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The challenge of tackling *Campylobacter* in Belgium



May 7th 2014

DG SANCO workshop on the control of *Campylobacter* in poultry

Isabel De Boosere



Content

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- Background
- National risk assessment
- National legislation on microbiological criteria for *Campylobacter*
- Further research
- Future perspectives

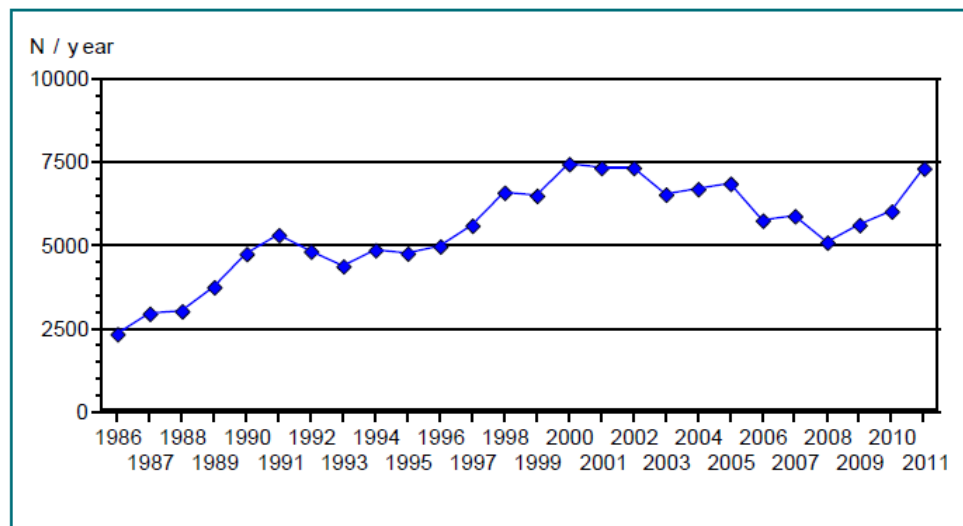


Background

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- *Campylobacter* is most commonly reported gastrointestinal bacterial pathogen in humans in Belgium since 2005.
- Monitored by the FASFC since 2000. The incidence of positive poultry samples is high and remains stable.



Trends & Sources
2010-2011, FASFC

Number of Campylobacter infections in humans by year

(1986-2011). Source: Sentinel Laboratory Network

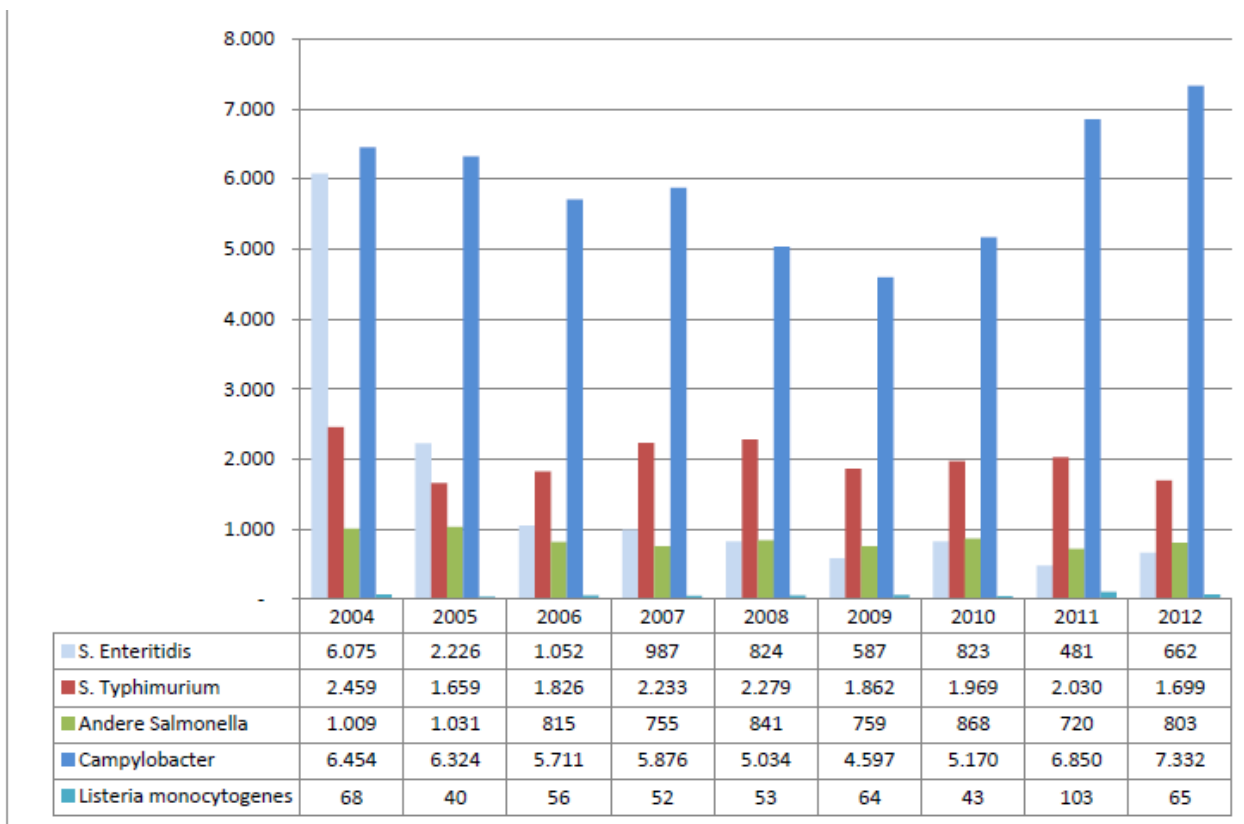


Background

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Number of reported human cases of the 3 most important food bacterial pathogens (source: IPH)



Annual report
FASFC, 2012



Background

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European legislation



- General Food Law, R 178/2002
- Since 2004: European hygiene package (R 852/2004 & R 853/2004) & R 2073/2005 → *necessity to review national legislation*

Belgian legislation



- Law of 24 Jan 1977 concerning the health protection of consumers regarding food and other products

Art. 5 obligation of risk assessment by Superior health Council before setting legislation on contaminants



National risk assessment

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Conseil Supérieur d'Hygiène  Hoge Gezondheidsraad

Advies van de Hoge Gezondheidsraad
April 2005

BIJDRAGE TOT EEN RISICOBEOORDELING
VAN *CAMPYLOBACTER* SPECIES IN VLEESBEREIDINGEN
OP BASIS VAN PLUIMVEEGEHAKT IN BELGIË

HGR n° 7947

Contribution to a risk
assessment - *Campylobacter*
spp. in meat preparations on
the basis of poultry minced
meat in Belgium

<http://www.gezondheid.belgie.be/portal/Aboutus/relatedinstitutions/SuperiorHealthCouncil/publications/4492397?ie2Term=&ie2section=9744>



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National risk assessment



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Why MC at retail level?

Primary production

Biosecurity on farms: no guarantee, difficult to maintain during long period

Vaccination, probiotics: no options

Research was initiated

Slaughter

Drastic reduction is not possible, only limited reduction

Retail

Often contaminated (67 %). Contamination levels are mostly unknown. Estimated that 9 % is contaminated with > 100 cfu/g



National risk assessment



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Preliminary probabilistic approach

6 scenario's, e.g.

Scenario 1 (estimated current situation)

9 % > 100/g , 24 % > 1/25 g & 67 % < 1/25 g

Scenario 3

0 % > 100/g , 9% > 10/g , 24 % > 1/25 g and < 10/g , & 67 % < 1/25g

Scenario 5

0 % > 10/g , 9% > 1/g , 24 % > 1/25 g and <1/g , & 67 % < 1/25g



National risk assessment



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Preliminary probabilistic approach

	Normal distribution of the contamination level (%)				Mean	Maximal probability of infection			
	> 1/g	>10/g	>100/g	>1000/g		P50	P75	P95	Max
Scenario 1	19.99	12.66	7.45	4.06	1.33 e-02	4.17 e-08	9.28 e-06	2.16 e-02	0.942
Scenario 3	11.47	5.15	1.97	0.64	2.22 e-03 : 6	4.41 e-08 =	2.06 e-06 : 5	6.27 e-04 : 35	0.861 : 1.1
Scenario 5	6.3	1.89	0.44	0.08	4.49 e-04 : 30	4.09 e-08 =	9.79 e-07 : 10	1.31 e-04 : 165	0.392 : 2.4



National risk assessment



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Quantitative risk assessment of *Campylobacter* spp. in poultry based meat preparations as one of the factors to support the development of risk-based microbiological criteria in Belgium

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National risk assessment



Table 5
Overview of the results (exposure, probability of infection, % infected) for the different tested situations

Situation	Exposure (cfu per 100 g serving)			Approach 2 (probability of infection)			Approach 3 (% infected)
	Mean	95% percentile	100% percentile	Mean	95% percentile	100% percentile	
1 ^a	2.02E+ 07	7.75E- 01	1.63E+ 13	2.38E- 03	7.55E- 05	3.66E- 01	0.0353
1 ^a (raw)	1.45E+ 10 (sit 1 × 718) ^b	4.35E+ 04 (sit 1 × 56180)	1.30E+ 16 (sit 1 × 802)	4.98E- 02 (sit 1 × 21)	4.14E- 01 (sit 1 × 548)		1.0155 (sit 1 × 29)
2	1.83E+ 05 (sit 1:110)	2.63E- 01 (sit 1:3)	1.35E+ 11 (sit 1:120)	1.38E- 03 (sit 1:2)	2.74E- 05 (sit 1:3)	3.55E- 01 (sit 1:1)	0.0089 (sit 1:4)
3	1.77E+ 03 (sit 1:11390)	9.47E- 02 (sit 1:8)	1.12E+ 09 (sit 1:14469)	6.72E- 04 (sit 1:4)	1.07E- 05 (sit 1:7)	3.38E- 01 (sit 1:1)	0.0016 (sit 1:22)
4	1.98E+ 01 (sit 1:1.0 × 10 ⁶)	3.70E- 02 (sit 1:21)	9.35E+ 06 (sit 1:1.7 × 10 ⁶)	2.42E- 04 (sit 1:10)	4.42E- 06 (sit 1:17)	3.16E- 01 (sit 1:1)	0.0003 (sit 1 : 118)
5	3.26E- 01 (sit 1:6.2 × 10 ⁷)	1.62E- 02 (sit 1:48)	7.77E+ 04 (sit 1:2.1 × 10 ⁸)	5.50E- 05 (sit 1:43)	2.00E- 06 (sit 1:38)	2.87E- 01 (sit 1:1)	0
6	1.23E- 02 (sit 1:1.6 × 10 ⁹)	8.11E- 03 (sit 1:95)	6.46E+ 02 (sit 1:2.5 × 10 ¹⁰)	6.33E- 06 (sit 1:376)	9.76E- 07 (sit 1:77)	1.78E- 01 (sit 1:2)	0
7	1.63E- 03 (sit 1:1.2 × 10 ¹⁰)	4.93E- 03 (sit 1:157)	8.67E+ 00 (sit 1:1.9 × 10 ¹²)	6.75E- 07 (sit 1:3525)	5.11E- 07 (sit 1:148)	6.91E- 03 (sit 1:53)	0
8	2.02E+ 06 (sit 1:10)	7.77E- 02 (sit 1:10)	1.63E+ 12 (sit 1:10)	9.32E- 04 (sit 1:3)	5.62E- 06 (sit 1:13)	2.73E- 01 (sit 1:1)	0.0143 (sit 1:2)
9	1.84E+ 04 (sit 1:1098)	2.63E- 02 (sit 1:29)	1.36E+ 10 (sit 1:1200)	4.44E- 04 (sit 1:5)	1.92E- 06 (sit 1:39)	2.48E- 01 (sit 1:1)	0.0024 (sit 1:15)
10	1.78E+ 02 (sit 1:113603)	9.49E- 03 (sit 1:82)	1.13E+ 08 (sit 1:144312)	1.60E- 04 (sit 1:15)	6.88E- 07 (sit 1:110)	2.18E- 01 (sit 1:2)	0.0004 (sit 1:88)

(raw) Indicates raw consumption of the product (no effect of cross-contamination or cooking included in the model).

^a Situation 1 is the original situation in Belgium with regard to the distribution of the *Campylobacter* contamination level (19.68% > 1 cfu/g; 12.44% > 10 cfu/g; 7.28% > 100 cfu/g; 5% > 1000 cfu/g).

^b (sit 1 × 718) indicates that the exposure is 718 times higher for sit 1 (raw) than for sit 1.

National risk assessment



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- Risk of human infection and disease ↓ when level of *Campylobacter* are better controlled and presence of high contamination levels is limited
- When elimination of preparations with $> 1000/g$ ($<1\%$) and reduction of $> 100/g$ (max. 2%) & $> 10/g$ (max. 5%)
→ reduction of probability of infection by a factor 6
- When elimination of preparations with $> 100/g$ ($<1\%$) and reduction of $> 10/g$ (max. 2%)
→ reduction of probability of infection by a factor 30
- Communication needed to point out hazards of consumption of raw meat and necessity to heat thoroughly



National legislation on microbiological criteria for *Campylobacter*

RD of 26 April 2009



Annexe à l'arrêté royal du 26 avril 2009 concernant des critères microbiologiques applicables aux denrées alimentaires

Critères d'hygiène des procédés

Catégorie de denrées alimentaires	Micro-organisme/ Métabolite	Limites (1)		Plan d'échantillonnage (2)		Point d'application du critère	Actions correctives
		m	M	n	c		
1. Viandes hachées de volaille et préparations de viande à base de viande de volailles, destinées à être consommées cuites (3)	<i>Campylobacter</i> spp. thermotolérants	100 ufc/g		5	0	Fin du processus de production	Améliorations de l'hygiène de la production, de la sélection et/ou de l'origine des matières premières



Further research



- EFSA's analysis of the baseline survey (2008), conclusions:
“These findings indicate that certain slaughterhouses are more capable than others in preventing Campylobacter contamination and in controlling the contamination and/or the Campylobacter counts on the carcasses. This implies that slaughterhouse processing offers an opportunity for Campylobacter risk mitigation.”
- Analysis of Belgian data **confirmed** the EFSA observation



CAMPYVAR, CAMPYTRACE



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Campylobacter contamination in broiler carcasses and correlation with slaughterhouses operational hygiene inspection

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Further research: CAMPYVAR & CAMPYTRACE

Some preliminary results

Slaughterhouses operate high risk raw material - almost 60 % of batches are *Campylobacter* positive and usually broilers are colonised with high numbers ($> 7,5 \log \text{ cfu/g}$)

Both *Campylobacter* colonisation level in the caecal content and especially the carriage of *Campylobacter* on feathers differs between batches.

Breast skin can be highly contaminated with *Campylobacter*.

Campylobacter contamination on feathers and on breast skin mostly increased significantly during transport and holding time.



Further research: CAMPYVAR & CAMPYTRACE

High variability in *Campylobacter* carcass contamination
within batches
between batches in slaughterhouse
between slaughterhouses

High risk material - *Campylobacter* colonisation level in the caecal content
and the carriage of *Campylobacter* on feathers

Certain slaughterhouses are able to produce lower numbers of highly
contaminated carcasses than others

Campylobacter contamination is mainly influenced by the following
processes:

Plucking and evisceration ↑

Washing and chilling (combined effect) ↓ (BUT water immersion)



Further research: CAMPYVAR & CAMPYTRACE

If only *Campylobacter* negative batches are slaughtered: non-contaminated carcasses (i.e. no enumerable levels of > 10 cfu/g)

The slaughter of positive batches results in immediate contamination of carcasses across the slaughter line.

When only positive flocks are slaughtered, *Campylobacter* carcass contamination remains at the same level during the process day.

Campylobacter is transmitted from a positive to a subsequent negative batch, but the transmission is restricted and decreases quickly to non-enumerable numbers over time.

If the preceding positive batch is colonised at a low level no carcass contamination occurs in the following negative batch.



Further research



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This resulted in a change of the action limits for broiler & laying hen carcasses & fresh meat with skin at slaughterhouse, cutting & processing plants & retail to a level of 1000 cfu/g

[Advice 10-2012](#) The evaluation of the document "Action limits for microbiological contaminants in food" (dossier Sci Com 2011/21)

<http://www.favv.be/thematischepublicaties/actiegrenzenvoormicrobiologischecontaminanteninlevensmiddelen.asp> (NL)

<http://www.favv.be/publicationsthematiques/Limitesdactionpourlescontaminantsmicrobiologiquesdanslesdenreesalimentaires.asp> (FR)



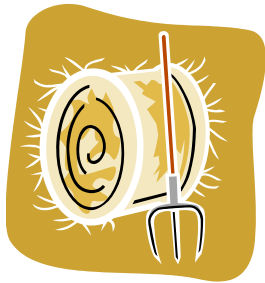
Future perspectives

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Further research at primary production level

- **CAMPYNANOCURE**



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www.health.belgium.be

