

Campylobacter Risk Assessment in the EU: Past, present and future.

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Overview

- Past

- MedVetNet WP 24: a comparison of "*Campylobacter* in broiler meat risk assessments" in Europe

- Present

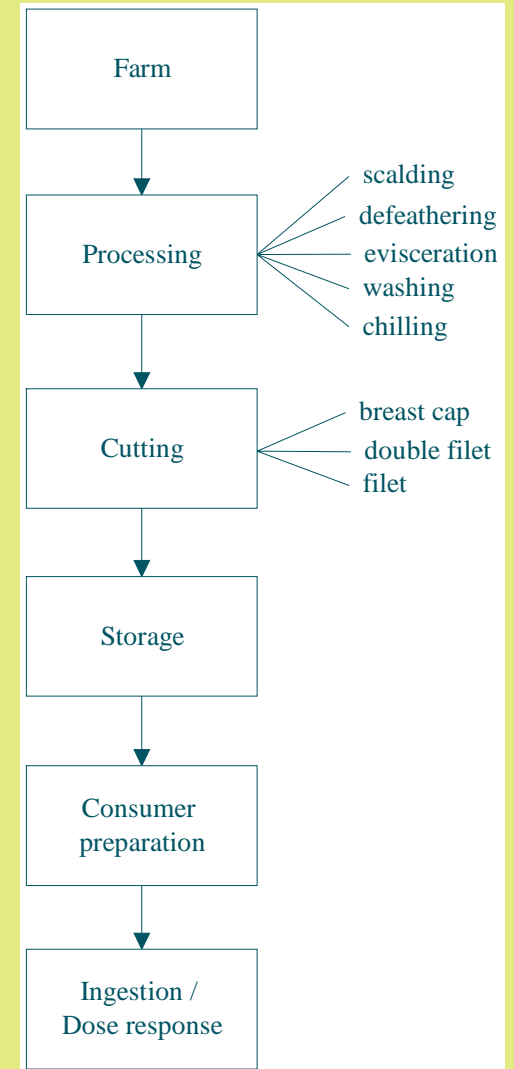
- A consensus framework: CRAF

- Future

- Challenges of European *Campylobacter* QMRA

Risk assessment: what do I mean?

- Food chain risk assessment
- Model describes transmission and survival of *Campylobacter* in the broiler meat chain: changes in distribution of concentrations
- Exposure assessment
+ Dose response = risk
- Quantitative Microbiological Risk Assessment (QMRA) is still developing!



Why we need risk assessment

- **Relative** risk estimates
 - The effects of control measures
 - Comparison of interventions all over the food chain
- Added value
 - food chain data
 - epidemiology
 - below the detection limit
 - check our understanding
- Indispensable for PO / target setting

***Campylobacter* in broiler meat risk assessments in Europe**

• UK	Hartnett	2001
• Denmark	Rosenquist, Christensen	2003
• Netherlands	Havelaar, Nauta	2005
• Germany	Brynstad	2006
• Belgium	Uyttendaele	2006
	Gellynck, Messens	2008
• Sweden	Lindqvist, Lindblad	2008
• Italy	Calistri, Giovannini	2008

MedVetNet Workpackage 24

March 2006 - June 2009



- Objective: consensus on *Campylobacter* QMRA?
- UK, DK, GE, NL models compared
 - Input from New Zealand and FAO/WHO risk assessment

- Differences
 - objectives
 - approach
 - models
 - results



- Similar conclusions

Different objectives



- Gain risk assessment modelling experience
- Human incidence estimation
- Evaluation of risk reduction after intervention and control
 - general
 - specific interventions
 - incl. economic analysis
- Interaction with risk management

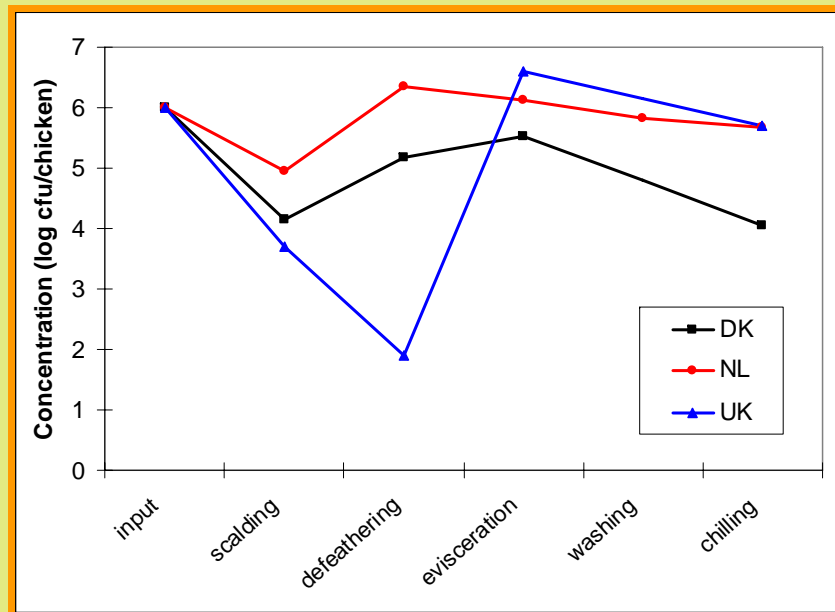
Differences between models

- objectives
- expertise of modellers
- national differences
- data and/or expert opinion
- statistical description and/or dynamic model
- details included in the models
- channel assignment
- end product
 - whole carcass
 - specific product
 - side dish
- but all use quantitative risk assessment
 - probabilistic models



Differences between model results

example



Three chicken processing models with the same input

- different dynamics
- similar end results
- similar effects of interventions (?)

Different end results ?

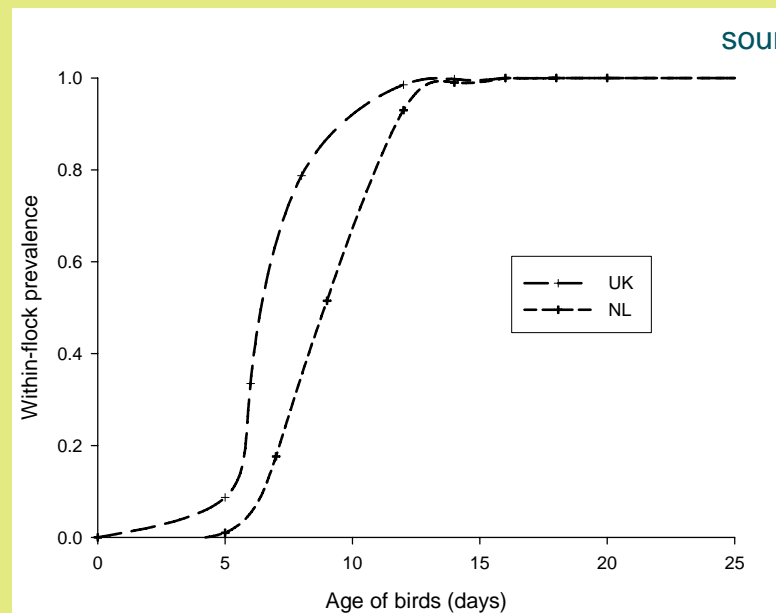


- Varying human incidence estimates
 - Differences in models, (national) data and assumptions
 - Risk estimates are uncertain
 - Not easy to decide what is the "main cause" of differences in results
- Evaluation of risk reduction after intervention
 - In general similar, despite quantitative differences
 - Relative risk estimates are less uncertain

Similar conclusions (1)



- Farm models predict many low prevalent flocks at the farm that may not be detected



- False negative flocks occur frequently

Similar conclusions (2)

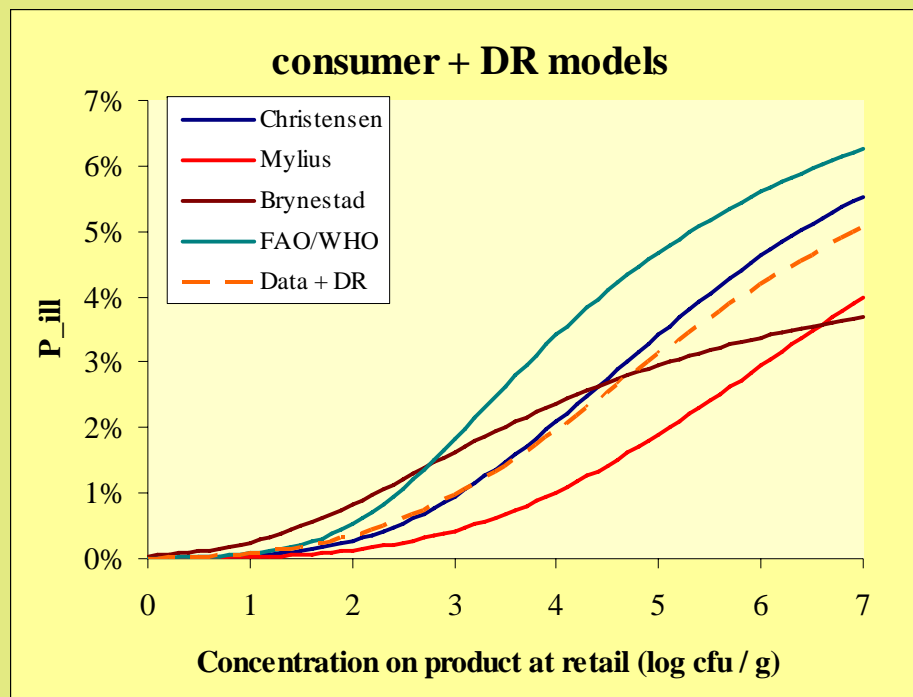


- "Logistic slaughter" has little effect
 - No growth of *Campylobacter* in processing environment
 - Each model MUST predict that concentrations on carcasses of cross contaminated flocks are lower
- Data:
 - Typing shows *Campylobacters* are transmitted from one flock to the other (e.g. Miwa et al. 2003)
 - Transferred quantities are small (Johannessen et al. 2007)

Similar conclusions (3)



- High concentrations pose the largest risks



Nauta *et al*, unpublished results

- targetting high concentrations is an effective intervention
- get data on *distributions* of concentrations, not just *means*
- confirmed by Callicott *et al*. (2008)

Conclusions from WP 24



- QMRA model must be fit for purpose
 - *different purposes require different models*
 - *balance between simple and complex*
- Many modelling methods explored
 - *try to combine the good qualities of different models*
- Similar conclusions !
 - *useful insights for risk managers*
- No consensus European Risk assessment Model
 - *no single purpose, many national differences*
- Towards a consensus Approach
 - *development of Campylobacter Risk Assessment Framework (CRAF)*

Campylobacter Risk Assessment Framework CRAF



- Software tool for risk assessors
- Structured information on five Campy QMRAs
- Compare and link models for modules
- An aid to make your own *Campylobacter* QMRA

19 February 2009 :

The Final General Meeting of MedVetNet Workpackage 24

20 February 2009 :

Campylobacter Risk Assessment Framework (CRAF)
Training Course

hosted by BfR in Berlin

INVITATION

rivm

National Institute
for Public Health
and the Environment

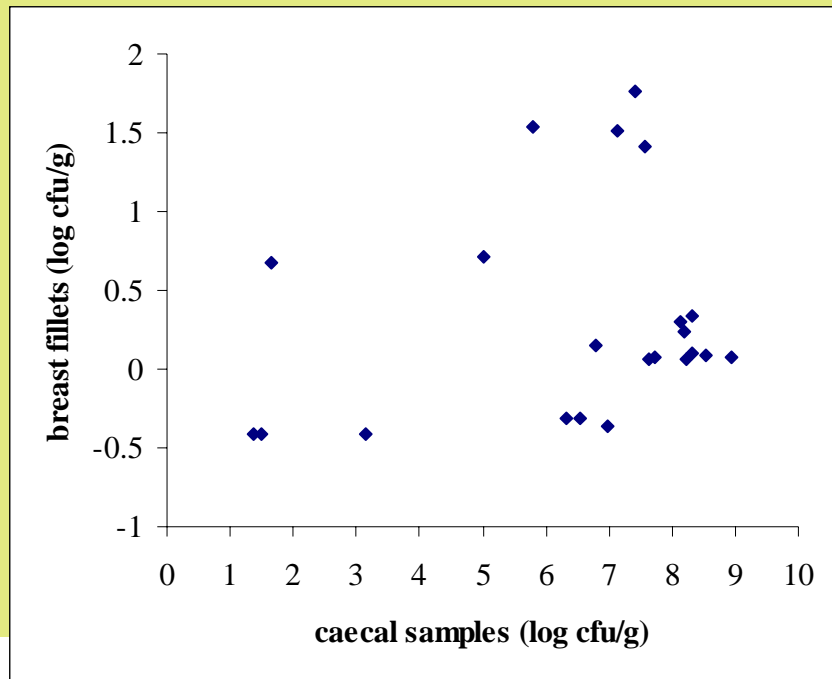


New *Campylobacter* QMRA in Europe (1)

- Baseline data from caeca and neck skins
- Challenge: how to relate those data to risks?
 - QMRA models don't have either of them as inputs
 - Data don't always show a good link caecal samples - meat products; why not?

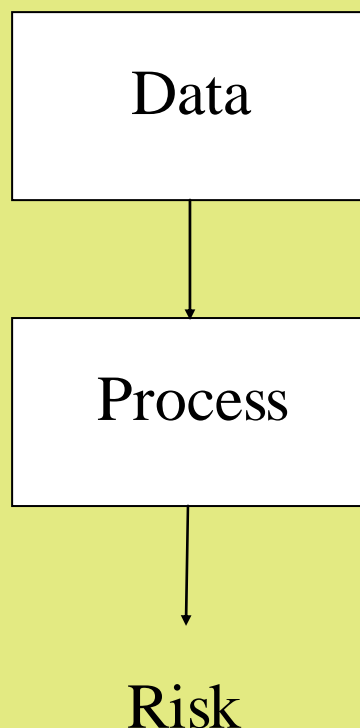
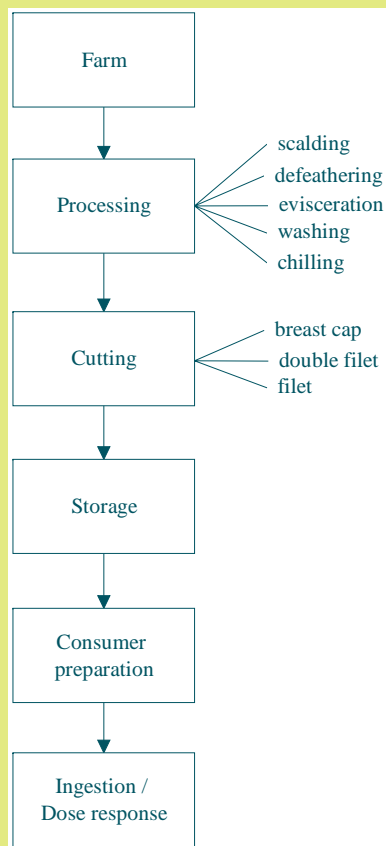
product	caeca		tot
	-	+	
	-	+	tot
-	17	19	36
+	4	22	26
tot	21	41	62

source: Nauta, Bolder *et al*, in prep.



New Campylobacter QMRA in Europe (2)

- Target setting: link with human health risks



Quantitative distribution

Differences between member states

Challenge: How to model the differences for each MS?
How important are those?

Take away messages

- Much *Campylobacter* QMRA experience in Europe, but

- different objectives and approaches
- different results

still

- similar and useful conclusions

- European *Campylobacter* QMRA needs

- a clear objective
- further development of QMRA modelling
 - integration of good ideas
 - balance between complexity and simplicity
 - incorporation of differences between MS