

UNSW

Monte Carlo Pathogen Risk Assessment Modeling in the Water Industry Using @ Risk

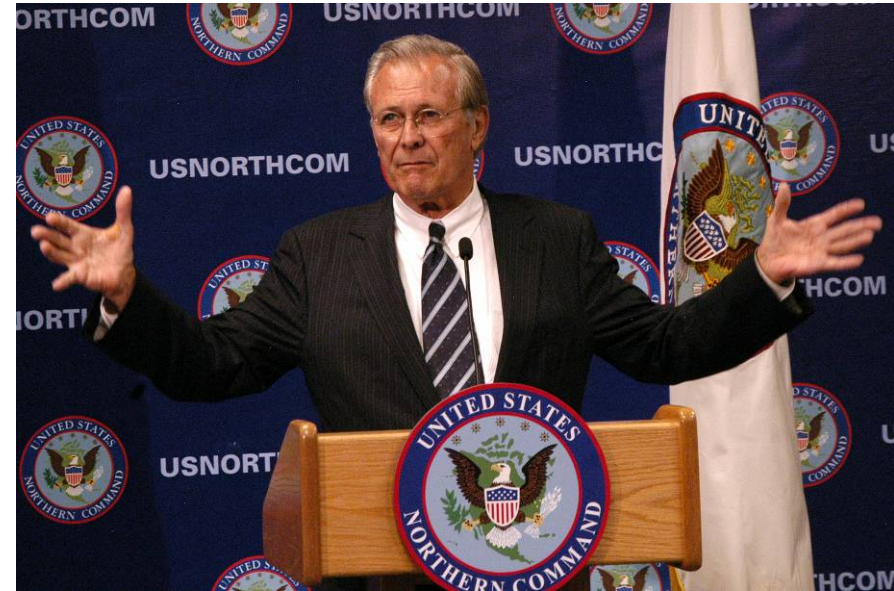
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A Philosophy for @Risk Modellers?

Rumsfeld's Postulates

- Known knowns
 - Contaminants we know we know
(*Cryptosporidium*)
- Known unknowns
 - Contaminants we know we don't know
(*Helicobacter*)
- Unknown unknowns
 - Contaminants we don't know we don't know
(Prions??)
- Unknown knowns?
 - Contaminants we don't know we know
(emergent pathogens)



Quantitative Microbial Risk Assessment (QMRA)

- Estimation of risks arising from exposure to pathogens
- Calculated for multiple exposure pathways
 - *based on initial concentration, barrier reductions, consumption rate, & dose response for population/pathogens*
- Calculations use various PDFs to describe each stage in exposure path)
- Linked in series to get final risk estimate:
 - *Disability Adjusted Life Years (1 μ DALY)*
 - *Infection probability (drinking 'benchmark' $10^{-4} \cdot \text{person}^{-1} \cdot \text{y}^{-1}$)*
 - *Concentration of pathogen*

Paper Aims

For theoretical Basis for QRA/QMRA see Haas, Rose and Gerba 1999

Presentation focuses on applying QRA operationally Using @Risk as a TOOL within a larger scheme



1. Outline Infectious Disease and Water Issue
2. Why Quantitative modelling of water supply with @Risk



3. Application of Quantitative Risk Technology



4. Strategies for Addressing Use Challenges - making @Risk a Useful Day to Day Tool



5. @Risk Programming Nuts and Bolts



1. Infectious Diseases and Water

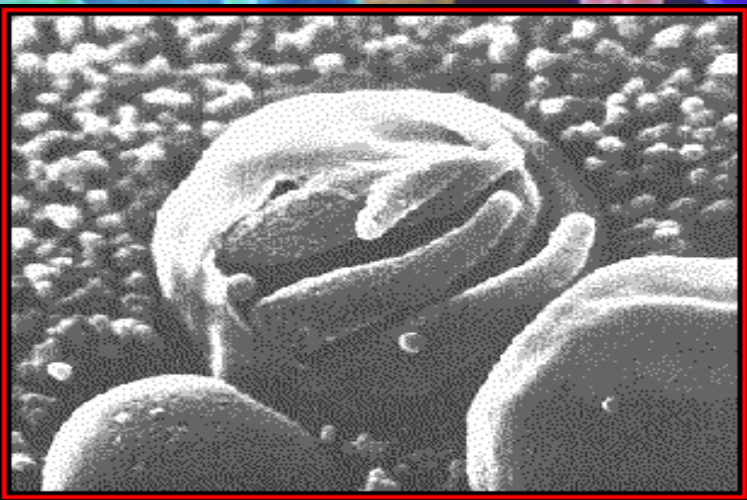
- The Culprits
- Past and Present Situation



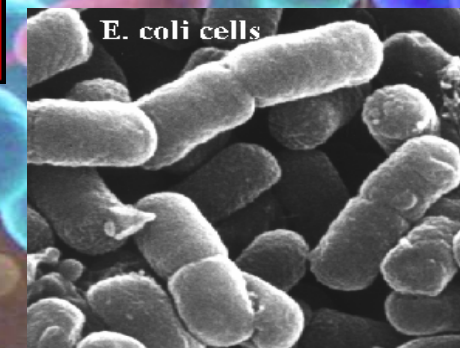
MONSTER SOUP commonly called THAMES WATER. (being a correct representation of that precious stuff doled out to us)

William Heath, 1828

Microbial Pathogens



Cryptosporidium oocysts – 3 μm



E. coli cells

Escherichia coli O157H7 – 1 μm cells



Viruses – bar = 0.1 μm

Robert Koch about his observations in India (1884):

cholera was endemic in certain regions in Bengal, and localized around ponds and swamps with multiple uses by the inhabitants

Today still 1.1 billion with not clean drinking water

source of drinking water



cleaning of household utensils

bathing

Today still 2.4 billion with no sanitation

washing of clothes, including those of persons who had died of cholera

input of wastewater, human excreta

Outbreak of EHEC in Walkerton, Canada

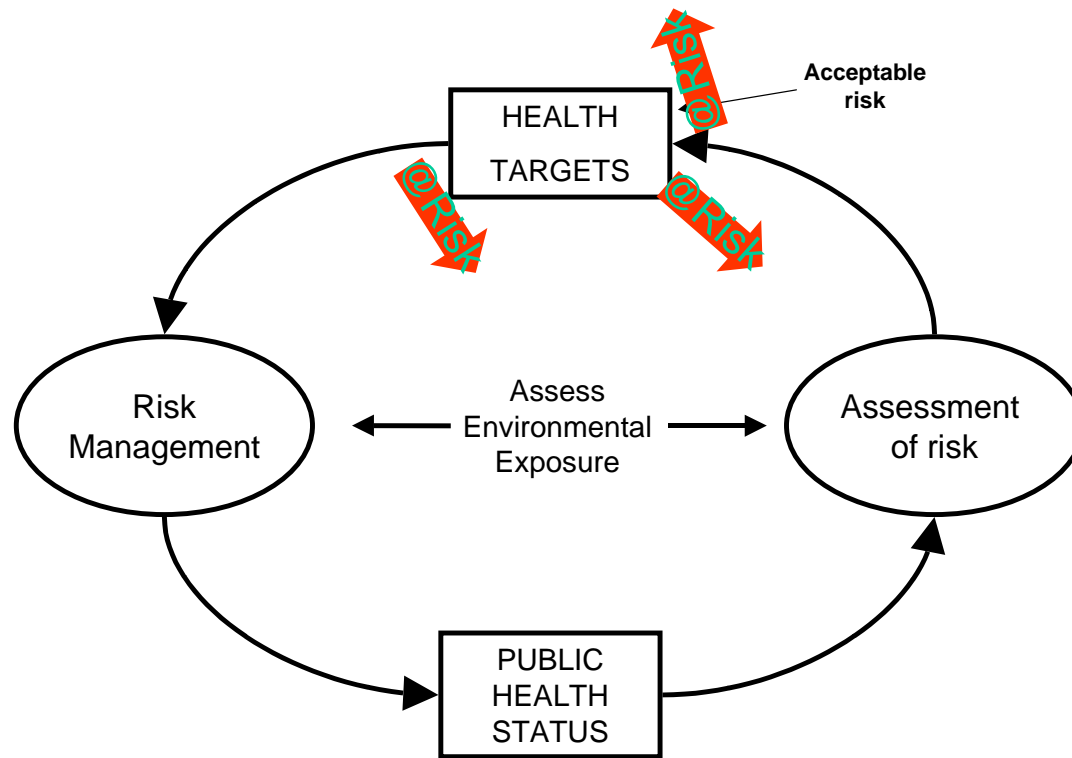
- Walkerton, Ontario (Canada) with 4,800 residents; in May 2000: more than 2,300 persons became ill, seven people died
- contamination of drinking water system with primarily *E. coli* O157:H7, but also *Campylobacter jejuni*
- primary source: manure that had been spread on a farm near a shallow well in an area of highly fractured bedrock
- outbreak 14 days after heavy rainfalls, which promoted contamination of wells
- failure of continuous chlorine and turbidity monitoring of water
- delay in reporting positive *E. coli* results in water samples



2. Why Quantitative Modelling Using @Risk

1. Guideline promotion of Risk Assessment based Management
2. Risk Principles and the Drive for Quantification
3. The Complexity problem

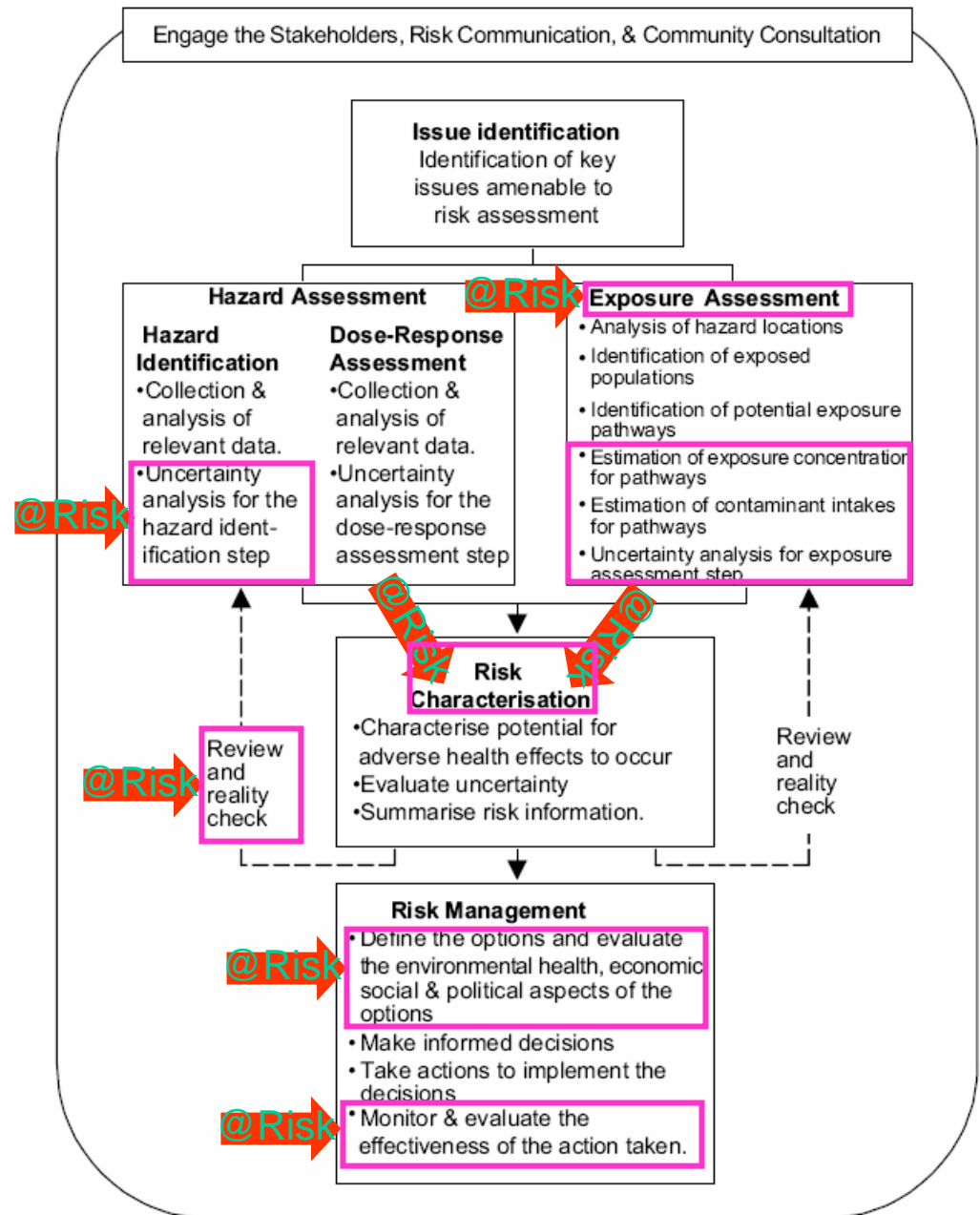
WHO Scheme for Water Quality Assessment and Management



From: Fewtrell & Bartram, 2001. in Water Quality Guidelines, Standards and Health (WHO). IWA publishing.

Health Risk Assessment (EnHealth 2002)

- Use scheme to systematise work
- Data mining + gap filling (CWWT work)
- Requires:
 - Hazard assessment (research lit review)
 - Dose response (literature)
 - Exposure Pathway Assessment (Initial provides basis for research program)



Example > Australian Drinking Water Guidelines

<http://www.nhmrc.gov.au/publications/synopses/eh19syn.htm>

The Six Fundamental Principles (strategic)
Twelve elements (operations)

Framework Includes:

-  • HACCP - Hazard Analysis Critical Control Point
-  • AS/NZS 4360 - Risk Management

Principles 1 & 2



- *The greatest risks to consumers of drinking water are pathogenic microorganisms.*

Protection of water sources and treatment are of paramount importance and must never be compromised.

- *The drinking water system must have, and continuously maintain, **robust multiple***



 ***barriers** appropriate to the level of potential contamination facing the raw water supply.*



Principles 3, 4 & 6

- *Any sudden or extreme change in water quality, flow or environmental conditions (eg extreme rainfall or flooding) should arouse suspicion that drinking water might become contaminated.*
- *System operators must be able to respond quickly and effectively to adverse monitoring signals*
- *Ensuring drinking water safety and quality requires the application of a considered risk management approach*

The 12 Elements of The Framework

1. Commitment to Drinking Water Quality Management
-  2. Assessment of the Drinking Water Supply System
-  3. Preventive Measures for DWQ Management
-  4. Operational Procedures and Process Control
-  5. Verification of Drinking Water Quality
-  6. Management of Incidents and Emergencies
7. Employee Awareness and Training
8. Community Involvement and Awareness
-  9. Research and Development
10. Documentation and Reporting
-  11. Evaluation and Audit
-  12. Review and Continual Improvement

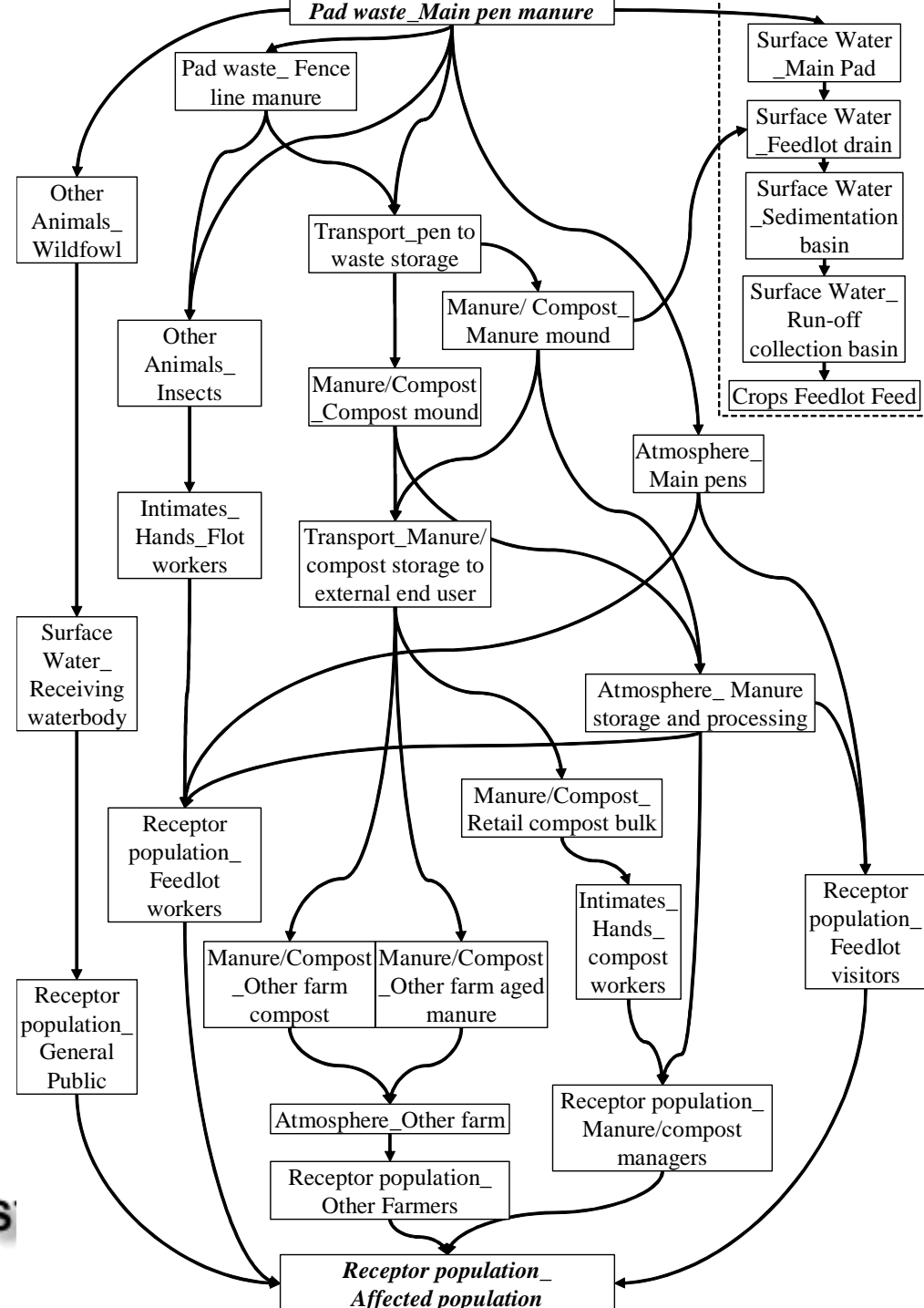
Definition of Risk

Risk = likelihood x
consequence

Qualitative risk analysis matrix: level of risk for each hazardous event

Likelihood	Consequences				
	1 Insignificant	2 Minor	3 Moderate	4 Major	5 Catastrophic
A (almost certain)	Moderate	High	Very high	Very high	Very high
B (likely)	Moderate	High	High	Very high	Very high
C (moderate)	Low	Moderate	High	Very high	Very high
D (unlikely)	Low	Low	Moderate	High	Very high
E (rare)	Low	Low	Moderate	High	High

Exposure Pathway Complexity (feedlot example)



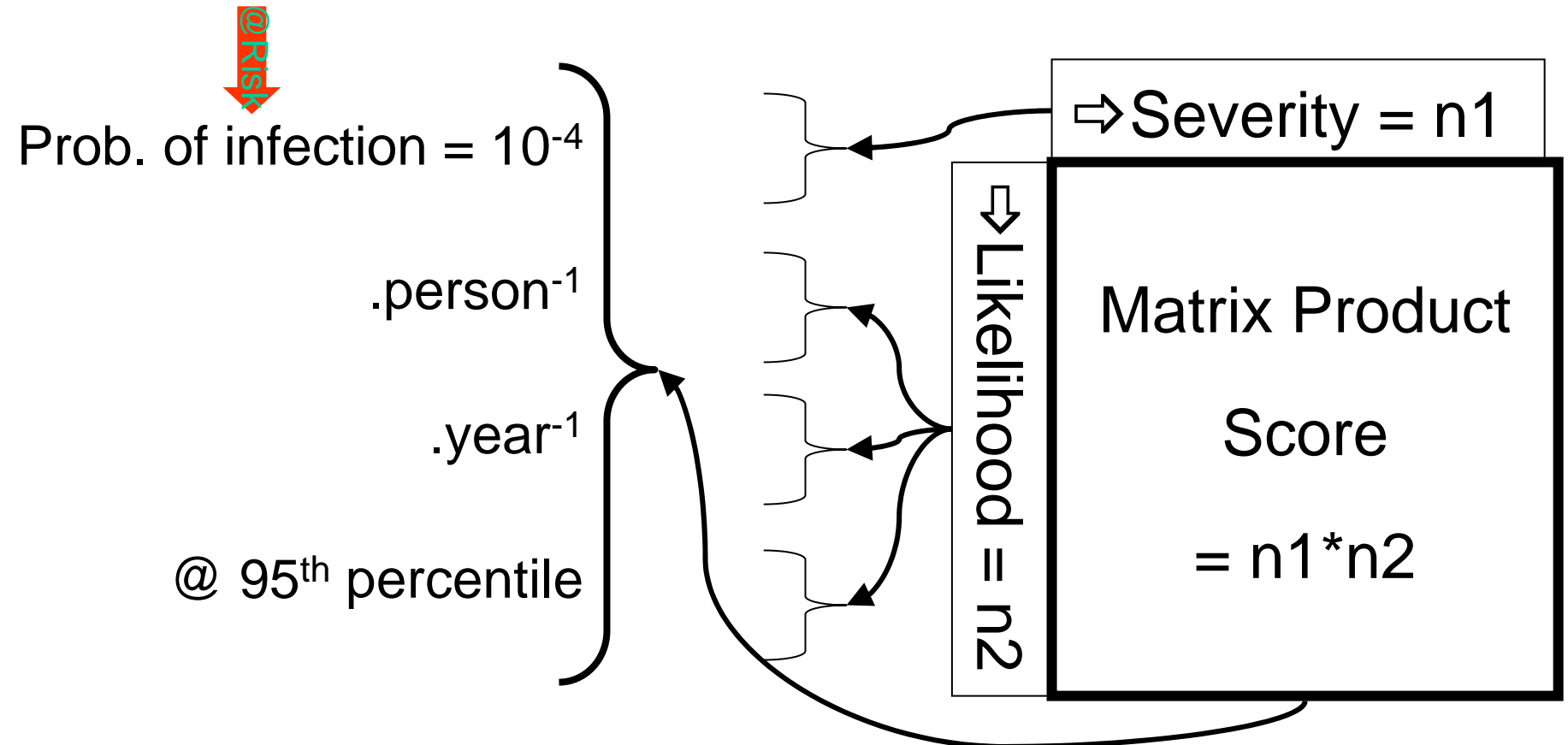
QMRA

(Decision Analysis)

Qualitative Risk

Assessment

(Decision Intuition)



Other drivers for going 'Quantitative'

- Can't maintain zero risk fiction— need more objectivity & precision.
- More knowledge (pathogens, impacts, environment) and need to better utilise expensive data/study outcomes
- Need to define/quantify 'safe' in auditable fashion
- Ecological Sustainability Philosophy
 - Base natural resource use limits and 'sustainability' of exploitation on mathematics + science (e.g. Malthus, Ehrlich, Diamond)

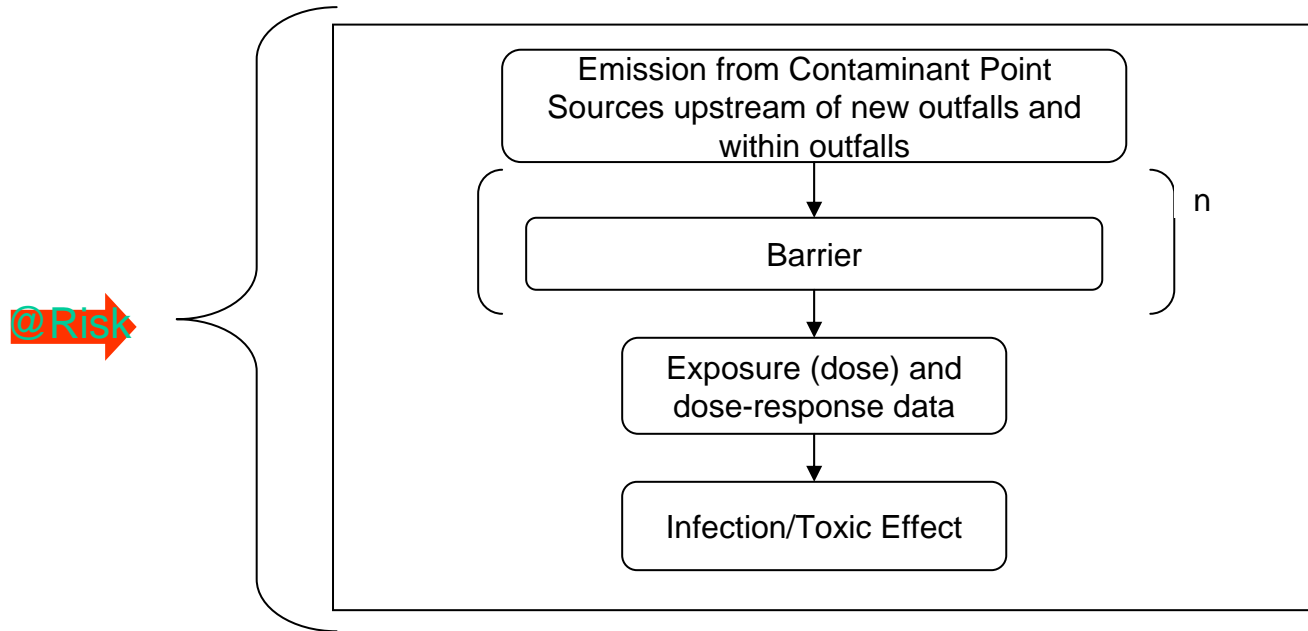
3. Application of Quantitative Risk Technology in the Water Industry

- Examples of Use
- The Barrier Concept
- Modelling challenges posed when attempted

Examples of Use

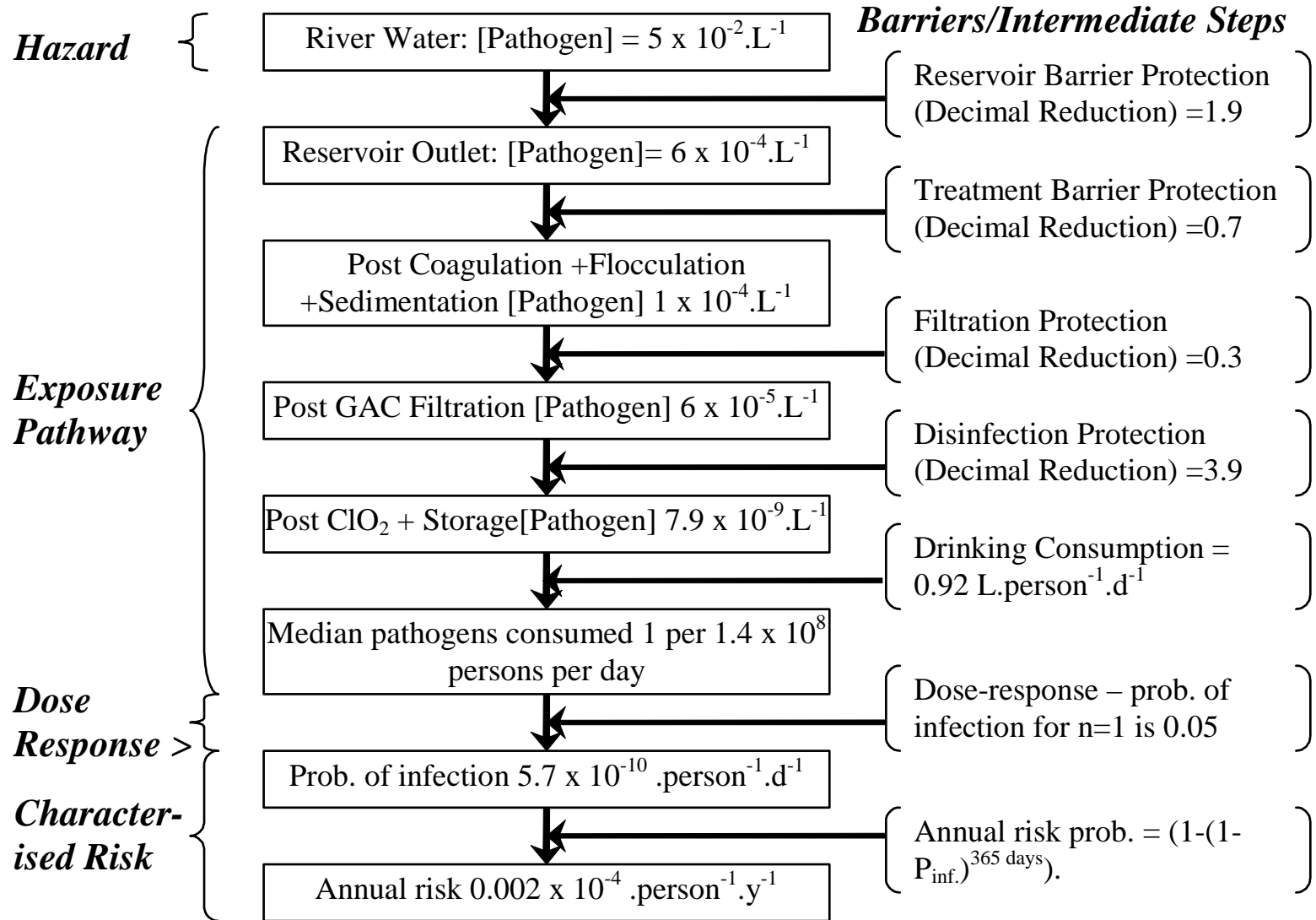
- Water Supplies supplies - EU MicroRisk (www.microrisk.com) - 14– *So many – need to 'Industrialise' QMRA?*
- Water Reuse - Flow Replacement = *Flow Augmentation*
- Recreational risk exposure - Lake Parramatta = *Bathing in stormwater/run-off*

The Barrier/Exposure Pathway Concept





Integrated (“Holistic”) Exposure Pathway





Quantitative Risk Modelling Challenges

- Logistics (next slide)
- Specialisation
 - Institutional environmental assessors cant be experts in all software
- ‘Risk Assessment not Industrialized (MicroRisk)
 - Documentation of assumptions
 - Coping with multiple scenarios
 - Large numbers of similar but unique projects
- The allergy of senior managers to numbers
 - Detail distracting and often not informative
 - Opacity of high level models





Logistics

~Infinite Permutations
and Combinations

- Pathogens
- Exposure Scenarios
- Hazardous Events (multiple?)
- Barrier functions
- Environment (season, load)
- Populations
- Hydrology
- Multiple Exposure Pathways



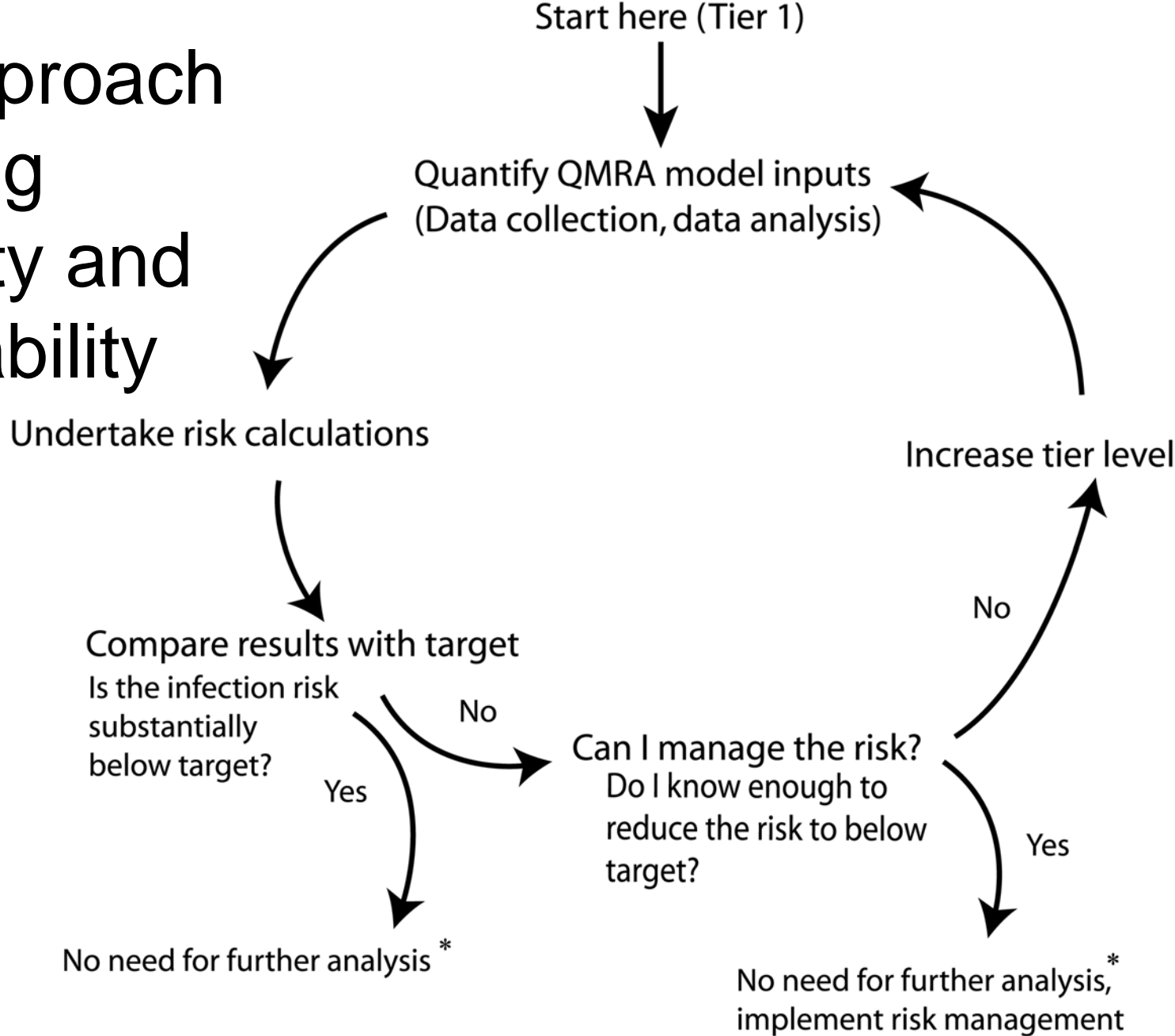
4. Strategies for making @Risk a Useful Day to Day Tool

- Programming strategy
- Tiered Approach
- Linking Risk Characterisation to Risk Management Information Needs (HRA scheme)

Programming strategies

- **Use of Metamodel Approach** (central calculation engine + library of @Risk submodels – accessed like lookup tables)
- **Construction of Informative** (easily revised) **Scenarios**
- **Separation of Baseline and ‘Hazardous Events’** (deal with one bit at a time)
- **Scenarios based on Exposure Pathway/Barrier Assessment** > drives input parameter data mining

Tiered approach in reducing uncertainty and data variability



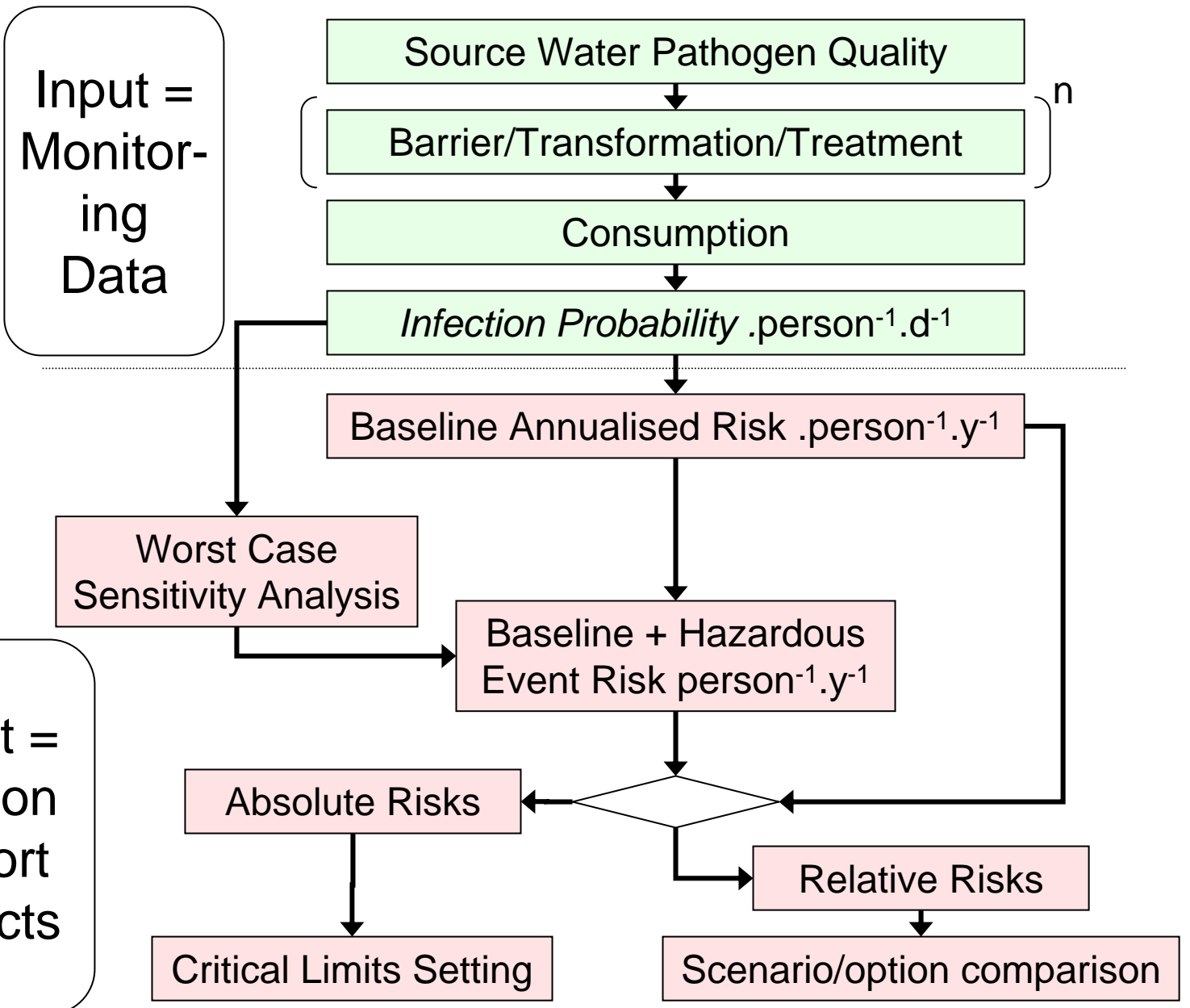
Managing the Risk Assessment

- Use style
 - *For Decision Support*
 - *Address questions/hypotheses/what if?*
 - *Critical limit development (from HACCP)*
- Use of reference/baseline scenarios
 - *High ('event') Risk/Concurrent Failure Modelling*
- High Priority Foci
 - *large population, worst environment, major exposure points, priority pathways*
- 'Industrialise' Risk Assessment
 - *hydrology example - what is possible / what not to do*
- Develop Operational Assessment System

5. @Risk Programming – the Nuts and Bolts

- Inputs and Outputs Scheme
- Baseline > Sensitivity > Events > Critical Limits
- Programming with Excel and @ Risk
- Program illustration
- Cautionary notes

Linking Data and Decisions



Output =
Decision
Support
Products

Campylobacter Risk Sweden 2

Critical Stage	Stage Transformation Units and Modal Value	Output after Transformation	
Source Water	-	Pathogen. L ⁻¹	4.7 x 10 ⁻²
Reservoir	Decimal Reduction = 1.87		6.3 x 10 ⁻⁴
Coag./Floc./Sediment	Decimal Reduction = 0.73		1.2 x 10 ⁻⁴
GAC Filtration	Decimal Reduction = 0.28		6.3 x 10 ⁻⁵
ClO ₂ + Storage	Decimal Reduction = 3.9		7.9 x 10 ⁻⁹
Consumption	L.d ⁻¹ consumed = 0.92		7.4 x 10 ⁻⁹
Dose Response	(infectious dose <i>ca</i> 10 organisms)		Infection prob.
Annualized Rate (@Risk simulated values)	Annualization (1-(1-P _{inf.}) ^{365 days})	2 x 10 ⁻⁷ .person ⁻¹ .y ⁻¹	

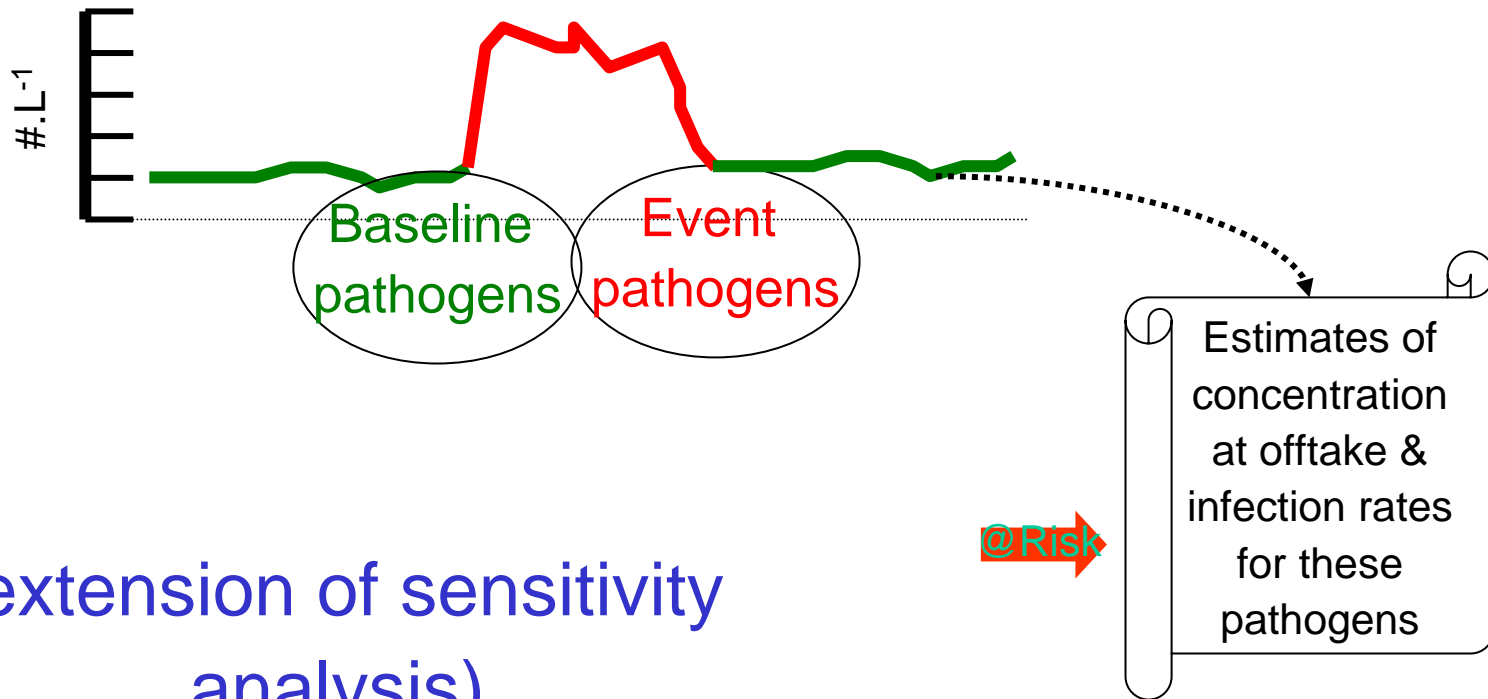
Baseline Infection Rates

Measure	Water Supply system	Pathogen					
		<i>Crypto-sporidium</i>	<i>Giardia</i>	<i>Campylo-bacter</i>	<i>E. coli</i> O157	<i>Norovirus</i>	Enterovirus
95 th %ile Annualized Probability (10 ⁻⁴ .person ⁻¹ .y ⁻¹)	Australia 1	0.68	6.7E-06	0.29		-	-
	UK 1	0.16	1.7E-06	1.21E-07	1.9E-08	-	1.51E-06
	Sweden 1	<u>2200</u>	0.44	0.84	-	(19)	(3.0)
	Sweden 2	<u>334</u>	1.40	0.012	-	(0.66)	(0.082)

Notes: 1. Infection rates in brackets are based on 95th percentile source water concentration data
 2. Bold + underlined = of high concern, Bold + Italics marginal concern

Assigning the origin of Pathogens to Baseline v. Event Conditions

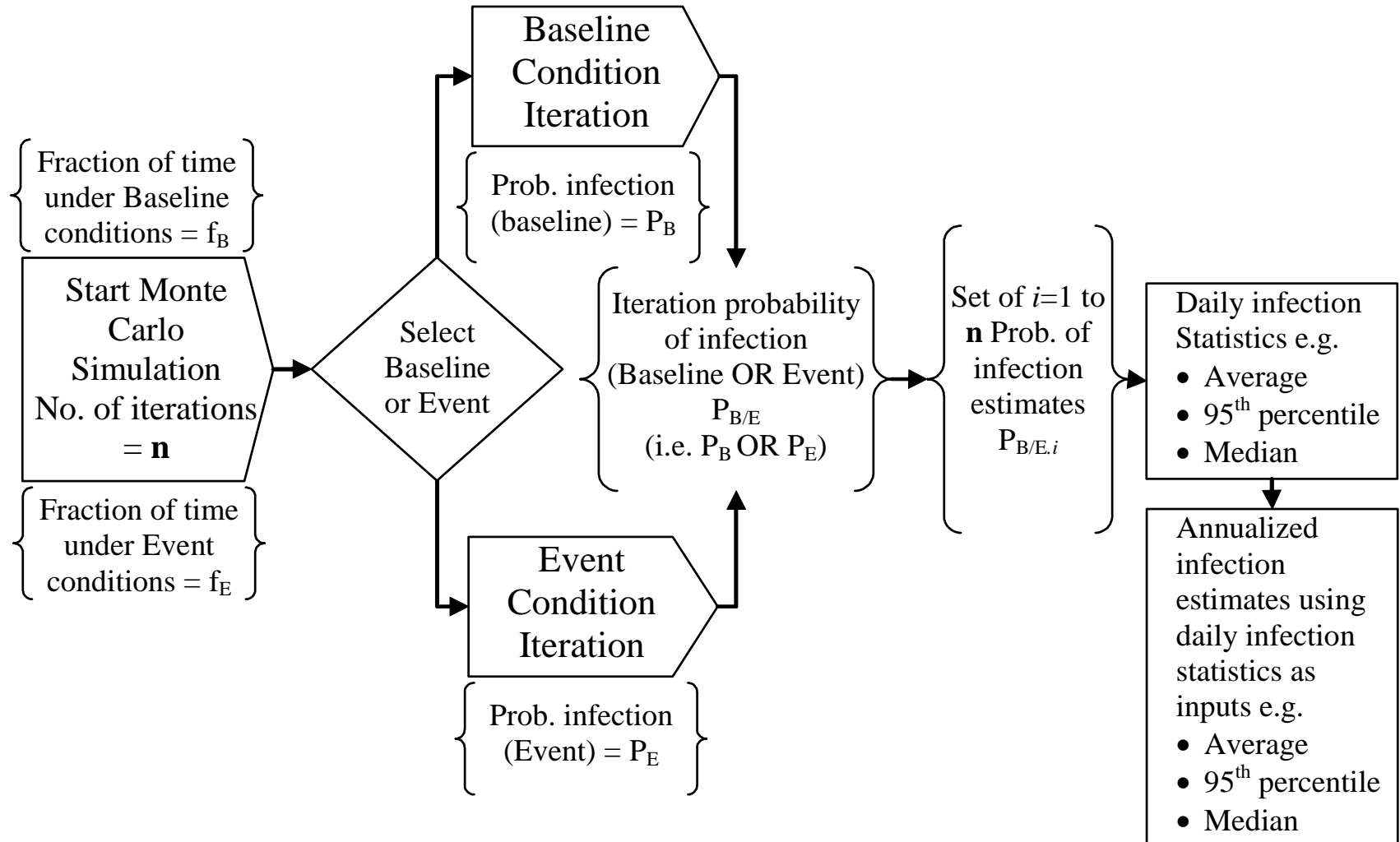
Small
River



(extension of sensitivity
analysis)



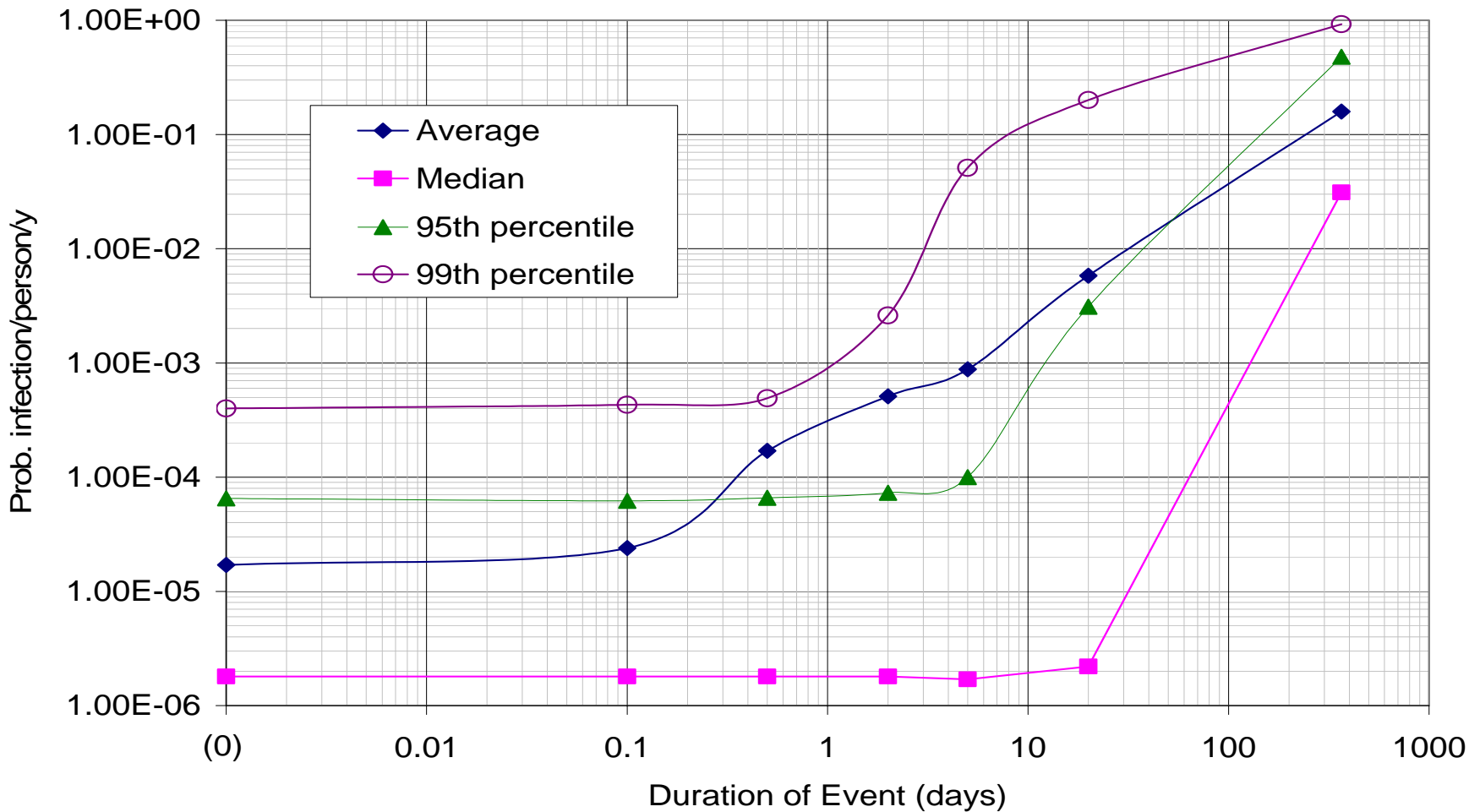
Concurrent Baseline and Event Simulation with @Risk



Average Annualised Risk person⁻¹.y⁻¹ for swimming (illustrates hazardous events issue)

Scenario Summary Description	High Flow Penrith Recreation (26 days)	Low Flow Penrith Recreation (26 days)	Low Flow Penrith Recreation (26 days) + <u>MF/RO failure</u>
<i>Cryptosporidium</i>	2.94E-13	6.31E-12	-
<i>Campylobacter</i>	2.41E-09	5.29E-08	7.49E-02
Rotavirus	2.46E-08	5.30E-07	9.05E-02

Infection risk in response to simulated chlorination failure (illustrates critical limit issue)



The Metamodel Library

@Risk Probability Density function

Model details

Model coefficients

PDF Code	PDF	CTC Simulations	PDF Summary	Locality	Study	Glasses	Expression form	Expression	Reference	Units	Spare	Coefficient 1	Coefficient 1 value	Coefficient 2	Coefficient 2 value	
43	CNSMP_40	0.88056	Sweden Transtr	Sweden	Transtrand 60-69 y	Risk function	Lognormalalt	Westrell et al. Tal	Litres/day	10th perc	0.4	Median	0.8	90th pe	1.46	
44	CNSMP_41	0.89609	Sweden Transtr	Sweden	Transtrand 70+ y	Risk function	Lognormalalt	Westrell et al. Tal	Litres/day	10th perc	0.48	Median	0.89	90th pe	1.32	
45	CNSMP_42	0.386	Canada cold tap	Canada	cold tap water	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	0.386	Std Dev	0			
46	CNSMP_43	0.76	Canada cold tap	Canada	cold tap water + bottled	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	0.76	Std Dev	0			
47	CNSMP_44	0.506	USA cold tap w	USA	cold tap water	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	0.506	Std Dev	0			
48	CNSMP_45	0.667	USA cold tap w	USA	cold tap water + bottled	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	0.667	Std Dev	0			
49	CNSMP_46	0.77	France winter c	France	winter cold tap water	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	0.77	Std Dev	0			
50	CNSMP_47	1.61	France winter c	France	winter cold tap water +bottled	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	1.61	Std Dev	0			
51	CNSMP_48	0.9	France Spring c	France	Spring cold tap water	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	0.9	Std Dev	0			
52	CNSMP_49	1.98	France Spring c	France	Spring cold tap water + bottled	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	1.98	Std Dev	0			
53	CNSMP_50	0.15	Netherlands col	Netherlands	cold tap water Teunis	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	0.153	Std Dev	0			
54	CNSMP_51	0.19	England +Wale	England +Wal	cold tap water	Risk function	Normal	Westrell et al. Tal	Litres/day	Average	0.19	Std Dev	0			
55	CNSMP_52	3	Sweden Westre	Sweden	Westrell thesis work+3 L sim	Risk function	Lognormal		Litres/day	Mu	3	Sigma	0.2			
56	CNSMP_53	1	Sweden Westre	Sweden	Westrell thesis work	Risk function	Lognormal	Westrell al 2003 t	Litres/day	Mu	1.1	Sigma	0.63			
57	CNSMP_54	6	Point Estimate	Point Estimate	Conceptual				Litres/day							
58	CNSMP_55	5	Point Estimate	Point Estimate	Consumption for Sensitivity at	High consum	Point value	Mons et al 2005	Litres/day	95th %	1.5					
59	CNSMP_56	0.03	SWHN	Recreation cons	Recreation co	Typical values	Recreation ci	Triangular	SWHN Table	Litres/day	Minimum	0.01	Mode	0.03	Maxim	0.05
60	CNSMP_57	0.1	SWHN	Maximum recre	Maximum recr	Maximum estimate	Recreation ci	High value	SWHN Table	Litres/day	Maximum	0.1				
61	CNSMP_58	0.005	SWHN	Typical serving	Typical serving	Draft Aust. Guidelines	Lettuce	Point value	National draft 200	Litres/day	Point	0.005				
62	CNSMP_59	0.011	SWHN	Maximum servir	Maximum serv	Draft Aust. Guidelines	Lettuce	Point value	National draft 200	Litres/day	Point	0.011				
63	CNSMP_60	0.06	SWHN	US range of Ovs	US range of O	Oyster impact study - model for mussels		Uniform	AWT 1997	Litres/day	Minimum	0.06	Maximur	0.24		
64	CNSMP_61	0.24		n	Oyster impact study - model for mussels			Point value	AWT 1997	Litres/day	Point	0.24				

Selected Model

EXCEL Scenario MetaModel

1. @Risk Add in for Monte Carlo Simulations

Pathogen Removal>Consumption>Infection Tree Simulator											
Date modelled 3/Aug/2006 9:57											
Simulator version 6_3_dist_acute.xlsScenario_baseline											
Barrier No.	1	2	3	4	5	6	7	8	9	10	
Stage process	Input	Transformation	Transformation	Transformation	Transformation	Transformation	Transformation	Transformation	Consumption	Dose Response	
Process Library Table (wksheet)	R1_INPUT	R2_DECIM	R2_DECIM	R2_DECIM	R2_DECIM	R2_DECIM	R2_DECIM	R2_DECIM	R3_CNSMP	R4_DSRSP	
Process Code No.	58	79	70	74	1	1	1	1	28	5	
Process Class	Initial input concentration	Change in Concentration based on DEC PDF	Change in Concentration based on DEC PDF	Change in Concentration based on DEC PDF					Consumption arising from intake rate	Infections in response to intake	
Process Summary Description	Myponga River Year Average_Cryptosporidium modelled using Cryptosporidium data	Cryptosporidium removal by Myponga Reservoir Reduction Factor modelled using Cryptosporidium data	Giardia and Cryptosporidium removal by Myponga DAF Optimal modelled using Yeast data	Giardia and Cryptosporidium removal by Myponga RSF Optimal modelled using Yeast data					Adelaide Discrete general 0.25 L glass Poisson	Teunis pooled data_Cryptosporidium modelled using Cryptosporidium data	
Process PDF Modal Value	1.07	1.47	1.72	3.57					0.75	2.7368E-08	
Primary Literature Reference/Data source	Signor data	Signor et al. in prep/PhD thesis	Signor et al. in prep/PhD thesis	Signor et al. in prep/PhD thesis					Mons et al 2005	Teunis et al 2002	# Simulations 1
Tier Rating - Input and Transformation PDFs	3	2	2	2.5					3	2.5	sd=0.44, min. tier
No. organisms @ end of stage(statistics)	No. .L ⁻¹	No. .L ⁻¹	No. .L ⁻¹	No. .L ⁻¹					No. ingested	prob. inf. person ⁻¹ .d ⁻¹	Days 365
Modal @Risk default value	1.073740	3.63E-02	6.91E-04	1.86E-07					1.39E-07	2.74E-08	9.99E-06
Simulation Average	1.09111	3.49E-02	1.62E-03	4.76E-07					3.68E-07	7.21E-08	2.63E-05
0.95 output percentile	2.837576463	1.31E-01	7.07E-03	2.70E-06					1.99E-06	3.90E-07	1.42E-05
0.5 output percentile	0.655324696	1.88E-02	2.52E-04	8.97E-08					5.09E-08	9.99E-09	3.65E-05
0.05 function percentile	0.11	1.15	0.71	2.83					0	0	
0.95 function percentile	2.84	2.00	2.71	4.30					1.5	3.90201E-07	

2. Create Scenario by Selecting Input Functions from PDF Model library

4. Estimate concentration after each barrier

3. Run Simulations

5. Output risk probability statistics and pathogen numbers

Extracting Library Submodels

Selecting the submodel

Model name & Reference Cell

Model information

Extracted Model Monte Carloed Value

Barrier No.	1	2
Stage process	Input	Transformation
Process Library Table (wksheet)	R1_INPUT	R2_DECIM
Process Code No.	106	323
Process Class	Initial input concentration	Change in Concentration based on DEC PDF
Process Code	INPUT_106	DECIM_323
Process Summary Description	Raw sewage content based on literature review_Cryptosporidium modelled using Cryptosporidium data	All pathogens removal by Hunts Ck dilution sewage suboptimal modelled using Indicator dilution data
Process PDF Modal Value	3162.27766	3.18708664
Primary Literature Reference/Data source	Medema et al 2001	LP Stage 3 Report

B4 >
B5 >

B7 fx =OFFSET(INDIRECT(RIGHT(B4,5)&"_"&B5),0,-1)

B8 fx =OFFSET(INDIRECT(B7),0,2)

B9 fx =INDIRECT(B7)



Other Input/Outputs

9	10	Point Simulations		
Consumption	Dose Response	Point 'Input' Value	Point Decimal Reduction	Point Consumption (L)
R3_CNSMP	R4_DSRSP	Point 'Input' Value	Point Decimal Reduction	Point Consumption (L)
54	36	2488	4.754466	6
Consumption arising from intake rate	Infections in response to intake	6. Point value Simulations: The 3 white boxes allow inputting of point values for 'input', 'transformation' and consumption PDFs. Values entered are each mirrored in a reference table PDF and from there can be selected for a		
CNSMP_54	DSRSP_36	ed either for simple point value lation.		
Point Estimate Consumption for Sensitivity and Goal Seek Conceptual L	Teunis pooled data_Cryptosporidium modelled using Cryptoc	Or using the @Risk 'Goal Seek' tool they can be altered to find that input value which yields a g. a 95th pc of persons per me is shown in usual way.		
6	0.005185741	7. Tier of Assessment indicates input data robustness		
0	Teunis et al 2002	# Simulations	# iterations	
		1	1000	
3	2.5	mean=2.5, sd=0.44, min. tier = 2 (n=6)	<<<<< Tier Score	

Simulation Title		
Lake Parramatt Risks		
System name	Pathogen name	Short CTC name (4 characters) to highlight PDFs in reference tables
CTS-5	Cryptosporidium	LP_risk
8. Simulation Details		
5. To compare one simulation with another copy the values in L15:L22 and PASTE VALUES them into M15:M22. Then run a new simulation. The log10 FS values will appear in column N		



Baseline + Hazardous Event Simulation Outputs

12. @Risk Function Labels

9. Event and Baseline Risk Probabilities

10. Aggregate Risk

11. Event details

		Event scenario infection rate - Pinf 1 d	Baseline (values automatically copied)		Total Baseline Period (days)		365	
					Total Event Period (days)		50	
					Proportion time Baseline State		0.863014	
					Proportion Time as Event [1 in 6]		0.136986	
			Reference Inf. person ₁ .d ⁻¹	Baseline scenario infection rate - Pinf 1 d	Total P inf. person ⁻¹ .d ⁻¹	P. of inf. person-1 .365 days-1	Event Increase Factor	
Modal @Risk default value			3.95E-03	2.66E-05	2.66E-05	2.66E-05	9.66E-03	1.00000
Simulation Average			4.84E-03	3.31E-05	3.31E-05	4.22E-04	1.43E-01	11.25273
0.95 output percentile			1.20E-02	8.39E-05	8.39E-05	4.29E-03	7.92E-01	145.13301
0.5 output percentile			3.62E-03	2.43E-05	2.43E-05	2.51E-05	9.13E-03	1.00000
0.99 output percentile			1.44E-02	1.02E-04	1.02E-04	8.85E-03	9.61E-01	148.40449
0.975 output percentile			1.30E-02	9.12E-05	9.12E-05	5.36E-03	8.59E-01	148.38993
No. organisms @ end of stage (statistics)		1						
3. Values from B20 to K27 are input and output statistics for each stage simulation (rows 9 and 20). The		1						

Reporting

BASELINE BASICS

Software

Program QPES v 1.0 using @Risk 4.5 Professional Add ins
 Main Scenario version C:\Documents and Settings\djroser\My Documents\QMRA\Swedish students\[\Event Tree Risk
 worksheet Estimation Simulator version 6_5 - at risk conference.xls\Scenario_baseLine
 Date modelled 6-Sep-2007 16:25

Model

Model Title Lake Parramatta Risks
 System modelled CTS-5
 Pathogen modelled Cryptosporidium
 Short code for hi-light LP_risk
 No. of iterations 1000
 No. of Simulations 1

BASELINE SCENARIO

Inputs (Summary)

Barrier codes

Barrier No.	Barrier Type	Worksheet Library	Barrier name	Tier	Barrier Class
1	Input	R1_INPUT	INPUT_106	3	Initial input concentration
2	Transformation	R2_DECIM	DECIM_323	2	Change in Concentration based on DEC PDF
3	Transformation	R2_DECIM	DECIM_331	2	Change in Concentration based on DEC PDF
4	Transformation	R2_DECIM	DECIM_328	2.5	Change in Concentration based on DEC PDF
5	Transformation	R2_DECIM	DECIM_1	0	Change in Concentration based on DEC PDF
6	Transformation	R2_DECIM	DECIM_1	0	Change in Concentration based on DEC PDF
7	Transformation	R2_DECIM	DECIM_1	0	Change in Concentration based on DEC PDF
8	Transformation	R2_DECIM	DECIM_1	0	Change in Concentration based on DEC PDF
9	Consumption	R3_CNSMP	CNSMP_56	3	Consumption arising from intake rate
10	Dose Response	R4_DSRSP	DSRSP_36	2.5	Infections in response to intake

Overall Simulation Tier Level

mean=2.5, sd=0.44, min. tier = 2 (n=6)

Transformation Key information

Barrier PDF

Barrier No.	Mode	Barrier Summary Description
1	3162.27766	Raw sewage content based on literature review_Cryptosporidium modelled using Cryptosporidium data
2	3.187086643	All pathogens removal by Hunts Ck dilution sewage suboptimal modelled using Indicator dilution data
3	2.180456064	All pathogens removal by Lake Parramatta stormwater dilution modelled using Management Trigger point data
4	1	Virus and protozoa removal by LP - 3 day other pathogen inactivation modelled using Bacterial indicators data
5	0	Null transformation - Used where < 7 transformations to fill blanks or to omit a process step - sensitivity analysis
6	0	Null transformation - Used where < 7 transformations to fill blanks or to omit a process step - sensitivity analysis
7	0	Null transformation - Used where < 7 transformations to fill blanks or to omit a process step - sensitivity analysis
8	0	Null transformation - Used where < 7 transformations to fill blanks or to omit a process step - sensitivity analysis
9	0.03	Recreation consumption range Typical values L
10	2.65894E-05	Teunis pooled data_Cryptosporidium modelled using Cryptosporidium data

Calculation Outputs

No. organisms @ end of stage (statistics)

Barrier No.	Units	Outputs					
		Modal @Risk default value	Simulation Average	95th %ile	50th %ile	99th %ile	97.5th %ile
1	No. .L-1	3162.27766	3908.012621	8843.546921	3098.77502	9625.47081	9334.4172
2	No. .L-1	2.055480479	2.540208204	5.748305499	2.01420376	6.25655603	6.0673712
3	No. .L-1	0.013566171	0.016765374	0.037938816	0.01329374	0.04129327	0.0400446
4	No. .L-1	0.001356617	0.001676537	0.003793882	0.00132937	0.00412933	0.0040045
5	No. .L-1	0.001356617	0.001676537	0.003793882	0.00132937	0.00412933	0.0040045
6	No. .L-1	0.001356617	0.001676537	0.003793882	0.00132937	0.00412933	0.0040045
7	No. .L-1	0.001356617	0.001676537	0.003793882	0.00132937	0.00412933	0.0040045
8	No. .L-1	0.001356617	0.001676537	0.003793882	0.00132937	0.00412933	0.0040045
9	No. ingested	4.06985E-05	5.06279E-05	0.00012843	3.7221E-05	0.00015657	0.0001397
10	prob. inf. person-1,d-1	2.65894E-05	3.30726E-05	8.38834E-05	2.4318E-05	0.00010225	9.123E-05

Time Period Adjusted Risk (generally annualized)

Persons	Days	Statistic >					
		Modal @Risk default value	Simulation Average	95th %ile	50th %ile	99th %ile	97.5th %ile
1	1	2.66E-05	3.31E-05	8.39E-05	2.43E-05	1.02E-04	9.12E-05

Run Simulation Scenario (time permitting)

- Lake Parramatta Bathing
- Infection risk probability:
 - normal
 - & with contaminated water *ca* 20% of the time

QMRA Limitations v. Strengths

- Strengths (some of many)
 - Easily conceptualised outputs
 - Hypothesis testing possible
 - Exploration of scenarios, Events, management actions
 - Numerical Critical Limit Definition
 - Whole of system analysis possible
 - Promotes use in management of objective scientific data and systematic identification of knowledge gaps
- Limitations (not limitations of @Risk but output interpretation)
 - Not reality
 - PDF selection can be challenging
 - Various ‘Known unknowns’ e.g. pathogen viability and subpopulations
 - Order of magnitude precision only – may overestimate risk?

QUESTIONS

