

MINIMUM AND OPTIMAL REQUIREMENTS FOR NPHRL

SURVEY RESULTS ON CAPACITY FOR TESTING AND SURVEILLANCE OF AMR

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Network meeting 2021 November 30 – December 1





- All Member States (MS) are obliged to collect relevant and comparable data on Salmonella and Campylobacter:
 - infections in humans,
 - food-related outbreaks, and
 - the occurrence of resistance to antimicrobials (AMR) relevant for the treatment of human infections with these bacteria
- The data must be reported to ECDC, however the it is not always comparable due to:
 - dissimilar organisation and operation of the national public health reference laboratories (NPHRLs) in different MSs
 - variation in data quality level
 - differences in the case definitions and methodologies used

CORE FUNCTIONS OF NPHRL





European Centre for Disease Prevention and Control. Core functions of microbiology reference laboratories for communicable diseases. Stockholm: ECDC; 2010

MONITORING OF NPHRL MICROBIOLOGY SYSTEM



European Centre for Disease Prevention and Control. EU Laboratory Capability Monitoring System (EULabCap) – Report on 2018 survey of EU/EEA country capabilities and capacities. Stockholm: ECDC; 2020.

Substantial variation in capacities and capabilities between the laboratories

Some laboratories:

- Had an insufficient level of laboratory capacity and capability to conduct effective public health surveillance

- Were unable to provide an adequate level of disease threat response

RECOMMENDATIONS OF MINIMUM AND OPTIMAL REQUIREMENTS OF THE REFERENCE LEVEL FUNCTIONS FOR THE NATIONAL SURVEILLANCE SYSTEMS FOR AMR IN SALMONELLA AND CAMPYLOBACTER IN HUMANS

1. Support to primary diagnostic testing

2. Laboratory-based surveillance of AMR, alert and response

- Support to national surveillance networks
- National outbreak response support
- Participation in EU disease networks and research
- The design of AMR surveillance system, sampling and testing frequency
- Referral, timing and storage of *Salmonella* and *Campylobacter* isolates

3. National Public Health Reference Laboratory Services

- Internal and external quality control
- Reference diagnostics and characterisation of Salmonella
- Reference diagnostics and characterisation of *Campylobacter*

MINIMAL VS. OPTIMAL REQUIREMENTS



- In the same laboratory, both minimum and optimal requirements may be applied, e.g.
 - minimum requirements for the majority of clinical samples/isolates
 - optimal requirements on a subset of samples/isolates

- If, NPHRLs do not have the capacity and capability to operate according to at least minimum requirements
 - service level agreement(s) with other expert or reference level
 laboratories should be in place



Salmonella

Requirements	Serotyping	Antimicrobial resistance	Cluster detection
Minimum	Phenotypic or genotypic: common serovars	Phenotypic AST or genotypic AMR prediction	High resolution molecular typing (e.g. MLVA)
Optimal	Phenotypic or genotypic: all serovars	Phenotypic AST and WGS-based AMR prediction*	WGS-based (e.g. cgMLST, wgMLST, SNP)

Campylobacter

Requirements	Species	Antimicrobial resistance	Cluster detection
Minimum	Phenotypic or genotypic: <i>C. jejuni,</i> <i>C. coli</i>	Phenotypic AST or genotypic AMR prediction	Not applicable**
Optimal	Phenotypic or genotypic: all species	Phenotypic AST and WGS-based AMR prediction*	WGS-based (e.g., cgMLST, wgMLST, SNP)

* a defined proportion of isolates or selected isolates are periodically tested phenotypically to ensure detection of novel resistance mechanisms

** Some laboratories may use Pulsed-field gel electrophoresis (PFGE) for cluster detection but this is not considered as a minimum requirement



- to identify capacity and capability gaps in technical and analytical skills of AMR testing and strain subtyping at national level in all countries
- identify countries with the greatest needs for capacity building in phenotypic and genomic testing of AMR ('priority countries')
 - provide tailored technical and operational support for implementation of phenotypic and genomic testing of AMR in human Salmonella and Campylobacter
 - provide direct advice and support for each country to create an action plan to strengthen its national reference laboratory capacities in line with standards set out in EULabCap documentation
- identify 'additional countries' facing challenges to identify and manage outbreaks of Salmonella spp. and Campylobacter spp.

SURVEY OUTCOME

- Sent to 37 countries (32 EU/EEA/EU Health programme countries and 5 EU-candidate countries)
- Responses received from 31 EU/EEA/EU Health programme countries and all 5 EU-candidate countries (n=36)
 - In 29/36 countries responses from the same laboratory covering both Salmonella and Campylobacter were received
 - In 6/36 countries responses from two laboratories covering either *Salmonella* or *Campylobacter* were received
 - One country responded for *Salmonella* spp. only

SALMONELLA

- AMR testing overview
- Data and sample collection for surveillance of AMR
- National AMR surveillance system coverage
- Molecular subtyping of *Salmonella* isolates

AMR TESTING OVERVIEW - SALMONELLA





The purpose and the number of antimicrobials tested (phenotypic and genotypic)

N=30

DATA FOR AMR SURVEILLANCE - SALMONELLA



Collection of laboratory data



submitting laboratory data and/or sample material % of the local/regional clinical laboratories



% of the total number of Salmonella enteritis cases tested for AMR annually

		76-100	51-75	26-50	0-25	Unknown	N=23
R	76-100	N=8	N=3	N=2			EU/EEA/EU Health
nce of AN	51-75		N=3		N=1		
surveilla	26-50					N=1	
national	<mark>0-25</mark>		N=1	N=1			
for	Unknown				N=1	N=2	

MOLECULAR SUBTYPING OF SALMONELLA ISOLATE



Objectives of molecular subtyping



N=22 EU/EEA/EU Health

AMR testing on outbreak isolates



SALMONELLA AMR SURVEILLANCE IN NON-EU COUNTRIES





CAMPYLOBACTER

- AMR testing overview
- Data and sample collection for surveillance of AMR
- National AMR surveillance system coverage
- Molecular subtyping of *Campylobacter* isolates

N=28

The purpose and the number of antimicrobials tested in all laboratories

EU/EEA/EU Health 4 Surveillance Other 3 2 1 0 _ab 10 Lab 23 9 ∞ 6 Lab 11 Lab 12 Lab 13 Lab 14 Lab 15 Lab 16 Lab 18 Lab 19 Lab 20 21 Lab 24 25 26 28 Lab 17 22 27 Lab Lab Lab Lab Lab Lab ab. Lab Lab Lab Lab Lab Lab Lab Lab C. jejuni + C.coli 96 % SEROVARS tested Phenotypic AMR 93 % Ρ Р **METHODS** GG G G G G G G **Genotypic AMR** 29 % **Disk diffusion** 77 % 31 % **Broth microdilution AST METHODS** 42 % **Gradient strip** Ο Automated system 4 % AST BREAKPOINTS EUCAST CBP 96 % 00 000 0 EUCAST EBP 0 0 31 % **CLSI CBP** 8 % **AST Performance level** •



Collection of laboratory data







MOLECULAR SUBTYPING OF CAMPYLOBACTER ISOLATES

STATENS SERUM INSTITUT

Objectives of molecular subtyping



N=16

EU/EEA/EU Health

AMR testing on outbreak isolates



CAMPYLOBACTER AMR SURVEILLANCE IN NON-EUS SERUM INSTITUT





SUMMARY



- Nearly all NPHRLs have a capacity to perform testing of AMR in Salmonella and Campylobacter isolates from humans using either phenotypic of genotypic methods, however:
 - Approx. 2/3 of NPHRLs perform AMR surveillance and strain subtyping of *Salmonella* isolates
 - Approx. 1/3 of NPHRLs perform AMR surveillance and strain subtyping of *Campylobacter* isolates





ACKNOWLEDGEMENTS



Team members at SSI

- Eva Møller Nielsen
- Susanne Schjørring
- Malgorzata Ligowska-Marzeta
- Jeppe Boel
- etc.

- EURGen-RefLabCap team at SSI:
 - Valeria Bortolaia
 - Camilla Wiuff Coia

Everyone who responded to the survey!