



FWD AMR·  
RefLabCap

# SURVEY RESULTS ON WGS CAPACITY AND METHODS

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## *Salmonella*

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

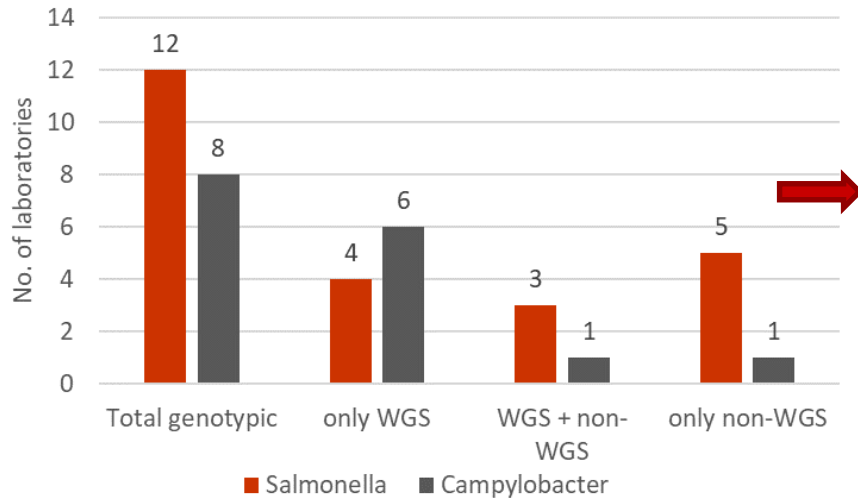
## *Campylobacter*

Requirements	Species	Antimicrobial resistance	Cluster detection
<b>Minimum</b>	Phenotypic or genotypic: <i>C. jejuni</i> , <i>C. coli</i>	Phenotypic AST or genotypic AMR prediction	Not applicable**
<b>Optimal</b>	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

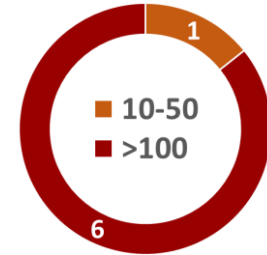
## Molecular methods in use for testing of AMR



## No. of isolates tested using WGS

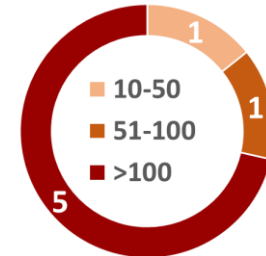
### Salmonella

- 7 WGS-based
- 8 non - WGS



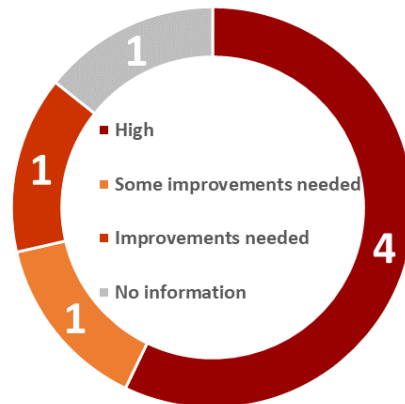
### Campylobacter

- 7 WGS – based
- 2 non - WGS

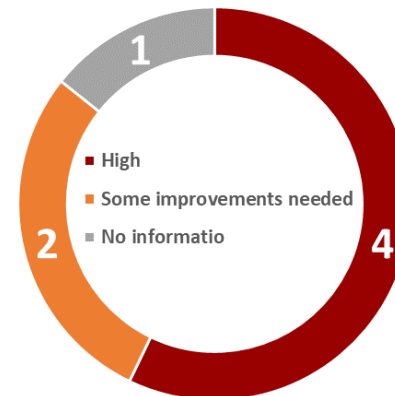


## Perceived performance of AMR prediction using WGS

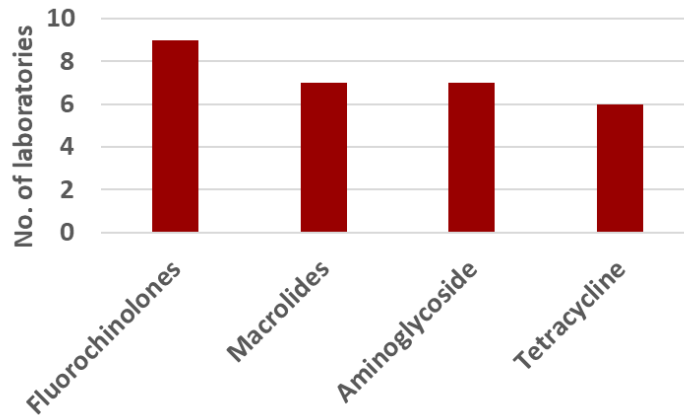
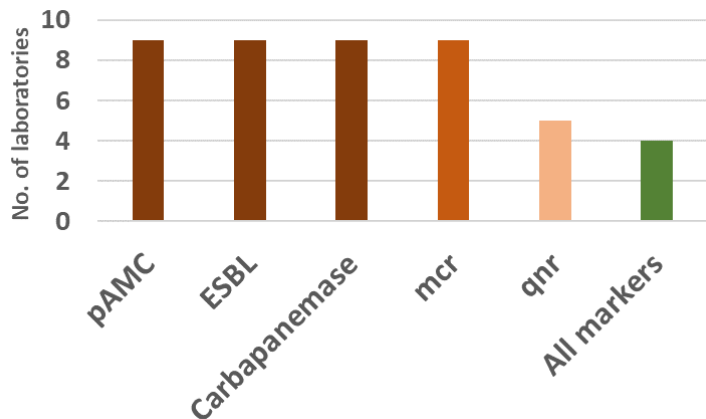
### Salmonella



### Campylobacter

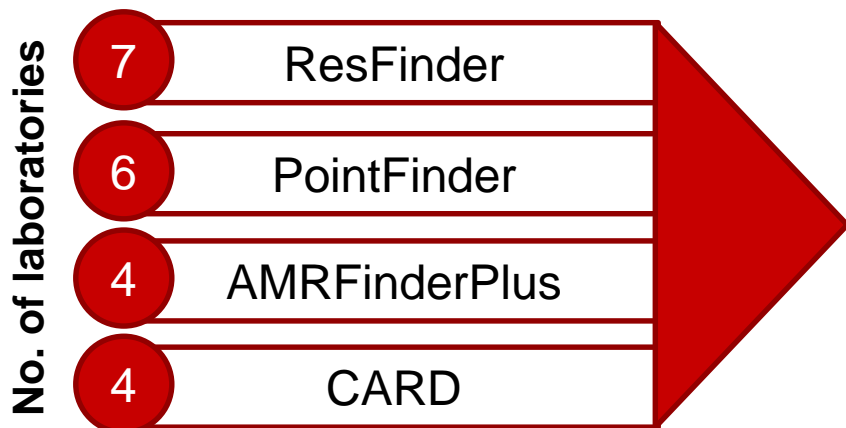


## Genetic markers tested routinely in *Salmonella* and *Campylobacter*\*



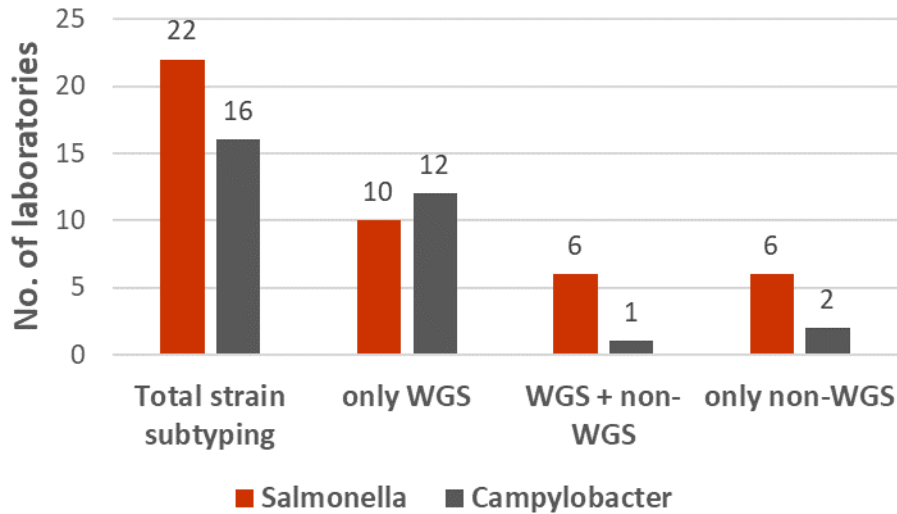
\*at least one genetic marker from each class is tested

## Databases in use for prediction of AMR in *Salmonella* and *Campylobacter*



All laboratories use a combination of databases, and often they are incorporated into certain tools or in-house build pipelines

## Molecular methods in use for strain subtyping



### *Salmonella*

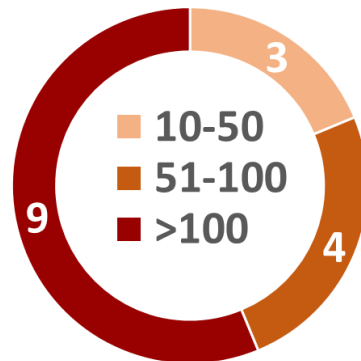
- 16 WGS-based
- 12 non – WGS (PFGE, MLVA)

### *Campylobacter*

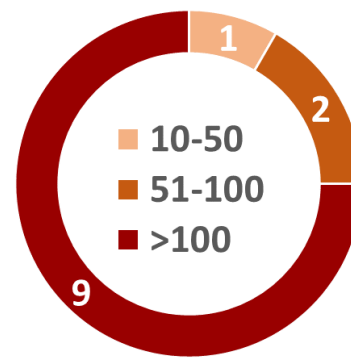
- 12 WGS – based
- 3 non – WGS (PFGE, *flaA*)

## No. of isolates tested using WGS annually

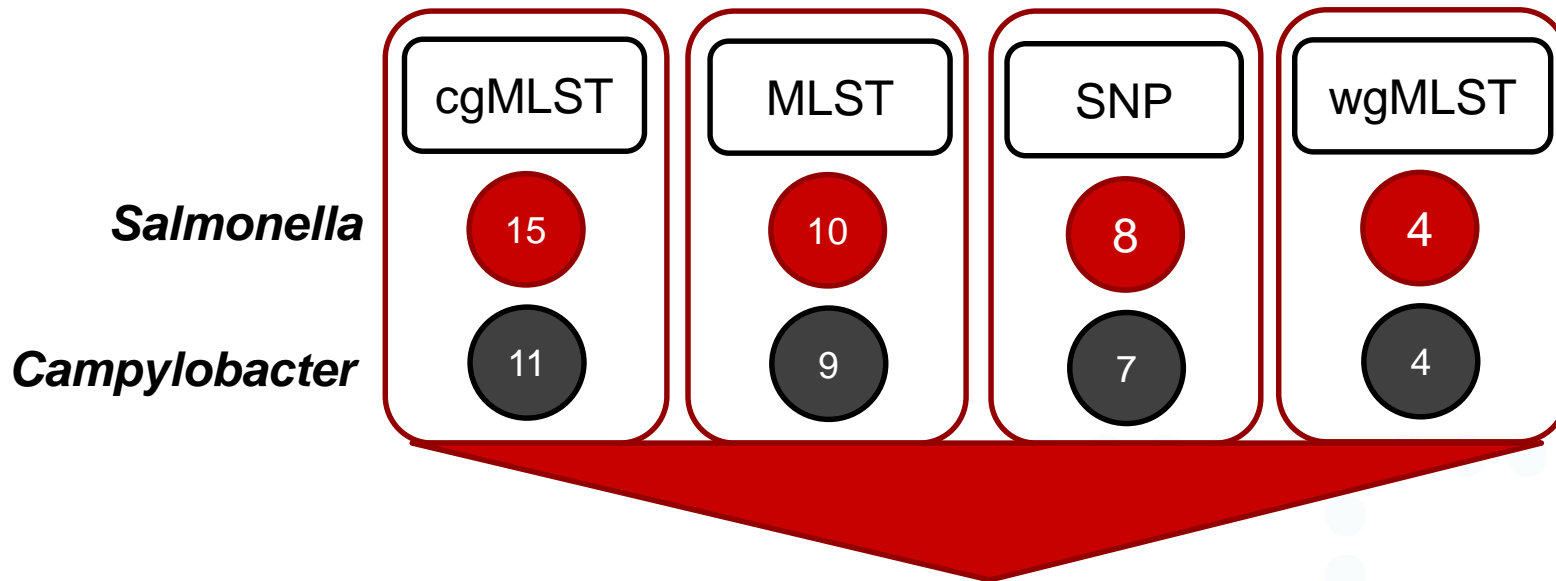
### *Salmonella*



### *Campylobacter*



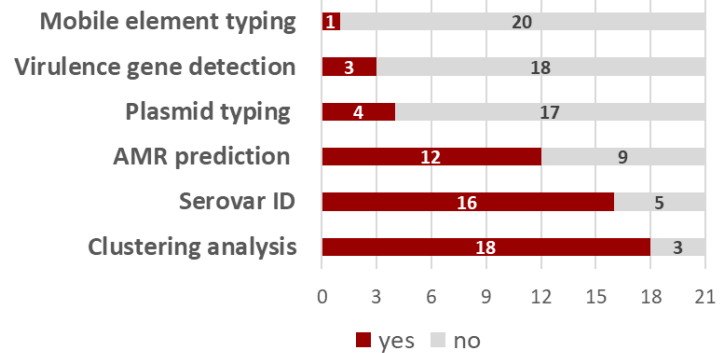
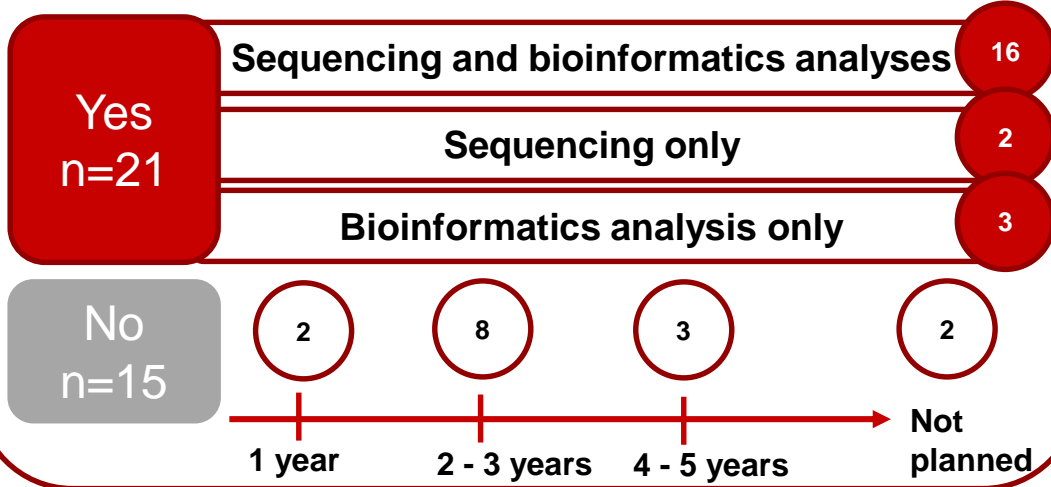
## WGS- based methods in use for strain subtyping



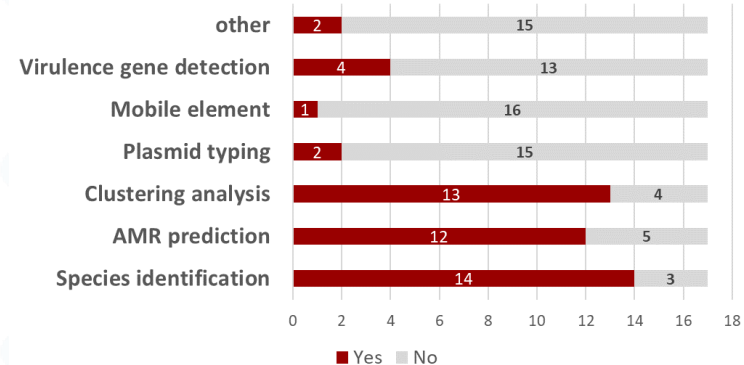
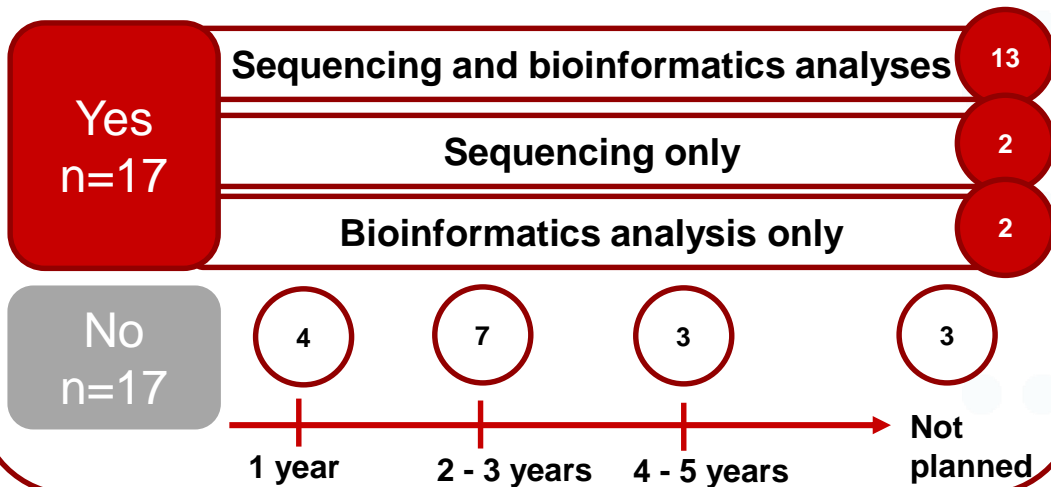
Often a combination of methods are used

# WGS CAPACITY AND UTILISATION

## Current WGS status in 36 countries for *Salmonella*



## Current WGS status in 34 countries for *Campylobacter*



## Commercial softwares in use

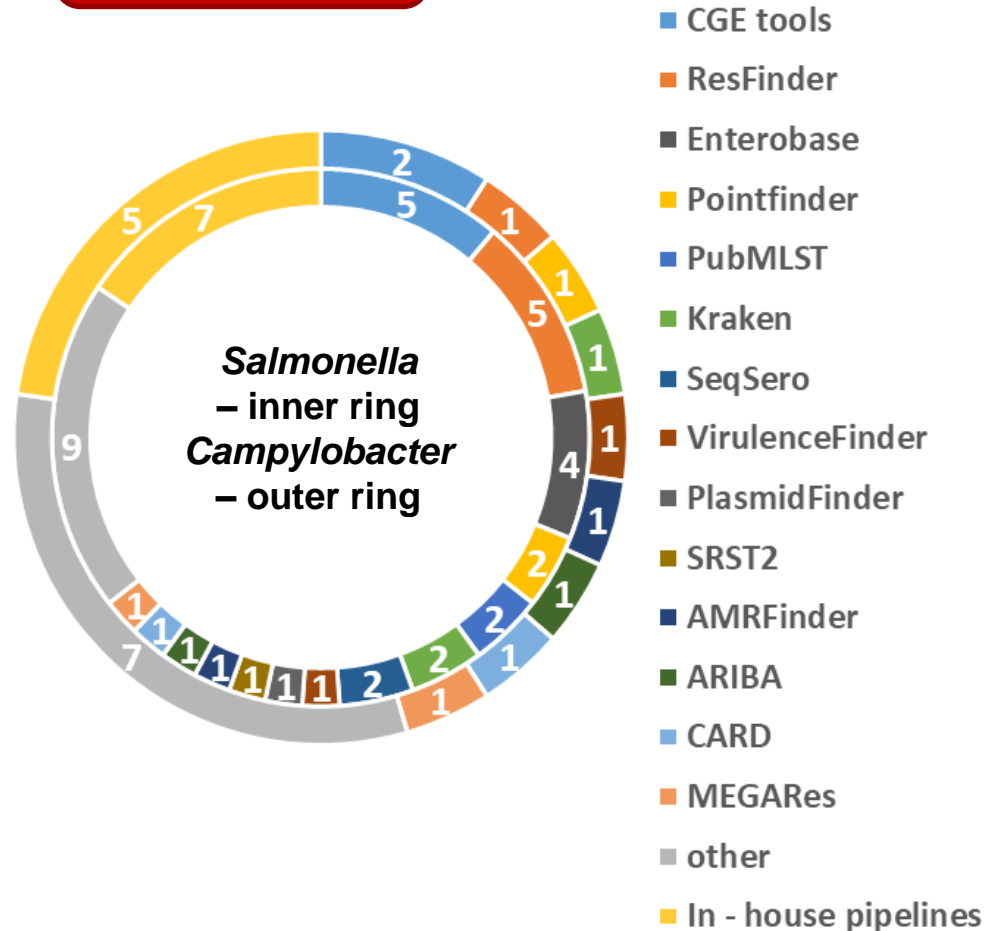
### *Salmonella*

- SeqSphere 10
- Bionumerics 4
- CLC Genomic Workbench 4

### *Campylobacter*

- SeqSphere 6
- Bionumerics 2
- CLC Genomic Workbench 3
- Genious 1

## Open source tools in use





- Up to 61% of countries have WGS capacity, however:
  - Up to 52% use WGS for strain subtyping routinely
  - 23% use WGS for prediction of AMR routinely
- High variation of bioinformatics methods, tools and softwares are used in different countries performing WGS – based prediction of AMR and strain subtyping of *Salmonella* and *Campylobacter*

