EFSA TSE activities 2021-2022

EURL on TSE 19th Annual meeting 17-18 October 2022







CONTENT

Finished:

ABP renewable fuels 2022

Ongoing:

- CWD monitoring
- 2021 EU summary report 2021
- Pig PAP

Pipeline:

Negligible risk classical scrapie CZ





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Pipeline:

- Negligible risk classical scrapie CZ
- Atypical CWD



ABP APPLICATION

multi-step catalytic co-processing hydro-treatment for the production of renewable fuels using **Category 3** animal fat and used cooking oils **BpRR (British Petroleum)**

Animal fat: Standard processing method: Processing method 1 (pressure sterilization) 133°C, 3 bar, 20min Used cooking oil

Catalytic co-processing hydro-treatment using a middle distillate followed by a stripping step. pressure > 60 bars temperature >270°C >4.7 minutes





ABP APPLICATION

Most resistant hazard: hazard identification. Indicator pathogens. Not prions

Approach:

spores from non-pathogen ber 2 bacterial species (*Bacitetibet* an *Desulfotomaculum* geocton) andard appli 28 orming indicator and

Standard appli

reduction tediedi subtilion profishedi br Adoptishedig \sim 5 log₁₀ in the spores of *B*. \log_{10} reduction in the spores of C.





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CWD MONITORING

- Detection of CWD in Norway 2016
- EFSA opinion (I) December 2016: proposal 3-year monitoring programme
 - ToR1: "aimed at detecting CWD and/or estimating the prevalence of CWD in Norway, Sweden, Finland, Iceland, Estonia, Latvia and Poland, which are the EU and EEA countries with reindeer and/or moose populations, depending on the level of prevalence which is wished to be detected".
- Com. Reg. (EU) 2017/1972: Estonia, Finland, Latvia, Lithuania, Poland and Sweden. Compulsory monitoring following EFSA opinion 2018-2020.
- Q4 2017 and Sweden (2021/2022 surveillance)
- Prion protein genotype for positives and negatives (or a sample).



	CWD MONITORING						
		Norway		Sweden	Finland		
		Reindeer	Moose	Red deer	Moose	Moose	
	2016	4	2				
	2017	9	1	1			
	2018	6	1			1	
	2019		2		3		
	2020	1	1		1	1	
	2021		2	1			
	2022	1	2	1			
	TOTAL	21	11	3	4	2	



EFSA is requested to provide a scientific opinion on the monitoring of CWD, based on the **results** of the abovementioned **monitoring programme** including the **statutory** data available in the EFSA database, **and any other monitoring data collected with the same epidemiological objective** and having become available since the publication of previous EFSA opinions on CWD

ToR1

To analyse the results of the monitoring programme carried out in Norway, Sweden, Finland, Iceland, Estonia, Latvia, Lithuania and Poland between 1 September 2017 and 28 February 2022 and, in particular, to assess if the two objectives as set in the 2016 EFSA opinion on CWD in cervids have been met





EFSA proposal (2017) and legal requirements (2001/999). Summary of the country-specific implementation

The analysis of the surveillance data.

Description of general and intensified surveillance Description surveillance data (EFSA database) Evaluation of objectives (representativeness and sensitivity) Description of the outcomes: caseload and prevaler

Description of the outcomes: caseload and prevalence (Ly- & Ly+)

- Estimation of the minimum detectable prevalence (design prevalence)
- Estimation of the sensitivity of the surveillance system





ToR2

- To describe any new knowledge on the epidemiology of CWD in Europe/European Union.
- Available epidemiological knowledge until the last EFSA CWD opinion (2019) Summary previous Opinions (CWD I, II and III) North America vs. Europe
- New epidemiological knowledge since the last EFSA CWD opinion

Mostly based on the Norwegian experience from 2019





ToR3

- To recommend, if considered appropriate, future CWD monitoring activities for the EU based on an assessment of the epidemiological situation
 - Framing through SWOT (strengths, weakness, opportunities, threats).
 - Defining different objectives of the future surveillance (animal health (introduction risk, status recognition), environmental issues, epidemiological knowledge, spillover risk, public health/zoonotic potential)
 - Provisions of general surveillance recommendations





ToR4

Based on what is known about the epidemiology of CWD in Europe/European Union, to describe the criteria relevant for considering an area not to be infected with CWD

- Need of defining the concept of area
- Combination of criteria: multiple requirements
- Multistep progressive strategy to achieve status of area free from infection





ToR5

- To provide the design of a **genotyping protocol** for positive samples, and for the negative samples of the 3-year monitoring programme stored as per point 3.3, section III.A of Annex III of Regulation (EC) No 999/2001, specifying which negative samples should be genotyped, the codons of the PRNP gene to be genotyped and **recommending genotyping assay/s** for the implementation of the requirement by the NRLs
 - Defining different objectives: detection, frequency, res/sus
- Focus on polymorphic species (by country)
- Multi-aim sample size
- Recommendations, not protocol.





CWD MONITORING

Mandate received: Assigned mandate: Charter approved: Acceptance mandate: Working group established (DoI?): Meetings:

Deadline for submission of opinion:

31 March 2023





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8 non – EU RC: Bosnia and Herzegovina, Iceland, Montenegro, North Macedonia, Norway, Serbia, Switzerland

Turkey (first time)

Albania: no surveillance

CATTLE

- No change in tested animals 1,021,252 (-9%)
 2 H-BSE in France & Spain
 - 4 L-BSE in France (2), Germany & Spain
 - 1 C-BSE in England. dairy cow (FS) Somerset, 6.5y
 - 2 H-BSE Brasil and 1 H-BSE Canada





SHEEP

- No change in tested animals 311,174 (-6.4%)
 - 551 cases (19.8%)
 448 (81.3%) CS from 6 MS ES (184), IT (148), EL(74)
 55 CS from IS
 103 (18.7%) AS from 13 MS PT, HU
 9 AS: IS (1) NO (8)

GOATS

- No change in tested animals 118,457 (-1.8%)
- 224 cases (-31.7%)
 219 (97.8%) CS from 6 MS. CY (135), ES (45), IT (23)
 5 (2.2%) AS



2021 EUSR TSE

SHEEP GENOTYPE

- 98.6% genotyped in NSP3, NSP3O, NSP4 and NSP5
- 5,411 random genotype, 9 MS. Excluding CY, 7.9% susceptible (IT 21.2%)

GOAT GENOTYPE

- 152 cases genotyped (2 AS, 149 CS and 1 inconclusive)
 - N146 and Q222 when available





CERVDIS

- **5,854** by 8 MS (82.1% from SE and RO)
 - 70.2% from the hunted/slaughtered fit for human consumption (HSHC)
 - No cases
 - NO: 21,670
 - 2 cases in moose, 1 case in red deer

https://www.efsa.europa.eu/en/topics/topic/tra nsmissible-spongiform-encephalopathies-tses





2021 EUSR TSE

STORYMAP



https://storymaps.arcgis.com/stories/43e7cf4146534e2c8f7c5dce66fe1e56



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2021 EUSR TSE

DASHBOARD



https://www.efsa.europa.eu/en/microstrategy/cattle-bse https://www.efsa.europa.eu/en/microstrategy/sheep-goats-scrapie https://www.efsa.europa.eu/en/microstrategy/cervids-cwd





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Commission Regulation (EU) No 2021/1372:

PAP from pigs and insects in poultry feed;
 PAP from poultry and insects in pig feed;
 Ruminant C&G in non-ruminant







Processed **A**nimal **P**rotein (PAP): animal protein derived entirely from **Category 3** material or products derived from them (except hides and skins, hooves, feathers, wool, horns, hair and fur, adipose tissue, catering waste not from international means of transport)

Annex X Com Reg (EU) 2011/142: "Processed animal protein of mammalian origin must have been submitted to processing method 1 (pressure sterilisation)"

Derogations using methods 2-5 or 7 for :

- Petfood
- Organic fertilisers and soil improvers
- Fuel

But not for feed for farmed animals!





EFPRA (European Fat Processors and Renderers Association): "no EU operator applies method 1 for the processing of PAP of porcine origin"

ToR

The Commission requests EFSA to provide a scientific opinion concerning the **efficacy of methods 2 to 5 and method 7** to inactivate relevant pathogens when producing processed animal protein (PAP) of porcine origin intended to feed poultry and aquaculture animals.

In particular, the scientific opinion should comprise an assessment of the **level of inactivation of relevant pathogens that could be present in processed animal protein of porcine origin intended to feed poultry and aquaculture animals**





PIG PAP







PIG PAP

Draft mandate received: Final mandate received: Mandate number: Question number: Charter approved: Working group established:

3 June 2022 8 July 2022 M-2022-00135 EFSA-Q-2022-00455 12 July 2022 2 September 2022

Deadline for submission of opinion:

30 June 2023





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 Annex VIII, Chapter A, Section A, point 2 to Regulation (EC) No 999/2001,

> a Member State can submit a request to the Commission to be recognised as a Member State, or zone of a Member State, with a negligible risk of classical scrapie.

Czech Republic submitted a request to the Commission to be recognised a Member State with negligible risk of classical scrapie on 12 May 2022

> Part of the dossier evaluated by the EC Part commission to EFSA. Art 31. Scientific report Level of testing Future surveillance plan





CZ NEGLIGIBLE RISK

- EFSA: Evaluation of the application of **Sweden, Finland, Denmark** to be recognised as having a negligible risk of classical scrapie (2015)
- CZ application: mid-October





Thank you Questions?









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