

# Hypothetical examples of FSO-derived Microbiological Criteria *Campylobacter* and *Salmonella* in raw poultry

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## Contents

ALOP/FSO/MC  
Dose-response  
MC for prevalence  
preventing the extremes ?

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## number of cases (ALOP)

$$\# \text{ of cases} = P \cdot C \cdot M \cdot S \cdot r$$

*P*: Prevalence

*C*: not log *C*!

*M*: mass per serving

*S*: amount of servings/ year

*r*: virulence, state

FSO based on *P* and *C*  
also compliance ?

## Dose response: *Salmonella*

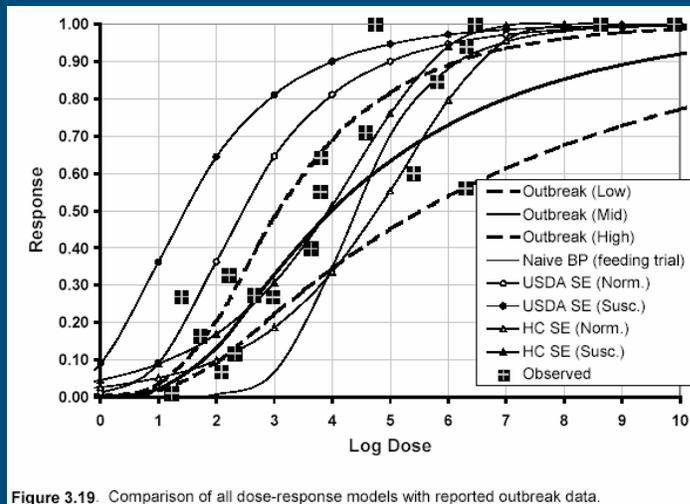


Figure 3.19. Comparison of all dose-response models with reported outbreak data.

## Dose response: *Salmonella*

Risk assessments of *Salmonella* in eggs and broiler chickens

89

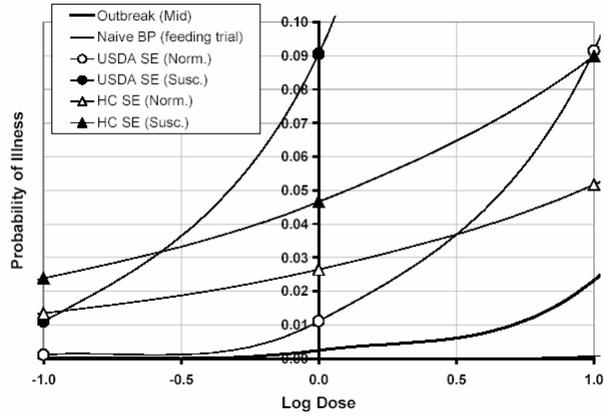


Figure 3.21. Comparison of alternative dose-response models in the -1.0 to 1.0 mean log dose interval.

## Dose response *Campylobacter*

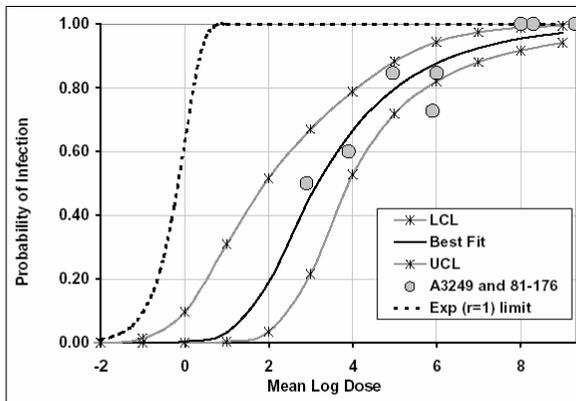


FIGURE 5.1 Beta-Poisson dose-response relationship for the probability of infection for *C. jejuni* based on human feeding trial data and two strains (A3249 and 81-176) and model parameters,  $\alpha = 0.21$ ,  $\beta = 59.95$   
LCL - Lower confidence limit      UCL - Upper confidence limit

## Dose response

Large uncertainty (“impossible” to gather real data)

Often relative risk can be estimated more accurately

*P* from 10% to 5%: factor 2 reduction in illness

*C* from 2 logs to 1 log: in certain cases 90% reduction of illness, sometimes less

## From FSO to PO

P/C at slaughter

storage

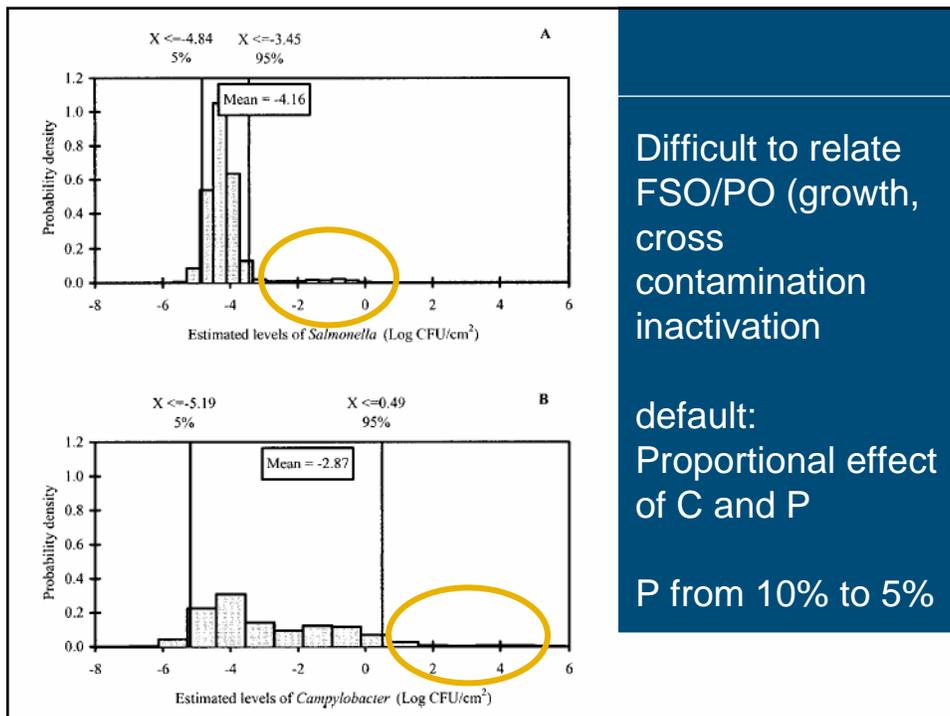
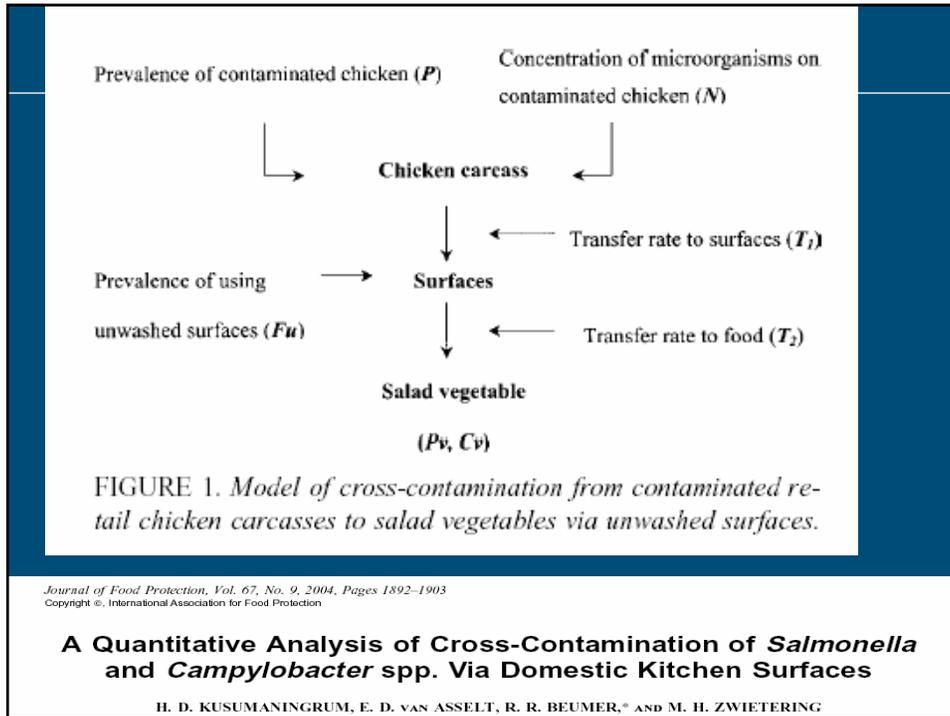
transport

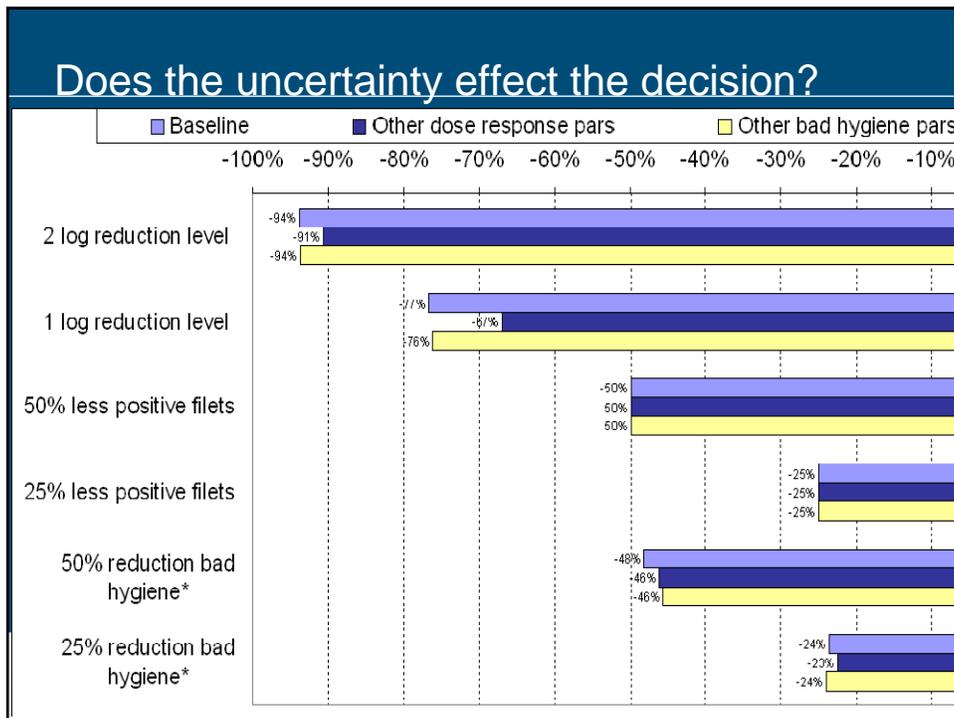
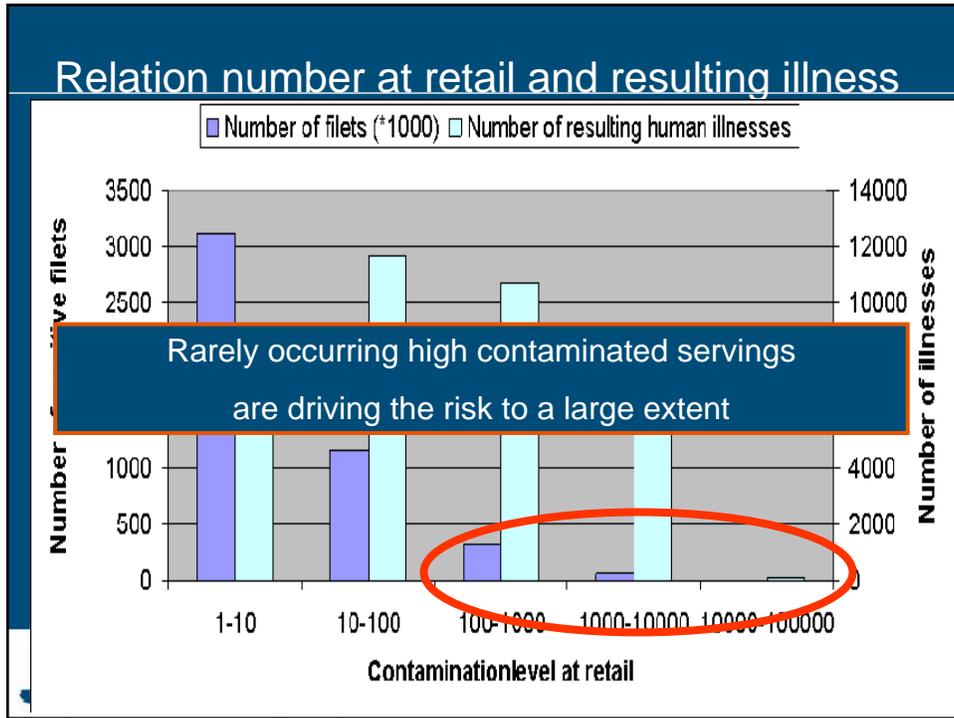
storage

preparation

survivor of (inadequate) cooking

recontamination (hands, tools, surfaces)





## From PO to MC ?

Objective that must (100%?) be reached

Standardised Method

Measurement inaccuracy

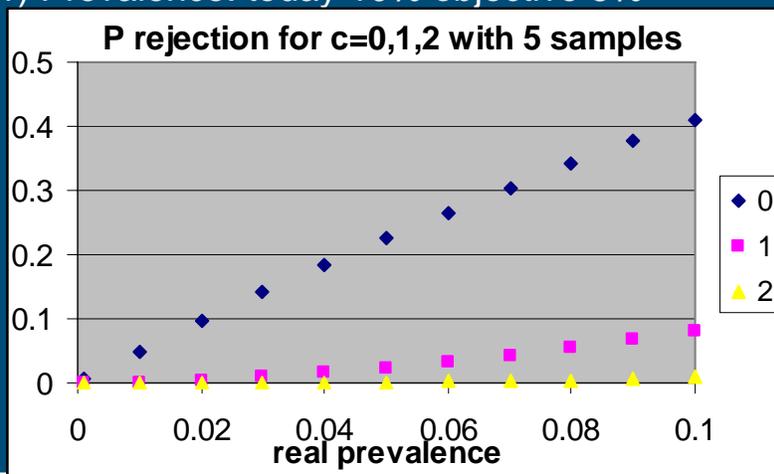
Sampling frequency: false positives/false negatives

- fresh product, only way to make “virtually” Salmonella- or Campylobacter-free is heating and irradiation

- decontamination might have relevant effect

## Criteria: presence/absence in 25 g

1) Prevalence: today 10% objective 5%



## What would be the result of $c=0$

$C=0$ :

real  $P=0.05$  OK:  $P_{\text{reject}}=0.23$   $(1-0.95^5)$

real  $P=0.10$  not OK:  $P_{\text{accept}}=0.59$   $(0.9^5)$

test is not discriminative

even with 20 samples 12% of not detecting  $P=0.1$  !

Better to focus on record of safety ?

Force by ratio  $P_{\text{detection}}/\text{costs}$

## How are criteria related to interventions

$P_{\text{detection}}$

fine:

- 100€	100€
- rework batch	1.000€
- destroy batch	10.000€
- blame and shame	100.000€
- out of business	10.000.000€

It is not the sampling scheme as such that determines policy and intentions producers

## Approach

- 1) Set level for P
  - 2) If more known about effects P/C set combined level P/C
- 
- 1) method to prove compliance
  - 2) method to reject highly contaminated batches
- 
- 1) For example record of safety  $P=0.05$  (yearly basis)
  - 2) Batch where  $>100$  cfu/g is rejected

## Conclusions

Difficult to structurally relate ALOP/FSO/MC for fresh products where the main route is by mishandling /recontamination

RA can give clues for relative effects of interventions

record of safety / batch sampling only to detect/reject extremes

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