{t_{end}} \sum_{i=1}^{n} I_i(r_i(t))$$
  
s.t. 
$$\sum_{i=1}^{n} r_i(t) = m_t, t = t_0, t_0 + 1, \dots, t_{end}$$
$$r_i(t) \in \{0, 1, \dots, m_t\}, t = t_0, t_0 + 1, \dots, t_{end}$$

The number of new infections at any time instance can be calculated using as input any existing disease transmission model (from compartmental modeling to agent-based modeling approaches).

**Epidemics Logistics** 

# Solution methodology

Step 1: Allocate resources to populations according to the incremental algorithm (Shih, 1974) for solving the respective static discrete resource allocation problem. The vaccination time duration under the current assignment is calculated.

Step 2: Check whether the current resource allocation should be altered. The resource allocation changes in two cases: a) arrival of additional resources, b) the region's vaccination with the shortest completion time finishes. If yes, move to Step 3. If not, then the vaccination campaign is completed (time  $t_{end}$  is reached) and the algorithm ends calculating the total number of infected people. Step 3: Calculate new populations' susceptibles numbers and return to

Step 1.



# Example: A case of influenza in Greece



The SVEIR influenza epidemic model (Samsuzzoha et al., 2012).

Population is divided into five subgroups: susceptible (S), vaccinated (V), exposed (E), infective (I) and recovered (R).



# **Model's parameters**

- $\beta$ : Contact rate
- $\beta_E$ : Ability to cause infection by exposed individuals
- $\beta_I$ : Ability to cause infection by infectious individuals
- 1- $\beta_V$ : Vaccine effectiveness
- $\sigma^{-1}$ : Mean duration of latency
- $\gamma^{-1}$ : Mean recovery time for clinically ill
- $\delta^{-1}$ : Duration of immunity loss
- $\mu$ : Natural mortality rate
- r: Birth rate
- *κ*: Recovery rate of latent
- $\alpha$ : Flu induced mortality rate
- $\theta^{-1}$ : Duration of vaccine-induced immunity loss
- CSR: the mobile medical teams' constant service rate

 $\varphi_t$ : Rate of vaccination. It is  $\varphi_t = r(t)CSR$ , which differs from the common SVEIR models' assumption that the vaccination rate is constant during the control effort.



# **Model's equations**

The model is represented by the following system of ordinary differential equations:

$$S'(t) = -bb_E \frac{ES}{N} - bb_1 \frac{IS}{N} - j_T S - mS + dR + qV + rN$$

$$V'(t) = -bb_E b_V \frac{EV}{N} - bb_1 b_V \frac{IV}{N} - mV - qV + j_T S$$

$$E'(t) = bb_E \frac{ES}{N} + bb_1 \frac{IS}{N} + bb_E b_V \frac{EV}{N} + bb_1 b_V \frac{IV}{N} - (m + k + S)E$$

$$I'(t) = SE - (m + a + g)I$$

$$R'(t) = kE + gI - mR - dR$$

The basic reproduction number for the previous model is provided by the next formula:

$$R_{0} = \frac{b(rb_{E} + ab_{E} + gb_{E} + Sb_{I})(r + q + b_{V}j_{t})}{(r + a + g)(r + k + S)(r + q + j_{t})}$$



**Epidemics Logistics** 

# **Greece's Health Districts and targeted populations**



![](_page_25_Picture_2.jpeg)

# **Scenarios and sensitivity analysis**

Intervention strategy	Resource allocation policy	Scenario
No intervention	-	Baseline
	Allocation of a single resource to all sub-populations	Fixed strategy
Reactive mass vaccination starting at day 7, 14, 21, 28 and 60 from the onset of the outbreak	Allocation of a constant amount of resources by using the size of each sub-population as the main driver	Maximum resources
	Dynamic reallocation of resources	Heuristic

![](_page_26_Picture_2.jpeg)

# Results

Vaccination at day 7

Vaccination at day 14

![](_page_27_Figure_3.jpeg)

Results

![](_page_28_Picture_1.jpeg)

![](_page_28_Figure_2.jpeg)

Heuristic algorithm's solution Gantt chart for d=7

- The maximum resources scenario outperforms the no vaccination scenario (the percentage difference of total infective cases ranges from 31.8% to 52.7% respectively, increasing when the vaccination campaign initiates earlier).
- The heuristic algorithm's solution also outperforms the maximum resources scenario where the percentage difference of total infected cases ranges from 1.1% to 2.3% respectively, translated into 15-20 less deaths per 1,000 infective cases averted.

# Smallpox outbreak: The case for regional mass vaccination (Dasaklis et al., 2017)

Smallpox is considered one of the most feared bioterrorist agents with a case-fatality rate ranging from 15 to 30%.

![](_page_29_Picture_2.jpeg)

# **Problem's description**

Two types of supplies should be transported from a central warehouse to RSCs and then to local PODs:

- Crucial medical supplies like vaccines. For these supplies certain transportation protocols should be followed and separate vehicles should be used.
- Vaccine administration supplies (smallpox vaccine coolers/refrigerators, vaccine diluents, sterilized bifurcated needles, etc.), general supplies and equipment (tables, chairs, water and cups, paper, telephones, fax machines etc.) and other emergency supplies (blankets, food meals, etc.). These supplies could be bundled together.

Different types of vehicles (in terms of capacity) and different modes of transportation should be used (according to the type of commodity transported/distributed). The objective is the minimization of the total amount of unsatisfied demand over all types of commodities, final demand points (Points of Dispensing), for all periods.

![](_page_30_Picture_5.jpeg)

# **Problem's formulation**

The modelling approach consists of two parts. The first part is the compartmental modelling approach related to the disease's progression. The second part relates to the epidemics control logistics network configuration model.

![](_page_31_Picture_2.jpeg)

# **Modelling smallpox's progression**

![](_page_32_Figure_1.jpeg)

- Infected but asymptomatic, non-infectious, and vaccinesensitive (I<sub>1</sub>)
- Infected but asymptomatic, non-infectious, and vaccineinsensitive (I<sub>2</sub>)
- Infected but asymptomatic and infectious (I<sub>3</sub>)
- Symptomatic and isolated (I<sub>4</sub>).

![](_page_32_Picture_6.jpeg)

# Notation

![](_page_33_Picture_1.jpeg)

T: be the planning horizon. t: time period (day), t-1,...,T.  $C_M$ : be the set of different essential medical commodities (vaccines)  $C_S$ : be the set of different ancillary supplies (bifurcated needles, food, blankets etc) C: be the set of commodities in total. It is  $C=C_M \cup C_S$ NSC: be the set of National Stockpiling Centres RSC: be the set of Regional Stockpiling Centres POD: be the set of local Points of Dispensing N=NSC+RSC+POD: be the union of nodes in the network

#### **Parameters**

#### Let

Let

 $S_{ic}(t)$ : be the supply of commodity type c in time period t at the National Stockpiling Centre i,  $i \in NSC$ ,  $c \in C$ , t=1,...,T

M(t): be the set of kinds of transportation means  $\{M_1(t), ..., M_K(t)\}$  in time period t, i.e. there are K different kinds (subsets) of non-identical transportation means with  $|M_k(t)|$  identical means of each kind, k=1,...,K

 $V_{kc}$ : be the capacity of transportation mean k for commodity type c, k=1,...,K, c  $\in$  C.

 $v_c$ : be the volume of commodity type c,  $c \in C$ 

G(N,E): be a graph, where E is the set of edges (i,j,k),  $i,j \in N$ , k=1,...,K

 $d_{ic}(t)$ : the demand for commodity type c in time period t at dispensing point i, i  $\in$  POD, c  $\in$  C, t=1,...,T

 $U_{ik}(t)$ : Unloading capacity for the facility in node i for transportation mean of kind k in time period t,  $i \in N$ , k=1,...,K, t=1,...,T

 $SC_i(t)$ : Storage capacity for the facility in node i in time period t,  $i \in N$ , t=1,...,T

LC<sub>ik</sub>(t): Loading capacity for the facility in node i for transportation mean k in time period t, i∈N, k=1,...,K, t=1,...,T

#### **Decision variables**

 $x_{ijck}(t)$ : amount of commodity type c transported from node i to node j by the k-th kind of transportation in period time t,  $i, j \in N$ ,  $i \neq j$ ,  $c \in C$ , k=1,...,K, t=1,...,T.

 $u_{ic}(t)$ : unsatisfied demand of commodity type c at node i in period time t,  $i \in POD$ ,  $c \in C$ , t=1,...,T

 $I_{ic}(t)$ : Inventory of commodity type c at node i in period time t,  $i \in N$ ,  $c \in C$ , t=1,...,T

### **Epidemics Logistics**

# **Smallpox progress model equations**

$$\begin{aligned} \frac{dS_{1}(t)}{dt} &= -\beta S_{1}(t)I_{3}(t) \\ \frac{dS_{2}(t)}{dt} &= -\beta S_{2}(t)I_{3}(t) - v_{s}\alpha(t)\sum_{i=1}^{n}I_{ic}(t)S_{2}(t) \\ \frac{dI_{1}(t)}{dt} &= \beta S_{1}(t)I_{3}(t) + \beta S_{2}(t)I_{3}(t) - r_{1}I_{1}(t) - v_{1}\gamma(t)\sum_{i=1}^{n}I_{ic}(t)I_{1}(t) \\ \frac{dI_{2}(t)}{dt} &= r_{1}I_{1}(t) - r_{2}I_{2}(t) \\ \frac{dI_{3}(t)}{dt} &= r_{2}I_{2}(t) - r_{3}I_{3}(t) \\ \frac{dI_{4}(t)}{dt} &= r_{3}I_{3}(t) - r_{4}I_{4}(t) \\ \frac{dR(t)}{dt} &= r_{4}I_{4}(t) + v_{s}\alpha(t)\sum_{i=1}^{n}I_{ic}(t)S_{2}(t) + v_{1}\gamma(t)\sum_{i=1}^{n}I_{ic}(t)I_{1}(t) \end{aligned}$$

![](_page_34_Picture_2.jpeg)

# **Smallpox progress model equations**

$$\alpha(t) = \frac{S_2(t)}{S_2(t) + I_1(t) + I_2(t) + I_3(t)}$$

$$\gamma(t) = \frac{I_1(t)}{S_2(t) + I_1(t) + I_2(t) + I_3(t)}$$

$$\sum_{i=1}^{n} d_{ic}(t) = S_2(t) + I_1(t) + I_2(t) + I_3(t)$$

As the vaccine provides protection to those susceptible to the disease and to those at stage 1 of infection there will be a quantity  $V_w(t)$  of vaccine wasted per period:

$$V_{w}(t) = \left[\frac{I_{2}(t) + I_{3}(t)}{S_{2}(t) + I_{1}(t) + I_{2}(t) + I_{3}(t)}\right] \sum_{i=1}^{n} I_{ic}(t)$$

![](_page_35_Picture_6.jpeg)

# Logistics network configuration model

### Assumptions:

- NSC, RSC and POD are disjoint sets, e.g. a RSC cannot serve as local POD, etc.
- The time to transfer commodities from a transportation mean to another is considered negligible.
- Travelling times between the two most distanced nodes of the network do not exceed the period of one day (24 hours).

![](_page_36_Picture_5.jpeg)

# Logistics network configuration model

 $min \sum_{i \in POD} \sum_{c \in C} \sum_{t=1}^{T} u_{ic}(t)$ 

#### Constraints

$$\begin{split} &I_{ic}(t) = I_{ic}(t-1) + S_{ic}(t) - \sum_{j \in RSC} \sum_{k=1}^{K} x_{ijck}(t) \text{, } i \in \text{NSC, } c \in \text{C, } t=1,...,\text{T} \\ &I_{ic}(t) = I_{ic}(t-1) + \sum_{\substack{j \in NSC \\ j \neq i}} \sum_{k=1}^{K} x_{jick}(t) - \sum_{\substack{j \in \text{POD} \\ j \neq i}} \sum_{k=1}^{K} x_{ijck}(t) \text{, } i \in \text{RSC, } c \in \text{C, } t=1,...,\text{T} \\ &I_{ic}(t) - u_{ic}(t) = I_{ic}(t-1) - u_{ic}(t-1) - d_{ic}(t) + \sum_{\substack{j \in RSC \\ j \neq i}} \sum_{k=1}^{K} x_{jick}(t) \text{, } i \in \text{POD, } c \in \text{C, } t=1,...,\text{T} \end{split}$$

$$\sum_{c \in C} I_{ic}(t) \leq SC_i(t), i \in \mathbb{N}, t=1,...,T$$

$$\sum_{\substack{i,j \in N \\ i \neq j}} v_c x_{ijck}(t) \leq V_{kc} \left| M_k(t) \right|, k=1,...,K, c \in C, t=1,...,T$$

$$\sum_{i \in RSC+POD \atop i \neq i} \sum_{c \in C} x_{ijck}(t) \leq LC_{ik}(t), i \in N, k=1,...,K, t=1,...,T$$

$$\sum_{\substack{j \in N \\ i \neq i}} \sum_{c \in C} x_{jick}(t) \leq UC_{ik}(t), i \in \mathsf{RSC+POD}, k=1,...,K, t=1,...,T$$

![](_page_37_Picture_8.jpeg)

# **Numerical experiments: Athens**

Regional units	Population	Intervention policy	Scenario	Resource allocation policy
Central Athens North Athens West Athens South Athens Piraeus East Attica West Attica Total	1,029,520 591,680 489,675 529,826 448,997 502,348 160,927 3,752,973	Case isolation Reactive regional mass vaccination within 4 days starting at days 22, 29, 36, 43 and 50 from the onset of the outbreak. Reactive regional mass vaccination within 9 days	Baseline A <sub>1</sub> A <sub>2</sub> A <sub>3</sub> A <sub>4</sub> A <sub>5</sub> B <sub>1</sub> B <sub>2</sub>	Unconstrained
Number of VS Number of vaccinators Vaccination rate Hours of operation for VC Estimated overall vaccination rate per day/POD	8 per VC 1 per VS 30–60 vaccinations per VS/hour 24 hours/16 hours 8640/5900	starting at days 22, 29, 36, 43 and 50 from the onset of the outbreak. Reactive regional mass vaccination starting at days 22, 29, 36, 43 and 50 from the onset of the outbreak with limited vaccine supply. Reactive regional mass vaccination starting at days 22, 29, 36, 43 and 50 from the onset of the outbreak where limited transportation capacity as well as limited vaccine supply are	$B_{4} \\ B_{5} \\ C_{1} \\ C_{2} \\ C_{3} \\ C_{4} \\ C_{5} \\ D_{1} \\ D_{2} \\ D_{3} \\ D_{4} \\ D_{5}$	Constrained

![](_page_38_Picture_2.jpeg)

# **Numerical experiments: Athens**

Parameters	Value
Set of commodities	Vaccines
	<ul> <li>Three other supplementary commodities</li> </ul>
	(water, food and blankets)
Periods	25
Demand for commodities	5900/POD/period
Set of vehicles	<ul> <li>4 (for vaccines' distribution)</li> </ul>
	<ul> <li>10 (for supplementary commodities)</li> </ul>
Points of Dispensing	47
Regional Stockpile Centres	7
National Stockpile Centres	1
Storage capacity of NSC	5,000,000 Units
Storage capacity of RSC	3,000,000 Units
Storage capacity of POD	600,000 Units
Vehicles capacity (for vaccines' distribution)	21,000 lt
Vehicles capacity (for distributing the other commodities)	90,000 lt
Loading capacity of NSC (the same for all commodities and periods)	1,000,000 Units
Loading capacity of RSC (the same for all commodities and periods)	750,000 Units
Unloading capacity of RSC (the same for all commodities and periods)	750,000 Units
Unloading capacity of POD (the same for all commodities and periods)	500,000 Units
Volume of commodities	Vaccines: 0.2 lt
	Water: 0.5 lt
	Food: 2 It
	Blankets: 4 lt
Vaccine supply at the NSC	Forty per cent in the initiation of the
	vaccination campaign and 60% 10 days after
	(for all scenarios)

![](_page_39_Picture_2.jpeg)

![](_page_40_Picture_0.jpeg)

Table 5. Data when vaccination lasts for 4 days.

Vaccination's initiation day	Number of persons vaccinated per day	Number of PODs to open	Number of infected individuals
22	625,004	72	1915
29	624,284	72	3810
36	622,931	72	7568
43	620,247	72	14,983
50	614,747	72	29,484

### Table 6. Data when vaccination lasts for 9 days.

Vaccination's initiation day	Number of persons vaccinated per day	Number of PODs to open	Number of infected individuals
22	277,543	32	2408
29	277,071	32	4788
36	276,113	32	9503
43	274,187	31	18,790
50	270,287	31	36,876

![](_page_40_Figure_5.jpeg)

Figure 3. Number of infected individuals for the first set of scenarios.

![](_page_40_Figure_7.jpeg)

![](_page_40_Figure_8.jpeg)

![](_page_40_Picture_9.jpeg)

#### **Epidemics Logistics**

![](_page_41_Picture_0.jpeg)

South Athens	529,826	7
Piraeus	448,997	6
East Attica	502,348	6
West Attica	160,927	2
Total		47

#### Table 9. Results of the third set of scenarios.

Vaccination's initiation day	Vaccination's termination day	Day when effective reproduction number drops below 1	Infected individuals
22	39	35	2833
29	46	42	5623
36	53	49	11,119
43	60	56	21,826
50	67	62	42,238

### Table 10. Results of the fourth set of scenarios.

Vaccination's initiation day	Vaccination's termination day	Day when effective reproduction number drops below 1	Infected individuals
22	43	38	3325
29	50	45	6,592
36	57	52	13,006
43	64	59	25,422
50	71	63	39,091

![](_page_41_Figure_6.jpeg)

Figure 7. Number of infected individuals for the third set of scenarios.

![](_page_41_Figure_8.jpeg)

![](_page_41_Figure_9.jpeg)

![](_page_41_Picture_10.jpeg)

![](_page_42_Figure_0.jpeg)

Figure 9. Vaccine's stockpile in the National Stockpile Centre for the fourth set of scenarios (per day).

![](_page_42_Picture_2.jpeg)

**Results** 

![](_page_43_Picture_0.jpeg)

- Epidemics control supply chain literature is fragmented
- Most of the available frameworks have little correlation with real-case scenarios and, therefore, the applicability of the modeling approaches might be limited
- Several aspects of the nature of the outbreak or of the agent triggering the outbreak have not explicitly taken into consideration when relevant supply chain decisions are to be made.

![](_page_43_Picture_4.jpeg)

# **Research directions**

- Multidisciplinary synergies
- End-to-End approaches
- Evaluation of models and large scale exercises (applicability of existing modeling approaches)
- Performance metrics
- Cross-functional drivers
- Coordination issues.

![](_page_44_Picture_7.jpeg)

# Literature

1. T.K. Dasaklis, C.P. Pappis, N.P. Rachaniotis (2012). Epidemics control and logistics operations: A review. *International Journal of Production Economics*, **139**, **2**: 398-410.

2. T.K. Dasaklis, N. Rachaniotis & Costas Pappis (2017). Emergency supply chain management for controlling a smallpox outbreak: the case for regional mass vaccination. *International Journal of Systems Science: Operations & Logistics*, **4**, **1**: 27-40.

3. N.P. Rachaniotis, T.K. Dasaklis, C.P. Pappis (2012). A deterministic resource scheduling model in epidemic control: A case study. *European Journal of Operational Research*, **216**, **1**: 225-231.

4. N. Rachaniotis, T.K. Dasaklis, C. Pappis (2017). Controlling infectious disease outbreaks: A deterministic allocation-scheduling model with multiple discrete resources. *Journal of Systems Science and Systems Engineering*, **26**, **2**: 219-239.