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Bergion, V., Lindhe, A., Sokolova, E. et al (2018)

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Water Research, 132: 111-123

http://dx.doi.org/10.1016/j.watres.2017.12.054

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Accepted Manuscript

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PII: S0043-1354(17)31049-7

DOI: 10.1016/j.watres.2017.12.054

Reference: WR 13452

To appear in: Water Research

Received Date: 13 July 2017

Revised Date: 31 October 2017

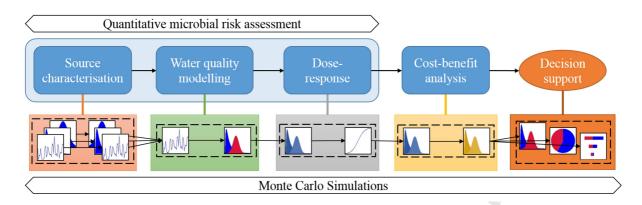
Accepted Date: 22 December 2017

Please cite this article as: Bergion, V., Lindhe, A., Sokolova, E., Rosén, L., Risk-based cost-benefit analysis for evaluating microbial risk mitigation in a drinking water system, *Water Research* (2018), doi: 10.1016/j.watres.2017.12.054.

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Risk-based decision model



Risk-based cost-benefit analysis for evaluating

² microbial risk mitigation in a drinking water

3 system

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8 Abstract

- 9 Waterborne outbreaks of gastrointestinal diseases can cause large costs to society. Risk
- management needs to be holistic and transparent in order to reduce these risks in an effective
- manner. Microbial risk mitigation measures in a drinking water system were investigated
- using a novel approach combining probabilistic risk assessment and cost-benefit analysis.
- Lake Vomb in Sweden was used to exemplify and illustrate the risk-based decision model.
- Four mitigation alternatives were compared, where the first three alternatives, A1-A3,
- represented connecting 25, 50 and 75 %, respectively, of on-site wastewater treatment
- systems in the catchment to the municipal wastewater treatment plant. The fourth alternative,
- 17 A4, represented installing a UV-disinfection unit in the drinking water treatment plant.
- Quantitative microbial risk assessment was used to estimate the positive health effects in
- terms of quality adjusted life years (QALYs), resulting from the four mitigation alternatives.
- 20 The health benefits were monetised using a unit cost per QALY. For each mitigation
- 21 alternative, the net present value of health and environmental benefits and investment,

maintenance and running costs	was calculated. The results showed that only A4 can reduce
the risk (probability of infection	a) below the World Health Organization guidelines of 10 ⁻⁴
infections per person per year (ooking at the 95 th percentile). Furthermore, all alternatives
resulted in a negative net presen	nt value. However, the net present value would be positive
(looking at the 50 th percentile u	sing a 1 % discount rate) if non-monetised benefits (e.g.
increased property value divide	d evenly over the studied time horizon and reduced microbial
risks posed to animals), estimat	ed at 800-1200 SEK (€100-150) per connected on-site
wastewater treatment system pe	er year, were included. This risk-based decision model creates
a robust and transparent decision	n support tool. It is flexible enough to be tailored and applied
to local settings of drinking wat	er systems. The model provides a clear and holistic structure
for decisions related to microbi	al risk mitigation. To improve the decision model, we suggest
to further develop the valuation	and monetisation of health effects and to refine the
propagation of uncertainties and	l variabilities between the included methods.

Keywords: cost-benefit analysis (CBA), decision support, drinking water, quality adjusted life

37 year (QALY), quantitative microbial risk assessment (QMRA), water quality modelling

1 Introduction

38

39	Risk management of drinking water systems (DWSs) is an iterative process including risk
40	assessment and risk mitigation (i.e. risk treatment) (ISO 2009). To be effective in providing
41	safe drinking water supply, the risk management must comprise the entire system, from
42	catchment to consumer. If the risks are unacceptable, risk mitigation measures should be
43	implemented, and alternatives for risk mitigation evaluated. Water Safety Plans procedures,
44	developed by the World Health Organization (WHO), can serve as a risk management
45	strategy for water providers (Bartram et al. 2009). However, in order to allocate societal
46	resources for risk mitigation in an efficient manner, the economic dimension of risk levels and
47	possible risk mitigation measures must be considered (WHO 2011).
48	Risks related to DWSs have been extensively discussed in the literature (e.g. Beuken et al.
49	2008, Keller and Wilson 1992, WHO 2011). Health risks in DWSs can be related to chemical,
50	microbial and radiological hazards (WHO 2011). In this paper, the microbial risks are the
51	main focus. Microbial risks in the form of pathogenic microorganisms can originate from
52	faecal sources (Dufour et al. 2012, Ferguson et al. 2009) related to humans (municipal
53	wastewater treatment plants (WWTPs) or on-site wastewater treatment systems (OWTSs) on
54	private properties) or animals (wild animals, domestic grazing animals or use of manure on
55	cropland). Pathogens in DWSs can cause endemic waterborne illness (Payment and Hunter
56	2001) as well as waterborne outbreaks of gastrointestinal diseases, resulting in high costs for
57	the society (Corso et al. 2003, Larsson et al. 2014). The WHO pointed out that the societal
58	costs for endemic waterborne illness and related gastrointestinal disease are commonly
59	underestimated (WHO 2001).
60	Quantitative microbial risk assessment (QMRA) has been applied to DWSs in various settings
61	(Haas et al. 2014, WHO 2016) in order to assess the risk in relation to an acceptable or

62	tolerable risk level. The result from a QMRA is typically reported as probability of infection,
63	disability adjusted life years (DALYs) or quality adjusted life years (QALYs). Both DALYs
64	and QALYs are health metrics that combine mortality and morbidity. Drinking water
65	producers commonly look at the (WHO) for guidance and the suggested risk levels of an
66	annual probability of infection of 10 ⁻⁴ per person per year, and DALYs of 10 ⁻⁶ per person per
67	year (WHO 2011).
68	To make informed decisions on which risk mitigation measure to implement in order to use
69	societal resources effectively, the alternatives need to be compared. Comprehensive lists and
70	procedures for identifying risk mitigation measures (e.g. Åström and Pettersson 2010, NZMH
71	2014, Rosén et al. 2010) are available. Decision support systems or decision models such as
72	cost-effectiveness analysis (CEA) and multi-criteria decision analysis (MCDA) can aid
73	decision makers in comparing the alternatives. If there are no regulations regarding acceptable
74	risk levels, other evaluation methods might be needed in order to justify the implementation
75	of risk mitigation measures. Cost-benefit analysis (CBA) provides a robust well-established
76	decision support approach to investigate the measure that is the most profitable or least costly
77	(if a certain risk level is required) for society (Boardman et al. 2011, Cameron et al. 2011).
78	Comparing mitigation measures directed at different parts of the supply system and
79	identifying the options most profitable for society are key steps towards a holistic and
80	sustainable risk management approach. Adopting holistic risk management also enables the
81	multi-barrier approach emphasised by the WHO (2011). Using CBA as a basis for decision
82	support helps to allocate monetary resources in an efficient manner providing possibilities to
83	compare mitigation measures with interventions in other sectors (e.g. food, health care, traffic
84	and environmental risk management). CBA facilitates optimisation of the societal resources
85	by comparing economic metrics, such as net present value (NPV), and performing
86	distributional analysis (Cameron et al. 2011). CRA also helps highlight the societal benefits of

87	reducing microbial risks in DWSs and creates a systematic and transparent decision support
88	tool.
89	Different frameworks for combining risk management, decision making process and CBA in
90	the drinking water context have been investigated (e.g. Assmuth et al. 2016, Rizak et al.
91	2003). Despite the aforementioned implementations, there are few, if any, methods that use a
92	probabilistic quantitative risk-based approach to create decision support in the form of a CBA
93	for microbial risk management in DWSs. To include an economic dimension and to perform a
94	CBA in this way is uncommon, even though the need is emphasised by the WHO (WHO
95	2001).
96	Aim
97	In this study we develop a method for creating a systematic, holistic and transparent decision
98	support for microbial risk management in DWSs. We present a novel CBA approach from
99	catchment to consumer. More in detail, we perform a CBA using a combination of water
100	quality modelling and QMRA to compare microbial risk mitigation alternatives in a DWS.
101	The methodology is exemplified using Lake Vomb in the south of Sweden. Different
102	alternatives of removing OWTSs are compared to installation of an additional treatment step
103	in the drinking water treatment plant (DWTP). We also highlight the choices that needs to be
104	made in the CBA-model, and what implications these might have on the outcome of the CBA
105	2 Risk-based decision model
106	The suggested approach for combining the methods for QMRA and CBA is presented as a
107	decision model in Figure 1. The four major compartments are: (i) source characterisation, (ii)
108	water quality modelling, (iii) dose-response, and (iv) CBA. The source characterisation
109	provides input to the water quality modelling, and the water quality modelling provides input
110	to the dose-response. The OMRA framework, including (i), (ii) and (iii), describes the entire

risk chain in the DWS and provides input for the CBA. Epistemic uncertainties (associated with lack of knowledge) and aleatory uncertainties (associated with natural variations) in all compartments are incorporated into the model by means of Monte Carlo (MC) simulations. The combination of methods aims to enable an estimation of the microbial risk in the DWS as well as an estimation of the effect of risk reduction measures and their societal profitability. Hence, the decision model can serve as a tool within the water safety plan framework. When analysing different mitigation measures, each compartment of the decision model needs to be executed. Detailed method descriptions of each compartment are presented in sections 3.2-3.4. It should be noted that this decision model is generic, and the applied methods in each case study should be selected to fit the specific context of the analysed DWS.

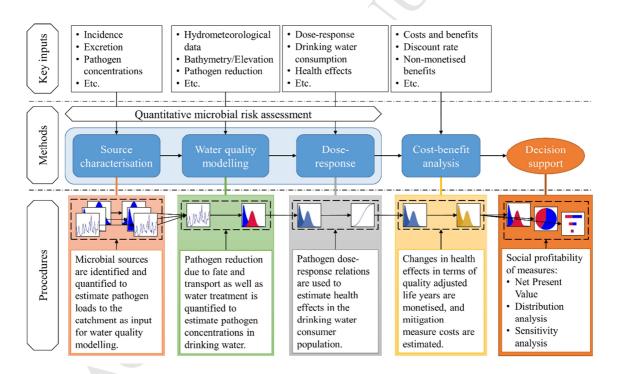


Figure 1. Risk-based decision model combining methods for evaluating and comparing microbial risk mitigation measures.

3 Methods

3.1 Lake Vomb

Lake Vomb is a small lake in Scania, the southernmost part of Sweden, providing 330,000 consumers with drinking water. The average water depth is 6.6 m, and the maximum depth is 16 m. Three major tributaries discharge into Lake Vomb: Borstbäcken, Torpsbäcken and Björkaån draining 26, 42 and 340 km², respectively. There are approximately 2800 OWTSs in the catchment (Norwegian Water BA 2009) posing a risk to the drinking water source. Other sources of microbial risks are e.g. WWTP, fertilisation using manure, grazing animals, wild animals. Raw water is extracted from Lake Vomb and artificially infiltrated into a glaciofluvial aquifer and then treated using conventional treatment consisting of rapid sand filtration and chlorination (Norwegian Water BA 2009). Figure 2 illustrates the case study area.

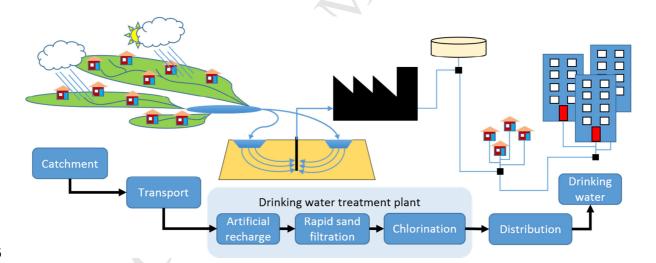


Figure 2 Schematic illustration of Lake Vomb drinking water system.

Microbial risk mitigation alternatives in different parts of the DWS were chosen to illustrate how the risk-based decision model can be used. The mitigation alternatives also reflect the contemporary trends in Sweden regarding OWTSs management and an increase in installation of UV-disinfection in DWTPs. Three of the analysed alternatives represent connection of

142	different proportions (25, 50 and 75 %, respectively) of the OWTSs in the catchment to the
143	municipal WWTP. The costs for the alternatives were based on connection of clusters of
144	closely located OWTSs. However, the pathogen load from these OWTSs was assumed to be
145	removed evenly across the different types of OWTSs and geographically across the catchment
146	area. This assumption was made because of the short transport time in the catchment (Sundahl
147	et al. 2008). The fourth alternative was to install UV-disinfection at the DWTP at Lake Vomb.
148	The four decision alternatives and one reference alternative were analysed:
149	• Reference alternative (A-Ref) – Continuation of the present state.
150	• Alternative 1 (A1) – Connecting 25 % (621) of the OWTSs to the local WWTP.
151	• Alternative 2 (A2) – Connecting 50 % (1240) of the OWTSs to the local WWTP.
152	• Alternative 3 (A3) – Connecting 75 % (1861) of the OWTSs to the local WWTP.
153	• Alternative 4 (A4) – An additional barrier, UV-disinfection, is installed at the
	DIVID
154	DWTP.
154 155	3.2 Quantitative microbial risk assessment (QMRA)
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155 156 157 158	3.2 Quantitative microbial risk assessment (QMRA) The QMRA methodology (Haas et al. 2014) was used for quantifying the health effects related to the reference alternative and the microbial risk mitigation alternatives. All inputs for the QMRA are listed in Table 1; input distributions represent both epistemic uncertainties and
155 156 157 158 159	3.2 Quantitative microbial risk assessment (QMRA) The QMRA methodology (Haas et al. 2014) was used for quantifying the health effects related to the reference alternative and the microbial risk mitigation alternatives. All inputs for the QMRA are listed in Table 1; input distributions represent both epistemic uncertainties and aleatory uncertainties.
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155 156 157 158 159 160 161	3.2 Quantitative microbial risk assessment (QMRA) The QMRA methodology (Haas et al. 2014) was used for quantifying the health effects related to the reference alternative and the microbial risk mitigation alternatives. All inputs for the QMRA are listed in Table 1; input distributions represent both epistemic uncertainties and aleatory uncertainties. 3.2.1 Source characterisation Human pathogens in wastewater from OWTS were quantified as described by Ottoson and
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pathogen concentration ($C_{Tributary}$, pathogens/L) of each reference pathogen in each tributary was calculated as:

$$C_{Tributary} = \frac{I_{Path} \cdot U \cdot D_{Path} \cdot FP \cdot C \cdot P}{365 \cdot 10^5 \cdot F_{Tributary}}$$
(1)

where I_{Path} (infections/year/ 10^5 inhabitants) was the incidence; U (no unit) was the factor for underreporting; D_{Path} (days) was the duration of excretion; FP (g) was the faecal production per person per day; C (pathogens/g) was the pathogen concentration in faeces when infected; P (persons) was the number of persons that are using OWTSs in the sub-basin; and $F_{Tributary}$ (L/day) was the average daily flow for each tributary.

Table 1 Inputs for the quantitative microbial risk assessment.

Red. in on-site sewage ^d	Input	Unit	Abbr.	Distr. s	Parameters
Factor of underreporting $^{\rm b}$ - U PV 38 Faecal production $^{\rm c}$ g FP N P05=115.7, P95=144.2 Red. in on-site sewage $^{\rm d}$ Log ₁₀ red. R_{OWTS} PV 0.72 Depth of unsat. zone $^{\rm c}$ m D_{PUZ} T Min=1.2, Mode=1.5, Max=2.4 Length of satur. zone $^{\rm c}$ m D_{PUZ} T Min=100, Mode=150, Max=250 Drinking water consumption $^{\rm f}$ L/day WI e $^{\rm N}$ N(μ =0.299, σ =0.57) Average flow Björkaân $^{\rm g}$ L/day F_{Biorka} PV 3.21 $^{\rm e}10^{\rm f}$ Average flow Torpsbäcken $^{\rm g}$ L/day F_{Biorka} PV 3.21 $^{\rm e}10^{\rm f}$ Average flow Borstbäcken $^{\rm g}$ L/day F_{Biorka} PV 3.46 $^{\rm e}10^{\rm f}$ Average flow Borstbäcken $^{\rm g}$ L/day F_{Biorka} PV 2.16 $^{\rm e}10^{\rm f}$ Incidence $^{\rm h}$ Inputs for norrovirus Incidence $^{\rm h}$ Log ₁₀ path/g C_{Noro} N P01=5, P99=9 Acd. lake transp. Björkaân $^{\rm k}$ Log ₁₀ red. $R_{Biork,Noro}$ N P1=5, p90=27, Loc=0 Acd in unsatur. zone $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ N P2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ N P2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ N P2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ N P2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ N P2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ N P2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ LN R2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ N P2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ LN P2, and an incidence $^{\rm h}$ Log ₁₀ red. $R_{Biork,Noro}$ PV Log ₁₀ P2, box PV Log ₁₀ P2				outs	<u></u>
Factor of underreporting $^{\rm b}$	Pers. connected to OWTSs ^a	#	P	PV	ARef=6215, A1=4661, A2=3107, A3=1554,
Faecal production $^{\circ}$ Red. in on-site sewage $^{\rm d}$ Log $_{10}$ red. R_{OWTS} PV 0.72 T Min=1.2, Mode=1.5, Max=2.4 Min=1.2, Mode=1.5, Max=2.4 Min=1.2, Mode=1.5, Max=2.50 Drinking water consumption $^{\rm f}$ L/day WI e $^{\rm N}$ N(μ=-0.299, c=0.57) Average flow Björkaån $^{\rm g}$ L/day F_{Bjorka} PV 3.46*10 $^{\rm 7}$ L/day F_{Bjorka} PV 3.46*10 $^{\rm 7}$ L/day F_{Bjorka} PV 2.16*10 $^{\rm 7}$ Min=1.2, Mode=1.5, Max=2.4 Min=1.2, Mode=1.5, Max=1.2, Mode=1.2,					A4=6215
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$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Faecal production ^c	g		N	P05=115.7, P95=144.2
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Average flow Torpsbäcken g L/day F_{Torps} PV $3.46*10^{7}$ L/day F_{Borst} PV $2.16*10^{7}$ Inputs for norovirus Incidence h Inf./10 5 inh/y I_{Noro} G α =6.25, β =10.6 Days excreting i days D_{Noro} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Noro} N P01=5, P99=9 Red. lake transp. Björkaån k Log ₁₀ red. $R_{Biork_{Noro}}$ N μ =4.5, σ =0.12, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. $R_{Borst_{Noro}}$ N μ =4.5, σ =0.11, Tr: Min=0 Red. in unsatur. zone l Log ₁₀ red. $R_{Borst_{Noro}}$ N μ =4.6, σ =0.10, Tr: Min=0 Red. in unsatur. zone l Log ₁₀ red. $R_{Borst_{Noro}}$ N μ =4.6, σ =0.10, Tr: Min=0 Red. in conv. treatm. n Log ₁₀ red. $R_{ST_{Noro}}$ T Min=0.05, Mode=0.3, Max=1 Red. in UV-treatm. 0 Log ₁₀ red. $R_{CT_{Noro}}$ T Min=0.4, Mode=0.5, Max=0.6 Red. in UV-treatm. 0 Log ₁₀ red. $R_{CT_{Noro}}$ PV 0.000716 QALYs per infection r QALY/inf D_{Noro} PV 0.0009 Incidence h Inf./10 5 inh/y D_{Noro} PV 0.0009 Incidence h Inf./10 5 inh/y D_{Camp} Log ₁₀ red. D_{Camp} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N μ =5.6, σ =0.40, Tr: Min=0 Red. lake transp. Björkaån D_{Camp} Log ₁₀ red. D_{Camp} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. D_{Camp} Log ₁₀ red. D_{Camp} N D_{Camp} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. D_{Camp} Log ₁₀ red. D_{Camp} N D_{Camp	Drinking water consumption ^f	L/day	WI	e^{N}	$N(\mu=-0.299, \sigma=0.57)$
Average flow Torpsbäcken g L/day F_{Torps} PV $3.46*10^{7}$ Average flow Borstbäcken g L/day F_{Borst} PV $2.16*10^{7}$ Logar Inputs for norovirus Incidence h Inf./10 5 inh/y I_{Noro} G $\alpha=6.25, \beta=10.6$ Days excreting i days D_{Noro} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log10 path./g C_{Noro} N P01=5, P99=9 Red. lake transp. Björkaån k Log10 red. $R_{Biork,Noro}$ N $\mu=4.5, \alpha=0.12, \mathrm{Tr}$: Min=0 Red. lake transp. Borstbäcken k Log10 red. $R_{Borst,Noro}$ N $\mu=4.5, \alpha=0.12, \mathrm{Tr}$: Min=0 Red. in unsatur. zone h Log10 red. $R_{Borst,Noro}$ N $\mu=4.6, \alpha=0.10, \mathrm{Tr}$: Min=0 Red. in atur. zone h Log10 red. $R_{Borst,Noro}$ N $\mu=4.6, \alpha=0.10, \mathrm{Tr}$: Min=0 Red. in in conv. treatm. h Log10 red. $R_{Borst,Noro}$ T Min=0.05, Mode=0.3, Max=1 Log10 red. $R_{CT,Noro}$ T Min=0.05, Mode=0.5, Max=0.6 Red. in CV-treatm. h Log10 red. $R_{CT,Noro}$ PV 4.2 Min=0.4, Mode=0.5, Max=0.6 Log10 red. $R_{CV,Noro}$ PV 0.000716 QALYs per infection f DALY/inf D_{Noro} PV 0.0009 PV	Average flow Björkaån g	L/day	F_{Bjorka}	PV	$3.21*10^8$
Average flow Borstbäcken g L/day F_{Borst} PV 2.16*10' Inputs for norovirus Incidence h Inf./10\$inh/y I_{Noro} G α =6.25, β =10.6 Days excreting i days D_{Noro} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Noro} N P01=5, P99=9 Red. lake transp. Björkaån k Log ₁₀ red. $R_{Bjork.Noro}$ N μ =4.5, σ =0.12, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. $R_{Bjork.Noro}$ N μ =4.5, σ =0.12, Tr: Min=0 Red. in unsatur. zone i Log ₁₀ red. $R_{Borst.Noro}$ N μ =4.6, σ =0.10, Tr: Min=0 Red. in satur. zone i Log ₁₀ red. $R_{Borst.Noro}$ N μ =4.6, σ =0.10, Tr: Min=0 Red. in satur. zone i Log ₁₀ red. $R_{UZ/m.Noro}$ T Min=0.0.5, Mode=0.3, Max=1 Log ₁₀ red. $R_{UZ/m.Noro}$ T Min=0.04, Mode=0.5, Max=0.6 Red. in Conv. treatm. o Log ₁₀ red. $R_{UV.Noro}$ PV 4.2 Infectivity p - M_{Noro} EBP σ =0.04, ρ =0.055 DALYs per infection f QALY/inf D_{Noro} PV 0.000716 QALYs per infection f QALY/inf D_{Noro} PV 0.0009 Inputs for Campylobacter Incidence h Inf./10\$inh/y I_{Camp} G σ =64.6, ρ =1.27 Days excreting i Log ₁₀ red. $R_{Bjork.Camp}$ N P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N P01=4, P99=10 Red. lake transp. Björkaån k Log ₁₀ red. $R_{Bjork.Camp}$ N μ =5.5, σ =0.40, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. $R_{Bjork.Camp}$ N μ =5.5, σ =0.29, Tr: Min=0 Red. in unsatur. Zone l Log ₁₀ red. $R_{Bjork.Camp}$ N μ =5.6, σ =0.38, Tr: Min=0 Red. in satur. Zone l Log ₁₀ red. $R_{Bjork.Camp}$ N μ =5.6, σ =0.38, Tr: Min=0 Red. in satur. Zone l Log ₁₀ red. $R_{Bjork.Camp}$ N μ =5.6, σ =0.38, Tr: Min=0 Red. in satur. Zone l Min=0.001, Mode=0.05, Max=1 Red. in conv. treatm. n Log ₁₀ red. $R_{IUZ/m.Camp}$ T Min=0.02, Mode=0.5, Max=1 Red. in conv. treatm. n Log ₁₀ red. $R_{IUZ/m.Camp}$ T Min=0.02, Mode=0.25, Max=0.3	Average flow Torpsbäcken g	L/day		PV	$3.46*10^7$
Incidence h	Average flow Borstbäcken g	L/day		PV	$2.16*10^7$
$\begin{array}{cccccccccccccccccccccccccccccccccccc$			Inputs for nor	ovirus	
Days excreting i days D_{Noro} LN $P01=13, P99=27, Loc=0$ Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Noro} N $P01=5, P99=9$ Red. lake transp. Björkaån k Log ₁₀ red. $R_{Biork,Noro}$ N $\mu=4.5, \sigma=0.12, Tr: Min=0$ Red. lake transp. Borstbäcken k Log ₁₀ red. $R_{Torp,DNoro}$ N $\mu=4.5, \sigma=0.11, Tr: Min=0$ Red. lake transp. Borstbäcken k Log ₁₀ red. $R_{Borst,Noro}$ N $\mu=4.6, \sigma=0.10, Tr: Min=0$ Red. in unsatur. zone l Log ₁₀ red. $R_{Borst,Noro}$ T Min=0.05, Mode=0.3, Max=1 Red. in satur. zone m Log ₁₀ red. $R_{Torp,Noro}$ T Min=0.05, Mode=0.3, Max=1 Log ₁₀ red. $R_{Torp,Coro}$ T Min=0.4, Mode=0.5, Max=0.6 Red. in Conv. treatm. o Log ₁₀ red. $R_{UV,Noro}$ PV 4.2 Infectivity p - m_{Noro} EBP $\alpha=0.04, \beta=0.055$ DALY/inf D_{Noro} PV 0.000716 QALYs per infection q DALY/inf D_{Noro} PV 0.0009 Thusts for $Campylobacter$ Infectione h Days excreting i days D_{Camp} Log ₁₀ path./g C_{Camp} N P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N P01=4, P99=10 Log ₁₀ red. $R_{Bjork,Camp}$ N P01=4, P99=10 Log ₁₀ red. $R_{Bjork,Camp}$ N $\mu=5.5, \sigma=0.40, Tr: Min=0$ Red. lake transp. Borstbäcken k Log ₁₀ red. $R_{Bjork,Camp}$ N $\mu=5.5, \sigma=0.29, Tr: Min=0$ Red. in unsatur. Zone l Log ₁₀ red. $R_{Bjork,Camp}$ N $\mu=5.9, \sigma=0.29, Tr: Min=0$ Red. in satur. Zone l Log ₁₀ red. $R_{CT,Camp}$ T Min=0.001, Mode=0.05, Max=1 Red. in satur. Zone l Log ₁₀ red. $R_{CT,Camp}$ T Min=0.001, Mode=0.05, Max=1 Red. in conv. treatm. n Log ₁₀ red. $R_{CT,Camp}$ T Min=0.001, Mode=0.25, Max=0.3 Red. in UV-treatm. o Log ₁₀ red. $R_{CT,Camp}$ PV 5.3	Incidence h	Inf./10 ⁵ inh/y	I_{Noro}	G	α =6.25, β =10.6
Red. lake transp. Björkaån k	Days excreting i	days		LN	
Red. lake transp. Björkaån k Log ₁₀ red. R_{Bjork_Noro} N μ=4.5, σ =0.12, Tr: Min=0 Red. lake transp. Torpsbäcken k Log ₁₀ red. R_{Bjork_Noro} N μ=4.5, σ =0.11, Tr: Min=0 μ=4.5, σ =0.11, Tr: Min=0 μ=4.6, σ =0.10, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Borst_Noro} N μ=4.6, σ =0.10, Tr: Min=0 Red. in unsatur. zone n Log ₁₀ red. R_{UZm_Noro} T Min=0.05, Mode=0.3, Max=1 Log ₁₀ red. R_{UZm_Noro} LN μ=6.6587, σ =5.5366 Red. in conv. treatm. n Log ₁₀ red. R_{UV_Noro} PV 4.2 Log ₁₀ red. R_{UV_Noro} PV 4.2 Infectivity p - m_{Noro} EBP σ =0.04, β =0.055 DALYs per infection q DALYinf O_{Noro} PV 0.000716 QALYs per infection r QALYinf O_{Noro} PV 0.0009 Inputs for Campylobacter Incidence h Inf./10 ⁵ inh/y I_{Camp} G σ =64.6, σ =1.27 days excreting i days O_{Camp} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N P01=4, P99=10 Log ₁₀ red. R_{Bjork_Camp} N ρ =5.5, σ =0.40, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Bjork_Camp} N ρ =5.5, σ =0.40, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Bjork_Camp} N ρ =5.9, σ =0.29, Tr: Min=0 Red. in unsatur. Zone l Log ₁₀ red. R_{Bjork_Camp} N ρ =5.9, σ =0.29, Tr: Min=0 Red. in satur. Zone l Log ₁₀ red. R_{Bjork_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in satur. Zone l Log ₁₀ red. R_{Bjork_Camp} T Min=0.001, Mode=0.05, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3	Path. conc. in faeces if inf. j	Log ₁₀ path./g	C_{Noro}	N	P01=5, P99=9
Red. lake transp. Torpsbäcken k	Red. lake transp. Björkaån k	Log ₁₀ red.		N	μ =4.5, σ =0.12, Tr: Min=0
Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Borst_Noro} N μ=4.6, σ=0.10, Tr: Min=0 Red. in unsatur. zone 1 Log ₁₀ red./m R_{UZ/m_Noro} T Min=0.05, Mode=0.3, Max=1 Red. in satur. zone m Log ₁₀ red./m R_{SZ_Noro} LN μ=6.6587, σ=5.5366 Red. in conv. treatm. 0 Log ₁₀ red. R_{CT_Noro} T Min=0.4, Mode=0.5, Max=0.6 Red. in UV-treatm. 0 Log ₁₀ red. R_{UV_Noro} PV 4.2 Infectivity p - m_{Noro} EBP m_{Noro} PV 0.000716 QALYs per infection q DALY/inf m_{Noro} PV 0.0009 Inputs for Campylobacter Incidence h Days excreting i days m_{Noro} DCamp LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g m_{Noro} Red. lake transp. Björkaån m_{Noro} Red. lake transp. Borstbäcken m_{Noro} Red. lake transp. Borstbäcken m_{Noro} Red. lake transp. Borstbäcken m_{Noro} Red. in unsatur. Zone m_{Noro} Red. in satur. Zone m_{Noro} Red. in satur. Zone m_{Noro} Red. in satur. Zone m_{Noro} Red. in conv. treatm. m_{Noro} Log ₁₀ red. m_{Noro} Red. in conv. treatm. m_{Noro} Red. in UV-treatm. m_{Noro} Red. in UV-treatm. m_{Noro} Red. in UV-treatm. m_{Noro} Red. Red. in Conv. treatm. m_{Noro} Red. Red. in Conv. treatm. m_{Noro} Red. Red. in UV-treatm. m_{Noro} Red. Red. Red. Red. Red. Red. Red. Red.	Red. lake transp. Torpsbäcken k	Log_{10} red.		N	μ =4.5, σ =0.11, Tr: Min=0
Red. in satur. zone $^{\rm m}$	Red. lake transp. Borstbäcken k	Log_{10} red.		N	μ =4.6, σ =0.10, Tr: Min=0
Red. in satur. zone $^{\rm m}$	Red. in unsatur. zone ¹	Log ₁₀ red./m	$R_{UZ/m\ Noro}$	T	Min=0.05, Mode=0.3, Max=1
Red. in conv. treatm. n	Red. in satur. zone ^m	Log ₁₀ red./m		LN	μ =6.6587, σ =5.5366
Infectivity $^{\rm P}$ — m_{Noro} EBP α =0.04, β =0.055 DALYs per infection $^{\rm q}$ DALY/inf D_{Noro} PV 0.000716 QALYs per infection $^{\rm r}$ QALY/inf D_{Noro} PV 0.0009 Inputs for $Campylobacter$ Incidence $^{\rm h}$ Inf./10 ⁵ inh/y I_{Camp} G α =64.6, β =1.27 Days excreting $^{\rm i}$ days D_{Camp} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. $^{\rm j}$ Log ₁₀ path./g C_{Camp} N P01=4, P99=10 Red. lake transp. Björkaån $^{\rm k}$ Log ₁₀ red. R_{Bjork_Camp} N μ =5.5, σ =0.40, Tr: Min=0 Red. lake transp. Borstbäcken $^{\rm k}$ Log ₁₀ red. R_{Torp_Camp} N μ =5.6, σ =0.38, Tr: Min=0 Red. in unsatur. Zone $^{\rm l}$ Log ₁₀ red. R_{Borst_Camp} N μ =5.9, σ =0.29, Tr: Min=0 Red. in satur. Zone $^{\rm l}$ Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.005, Mode=0.5, Max=1 Red. in conv. treatm. $^{\rm n}$ Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. $^{\rm o}$ Log ₁₀ red. R_{UV_Camp} PV 5.3	Red. in conv. treatm. ⁿ	Log_{10} red.		T	Min=0.4, Mode=0.5, Max=0.6
Infectivity $^{\rm p}$	Red. in UV-treatm. °	Log_{10} red.	$R_{UV\ Noro}$	PV	4.2
QALYs per infection $^{\Gamma}$ QALYinf Q_{Noro} PV 0.0009 Inputs for Campylobacter Incidence h Inf./10 ⁵ inh/y I_{Camp} G α=64.6, β=1.27 Days excreting i days D_{Camp} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N P01=4, P99=10 Red. lake transp. Björkaån k Log ₁₀ red. R_{Bjork_Camp} N μ=5.5, σ=0.40, Tr: Min=0 Red. lake transp. Torpsbäcken k Log ₁₀ red. R_{Torp_Camp} N μ=5.6, σ=0.38, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Rorst_Camp} N μ=5.9, σ=0.29, Tr: Min=0 Red. in unsatur. Zone l Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in satur. Zone l Log ₁₀ red./m R_{SZ/m_Camp} T Min=0.01, Mode=0.25, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. 0 Log ₁₀ red. R_{UV_Camp} PV <td>Infectivity ^p</td> <td>-</td> <td>_</td> <td>EBP</td> <td>α=0.04, β=0.055</td>	Infectivity ^p	-	_	EBP	α =0.04, β =0.055
QALYs per infection $^{\Gamma}$ QALYinf Q_{Noro} PV 0.0009 Inputs for Campylobacter Incidence h Inf./10 ⁵ inh/y I_{Camp} G α=64.6, β=1.27 Days excreting i days D_{Camp} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N P01=4, P99=10 Red. lake transp. Björkaån k Log ₁₀ red. R_{Bjork_Camp} N μ=5.5, σ=0.40, Tr: Min=0 Red. lake transp. Torpsbäcken k Log ₁₀ red. R_{Torp_Camp} N μ=5.6, σ=0.38, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Rorst_Camp} N μ=5.9, σ=0.29, Tr: Min=0 Red. in unsatur. Zone l Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in satur. Zone l Log ₁₀ red./m R_{SZ/m_Camp} T Min=0.01, Mode=0.25, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. 0 Log ₁₀ red. R_{UV_Camp} PV <td>DALYs per infection q</td> <td>DALY/inf</td> <td>D_{Noro}</td> <td>PV</td> <td>0.000716</td>	DALYs per infection q	DALY/inf	D_{Noro}	PV	0.000716
Inputs for Campylobacter Incidence h Inf./10 ⁵ inh/y I_{Camp} G α=64.6, β=1.27 Days excreting i days D_{Camp} LN P01=13, P99=27, Loc=0 Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N P01=4, P99=10 Red. lake transp. Björkaån k Log ₁₀ red. R_{Bjork_Camp} N μ=5. 5, σ=0.40, Tr: Min=0 Red. lake transp. Torpsbäcken k Log ₁₀ red. R_{Torp_Camp} N μ=5.6, σ=0.38, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Borst_Camp} N μ=5.9, σ=0.29, Tr: Min=0 Red. in unsatur. Zone l Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in satur. Zone l Log ₁₀ red./m R_{SZ/m_Camp} T Min=0.001, Mode=0.05, Max=1 Red. in conv. treatm. l Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. l Log ₁₀ red. R_{UV_Camp} PV 5.3	QALYs per infection ^r	QALY/inf		PV	0.0009
Days excreting i days D_{Camp} LN $P01=13, P99=27, Loc=0$ Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N $P01=4, P99=10$ Red. lake transp. Björkaån k Log ₁₀ red. R_{Bjork_Camp} N $\mu=5.5, \sigma=0.40, Tr: Min=0$ Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Torp_Camp} N $\mu=5.6, \sigma=0.38, Tr: Min=0$ Red. in unsatur. Zone l Log ₁₀ red. R_{Borst_Camp} N $\mu=5.9, \sigma=0.29, Tr: Min=0$ Red. in satur. Zone l Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. o Log ₁₀ red. R_{UV_Camp} PV 5.3	·	Inp	outs for Camp	ylobacter	
Days excreting 1 days D_{Camp} LN $P01=13, P99=27, Loc=0$ Path. conc. in faeces if inf. j Log ₁₀ path./g C_{Camp} N $P01=4, P99=10$ Red. lake transp. Björkaån k Log ₁₀ red. R_{Bjork_Camp} N $\mu=5.5, \sigma=0.40, Tr: Min=0$ Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Torp_Camp} N $\mu=5.6, \sigma=0.38, Tr: Min=0$ Red. in unsatur. Zone 1 Log ₁₀ red. R_{Borst_Camp} N $\mu=5.9, \sigma=0.29, Tr: Min=0$ Red. in satur. Zone 1 Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. 0 Log ₁₀ red. R_{UV_Camp} PV 5.3		Inf./10 ⁵ inh/y	I_{Camp}	G	α =64.6, β =1.27
Path. conc. in faeces if inf. $^{\rm J}$ Log ₁₀ path./g C_{Camp} N P01=4, P99=10 Red. lake transp. Björkaån $^{\rm k}$ Log ₁₀ red. R_{Bjork_Camp} N μ =5. 5, σ =0.40, Tr: Min=0 Red. lake transp. Borstbäcken $^{\rm k}$ Log ₁₀ red. R_{Torp_Camp} N μ =5. 6, σ =0.38, Tr: Min=0 Red. in unsatur. Zone $^{\rm l}$ Log ₁₀ red. R_{Borst_Camp} N μ =5.9, σ =0.29, Tr: Min=0 Nin=0.05, Mode=0.5, Max=1 Red. in conv. treatm. $^{\rm n}$ Log ₁₀ red. R_{CT_Camp} T Min=0.001, Mode=0.05, Max=1 Red. in UV-treatm. $^{\rm o}$ Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. $^{\rm o}$ Log ₁₀ red. R_{UV_Camp} PV 5.3	Days excreting i	days	D_{Camp}	LN	P01=13, P99=27, Loc=0
Red. lake transp. Torpsbäcken k Log ₁₀ red. R_{Torp_Camp} N μ =5.6, σ =0.38, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Borst_Camp} N μ =5.6, σ =0.29, Tr: Min=0 Red. in unsatur. Zone 1 Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.001, Mode=0.05, Max=1 Red. in UV-treatm. 0 Log ₁₀ red. R_{UV_Camp} PV 5.3	Path. conc. in faeces if inf. j	Log ₁₀ path./g		N	P01=4, P99=10
Red. lake transp. Torpsbäcken k Log ₁₀ red. R_{Torp_Camp} N μ =5.6, σ =0.38, Tr: Min=0 Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Borst_Camp} N μ =5.6, σ =0.29, Tr: Min=0 Red. in unsatur. Zone 1 Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.001, Mode=0.05, Max=1 Red. in UV-treatm. 0 Log ₁₀ red. R_{UV_Camp} PV 5.3	Red. lake transp. Björkaån k	Log_{10} red.	$R_{Bjork\ Camp}$	N	μ =5. 5, σ =0.40, Tr: Min=0
Red. lake transp. Borstbäcken k Log ₁₀ red. R_{Borst_Camp} N μ =5.9, σ =0.29, Tr: Min=0 Red. in unsatur. Zone 1 Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in satur. Zone 1 Log ₁₀ red./m R_{SZ/m_Camp} T Min=0.001, Mode=0.05, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. 0 Log ₁₀ red. R_{UV_Camp} PV 5.3	Red. lake transp. Torpsbäcken k			N	μ =5.6, σ =0.38, Tr: Min=0
Red. in unsatur. Zone 1 Log ₁₀ red./m R_{UZ/m_Camp} T Min=0.05, Mode=0.5, Max=1 Red. in satur. Zone 1 Log ₁₀ red./m R_{SZ/m_Camp} T Min=0.001, Mode=0.05, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. o Log ₁₀ red. R_{UV_Camp} PV 5.3	Red. lake transp. Borstbäcken k	Log_{10} red.			μ =5.9, σ =0.29, Tr: Min=0
Red. in satur. Zone 1 Log ₁₀ red./m R_{SZ/m_Camp} T Min=0.001, Mode=0.05, Max=1 Red. in conv. treatm. n Log ₁₀ red. R_{CT_Camp} T Min=0.2, Mode=0.25, Max=0.3 Red. in UV-treatm. $^{\circ}$ Log ₁₀ red. R_{UV_Camp} PV 5.3	Red. in unsatur. Zone ¹	Log ₁₀ red./m	R_{UZ/m_Camp}	T	Min=0.05, Mode=0.5, Max=1
Red. in UV-treatm. $^{\circ}$ Log ₁₀ red. R_{UV_Camp} PV 5.3	Red. in satur. Zone ¹	Log ₁₀ red./m		T	Min=0.001, Mode=0.05, Max=1
Red. in UV-treatm. Log ₁₀ red. R_{UV_Camp} PV 5.3	Red. in conv. treatm. ⁿ	Log_{10} red.	R_{CT_Camp}	T	Min=0.2, Mode=0.25, Max=0.3
Infectivity p - m_{Comp} EBP α =0.024, β =0.011	Red. in UV-treatm. °	Log_{10} red.		PV	5.3
- Cump	Infectivity ^p	-	m_{Camp}	EBP	α =0.024, β =0.011

DALYs per infection ^q	DALY/inf	D_{Camp}	PV	0.00328
QALYs per infection ^r	QALY/inf	Q_{Camp}	PV	0.0163
	Inp	uts for Crypto	osporidium	ı
Incidence ^f	Inf./10 ⁵ inh/y	I_{Crypt}	G	α =5.43, β =0.228
Days excreting ^g	days	D_{Cryp}	LN	P01=5, P99=30, Loc=0
Path. conc. in faeces if inf. j	Log ₁₀ path./g	C_{Cryp}	N	P01=7, P99=9
Red. lake transp. Björkaån ^k	Log_{10} red.	R_{Bjork_Cryp}	N	μ =4.5, σ = 0.12, Tr: Min=0
Red. lake transp. Torpsbäcken k	Log_{10} red.	R_{Torp_Cryp}	N	μ =4.6, σ =0.10, Tr: Min=0
Red. lake transp. Borstbäcken k	Log_{10} red.	R_{Borst_Cryp}	N	μ =4.6, σ =0.095, Tr: Min=0
Red. in unsatur. zone ¹	Log ₁₀ red./m	R_{UZ/m_Cryp}	T	Min=0.05, Mode=0.5, Max=1
Red. in satur. zone ¹	Log ₁₀ red./m	R_{SZ/m_Cryp}	T	Min=0.001, Mode=0.05, Max=1
Red. in conv. treatm. ⁿ	Log_{10} red.	R_{CT_Cryp}	T	Min=0.4, Mode=0.5, Max=0.6
Red. in UV-treatm. °	Log_{10} red.	$R_{UV\ Cryp}$	PV	3
Infectivity p	-	m_{Cryp}	EBP	α =0.115, β =0.176
DALYs per infection ^q	DALY/inf	D_{Cryp}	PV	0.00267
QALYs per infection ^r	QALY/inf	Q_{Cryp}	PV	0.0035

- a) 2.5 persons (Åström and Johansson 2015) per on-site wastewater treatment system.
- b) Mead et al. (1999)

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- c) Wyman et al. (1978)
- d) Log₁₀ reduction in on-site sewage was estimated using standard values for four types of systems, achieving 100 % 99 %, 95 %, 50 %, respectively (SEPA 1991, 2002, 2003, 2008b), and the proportions of OWTS types.
- e) Personal communication with B.M. Pott at Southern Sweden Water Supply (Sydvatten)
- f) Westrell et al. (2006)
- g) SMHI (2017)
- h) Based on yearly incidence data 2006-2016 (PHAS 2017). Norovirus and *Campylobacter*: Best fit using Chi-Squared, *Cryptosporidium*: Best fit using Kolmogorov-Smirnov. For norovirus: incidence was based on per season incidence and data from 2006-2012 was based on national data due to lack of regional estimates.
- i) Norovirus: The Min/Median of the range of days shedding from Atmar et al. (2008) was chosen as P01/P99 respectively, *Campylobacter* and *Cryptosporidium*: Reported triangular Min/Max from Petterson et al. (2016) was chosen for P01/P99 respectively.
- j) Norovirus: From Marshall et al. (2001) as reported in (Westrell 2004), *Campylobacter* and *Cryptosporidium*: Reported triangular Min/Max from Petterson et al. (2016) was chosen for P01/P99 respectively.
- k) Distributions fitted from three years (2005-2007) of simulated daily Log_{10} reduction from the hydrodynamic modelling. Tr=Truncation
- 1) Estimation based on expert judgement.
- m) Reduction calculated using 10,000 MC iterations using a groundwater transport model estimating the removal due to attachment, inactivation and dilution (Åström et al. 2016, Schijven et al. 2006).
- n) Norwegian Water BA (2009)
- o) Calculated using Equation 7. UV-dose (fluence) was set to 400 J/m². The inactivation constant (*k*) and intercept of the fluence axis (b) were set to (*k/b*) (0.106/0), (0,293/0) and (0,225/1.087) for norovirus, *Campylobacter* and *Cryptosporidium* respectively (Hijnen et al. 2006). Not to exceed the experimental range (Hijnen et al. 2006), the maximum Log₁₀ reduction was used for norovirus and *Cryptosporidium*.
- p) Norovirus: Teunis et al. (2008), Campylobacter: Teunis et al. (2005), Cryptosporidium: Teunis et al. (2002).
- q) Norovirus and Cryptosporidium: based on a re-analysis from Kemmeren et al. (2006), Campylobacter: based on Havelaar and Melse (2003).
- r) Batz et al. (2014)
- s) PV=point value, N=Normal distribution, T=triangular distribution, e^N=exponential with a normal distribution in the exponent, G=Gamma distribution, LN=LogNormal distribution, EBP=Exact Beta-Poisson distribution.

3.2.2 Water quality modelling

The pathogen concentrations at the raw water intake (C_{RW} , pathogens/L) were calculated as:

$$C_{RW} = (C_{Borst} \cdot 10^{-(R_{OWTS} + R_{Borst})}) + (C_{Torp} \cdot 10^{-(R_{OWTS} + R_{Torp})}) + (C_{Bjorka} \cdot 10^{-(R_{OWTS} + R_{Bjork})})$$
(2)

where C_{Borst} , C_{Torp} and C_{Bjorka} (pathogens/L) were the pathogen concentrations in the tributaries; R_{OWTS} (no unit) was the Log₁₀ reduction in the OWTSs; and R_{Borst} , R_{Torp} and R_{Bjork} (no unit) were the Log₁₀ reduction due to transport in Lake Vomb from the tributary to the raw water intake. Reduction in the catchment from OWTS discharge until entering Lake Vomb was conservatively assumed to be negligible due to the longevity of pathogens and the rapid transport (Sundahl et al. 2008) in the catchment. Hydrodynamic modelling was performed to simulate the fate and transport of pathogens from the point of entering Lake Vomb from the three tributaries to the raw water intake. Due to the linearity of the hydrodynamic model, a constant load was used to estimate the pathogen reduction. Decay of the pathogens was calculated as:

$$C_t = C_0 \cdot e^{-wt} \tag{3}$$

where C_t (pathogens/L) was the concentration at t; C_0 (pathogens/L) was the initial concentration; w (1/day) was the decay rate; and t was the time step. The w value was set to 0.23 for Campylobacter and 0.03 for Cryptosporidium and norovirus, based on the estimates of half-life of pathogens in environment. For Campylobacter, the median half-life was estimated from various literature sources (Catalao Dionisio et al. 2000, Cook and Bolster 2007, Hendricks 1971, McGee et al. 2002, Medema et al. 1997, Nasser et al. 2003, Ottosson and Stenström 2003, Rhodes and Kator 1988, Terzieva and McFeters 1991). For Cryptosporidium and norovirus the same half-life was used, estimated as the median half-life given in literature (Medema et al. 1997, Nasser et al. 2003, Ottosson and Stenström 2003). Cryptosporidium was also reduced due to settling in the lake. In the MIKE 3 FM, the settling velocity for Cryptosporidium was specified as 0.03 m/day, which is the settling velocity previously suggested for free oocysts (Medema et al. 1998). It was conservatively assumed that Cryptosporidium oocysts released into the lake were not attached to particles.

Comparing the input pathogen concentration at each tributary with the resulting pathogen concentration at the raw water intake, the daily Log_{10} reduction due to transport in the lake was calculated. Three year time-series of daily Log_{10} reductions were used to estimate the variability in the daily Log_{10} reduction for the three different transport paths. The estimated Log_{10} reductions for each tributary are presented in Table 1.

The pathogen concentration in drinking water (C_{DW} , pathogens/L) was calculated as:

$$C_{DW} = C_{RW} \cdot 10^{-(R_{UZ} + R_{SZ} + R_{CT} + R_{UV})}$$
 (4)

where R_{UZ} (no unit) was the total Log₁₀ reduction in the unsaturated zone; R_{SZ} (no unit) was the total Log₁₀ reduction in the saturated zone; R_{CT} (no unit) was the Log₁₀ reduction by the conventional treatment at the DWTP; and R_{UV} (no unit) was the total Log₁₀ reduction by the UV-disinfection. The chlorination step was assumed not to contribute to the microbial removal because of a small dose and that chloramine was used as disinfection agent¹.

The Log₁₀ reduction in the unsaturated zone (R_{UZ} , no unit) was calculated as:

$$R_{UZ} = R_{UZ/m} \cdot Dp_{UZ} \tag{5}$$

- where $R_{UZ/m}$ (Log_{10}/m) was the Log₁₀ reduction per meter, and Dp_{UZ} (m) was the depth of the unsaturated zone.
- For *Campylobacter* and *Cryptosporidium*, the Log₁₀ reduction in the saturated zone (R_{SZ} , *no unit*) was calculated as:

$$R_{SZ} = R_{SZ/m} \cdot L_{SZ} \tag{6}$$

where $R_{SZ/m}$ (Log_{10}/m) was the Log₁₀ reduction per meter in saturated zone, and L_{SZ} (m) was the length of the saturated zone. For norovirus, the Log₁₀ reduction in saturated zone (R_{SZ} , no

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¹ Personal communication the Southern Sweden Water Supply (Sydvatten).

- 256 *unit*) was estimated using a groundwater transport model (Åström et al. 2016, Schijven et al.
- 257 2006).
- The estimated Log₁₀ reductions in conventional treatment (R_{CT}) are presented in Table 1.
- The Log₁₀ reduction by the UV-disinfection (R_{UV} , no unit) was described as a first order
- 260 disinfection model and calculated as:

$$R_{N} = -x \cdot f - b \tag{7}$$

- where $x (cm^2/mJ)$ was an inactivation constant; b (no unit) was the interception of the fluence
- 263 axis; and $f(mJ/cm^2)$ was fluence.
- 3.2.3 Dose-response
- To estimate the health effects in the form of infections due to the pathogens in the drinking
- water, the pathogen daily dose (D, pathogens/day) was calculated as.

$$D = C_{DW} \cdot W \tag{8}$$

- where WI(L) was the daily ingested volume of drinking water per capita in Sweden (Westrell
- et al. 2006). All three reference pathogens were assigned the Exact Beta-Poisson dose-
- 270 response function. An Exact Beta-Poisson function can be represented by an exponential
- function with a beta distribution in the exponent (Equation 9); this approach has been reported
- to be representative in infection studies (Teunis et al. 2005, Teunis et al. 2002, Teunis et al.
- 273 2008).

$$P_{inf} = 1 - e^{-mD} (9)$$

- where P_{inf} (probability) was the daily probability of infection for each pathogen; m (no unit)
- was the infectivity; and D was the simulated daily pathogen dose that was ingested.
- The annual probability of infection (P_{annual} , probability) was calculated as:

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$$P_{annual} = 1 - \prod_{1}^{365} (1 - P_{inf})$$
 (10)

- A bootstrap technique was used to sample 365 random P_{inf} values for each iteration
- calculating the annual probability (Equation 10). This is necessary since the daily probability
- of infection is not constant for one year. P_{annual} was used to calculate the QALYs lost (QALYs,
- 282 *QALYs lost per person per year*):

$$QALYs = P_{annual} \cdot Q \tag{11}$$

- where Q (QALYs/infection) was the amount of QALYs reported per infection for each
- pathogen. All infections were assumed to result in QALYs. P_{annual} was also used to calculate
- 286 DALYs per person as suggested by the WHO (Havelaar et al. 2000, Kemmeren et al. 2006,
- WHO 2001). The Swedish population age structure of 2010 from the European database (EU
- 288 2010) was used to characterise the drinking water consumer population.
- 289 Three separate probabilities of infection² for the three pathogens were summarised into the
- total probability of infection ($P_{annual tot}$, probability) calculated as:

$$P_{annual_tot} = 1 - \left(1 - P_{annual_noro}\right) \cdot \left(1 - P_{annual_camp}\right) \cdot \left(1 - P_{annual_cryp}\right) (12)$$

- where P_{annual_noro} , P_{annual_camp} and P_{annual_cryp} (probabilities) were the annual probabilities of
- infection due to norovirus, *Campylobacter* and *Cryptosporidium* respectively.
- 294 3.3 Cost-benefit analysis (CBA)
- 295 A CBA was performed to compare the economic negative effects (costs) with the positive
- effects (benefits) for each alternative. All inputs for the CBA are presented in Table 2; input
- 297 distributions represent both epistemic uncertainties and aleatory uncertainties. To enable a

² This implies that the different events are independent. Since pathogens often originate from faecal contamination, one could argue that the presence of one pathogen could increase the probability for the presence of another, resulting in a positive correlation that has not been accounted for.

comparison of the alternatives' societal profitability, the net present value (*NPV*) was calculated and a distributional analysis was performed for each alternative. The *NPV* (*SEK*) was calculated as:

$$NPV = \sum_{t=0}^{T} \frac{(B_t)}{(1+r)^t} - \sum_{t=0}^{T} \frac{(C_t)}{(1+r)^t}$$
 (13)

- where *B* (*SEK*) and *C* (*SEK*) were the benefits and costs for each year *t* during the time

 horizon *T* (*years*); *r* (%) was the discount rate used. *T* was set to 100 years, representing the

 expected life-time of the mitigation alternatives.
- The procedure used in this study for taking into account the project-specific costs and benefits, as well as externalities, follows the basic concept of CBA given by e.g. Boardman et al. (2011), among others. The total annual benefits (B_{tot} , SEK) were calculated as:

$$B_{tot} = B_{health} + B_{environmental} + B_{other}$$
 (14)

- where B_{health} (SEK) were the benefits estimated from reduced negative health effects to drinking water consumers; $B_{environmental}$ (SEK) were the benefits from reduced nitrogen (N) and phosphorous (P) discharge to recipient water bodies due to increased treatment efficiency; and B_{other} (SEK) were other benefits.
- Health benefits (B_{health}) for A1-A4 were calculated as:

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$$B_{health} = \Delta QALYS \cdot QALY_B \cdot DWC_t \tag{15}$$

where $\triangle QALYs$ (QALYs) were the QALYs gained per person in year t in relation to the reference alternative (QALYs A-ref) for each mitigation alternative; $QALY_B$ (SEK/QALY) was the monetary value per QALY; and DWC (persons) was the number of drinking water consumers in year t. The value of a QALY is further discussed in the sensitivity analysis, Section 3.4.

- Environmental benefits for A1-A3 (in A4, environmental benefits were assumed to be zero)
- 321 were calculated as:

$$B_{Environmental} = N \cdot SEK_N + Php \cdot SEK_P$$
 (16)

- where N(kg) was the increased nitrogen removal; $SEK_N(SEK/kg)$ was the monetary value per
- kg nitrogen removed; Php(kg) was the increased phosphorus removal; and $SEK_P(SEK/kg)$
- was the monetary value per kg phosphorous removed.
- Other benefits (B_{other}) were not monetised using quantitative measures. However, to illustrate
- 327 the importance of these benefits, an analysis of how large they need to be to produce a
- positive *NPV* was conducted within the sensitivity analysis.
- 329 Investment costs were added to the first year of the CBA. For A1-A3, the investment costs
- 330 ($C_{Investments}$, SEK) were calculated as:

$$C_{Investments} = C_{WWIP} + C_{Pump} \cdot Pumps + C_{Con_WWIP} \cdot OWISs + C_{Pipe} \cdot WP \quad (17)$$

- where C_{WWTP} (SEK) was the cost for expanding the WWTP; C_{Pump} (SEK) was the cost per
- pump; Pumps (#) was the number of pumps needed; $C_{Con\ WWTP}$ (SEK) was the connection cost
- per OWTS; OWTSs (#) was the number of OWTS connected; C_{Pipe} (SEK/m) was the cost per
- pipe meter; and WP(m) was the pipe length for each alternative.

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Table 2 Inputs for the cost-benefit analysis.

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Input	Unit	Abbr.	Distr. m	Parameters
		General in	put	
Time horizon	у	T	PV	100
Discount rate	%	r	PV	High value=3.5, Low value=1%
OWTSs connected	#	OWTSs	PV	A1=621, A2=1,240, A3=1864, A4=0
Persons per OWTS	#	P_{OWTS}	PV	2.5 (as reported in Table 1)
Wastewater pipe length (A1) ^a	m	WP_{AI}	T	$Min=8*10^3$, $Mode=8.8*10^3$, $Max=9.5*10^3$
Wastewater pipe length (A2) ^a	m	WP_{A2}	T	$Min=1.4*10^4$, $Mode=1.5*10^4$, $Max=1.6*10^4$
Wastewater pipe length (A3) ^a	m	WP_{A3}	T	Min=2.8*10 ⁴ , Mode=3*10 ⁴ , Max=3.2*10 ⁴
Drinking water consumers	#	DWC	PV	330,000
Population increase	#/year	PI	PV	2300
	-	Investment	cost	<u> </u>
Number of pumps required b	Pumps	Pumps	PV	A1=1, A2=10, A3=20, A4=0
Cost for expanding WWTP (A1) ^a	SEK	$C_{WWTP\ AI}$	LN	P05=5*10 ⁶ , P95=6*10 ⁶ , Location=0
Cost for expanding WWTP (A2) ^a	SEK	$C_{WWTP A2}$	LN	P05=10*10 ⁶ , P95=11*10 ⁶ , Location=0
Cost for expanding WWTP (A3) ^a	SEK	$C_{WWTP\ A3}$	LN	P05=1.6*10 ⁷ , P95=1.7*10 ⁷ , Location=0
Cost per pump installation ^b	SEK	C_{Pump}	N	P05=4.5*10 ⁴ , P95=5.5*10 ⁴ , Location=0
Cost per meter pipe ^b	SEK/m	C_{Pipe}	LN	P05=3,150, P95=3,850, Location=0
Cost for installing UV treatm. c	SEK	C_{UV}	LN	P05=5.3*10 ⁷ , P95=5.7*10 ⁷ , Location=0
Cost for connecting to WWTP i	SEK/OWTS	$C_{Con\ WWTP}$	PV	$1.29*10^5$
		Annual co	osts	
Water use per person per day ^e	m ³ /p/y	WU	PV	58.4
Cost per m ³ water used f	SEK/m ³	C_{Water}	PV	26.36
Cost for water services ^f	SEK/y/Con	$C_{Con\ Year}$	PV	2,792
Cost for OWTS maintenance g	SEK/y	C_{Sludge}	PV	1,118
Lifetime of an OWTS h	у	$OWTS_{Life}$	PV	25
Cost for re-investing in OWTS ^b	SEK	$C_{OWTS\ RI}$	PV	$\sim 1.3*10^5$
Cost for UV treatm. maintenance d	SEK/y	$C_{UVmaint.}$	LN	P05=4.2*10 ⁵ , P95=4.4*10 ⁵ , Location=0
	•	Benefit	S	
Benefit/QALY avoided (High) j	SEK/QALY	$QALY_{B\ H}$	PV	$1.22*10^6$
Benefit/QALY avoided (Low) j	SEK/QALY	$\widetilde{Q}ALY_{BL}$	PV	$7*10^5$
Benefit per kg N avoided k	SEK/N	\widetilde{SEK}_N	PV	22.91
Benefit per kg P avoided k	SEK/N	SEK_{P}	PV	53.06
Increase, N removal/year (A1) ¹	kg N/y	N_{AI}	T	Min=3,550, Mode=4,000, Max=4,450
Increase, N removal/year (A2) ¹	kg N/y	N_{A2}	T	Min=7,100, Mode=8,000, Max=7,900
Increase, N removal/year (A3) ¹	kg N/y	N_{A3}	T	Min=10,700, Mode=12,000, Max=13,300
Increase P removal/year (A1) ¹	kg P/y	Php_{AI}	T	Min=580, Mode=650, Max=720
Increase P removal/year (A2) ¹	kg P/y	Php_{A2}	T	Min=1,160, Mode=1,300, Max=1,440
Increase P removal/year (A3) ¹	kg P/y	Php_{A3}	T	Min=1,750, Mode=2,000, Max=2,150

- a) Total wastewater pipe length and cost for expanding the municipal WWTP for each alternative was derived from personal communication with P. Fröjd at Sjöbo municipality and by using expert judgements.
- b) Cost per pump, amount of pumps and the pipe cost per meter were based on Swedish literature (Kärrman et al. 2012).
- c) Cost for installing UV-treatment was based on personal communication with B.M. Pott at Southern Sweden Water Supply (Sydvatten)
- d) Based on the cost per litre for medium sized drinking water treatment plants (Cotton et al. 2001).
- e) Åström and Johansson (2015)
- f) Since stormwater is not included, the fee for connecting OWTS to the municipal WWTP was reduced (Sjöbo Municipality 2016b).
- g) Sludge removal cost (968 SEK/year) (Sjöbo Municipality 2016a) and electricity cost (150 SEK/year) (expert judgement).
- h) Wastewater guide (2016)
- i) Connection fee (101,450 SEK) (Sjöbo Municipality 2016b), application fee (2,550 SEK) (Sjöbo Municipality 2016c), and excavation and plumbing on own property (25,000 SEK) (expert judgement).
- j) Svensson et al. (2015)
- k) SEPA (2008a)
- 1) Based on: 41% non-functioning (zero reduction) OWTSs (SEPA 2004); triangular distributions representing the nitrogen (Min=20, Mode=30, Max=40) and phosphorous (Min=60, Mode=70, Max=80) percentage removal in OWTSs (SEPA 2015); point values estimating the nitrogen (70) (SEPA 2017) and phosphorous (96) (SEPA 2013) percentage removal in WWTP. Triangular distributions were derived using MC simulations.
- m) PV=point value, T=triangular distribution, LN=LogNormal distribution, N=Normal distribution

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- Annual costs (ΔC_{Annual} , *SEK*) for A1-A3 when connected to the municipal WWTP in relation to having an OWTS were calculated as:
- $\Delta C_{Annual} = C_{Annual \ WWIS} C_{Annual \ OWIS}$ (18)
- where $C_{Annual\ WWTP}$ (SEK) was the annual cost per property when connected to the municipal
- WWTP; and C_{Annual_OWTS} (SEK) was the annual cost per property when having an OWTS.
- 365 The $C_{Annual\ WWTP}$ was calculated as:

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$$C_{Annual\ WWIP} = C_{Water} \cdot WU \cdot P_{OWIS} \cdot OWISs + C_{Con\ Year} \cdot OWISs$$
 (19)

- where C_{Water} (SEK/ m^3) was the cost for water use; WU ($m^3/p/year$) was the water use per
- person and year; P_{OWTS} (persons) was the number of persons per OWTS; OWTSs (#) was the
- number of OWTSs connected to the WWTP; and $C_{Con\ Year}$ (SEK/year/OWTS) was the annual
- 370 connection fee per OWTS per year.
- 371 The $C_{Annual\ OWTS}$ was calculated as:

$$C_{Annual_OWTS} = C_{Sludge} \cdot OWTSs + \frac{C_{Reinv} \cdot OWTSs}{OWTS_{Life}}$$
 (20)

- where C_{Sludge} (SEK/OWTS) was the annual cost for sludge removal per OWTS per year; C_{Reinv}
- 374 (SEK) was the cost for re-investing in a new OWTS; and OWTS_{Life} (years) was the expected
- 375 life time of an OWTS.
- For A4, the investment cost was the installation of UV treatment (C_{UV} , SEK), and the annual
- 377 cost was the maintenance of the UV treatment ($C_{UVmaint}$, SEK).
- A distributional analysis was performed by assigning costs and benefits to private OWTS
- owners, drinking water consumers, or inhabitants/visitors of the catchment of Lake Vomb.

3.4 Uncertainty and sensitivity analysis

To acknowledge epistemic and aleatory uncertainties, MC simulations were used in the QMRA and the CBA calculations. The model was divided into the following compartments: source characterisation; water quality modelling in the DWTP; dose-response in the QMRA; and the CBA. An adaptation of the local sensitivity analysis, which investigates the change in output by varying one input variable at a time, keeping all other input variables constant, as suggested by Schijven et al. (2013), was used. For the compartments with monotonic behaviour i.e. source characterisation, water quality modelling in the DWTP, and the CBA, the Spearman's rank correlation analysis was used (Mokhtari and Frey 2005). For the compartment with non-monotonic behaviour, i.e. dose-response in QMRA, scatter plots were used (Frey and Patil 2002). All results from the uncertainty and sensitivity analysis are reported in the supplementary material.

For investigating the uncertainties not suitable to model using probability distributions, scenarios were defined. Scenarios were used for the different values of a QALY, discount

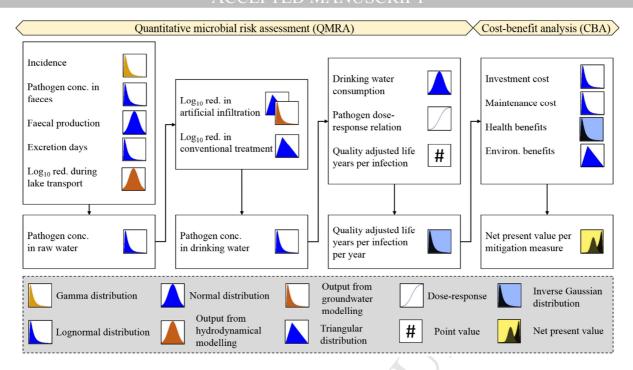
scenarios were defined. Scenarios were used for the different values of a QALY, discount rate, and the proportion of OWTSs contribution to the total pathogen load. The low and high values of a QALY were assumed to be 700,000 and 1,220,000 SEK based on the government implied willingness to pay for a QALY (Svensson et al. 2015). For the discount rate, two scenarios were used: 1 % or 3.5 %. The Swedish Transport Administration recommends 3.5 % for infrastructure projects in the traffic sector (STA 2016); this value is commonly used in other sectors as well. Since the time horizon was long and decisions affect several future generations, a low discount rate was also used. The pathogen load to Lake Vomb from OWTSs (L_{OWTS} , pathogens/day) was calculated as:

$$L_{OWTS} = C_{Tributary} \cdot F_{Tributary} \tag{21}$$

- The total pathogen load (L_{total} , pathogens/day) originates from many sources (e.g. WWTP,
- 404 fertilisation using manure, grazing animals, wild animals) and was illustrated as:

$$L_{total} = L_{owts} + L_{other}$$
 (22)

- where L_{other} (pathogen/day) was the load from all other pathogen sources. Since the OWTSs
- load to Lake Vomb in relation to the total pathogen load was unknown, two scenarios (L_{OWTS}
- 408 = 75 % and L_{OWTS} = 50 % of L_{total}) were investigated.
- To estimate in what range the non-monetised benefits in A1-A3 would have to be in order to
- render a positive NPV, a calculation of non-monetised benefits to reach break-even ($NPV \ge 0$)
- 411 was performed.
- 412 3.5 Software
- 413 For the source characterisation and CBA calculations, the MC simulations were performed
- using *Microsoft Excel*, @*RISK* version 7.5.1. For the drinking water treatment performance,
- virus groundwater transport model and the dose-response relationship, the MC simulations
- were performed using *Analytica* release 4.1.6.30. For the hydrodynamic modelling, the model
- 417 for Lake Vomb was developed using MIKE 3 FM (MIKE Powered by DHI), which is a
- deterministic three-dimensional numerical model that solves the incompressible Reynolds
- 419 averaged Navier-Stokes equations invoking the assumptions of Boussinesq and hydrostatic
- pressure (DHI 2011). The period 2005-2007 was simulated using the observed
- 421 hydrometeorological data.
- 422 Uncertainties were propagated between the different model compartments to calculate the
- probability distributions of the final results of the CBA. Using 10,000 MC iterations, the
- resulting probability distribution of the output of one model compartment was then used as an
- input in the next model compartment. The propagation of uncertainties and the combination
- of methods are illustrated in Figure 3.



428 Figure 3 Schematic illustration of how the different methods are combined in the model.

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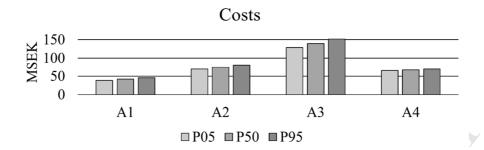
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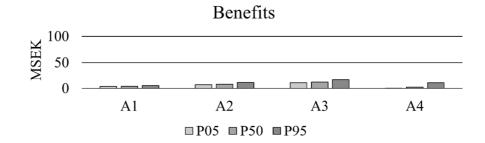
4 Results

- All results, unless stated otherwise, are from the scenario assuming that OWTSs are contributing 75 % of the total pathogen load to Lake Vomb. The complete results from the source characterisation (Table S1), dose-response (Table S2), CBA (Table S3), and uncertainty and sensitivity analysis (Tables S4-S8, Figures S1-S5) are presented in the supplementary material.
- 4.1 Cost-benefit analysis
- The costs, benefits and *NPV* for the scenario with a high value (1,220,000 SEK) of a QALY
- and a discount rate of 3.5 % are presented (Figure 4) for the 5th, 50th, and 95th percentiles.

a)



b)



c)

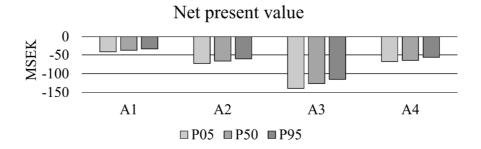


Figure 4. The costs (a), benefits (b) and NPV (c) for the scenario with the high value of a QALY (1,220,000 SEK) and the discount rate of 3.5 % are presented for the 5th, 50th, and 95th percentiles in million SEK (MSEK).

For A1-A3, the costs would be solely taken by the owner of the OWTS that will be connected to the WWTP (installation of pipes on their own property and a connection fee). For A4, the costs would be solely taken by the drinking water producers initially and eventually by the drinking water consumers through a higher drinking water fee.

447	The benefits were distributed between the drinking water consumers (health benefits) and the
448	inhabitants and visitors of the catchment area of Lake Vomb (environmental benefits). For
449	A1-A3, the drinking water consumers received 11 or 18 % and the inhabitants and visitors
450	received 89 or 82 % of the benefits, using a low or high valuation of a QALY, respectively.
451	For A4, the benefits were solely attributed to the drinking water consumers.
452	4.2 Uncertainty and sensitivity analysis
453	For A1-A3 using 3.5 % discount rate, the NPV sensitivity to the inputs in the CBA were (top
454	three in descending order): cost per pipe meter (C_{Pipe}), wastewater pipe length (WP), and
455	$\Delta QALYs$ (A1-A3). For A4 using 3.5 % discount rate, the NPV sensitivity to the inputs in the
456	CBA were (top three in descending order): $\Delta QALYs$ (A4), cost for installation of UV
457	treatment (C_{UV}) and cost for maintenance of UV treatment ($C_{UVmaint}$).
458	The concentration of norovirus in raw water (C_{RW_Noro}) was the most sensitive to the following
459	inputs (top three in descending order): concentration in faeces (C), incidence (I_{Noro}), and days
460	excreting (D_{Noro}) . The concentration of <i>Campylobacter</i> in raw water (C_{RW_Camp}) was the most
461	sensitive to the following inputs (top three in descending order): concentration in faeces (C) ,
462	Log_{10} reduction in Björkaån (R_{Bjork_Camp}), and Log_{10} reduction in Torpsbäcken (R_{Torp_Camp}).
463	The concentration of $Cryptosporidium$ in raw water (C_{RW_Cryp}) was the most sensitive to the
464	following inputs (top three in descending order): concentration in faeces (C), days excreting
465	(D_{Cryp}) , and incidence (I_{Cryp}) .
466	The concentration of norovirus in drinking water (C_{DW_Noro}) was the most sensitive to the
467	foellowing inputs (top three in descending order): Log ₁₀ reduction in saturated zone (R_{SZ_noro}),
468	raw water concentration (C_{RW_Noro}), and Log ₁₀ reduction per meter in unsaturated zone
469	(R_{UZ/m_Noro}) . The concentrations of Campylobacter (C_{DW_Camp}) and Cryptosporidium (C_{DW_Cryp})
470	in drinking water were the most sensitive to the following inputs (top three in descending

471	order): Log ₁₀ reduction per meter in saturated zone (R_{SZ_Camp} and R_{SZ_Crypp}), saturated zone
472	vertical length (L_{SZ}), and raw water concentration (C_{RW_Camp} and C_{RW_Cryp}).
473	The dose-response relationship between the concentrations of pathogens in drinking water
474	(C_{DW}) , infectivity (m) , drinking water consumption (WI) and the probability of infection (P_{inf})
475	was illustrated using scatter plots from the 10,000 MC simulations.
476	A scenario-based analysis was performed to analyse the effects on the final NPVs from
477	uncertainties regarding the QALY valuation, discount rate, and the OWTSs contribution to
478	the total pathogen load. The rank order of the 50^{th} percentiles for the NPV (A1>A4>A2>A3)
479	does not change depending on the level of OWTSs contribution to the total pathogen load nor
480	the QALY valuation. However, with a low discount rate (1 %), the rank order changes to
481	A1>A2>A4>A3.
482	Benefits that have not yet been monetised and included in the CBA that might alter the rank
483	order of NPV for the alternatives were identified. For alternatives A1-A3, non-monetised
484	benefits are:
485	• positive health effects for humans from improved water quality for recreational
486	activities in Lake Vomb;
487	• positive health effect for animals (both domestic and wild) from improved water
488	quality in the catchment and in Lake Vomb;
489	• perceived value for private OWTS owners not being responsible for treating their
490	wastewater;
491	• increased market value of the properties connected to the municipal water and
492	wastewater system;
493	• benefits of removing the possible risk of direct contamination of private wells by
494	OWTSs;

495	• possibility to recycle nutrients when wastewater is treated at the wwip;
496	• reduction of CO ₂ emission when sludge transportation trucks do not need to empty
497	closed tanks and three compartment septic tanks;
498	• reduced traffic accidents and related risks since heavy traffic is reduced in the
499	catchment area.
500	For alternative A4, non-monetised benefits are:
501	 less disinfection by-products due to lower dosage in chlorination;
502	 reduced handling and storage of chlorination chemicals.
503	For A1-A3, these additional benefits need to be 800-1200 SEK or 1800-2400 SEK per OWTS
504	per year for 1% and 3.5% discount rate, respectively, to give a positive NPV (50 th percentile).
505	These ranges apply for both the high and low valuation of a QALY.
506	5 Discussion
507	The aim of this study was to present an approach for comprehensive decision analysis using
508	CBA of microbial risk mitigation measures in DWSs, and including Lake Vomb as a case to
509	illustrate the assumptions needed and the associated variabilities and uncertainties. Below we
510	discuss the QMRA, the CBA, the uncertainties, and the overall applicability of the decision
511	model.
512	5.1 Quantitative microbial risk assessment
513	5.1.1 Source characterisation
514	Pathogen concentration in faeces and the pathogen excretion duration are subject to large
515	variability. In this study, it was assumed that the catchment was large enough to have
516	pathogens present at all times, evenly distributed geographically. However, if a smaller
517	catchment is to be described, it will be important to account for temporal and geographical

518	variations of the pathogen prevalence. If persons are infected, there will be high pathogen
519	concentrations in the OWTSs effluents, otherwise there will be no pathogens present. One
520	way forward is to combine the data on incidence with binary probability density functions.
521	This would capture the on/off characteristics of infections and enable the use of the decision
522	model on smaller systems, even on a single OWTS.
523	To acknowledge the ambiguity (a factor of 100 between values) and the lack of information in
524	the underreporting factor, it was assigned a point value and not included in the MC
525	simulations. However, the factor for underreporting is uncertain, and further investigations on
526	how to describe this input need to be conducted. The factor for underreporting is important,
527	since increased underreporting results in a corresponding increase of the estimation of the
528	pathogen concentration at the raw water intake.
529	The estimated pathogen load to Lake Vomb can be validated. The estimated concentrations of
530	Cryptosporidium in the tributaries (0.36-1.4 oocysts/L) in this study are in agreement with the
531	values reported by other studies, e.g. the mean of 0.62 oocysts/L in an Australian river
532	(Swaffer et al. 2014). The estimated concentrations could also be validated by monitoring the
533	local pathogen concentrations in the catchment; however, this is tedious and expensive.
534	Instead, based on the factor for underreporting and the incidence of norovirus (since it was the
535	pathogen causing the main part of the loss of QALYs), we made an estimate of the annual
536	infections in the drinking water consumer population, confirming that the waterborne
537	infections only represented a small proportion of the total infections calculated from the
538	incidence. Chosen values and associated probability distributions should be regarded as a
539	possible, but not necessarily the optimal, representation of the pathogen source characteristics.
540	5.1.2 Water quality modelling
541	The log ₁₀ reduction during transport in Lake Vomb was estimated using hydrodynamic
542	modelling encompassing several years of daily and sub-daily variation in

543	hydrometeorological data. Looking at the best fit, a normal distribution was reasonable to use
544	for describing the variability in pathogen reduction. Future development of water quality
545	modelling within the decision model is to include probabilistic modelling. Probabilistic
546	modelling will further facilitate risk-based modelling approaches, QMRA and holistic water
547	resource management (Oliver et al. 2016).
548	The model describing the artificial groundwater recharge system was highly simplified. The
549	pathogen reduction was based on a conceptual model describing the artificial groundwater
550	infiltration as one system, when in reality there are many smaller sub-systems with complex
551	flow and transport conditions between different infiltration ponds and abstraction wells.
552	Nonetheless, the model is assumed to give a good understanding of the key processes
553	affecting the level of reduction in the artificial infiltration.
554	Local investigations of the barrier efficiency at the DWTP would be preferred. Since it is not
555	ethical to use active DWSs to directly test the reduction of pathogens, surrogate organisms
556	can be used instead. It is also possible to use literature estimates. The Log_{10} reduction of
557	Cryptosporidium by the UV-disinfection was not allowed to be higher than 3 Log ₁₀ , in order
558	not to interpret results outside of the investigated range (Hijnen et al. 2006). Although this can
559	be considered a low reduction given the efficiency of UV-disinfection towards
560	Cryptosporidium, this approach is used in the QMRA-tool for drinking water producers in
561	Sweden. However, the truncation in UV-treatment needs to be further investigated and
562	thoroughly reviewed. Investigation of altering the UV-dose may also be of importance for
563	future implementation of the decision model.
564	5.1.3 Dose-response
565	The estimated annual probability of infection (P_{annual}) was slightly higher than the WHO
566	guidelines in the current situation (A-ref, 50 th percentile), while the <i>DALYs</i> were under the
567	threshold (A-ref, 5 th percentile), indicating that there is ambiguity whether the microbial risks

568	were acceptable or not. The large uncertainty and variability described in input probability
569	distributions should be taken under consideration when interpreting the results. To get below
570	the WHO guideline for P_{annual} looking at the 50 th percentile, A2, A3 and A4 are the possible
571	options, while A1 almost reaches the threshold. It is only A4 that meets the guideline level
572	with respect to the 95 th percentile. Even though no strict guideline level exists in Sweden,
573	drinking water producers should be aware of the discrepancy between meeting the DALY or
574	P_{annual} WHO guideline. The same pathogen concentration in drinking water can meet one
575	target and miss the other.
576	P_{annual} was calculated into to QALYs using standard unit values adopted from a study from
577	the U.S (Batz et al. 2014). It may be argued that results from the U.S. are not representative
578	for Swedish settings. Even so, to illustrate the methodology, it was assumed that the U.S.
579	values would be useful. However, further development of the model could use more detailed
580	health effect quantification and implement local studies for estimating the quantity of the
581	health risk reduction in the risk mitigation alternatives. To monetise the health effects, there
582	are other approaches which can be implemented into the model, e.g. information from
583	previous events, quality of life investigations, etc.
584	5.2 Cost-benefit analysis
585	None of the mitigation alternative rendered a positive NPV. However, the NPV results must
586	be interpreted using a wider perspective in combination with other results from the CBA, such
587	as distributional analysis and non-monetised benefits. From a socio-economic perspective, it
588	is important to identify the alternative with the least negative NPV (A1). In a situation where
589	decision makers are required to reduce the microbial risk, they will need to choose an
590	alternative. Such a situation would occur e.g. if there is a guideline or risk level that needs to
591	be achieved, such as the WHO recommendation of a maximum yearly probability of infection
592	per person of 10 ⁻⁴ or a maximum DALY of 10 ⁻⁶ (WHO 2011). If looking at the 50 th percentile

593	with 1 % discount rate, both A1 and A2 resulted in higher NPVs than A4. Nevertheless, as
594	noted above, only A4 would achieve the WHO recommendation of the P_{annual} with a high
595	degree of certainty (looking at the 95 th percentile).
596	When monetising health benefits, it is important to make sure that the underlying valuation
597	study represents the relevant health effects. The monetisation of health benefits was based on
598	a governmental implied willingness to pay for a QALY (Svensson et al. 2015). The values
599	used were estimated from a societal perspective, i.e. the effects both within the health care
600	sector (e.g. reduced medical and hospitalisation costs) and beyond the health care sector (e.g.
601	reduced discomfort from being ill and loss of production) were accounted for.
602	When decision makers choose an alternative, they also accept the distribution between
603	beneficiaries and payers associated with the decision. Even though the Kaldor-Hicks
604	criterion ³ can be argued, the distribution of the costs and benefits will need to be
605	communicated with stakeholders. Alternative A4 is the only alternative when the beneficiary
606	and the payer are the same stakeholder. In decision making, distributional analysis can be of
607	importance when applying the polluter pays principle.
608	5.3 Uncertainty and sensitivity analysis
609	The Spearman's rank correlation is inadequate (Ellouze et al. 2010) for measuring sensitivity
610	when analysing complex relationships such as the dose-response relation in QMRA. We have
611	used scatter plots to illustrate the relationships between drinking water pathogen
612	concentration, drinking water consumption and the infectivity. Future research needs to
613	investigate more advanced sensitivity analysis methods (see e.g. Mokhtari and Frey 2005).

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³ The Kaldor-Hicks criterion, simply put, state that: beneficiaries can compensate those that pay or experience negative effects. However, the compensation only needs to be possible and not realized, since it is argued that if a decision is societally profitable, the costs and benefits will eventually even out with regard to different stakeholders.

614	Since the total risk level in the drinking water system cannot be estimated, it was important to
615	investigate whether the results change if the OWTSs contribution to the total risk is altered.
616	Results showed that changing the OWTSs contribution to the total pathogen load did not
617	change the ranking of the alternatives.
618	After scrutinising the CBA results, decision-makers need to consider benefits that were
619	omitted from the monetised analysis. Even though the NPVs were negative for the A1-A3
620	alternatives, all alternatives could render a positive NPV (looking at the 50 th percentile) if
621	these other benefits could be valued in the range of 800-2400 SEK per connected OWTS per
622	year. The value of the benefits when using a 3.5 % discount rate need to be approximately
623	1000 SEK higher than when using a 1 % discount rate.
624	Some factors vary over time both within a year, e.g. incidence and water flow etc., and over
625	longer time periods, e.g. population increase, climate change etc., to mention a few. The
626	model included a population increase based on population projections for Sweden in general.
627	However, the inter-yearly variations have not been included. For further development of the
628	decision model, methods for including these temporal variations and uncertainties need to be
629	developed.
630	5.4 Risk-based decision model
631	Depending on the type of decision and the local settings, other methods than presented in this
632	paper can be more suitable to combine in the decision model. For decisions aiming at
633	reaching a certain guideline or threshold value, a CEA may be preferred, instead of a CBA.
634	CBA represents a strict anthropocentric and utilitarian context, only accounting for benefits
635	attributed to human values (Hutton 2001). If decision makers want to include intrinsic values,
636	they need to apply methods that can consider such values as well, such as multi-criteria
637	decision analysis (see e.g. DCLG 2009). In such multi-criteria decision models, the decision

support rendered from the CBA and the QMRA can be used as input for appropriate criteria. 638 To give some examples, the NPV can provide information to the economic dimension, 639 distributional analysis and QALY assessment can provide input to the social dimension, and 640 water quality modelling can provide input to the environmental dimension in a sustainability 641 assessment, see e.g. Rosén et al. (2015). 642 The focus of this study was to describe the methodology of comparing microbial risk 643 644 mitigation measures using CBA in combination with QMRA to estimate risk levels and the effect of possible mitigation measures. Benefits, in terms of the health risk reduction obtained 645 in each alternative were described in detail. Environmental benefits were included using a 646 647 more simplified approach. However, including the environmental benefits illustrates a key element of the CBA, i.e. the possibility to include other benefits, apart from the target risk 648 reduction. These additional benefits may be of substantial importance and heavily affect the 649 final decision. 650 The decision model incorporates both aleatory and epistemic uncertainties in the input 651 probability distributions. To further develop the model and to provide additional decision 652 support, these uncertainties can be divided. This separation would also facilitate additional 653 decision analyses, e.g. value of information analysis. 654 6 Conclusions 655 Results from the case study showed that the alternative to connect the smallest proportion 656 (25 %) of on-site wastewater treatment systems to the wastewater treatment plant (A1) at 657 Lake Vomb was the most societally beneficial. However, the only alternative that would 658 reduce the annual probability of infection to meet the WHO guidelines with a high degree of 659 certainty (95th percentile) was installing UV-disinfection (A4). In relation to the development 660 of the risk-based decision model, the following conclusions were drawn: 661

- The developed decision model is flexible and can be tailored to different drinking water systems and different types of decision problems.
 - To implement the decision model, a multitude of uncertainties and variabilities needs to be addressed. However, the model provides tools to include these variabilities and uncertainties in a structured manner.
 - Through the process of performing the cost-benefit analysis, aspects important for decision making that may otherwise easily be overlooked or ignored are openly displayed and assessed.
 - The combination of quantitative microbial risk assessment and cost-benefit analysis
 provides a novel decision model that creates transparent and holistic decision support
 tool for microbial risk mitigation.
 - For improvement of the decision model, we suggest to further develop the valuation and monetisation of health effects and the propagation of variability and uncertainty between the included methods.

Acknowledgement

This research has been performed as part of the project *Risk-Based Decision Support for Safe Drinking Water*, financed by the Swedish Water and Wastewater Association (project 13-102), and as part of DRICKS, a framework programme for drinking water research at the Chalmers University of Technology. The information provided by the local water utility (Southern Sweden Water Supply) and the municipality of Sjöbo was most appreciated. Input regarding the sensitivity analysis, provided from Tommy Norberg was highly appreciated.

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- A decision model for drinking water context, combining QMRA and CBA, was developed.
- This flexible model can be tailored to different systems and decision problems.
- The microbial risk mitigation measures were compared in a Swedish case-study.
- Microbial risk reduction was measured in QALYs and monetised.
- This novel decision model provides transparent and holistic decision support.

