Mapping and evaluation of national capacities in local and regional laboratories for the detection and characterization of Salmonella in humans in Serbia

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Mapping objectives

General

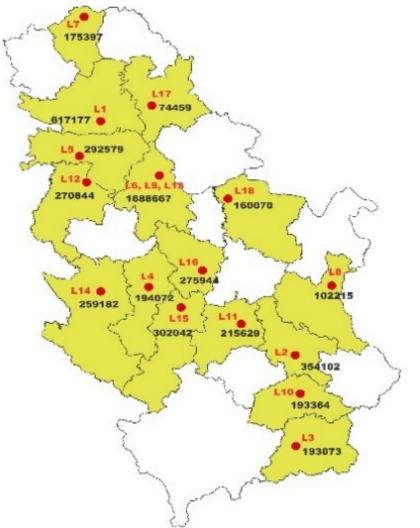
• To establish sentinel surveillance of AMR of human Salmonella in Serbia

Specific

- To improve refferal of isolates to NRL
- To identify how many laboratories can identify S. Enteritidis (most prevalent serotype)
- To harmonize the methodology used for AST in the laboratories according to EU protocol
- To identify how many laboratories perform culture-indipendent detection of Salmonella



Mapping methodology - Selection of laboratories



- All 4 regions including 16 districts
- Covering 82% of Serbian population
- Representative sample of 28% mapped laboratories in network (63 laboratories)
- 14 regional PH laboratories
 & 4 hospital (one pediatric)

Region	District/ Laboratory code
North Serbia	Severnobački L7
	Srednjobanatski L17
	Sremski L5
	Južnobački L1
Belgrade	L6, L9, L13
Central and West Serbia	Šumadijski L16
	Moravički L4
	Raški L15
	Rasinski L11
	Mačvanski L12
	Zlatiborski L14
South and East Serbia	Zaječarski L8
	Braničevski L18
	Nišavski L2
	Jablanički L10
	Pčinjski L3

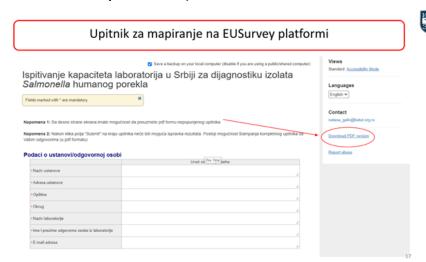


Mapping timeline

2022 2023

December:

- Email invitation to laboratories
- NRL_Questionnaire on EU Survey Platform (4 sections, 47 mandatory questions)



January:

- On –line meeting with participants mapping and questionnaire
- Disributed web link with questionnaire to participants

February:

Statistical analysis and evaluation of results

March:

• Summary report to RefLabCap Team

September:

 Detailed summary report to mapped laboratories

December:

Short report to decision makers,
 Ministry of Health & IPHS



National system for diagnostics of human Salmonella

Strengths/Weaknesses/Needs



- Good geographic coverage, but not a well-functioning laboratory network in relation to refferal of isolates
- Diagnostics of human *Salmonella* is mandatory according to the Serbian Law on Protection of Population from Communicable Diseases (all laboratories are performing diagnostic)



- National surveillance is based on species level, reporting of serotype is not mandatory
- National digital surveillance system for salmonellosis is not integrated with and AMR data of human Salmonella
- No national requirement for accreditation of the laboratories, only one accredited (ISO 15189)
- Number of laboratories who participaded in EQAs for human Salmonella in last three years is negligable
- No national guidelines for diagnostics and refferal of human Salmonella



- Support from PH authorities to implement accreditation in laboratories in PHIs and to issue national diagnostics guidelines
- NRL support through guidances, QC materials, implementation of EQA trails, consultancy to laboratories



Laboratory resources (human, equipment, diagnostics, funding) Strengths/Weaknesses/Needs



- Human resources (laboratory personel trained for diagnostic of Salmonella) highest grade
- Most of the laboratories are satisfied with availability and quality of laboratory equipment



- Deficit of labotratory staff (undercapacity)
- Limited resources for serotyping of *Salmonella* (low budget for purchase of antisera)
- Process of procurament of laboratory suplies and equipment maintenance situation



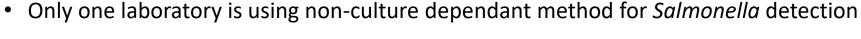
- Continous education of laboratory staff
- Budget improvement



Diagnostic methods for detection – Strengths/Weaknesses/Needs



• All laboratories are using enrichement method for Salmonella isolation



- Choice of culture media not harmonised
- Inadequate conditions and capacities for preservation of Salmonella isolates (only outbrake isolates)



- Issuing of national guidelines
- Implementation of molecular methods for detection of Salmonella in clinical samples







Characterization methods – Strengths/Weaknesses/Needs



• All laboratories have capacity for characterization of species and AST of Salmonella according to EUCAST



- In most laboratories capacities for serotyping of Salmonella are limited just to S. Enteritidis
- Just a few laboratories have Maldi-tof system
- Duplicate biochemical identification tests (manual and api 20E test),
- AST in majority of laboratories is performed for therapeutic purpose, only few for surveilance
- AST is not harmonised according to EU protocol
- Half laboratories are performing DDT and automated dilution sistem (increasing the expencess)
- Inconsistency in performing QC by EUCAST



- Laboratory network needs continuous work on methodology harmonisation with support of NRL
- Necessity of regular referall of isolates to NRL for a final characterisation



Refferal of isolates/samples to NRL – Strengths/Weaknesses/Needs



- Sample refferal to NRL is a legal obligation (for IPHs, not for clinical laboratories)
- Laboratories are reporting *Salmonella* to surveilance system (mostly on species level), without editing according to NRL result of serotypes



- No national guidance for handling and reffering samples/isolates (irregular)
- NRL's guidance for referral isolates are not used in each laboratory
- NRL sample refferal from usually filled with partial data related to the case (just gender and age, no epidemiological data, no travel history)
- 60% of laboratories are saving copies of reffered samples to NRL



- Harmonization of sample referal procedure to NRL methodology
- Raise awarness of importance of refferal of isolates/samples to NRL



Outcomes

- 1. Salmonella isolates refferal to NRL from local/regional laboratories is significantly improved
- 2. NRL has updated SOPs for detection and characterisation *Salmonella*, including AST in accordance with EU protocol
- 3. SOPs are uploaded on official site IPHS and distributed to Serbian Medical Society https://www.batut.org.rs/index.php?category_id=69
- NRL implemented a pilot sentinel Salmonella AMR surveilance with all mapped laboratories
- 5. Planning of first round of national EQA



Thank you!