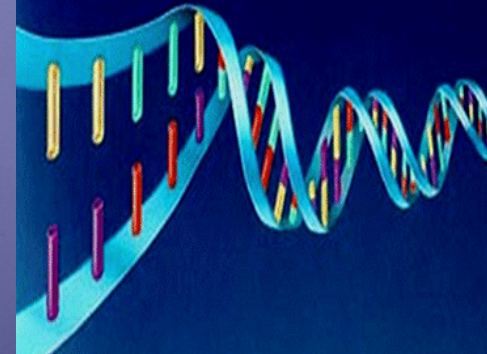


Modern techniques to assess shelf life and safety

TNO Quality of Life

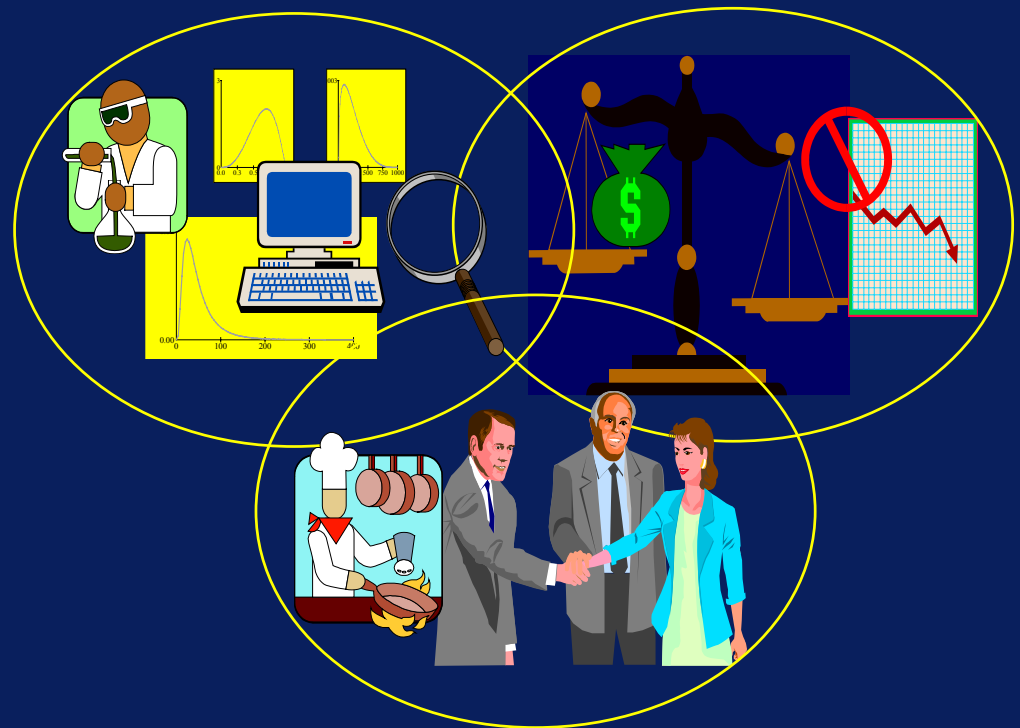


Erik Hoornstra



Introduction

- Principles of risk assessment
- Quantitative risk assessment
- Example
- Risk assessment in TTI
- Safety and shelf-life



TNO Department Microbiology

- **Research during product and process innovation**
- **Novel preservation strategies**

- **Testing and Validation**
- **European Hygienic Engineering and Design Group (EHEDG) certified test laboratory**

- **Trouble-shooting and emergencies (24 hour service)**
- **Rapid detection tools**
- **New developments: genomics & metabolomics**

The challenge for food companies

Safety
Freshness
Convenience
Shelf-life

Government FSO's
Customer requirements
Consumer demands
Company objectives

Criteria for:

- **Product**
- **Process**
- **Storage**

Question

What is safe food ?

Why define safe food

EU

- **“The free movement of safe and wholesome food ...”**
- **“Food shall not be placed on the market if it is unsafe”**
- **“A high level of protection of human life and health ...”**
- **“... free movement ... only if food safety requirements do not differ from Member State to Member State”**
- **“The Community has chosen a high level of health protection ...”**

Other aspects of defining safe food

EU

- **“It is necessary to ensure that consumers, other stakeholders and trading partners have confidence in the decision making process ...”**
- **“... risk analysis provides a systematic methodology for the determination of effective, proportionate and targeted measures to protect health ...”**
- **“... scientific risk assessment alone cannot provide all the information on which a risk management decision should be based ...societal, economic, traditional, ethical and environmental factors and the feasibility of controls”**
- **Precautionary principle: risk is identified but unknown / uncertainty**

Who should define food safety

EU

- “... the European Food Safety Authority should provide a comprehensive independent scientific view of the safety of the food ...”
- “ EFSA should be able to commission scientific studies ...”

Remember

- Defining food safety is a political issue

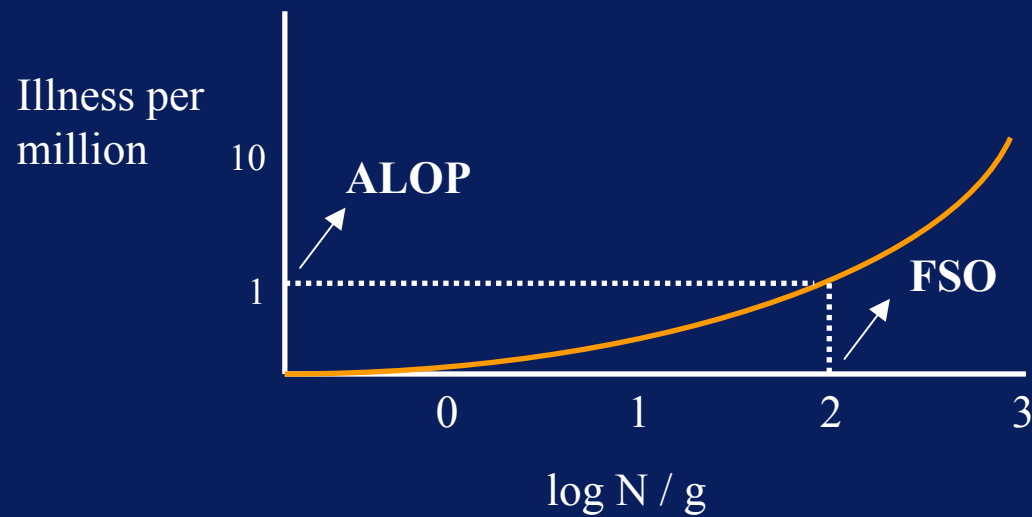
Safe food

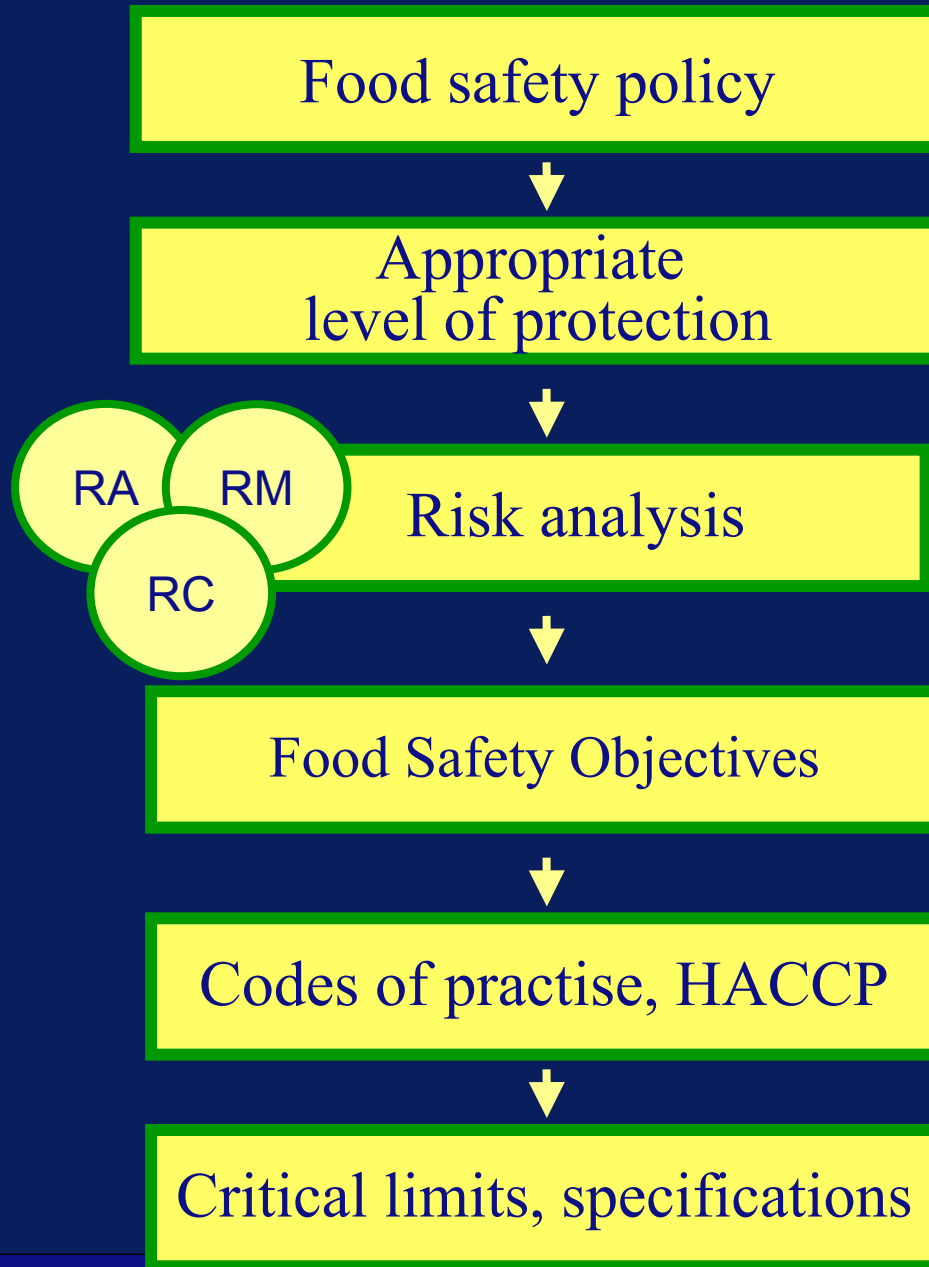
EU

- **Unsafe = “Injurious to health”**
- **Unsafe = “Unfit for human consumption”**
- **“... normal conditions of use of the food by the consumer and at each stage of production, processing and distribution”**
- **“... information provided to the consumer ... information on label ...”**
- **“... according to its intended use.”**
- **“... any food which is unsafe is part of a batch ... all food in that batch is unsafe, unless ... detailed assessment ...”**
- **“Traceability: Identify any person from whom they have been supplied ...”**

Safe food – ALOP and FSO

- **ALOP** = appropriate level of protection
- **FSO** is maximum level of a hazard at the moment of consumption





Safe food – Derived concepts from FSO

Performance objectives (in the food chain)

- “*Salmonella* shall not exceed 1 cfu / 10 ml before distribution”
- ‘*Listeria* absent per 25 gram in chilled foods (where it is able to grow)’

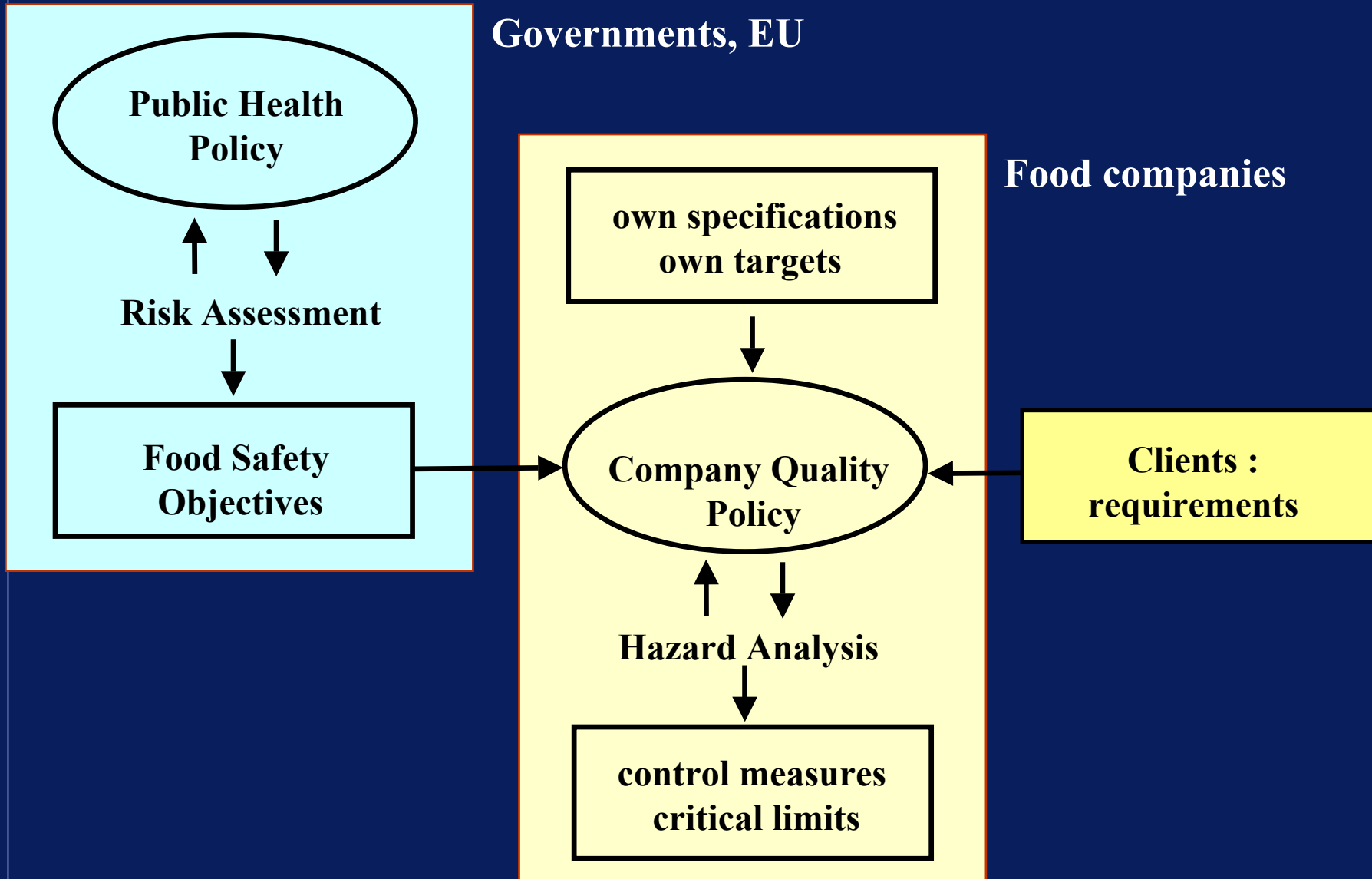
Performance criteria (change during a step)

- “Assure a 12 log reduction of *C. botulinum* in low acid canned foods”
- “Juice process should achieve 5 log reduction of *E. coli* and *Salmonella*”
- “Avoid more than 3 log increase of *S. aureus* during meat fermentation”

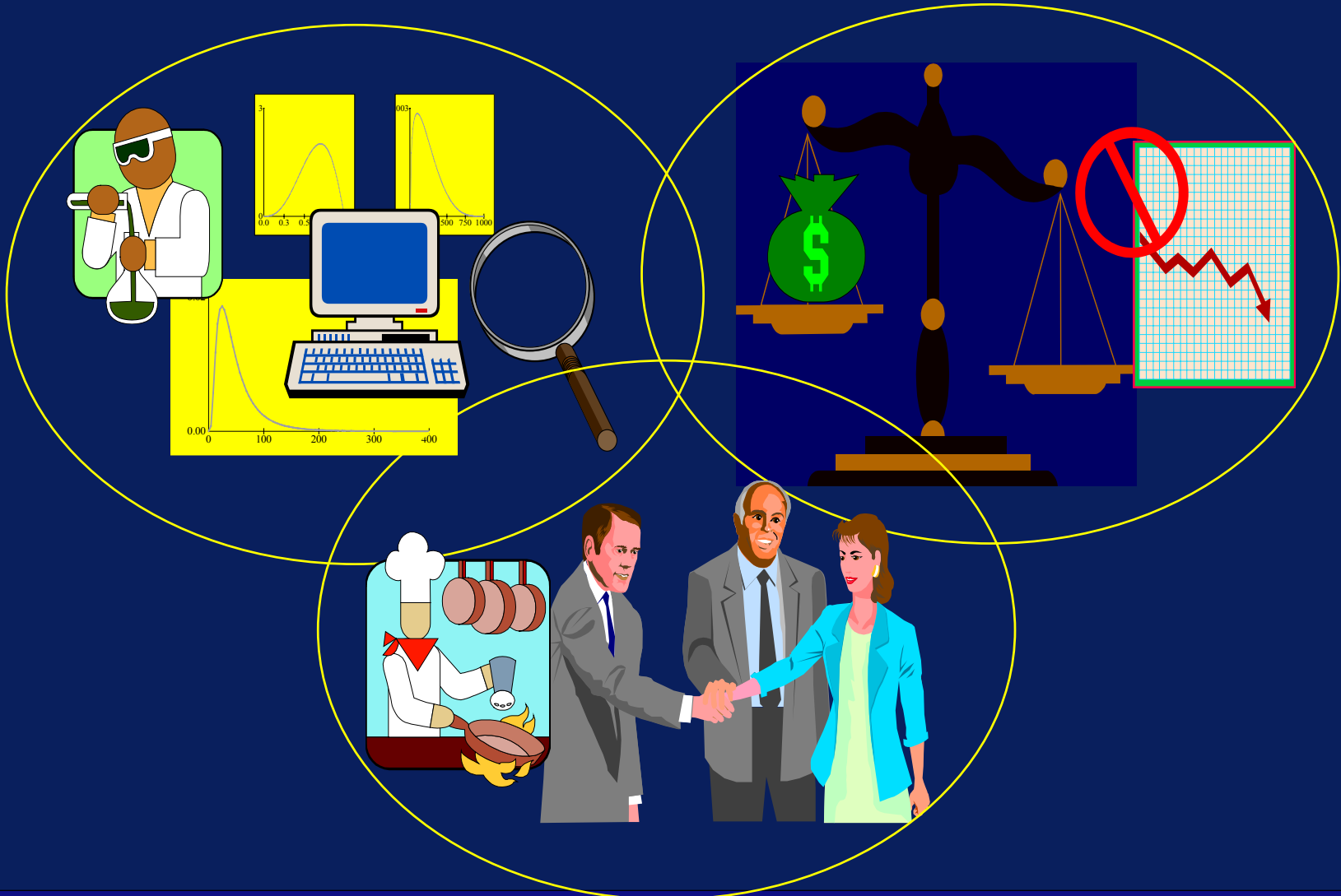
Process criteria

- 2.4 minutes 121°C for sterilisation
- 15 seconds 71°C for commercial pasteurisation of milk
- Storage temperature < 7 C

Risk assessment at two levels



Risk analysis



Hazard identification

Government

- Epidemiology
- (Public) opinion

Food company

- Raw materials
- Production process
- Product: complaints, history



Hazard characterization

Government

- **Dose-response relation**
 - Virulence of microorganism
 - Consumer sensitivity

Food company

- **Criteria**
 - Food Safety Objectives
 - Legislation
 - Specifications



Exposure assessment

- **Risk factors**
 - Occurrence in raw materials
 - Growth
 - Inactivation
 - Mixing, portioning
 - Recontamination
- **Quantification**
 - Single point: worst-case
 - Single point: what if
 - Probability distribution functions



Risk characterization

- Risk estimate: 0.....1
- Risk classes: f (occurrence, severity)
- Risk profile
- Prioritize control measures
- Scenario analyses
- Uncertainty and variability



Learnings from first years of experience using QRA (negative)

- Quantitative risk assessment may be time consuming
- Sometimes trivial results
- Uncertainty about risk factors
- Results difficult to understand by managers
- Lack of data
- Do we really understand what is going on



Learnings from first years of experience using QRA (positive)

- **Zero risk does not exist**
- **Quantification gives a lot of insight**
- **Uncertainty and variability**
- **List most important risk factors**
- **Scenario analysis for efficient improvements**
- **Meaningful sampling**



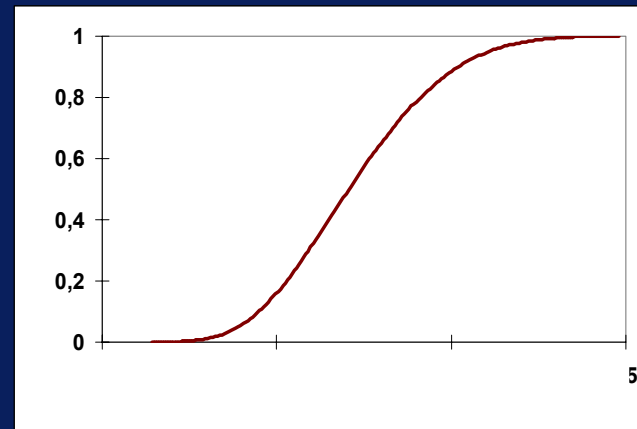
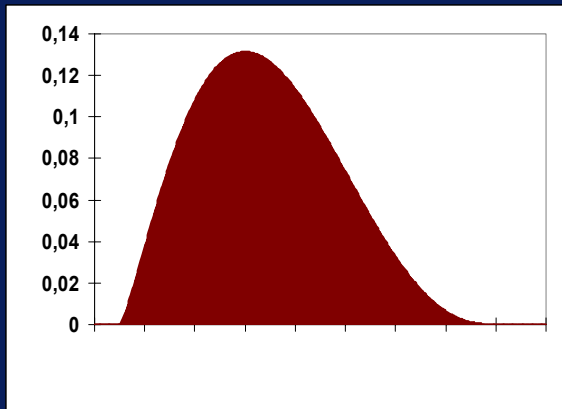
Single point estimation vs. Probability distribution functions

$$10 \times 10 = 100$$

$$1 \times 10 = 10$$




$$5 \times 10 = 50$$

$$20 \times 10 = 200$$



Single point calculations

Example: total travel time = **bicycle** + **train** + **bus**

	minimum	most likely	maximum
	5	10	20
	35	40	60
	10	15	20

Result

50 min.

65 min.

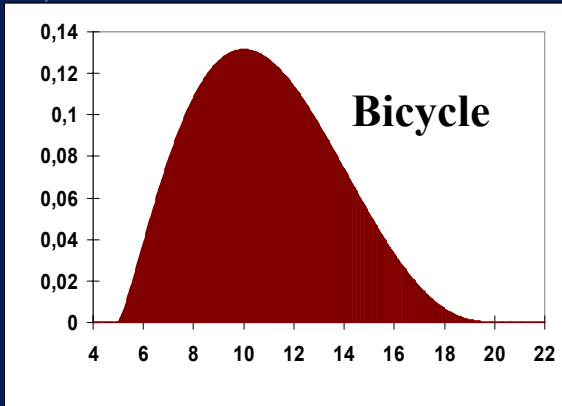
100 min.

best case

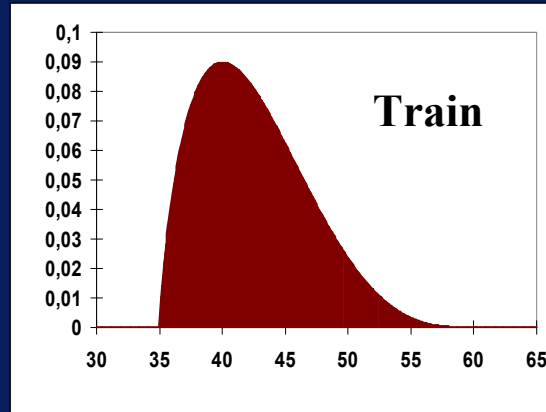
most likely

worst case

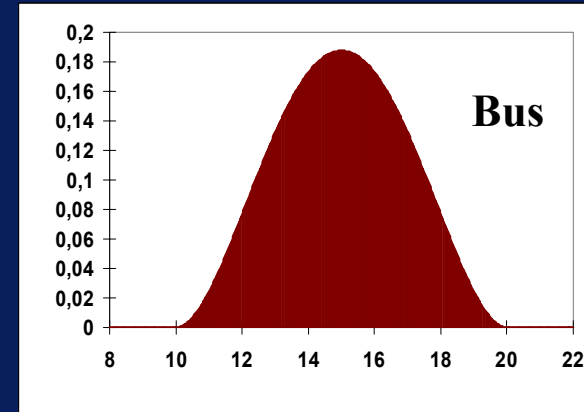
Probability distribution functions



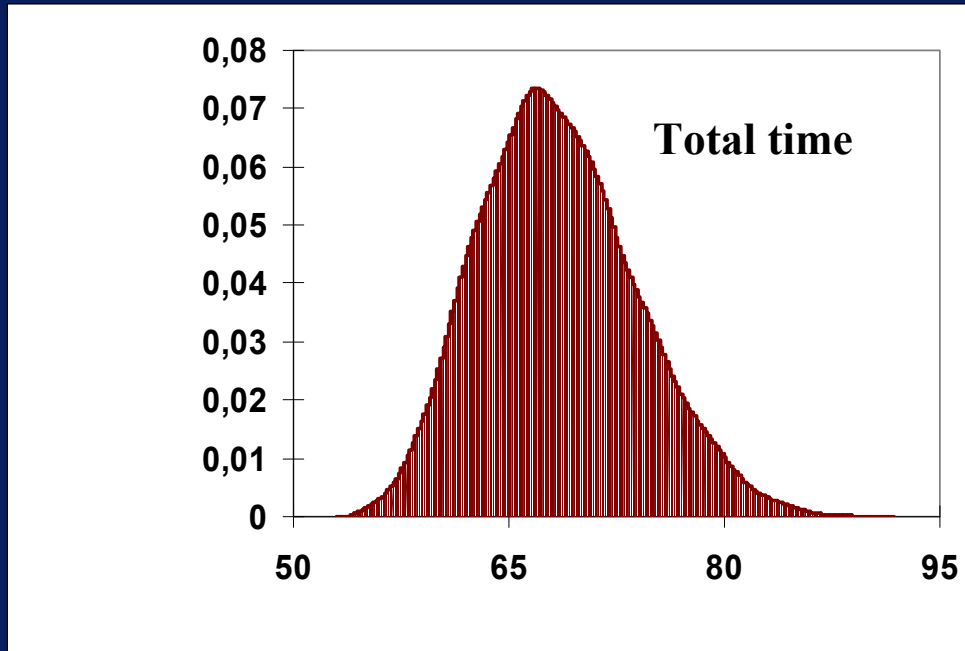
+



+



=



Average time 68 min

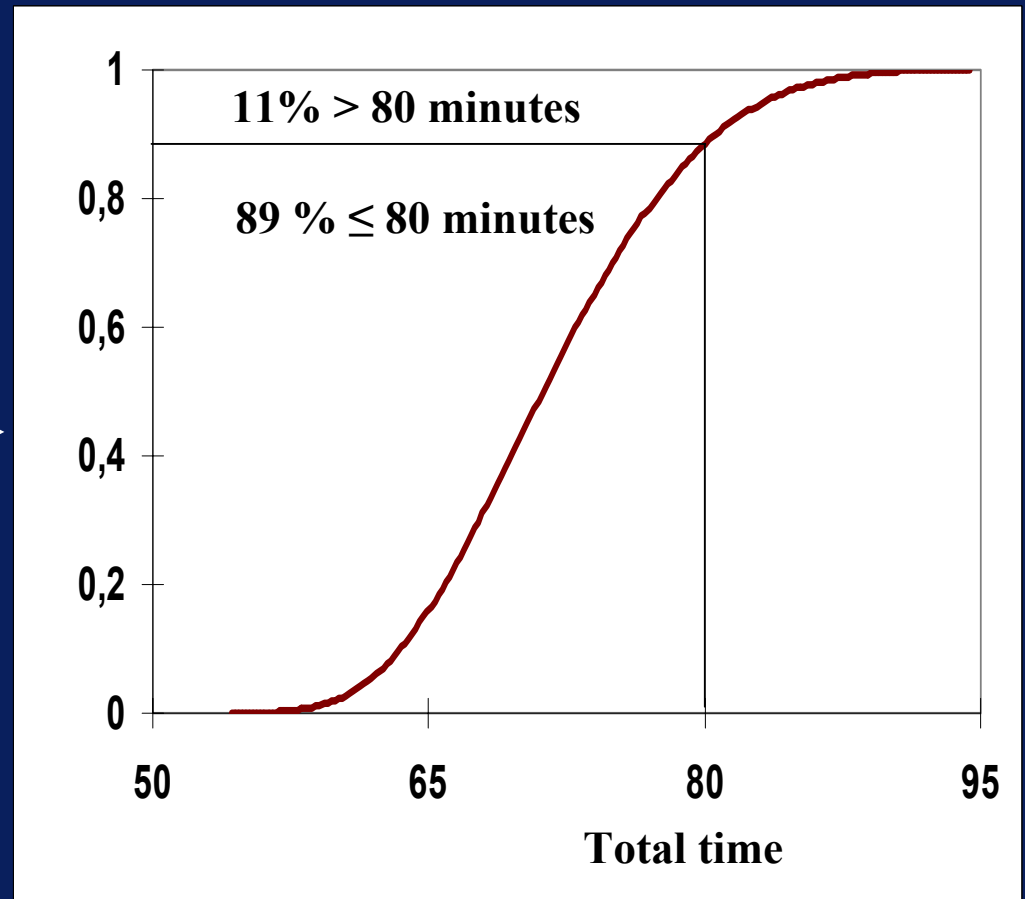
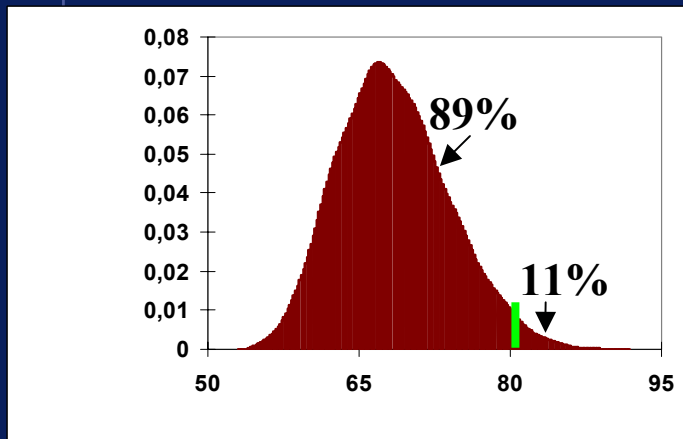
Time < 60 min 5.1 %

Time > 80 min 11 %

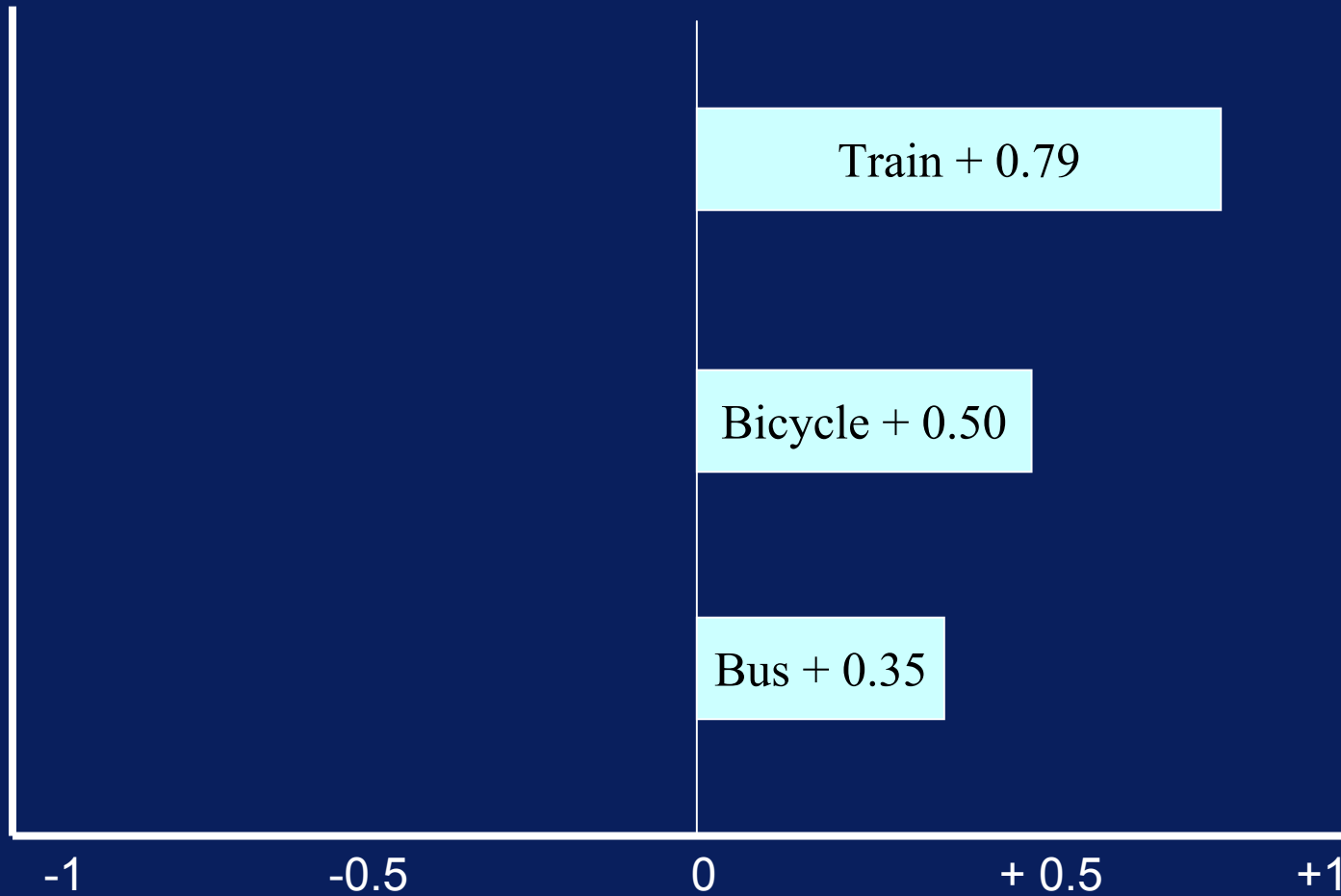
Time > 90 min 0.01 %

Probability distribution functions

Cumulative probability function



Sensitivity analysis



Monte Carlo simulation in @Risk

@RISK

File Edit Settings Variables Execute Results Window Help

Open Save Sim Sett +Output List Simulate Results Graph Summary Hide

Results

Simulation

Simulation #1:
Boerenmetworst.xls
Iterations= 100000 Simulations= 1
Input Variables= 19
Output Variables= 26
Sampling Type= Latin Hypercube
Runtime= 02:33:55
Run on 18-12-98, 15:21:15

Summary of Results

Cell	Name	Minimum	Mean	Maximum
E6	C feces / E	-1,276251	2,091336	5,559983
E7	overdracht / E	-9,669935	-5,100007	-1,21099
E9	C cm snippers / E	1,6506E-09	0,3658856	711,314
E12	oppervlakte / E	0,7504331	1	1,249238
E14	C g snippers / E	1,476751E-05	0,3683755	785,5647
E16	m / E	113,6647	135	157,1977

Statistics Data Sensitivity Scenarios

Simulation Statistics

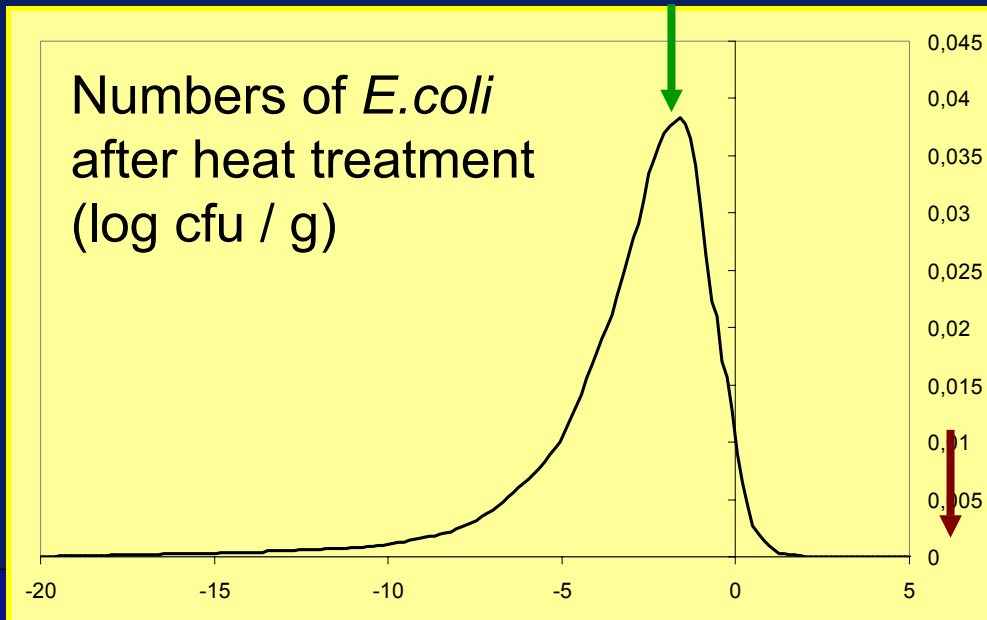
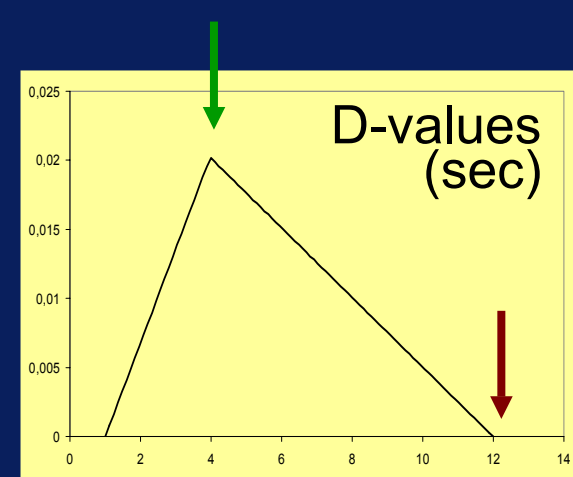
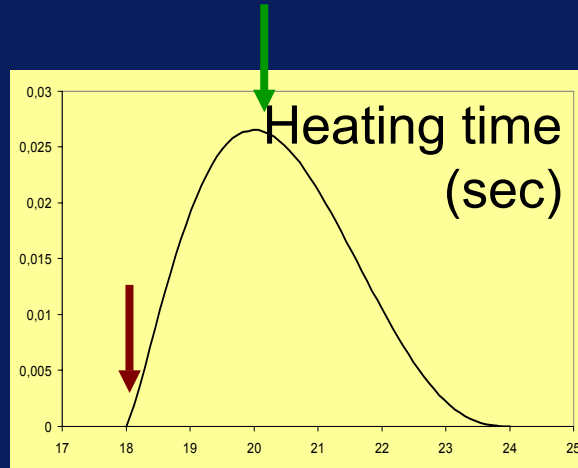
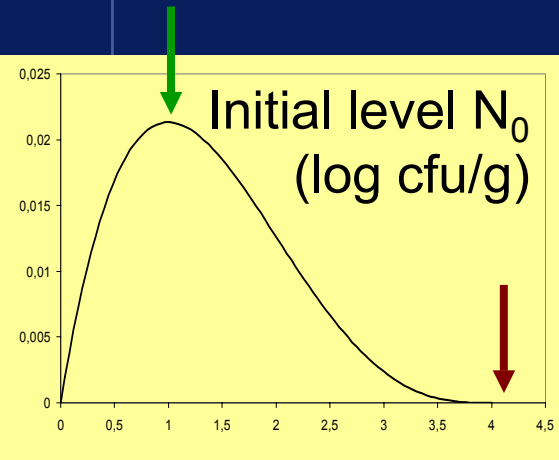
Name	overdracht / E	C cm snippers / E
Description	Output	Output
Cell	E7	E9
Minimum =	-9,669935	1,6506E-09
Maximum =	-1,21099	711,314
Mean =	-5,100007	0,3658856
Std Deviation =	0,8999839	6,473752
Variance =	0,809971	41,90947
Skewness =	-4,848558E-04	64,08714

Current Variables: Outputs=26, Inputs=19 Settings: Simulations=1 Iterations=1

Output Graph - Cell E7

Distribution for overdracht / E/E7

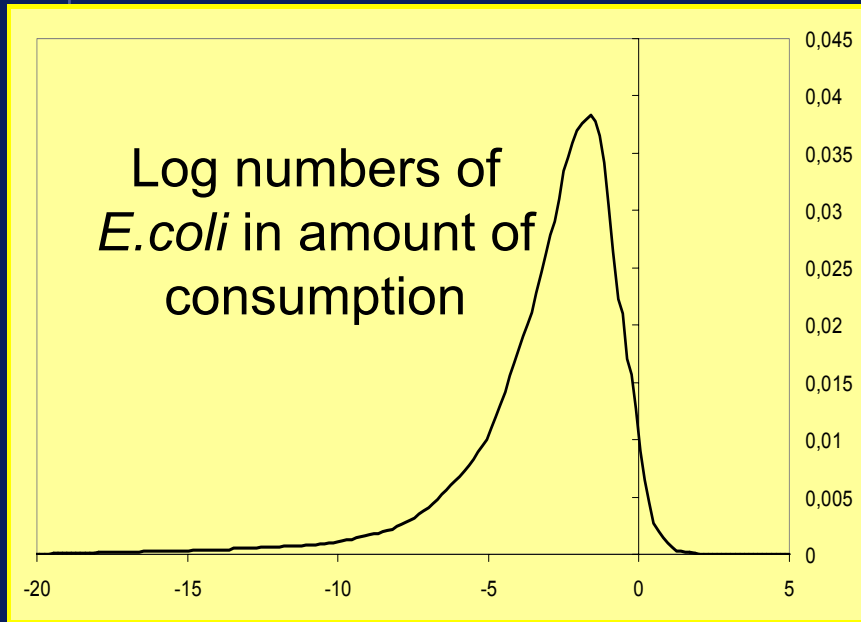
Exposure assessment of *E. coli* O157:H7



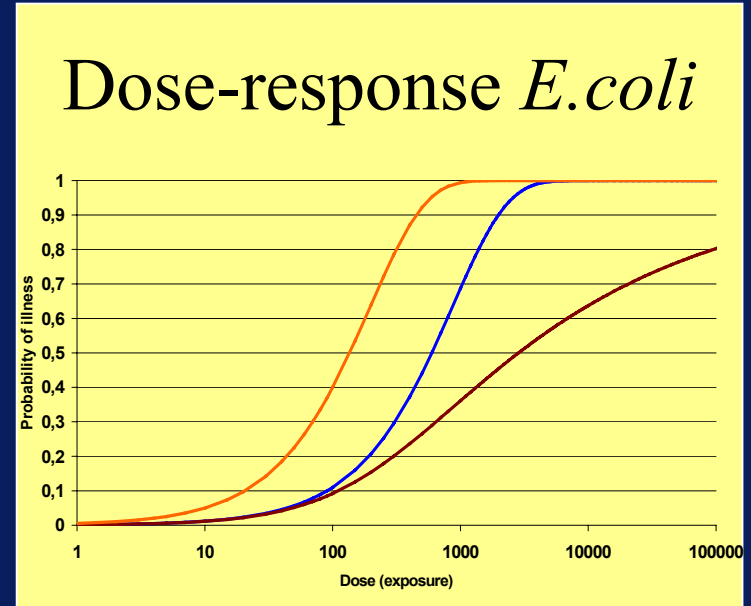
↓ Worst case situation

↓ 'Average' situation

Risk characterisation of *E. coli* O157:H7



X



=

Probability of illness after consumption

- Average, most likely, 90% confidence interval
- Probability of exceeding ALOP

Risk assessment of *E. coli* O157:H7 in raw fermented sausage

- Pathogens can be present in raw meat
- Fermentation results in reduction of pathogens
- No additional heating step



Risk assessment of *E. coli* O157:H7 in raw fermented sausage

	contamination	performance	end product
• worst-case	1000 / g	5 D	1 / 100 g
• “what if”	1 / g	2 D 3 D	1 / 100 g 1 on 10 with 1 / 100 g

If 1000 / g occurs in 0.1% of cases

If 1 / 100 g results in probability of illness of 0.1%

5D results in risk of 1 in million
ACCEPTABLE ???

3D results in risk of 1 in 10,000
ACCEPTABLE ???

Risk assessment in TTI

Slaughterhouse



contamination



contamination



contamination

Chill chain (distribution and retail)



growth

Consumer



growth



inactivation
recontamination



intake

Initial level

- % positive

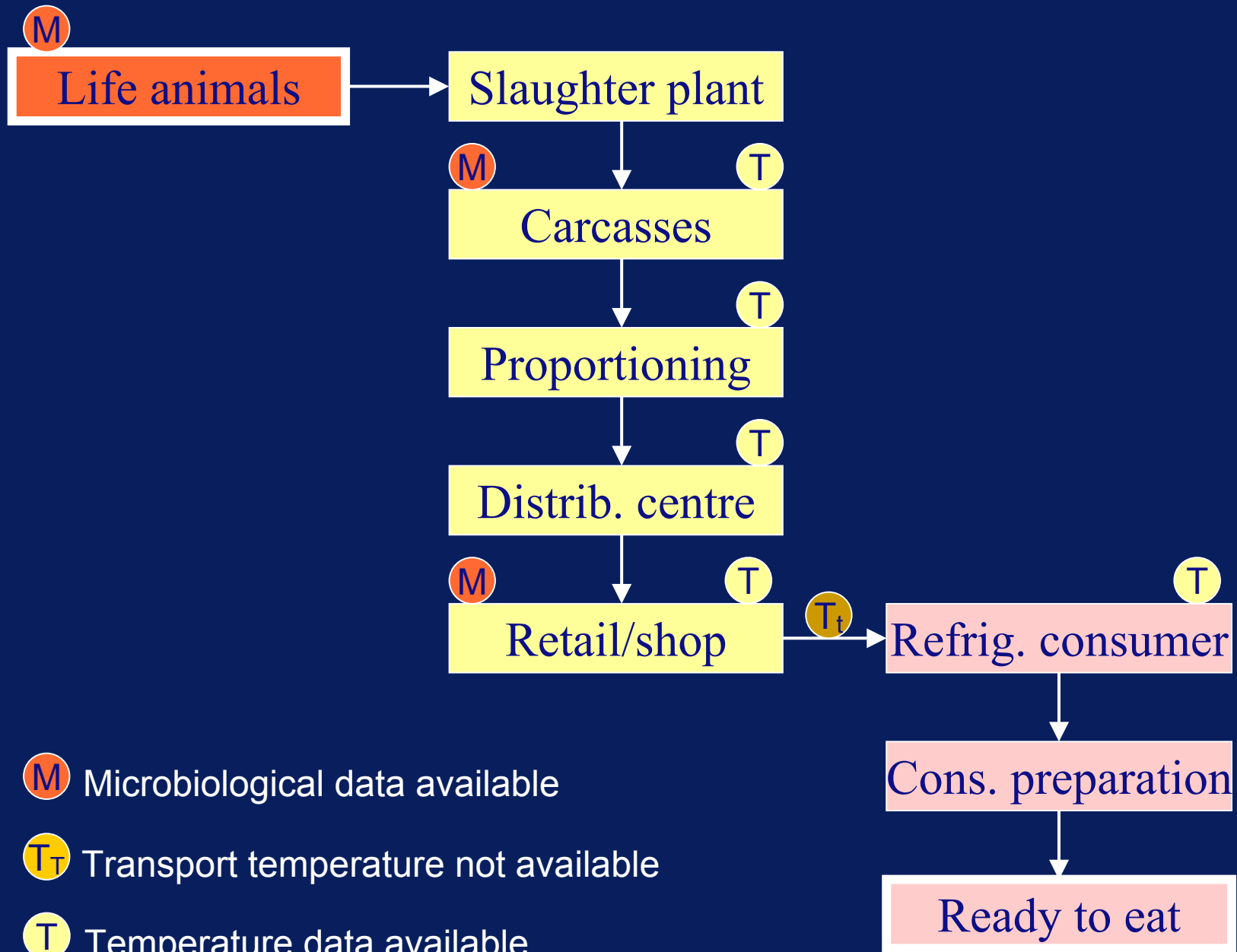
- numbers

Growth

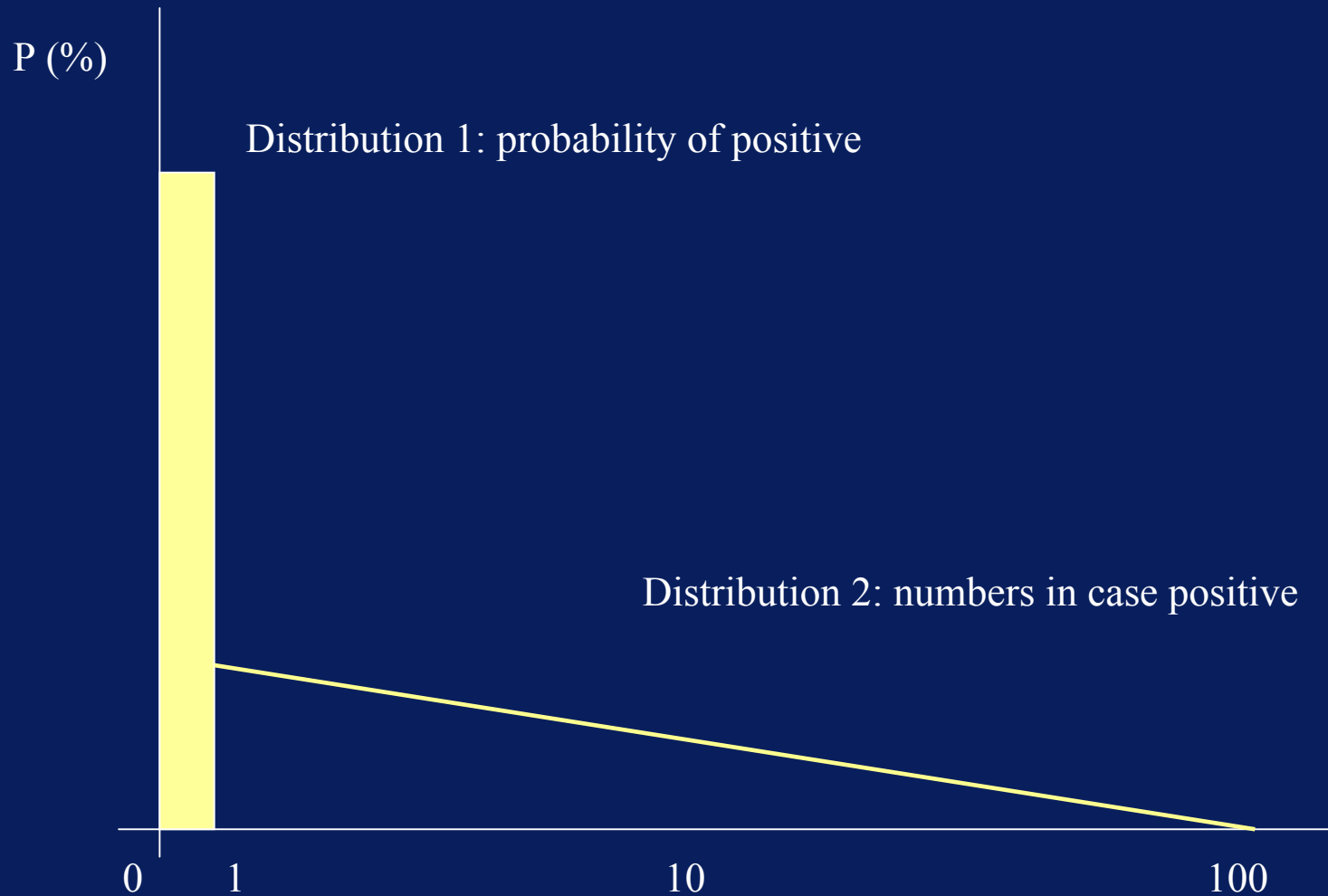
Level at time of consumption

- % positive

- numbers

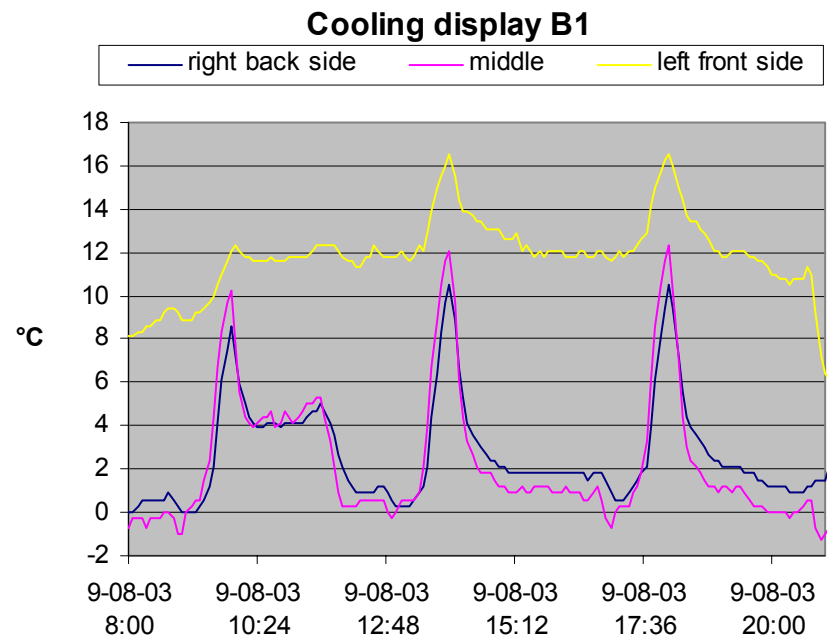
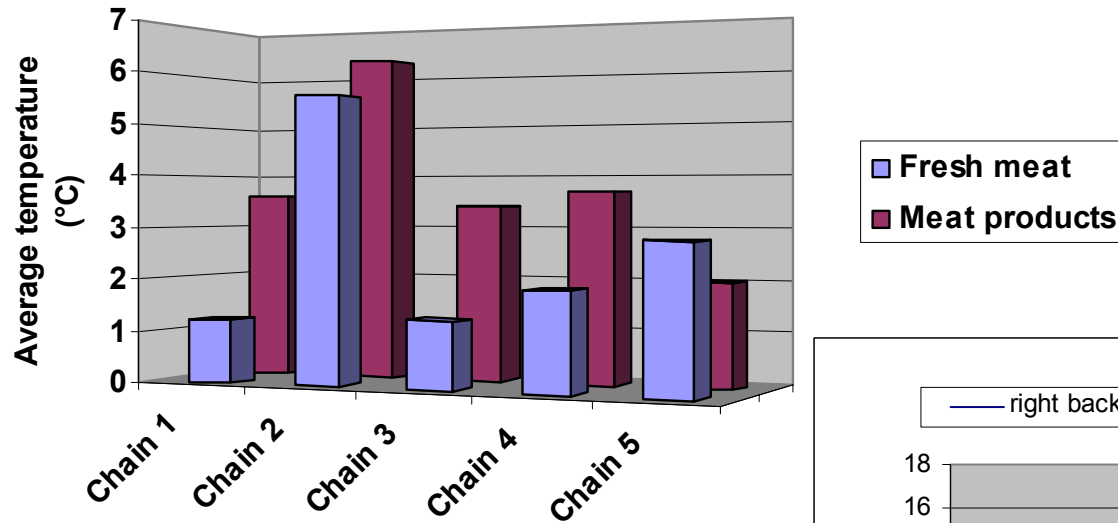


Occurrence of pathogens on raw meat

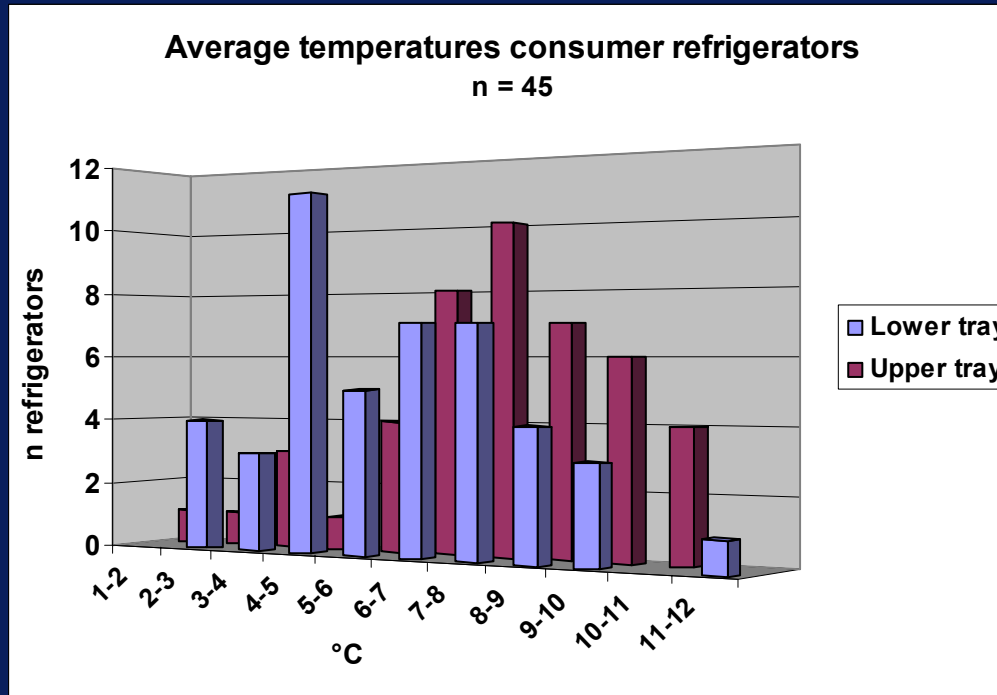


Temperature distribution in retail

Average temperatures in displays for fresh meat or meat products in different supermarket chains

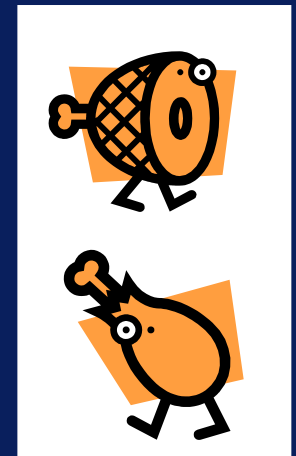


Temperature distribution in households

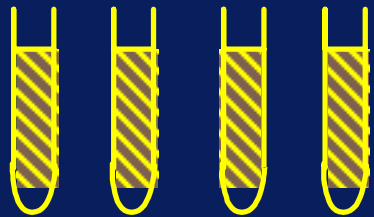


Refrigerators with average temperatures above 7°C

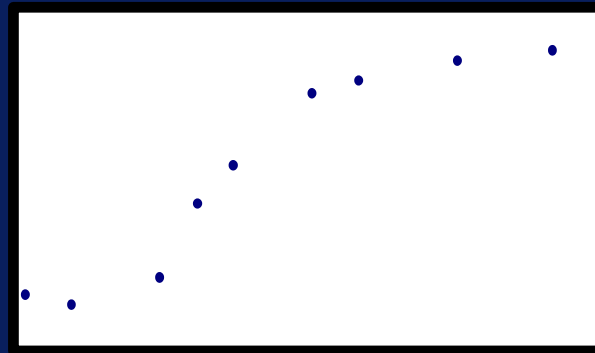
- Upper tray 60%
- Lower tray 33%
- Overall 42%
- Upper and lower tray 33%



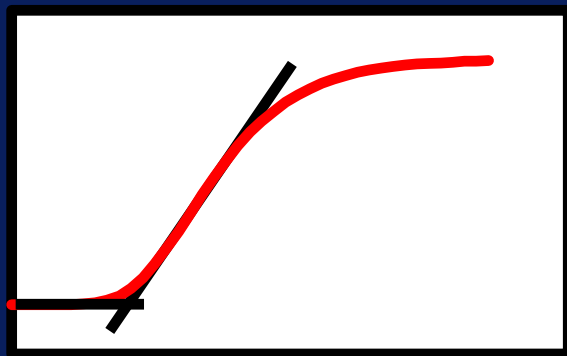
Modelling to predict safety and shelf-life



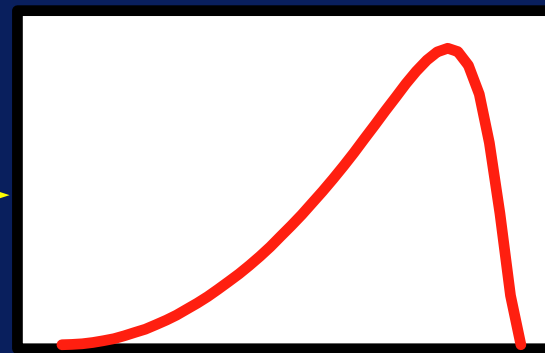
Experimental Design



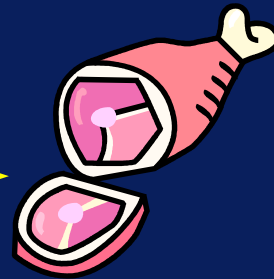
Generation of data



Curve fitting



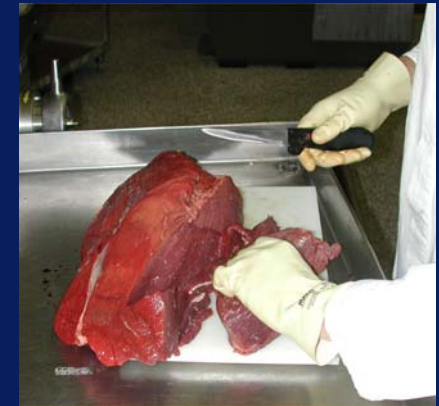
Modelling



From TTI to SMAS

TTI

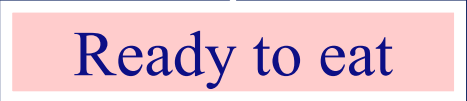
- Responds independent of initial level
- Prediction: SSO increase 0 log, 3 log, 5 log
- Prediction: pathogens increase not, factor 2, factor 10



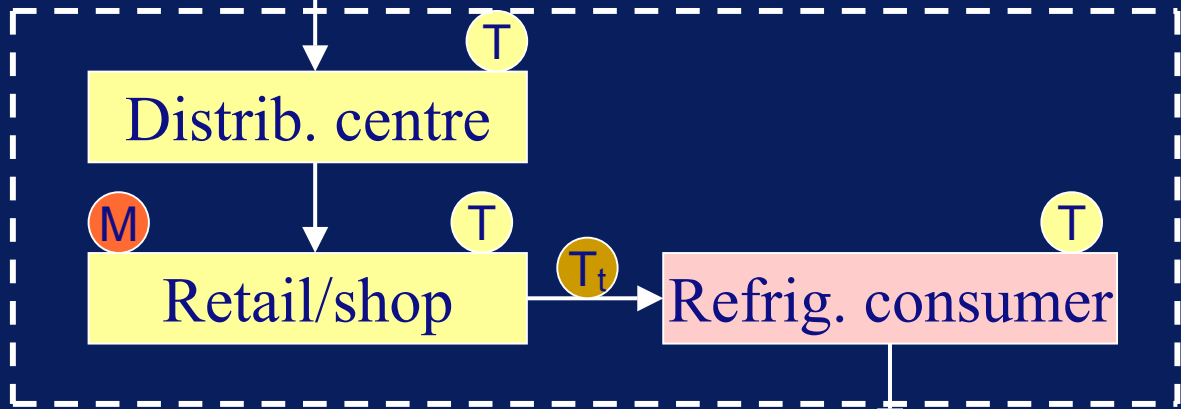
SMAS

- Remaining shelf-life is ...
- (Risk of spoilage is ... % (depending on initial level))
- Risk of illness is increased by factor ...
- (Risk of illness is ... % (depending on initial level and cooking, etc.))





Temperature/time controlled by TTI-system



Discussion about risk assessment

- **Define scope and objectives before risk assessment**
- **Use experts**
 - product/process, microbiology, statistics
- **Limit the models to relevant factors**
 - raw materials, process, storage, (consumer)
- **Verification of results with available data**
 - microbiology, epidemiology

