



# EFSA activities on Lumpy Skin disease: recent and ongoing activities

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Standing Group of Experts on Lumpy Skin Disease  
in the South East Europe region – Istanbul, 12-13  
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## TIMELINE OF EFSA ACTIVITIES ON LSD

1. Jan 2015, scientific opinion:  
focusing the problem
2. May 2016, EFSA-EC workshop,  
fostering cooperation
3. Aug 2016, Urgent advice : assessing  
effectiveness of control measures
4. 2016-2017, Current data collection:  
learning from epidemics



# FIRST EFSA OPINION – HIGHLIGHTING GAPS

## KNOWN

- Transmission by haematophagus arthropod vectors
- LSDV detectable in animal secretions
- Live vaccines
- PCR, SNT

## UNKNOWN

- Which vector species? Biological vectors?
- direct or indirect transmission?
- Milk products? Safe organs? Contaminated feed?
- ELISA? DIVA, safety and purity issues?

### Science has moved forward:

- DIVA qPCR (Menasherow et al., 2016)
- Full genome sequencing of vaccin strain (Mathjis et al., 2016)



# Simulating spread and impact



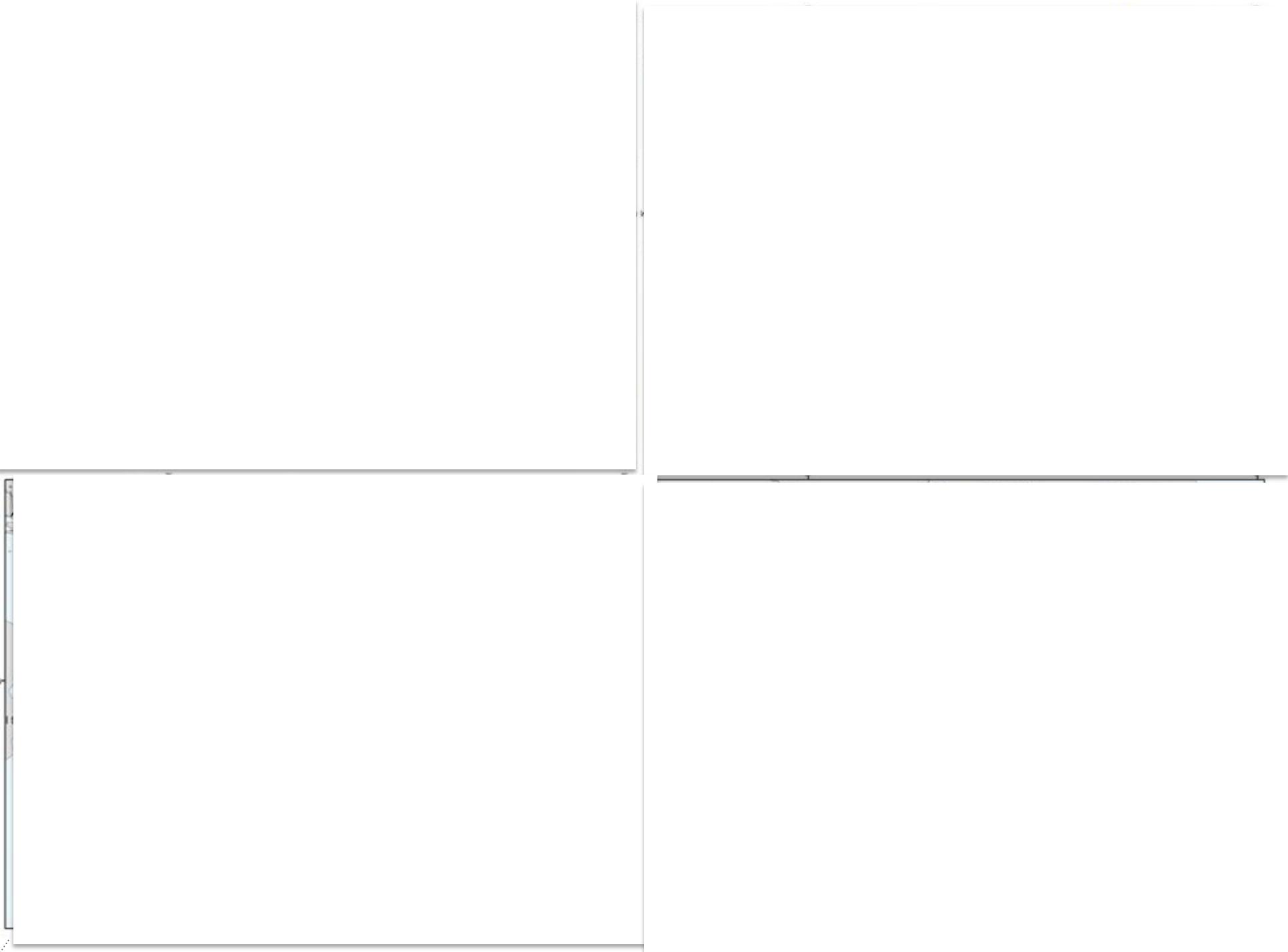
## METHODOLOGY

# Stochastic kernel-based model of LSD spread

➤ between-farms transmission

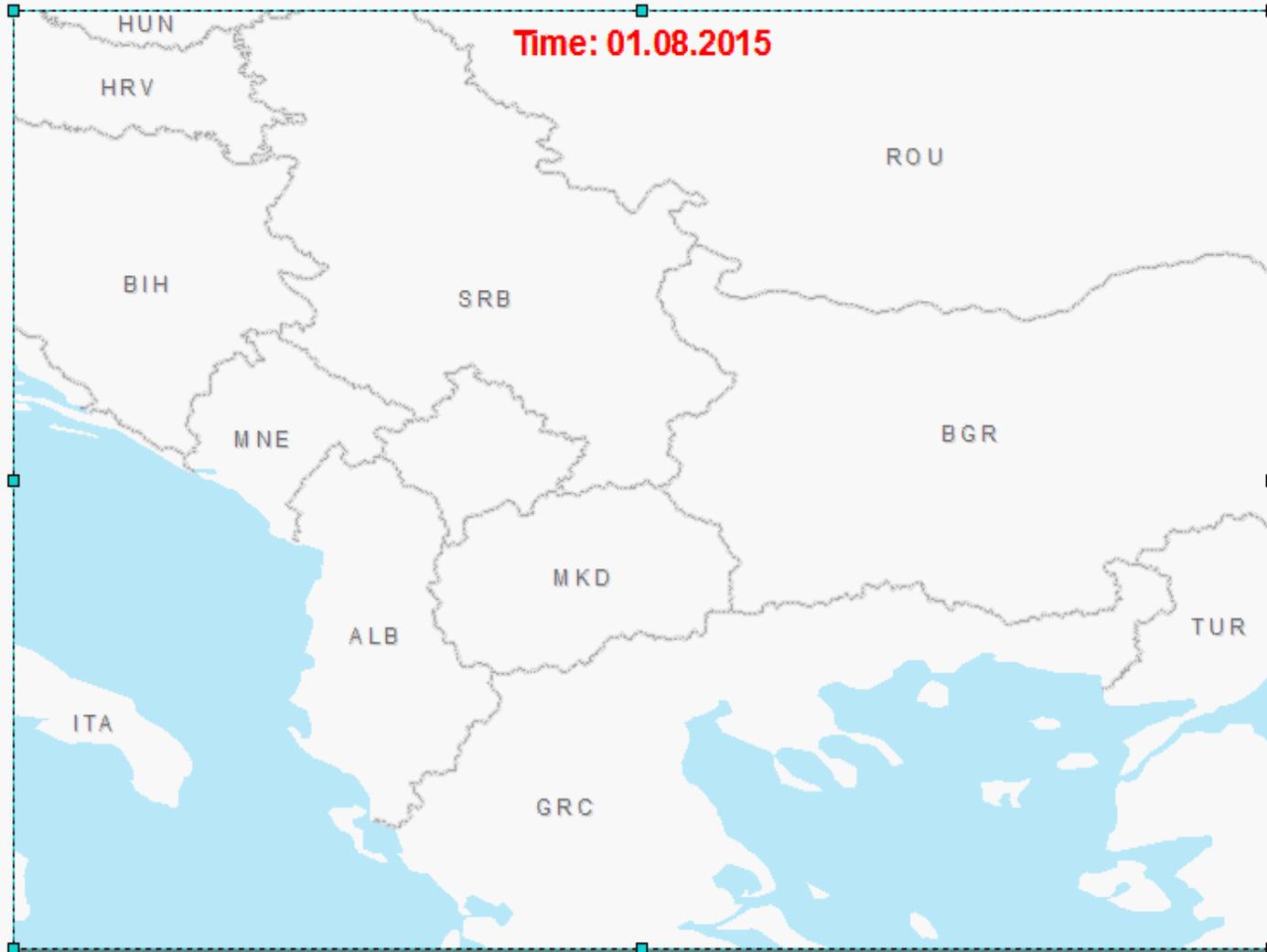
### Control strategy scenarios:

- i) the removal of clinical cases
- ii) whole-herd culling after 7, 15 or 28 days after infection.





# WHAT HAPPENED SINCE SUMMER 2015



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## SPREAD SCENARIOS EXPLORED IN GR AND BG

EC: assess the spread and persistence of a partial stamping-out policy compared to total stamping out





## ELEMENTS INCLUDED IN THE MODEL

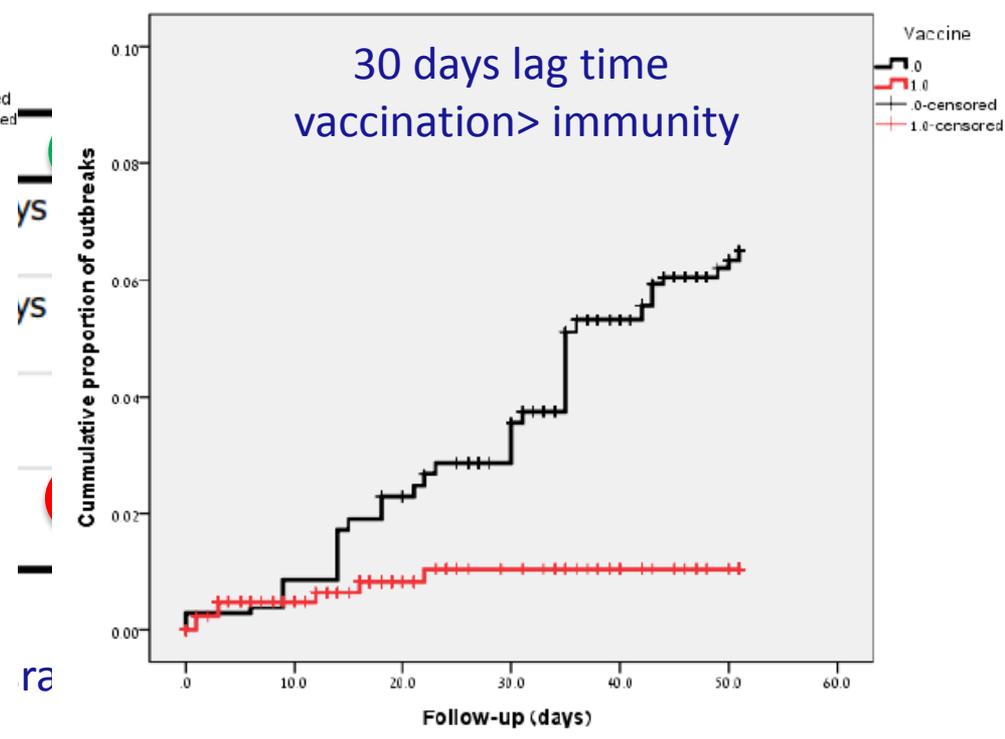
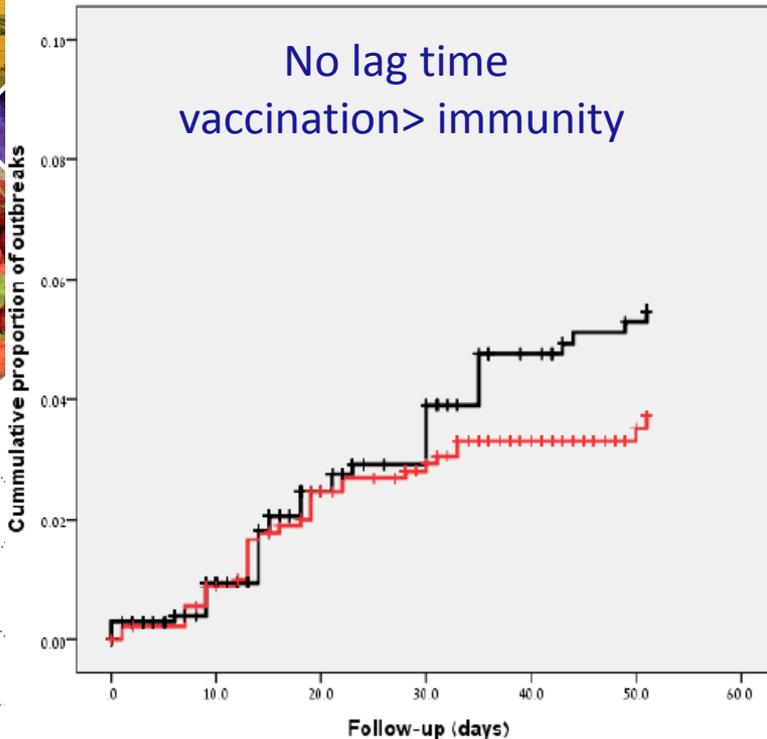
- **Delay between infection and report:** 1-2 weeks (mean 10.5 days, gamma dist.)
- **delay between report and stamping out:** based on the data from Greece and Bulgaria (mean 7.6 days, gamma distr.)
- **Partial stamping out:** i) by reducing outbreak duration; ii) by reducing the infectiousness; iii) by increasing the outbreak duration and by reducing the infectiousness
- **Total stamping out:** Removing the farm at a certain time (mean of 7.6 days after reporting)
- **Vaccination:** replacing herd sizes with the number of unprotected animals in each herd
- **vaccination effectiveness** 75% (Ben-Gera et al. 2015) and calculated from the data from Greece and Bulgaria, and 40%.
- **Preventive and reactive vaccination:** different time of vaccination start
- **Vaccination coverage:** 95% farms
- **Maximal protection:** after 21 days post vaccination

		VACCINATION		
		None	after virus entry	before virus entry
STAMPING OUT	TOTAL			
	NONE			
		PARTIAL		
		NONE		

# ESTIMATION OF VACCINATION EFFECTIVENESS IN THE FIELD

## survival analysis comparing LSD incidence

probability of infection in the **vaccinated** and **unvaccinated** farms



ys  
ys  
ra

## KEY MESSAGES

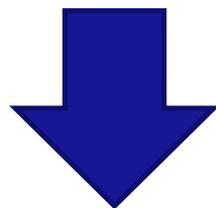
- 
- vaccination better than any stamping-out policy to reduce LSD spread
  - Performance of type of stamping out depends on effectiveness of coupled vaccination
  - partial stamping out leads to limited increase of spread compared to total stamping out
  - most effective vaccination policy:
    - protection developed at the time of virus entry
    - high coverage within and between farms

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## OBJECTIVES

- Learn from current outbreaks
- Strengthen collaboration between EFSA and MSs and neighbouring countries
- Harmonise and increase efficiency of data collection



**Improve RA quality**



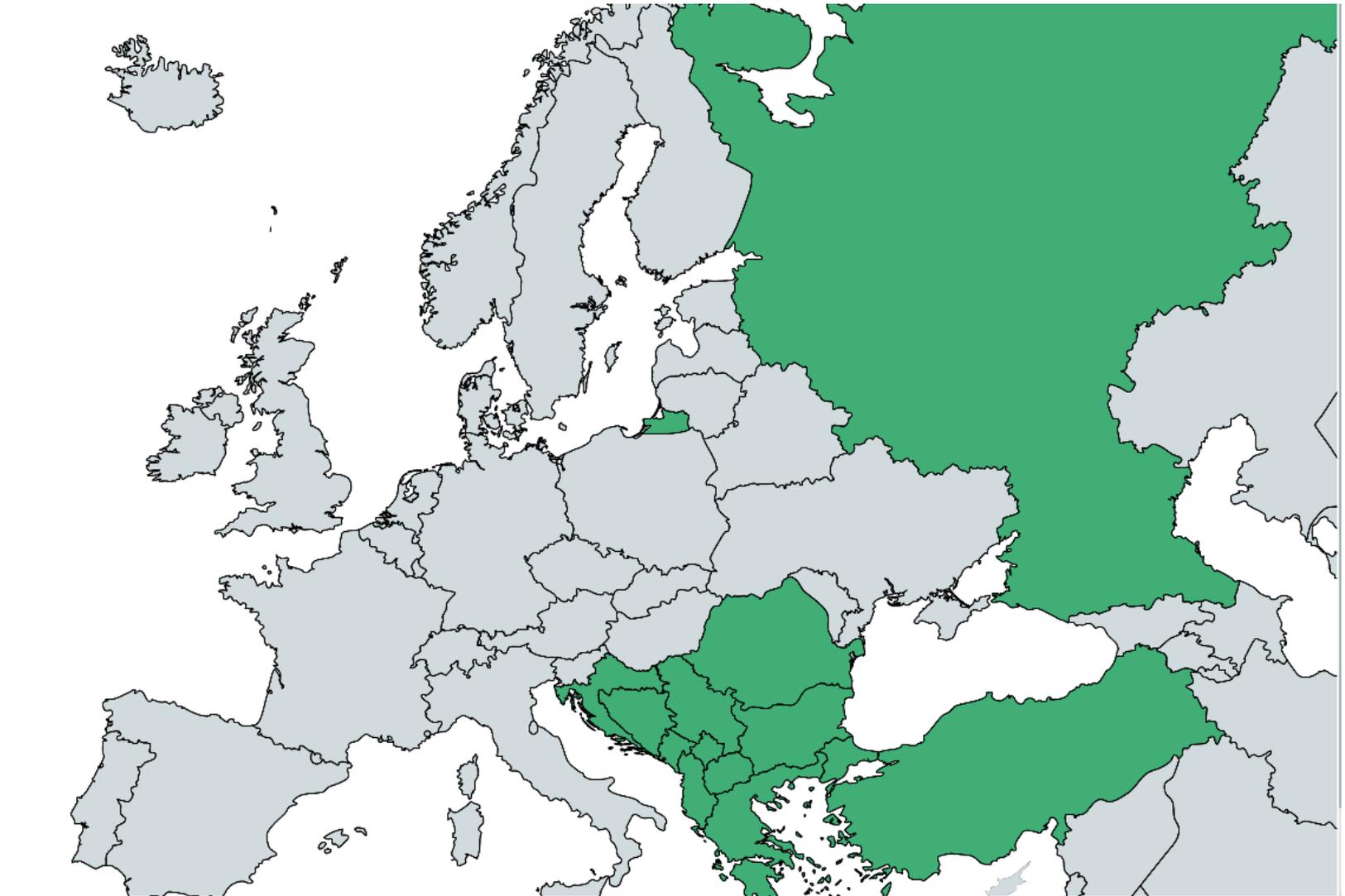
## DATA TO BE COLLECTED

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- Demography
  - Outbreaks
  - Vaccination
  - Movement of animal/people/feed
  - Laboratory test
- 
- Geo-climatic data and land cover
  - Vector presence/abundance

## POSSIBLE USE OF THESE DATA

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- **Spatial and temporal patterns of outbreak:** infection kernel, morbidity between farms, within farm
  - **Seasonality of the disease:** Relationship between outbreak and climatic issues and vector activity
  - **Risk factors;** type of farming, grazing, animal movements, farm/animal density, introduction of new animals/feed, land cover, season/climatic
  - **Estimation of subclinical disease**
  - **Vaccination effectiveness and safety**

# COUNTRIES INVOLVED



## WORKSHOP

# EFSA technical meeting, 20th Dec 2016

- what data would be useful (data model)
- What data are/could be available
- What else can be collected along 2017
- how to submit data by Data Collection Framework of EFSA

# OVERALL CONCLUSIONS

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- LSD as neglected disease with lots of **knowledge gaps**
  - **Research needed** for :
    - vector biology
    - diagnostics for mass screening, DIVA
    - Vaccine – safety, DIVA
  - New epidemiological situations: adapted **policies**
  - **Regional problem** > **regional cooperation** for enhancing preparedness, knowledge sharing
  - Learning from epidemics:
    - **ready data model for data collection**
    - **Coordinating centre for data collection**



# Thank you for your attention!

## Acknowledgements

- Representatives from National Authorities from affected and at risk countries
- WG experts

## EFSA staff:

- Andrey Gogin
- José Cortiñas Abrahantes

**Further info and all EFSA outputs at:**

**<https://www.efsa.europa.eu/en/topics/topic/lumpyskindisease>**