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Evaluation of a cross contamination model describing transfer of Salmonella spp. and Listeria monocytogenes during grinding of pork and beef



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Can a promising cross contamination model be successfully applied to any condition of meat grinding? To answer this question we performed different challenge tests and developed a set of evaluation approaches.

The cross contamination model (Møller et al. 2012) was evaluated to investigate its capability of describing transfer of *Salmonella* spp. and *L. monocytogenes* during grinding of pork and beef of varying sizes (50 – 324 g) and numbers of pieces to be ground (10 – 100), in two grinder systems.



Data from 19 trials were collected. Three different evaluation approaches were applied:

- ASZ an Acceptable Simulation Zone method which compared observed transfer with simulations using the proposed model
- ii) QMRA each trial was fitted to equation 1 and its respective parameter estimates were integrated in a Quantitative Microbiological Risk Assessment model (Møller et al. 2015) to compare risk estimates
- iii) TTP the Total Transfer Potential was calculated for each of the trials based on fitted parameter estimates

Møller et al. (2012) model $M_i = (1-a_1)(1-a_2)(1-c_2) P_i + (b_1 \operatorname{gr}_{1,i-1}) + (b_2 \operatorname{gr}_{2,i-1})$

Lessons learned:

- Results indicated that transfer estimates were not applicable for unlike processing
- QMRA risk estimates and TTP both revealed that risk attribution from grinding was influenced by:
 - 1) sharpness of grinder knife
 - 2) specific grinder
 - 3) grinding temperature
 - 4) specific pathogen was of minor importance

Explaining	Møller	et al.	(2012)	model	
	SOME MIDE				

	a1-transfer from piece of meat to E1
1	a2 - transfer from piece of meat to E2





		Classificatio	on of trials according agre	eement of results	obtained	l with three d	ifferent approaches		
		for evaluatin	ng performance of the monst during meat grinding in	odel proposed by n trials with at leas	Møller et st 15 piec	al. (2012) decess of meat.	escribing the transfer		
	DECISION MAKING WHEN:	Trial Si	rial Size of ASZ ^b to include entification 95 % of the predictions (± CFU/portion of meat)	Absolute Risk ^c Estimates x 10 ⁻³	TTP % ^d	Møller et al. (2012) model		A positive correlation was found	
	at least 2 evaluation approaches are IN AGREEMENT	identification				Applicability	Evaluation approaches in agreement	between QMRA risk estimat and TTP %	
	9.0 8.5 8.0 7.5 7.0 6.5	1	2.0	3.02	103	-	3		
		2	2.0	2.87	101		3	Evaluated parar estimates ar APPLICABI	
lioi		3	2.0	2.94	100	-	3		Evaluated parameter
U/port		4	0.8	1.01	21	+	3		estimates are
g10 CFL		5	1.4	1.56	36 🤇	+	2		APPLICABLE
evel (lo	6.0	6	1.2	1.10	24	+	2		(58 % of trials)
ogen	5.5 5.0 4.5 4.0 0 10 20 30 40	7	1.1	0.86	16	+	2		
4.5 4.0 0		8	1.2	0.62	11	+	2		
		9	2.0	2.21	60	-	3		
	number of portion	11	1.1	1.92	51	-	3		Evaluated parameter
_	12	12	1.4	3.04	102		3	>	
8.5 (hotion) (0.5	8.5 Most observations outside	13	1.2	1.44	35	+	2		(42 % of trials)
	ASZ = $\pm 1.0 \log_{10}$ CFU per portion	14	1.5	2.22	64	-	3		

1.54

2.24

1.59

1.41

1.14

0.96





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^b ASZ – Acceptable Simulation Zone.

1.1

1.1

2.2

0.6

0.6

0.5

15

16

17

18

19

20^a

^c Risk estimates from scenarios testing different sets of transfer parameter estimates, and using the QMRA of *Salmonella* in meatball processing model (Møller et al., 2015) at low concentration and prevalence of the pathogen.

^d Calculated with base on Equation 1. It indicates the percentage (%) of CFU of *Salmonella*, from the contaminated pieces of meat that ends up in the total portion of ground meat, assuming that the grinding process will continue forever.





