DR. MICKAEL CARGNEL (Orcid ID: 0000-0002-7490-7903)

PROF. CLAUDE SAEGERMAN (Orcid ID: 0000-0001-9087-7436)

Article type : Original Article

Prioritization of livestock transboundary diseases in Belgium using a multi-criteria decision analysis tool based on drivers of emergence

Juana BIANCHINI ¹, Marie-France HUMBLET ², Mickaël CARGNEL ^{1,3}, Yves VAN DER STEDE ^{3,4#}, Frank KOENEN ³, Kris DE CLERCQ ³, Claude SAEGERMAN ^{1*}

¹Research Unit in Epidemiology and Risk Analysis Applied to Veterinary Sciences (UREAR-ULiege), Fundamental and Applied Research for Animals & Health (FARAH) Centre, Faculty of Veterinary Medicine, Liege University, Liege, Belgium

²Department of Occupational Safety and Hygiene, Biosafety and Biosecurity unit, Liege University, Liege, Belgium

³Sciensano, Brussels, Belgium

⁴European Food Safety Authority, Parma, Italy

Disclaimer: Yves Van der Stede is currently employed with the European Food Safety Authority (EFSA) in the ALPHA Unit that provides scientific and administrative support to EFSA's scientific activities in the area of Animal Health and Welfare. The positions and opinions presented in this article are those of the authors alone and are not intended to represent the views or scientific works of EFSA.

This article has been accepted for publication and undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi: 10.1111/tbed.13356

*Corresponding author: claude.saegerman@uliege.be

ABSTRACT

During the past decade, livestock diseases have (re-)emerged in areas where they had been previously eradicated or never been recorded before. Drivers (i.e. factors of (re-)emergence) have been identified. Livestock diseases spread irrespective of borders, and therefore, reliable methods are required to help decisions makers to identify potential threats and try stopping their (re-)emergence. Ranking methods and multi-criteria approaches are cost-effective tools for such purpose and were applied to prioritize a list of selected diseases (N=29 including 6 zoonoses) based on the opinion of 62 experts in accordance with 50 drivers-related criteria. Diseases appearing in the upper ranking were porcine epidemic diarrhoea, foot-and-mouth disease, low pathogenic avian influenza, African horse sickness, and highly pathogenic avian influenza. The tool proposed uses a multi-criteria decision analysis approach to prioritize pathogens according to drivers and can be applied to other countries or diseases.

Keywords: Drivers; Transboundary diseases; Zoonoses; Prioritization; Ranking; Belgium; Multicriteria decision analysis (MCDA); Expert elicitation; Cluster analysis; Sensitivity analysis.

INTRODUCTION

The Food and Agriculture Organization of the United Nations (FAO) has defined transboundary animal diseases as "epidemic diseases which are highly contagious or transmissible and have the potential for very rapid spread, irrespective of national borders, causing serious economic and sometimes public health consequences" (Food and Agricultural Organization, 2018). Thus, livestock diseases may be responsible for negative social, economic and environmental impacts, at different levels (locally, nationally, regionally and internationally). Hence, the introduction of a new livestock disease not only has an impact on animal health, but it also affects international trade, food supply and if zoonotic, human health (Food and Agricultural Organization, 2018).

With the societal and technological changes occurring during the twentieth century, novel pathogens have appeared with countries experiencing human and animal diseases they have never

seen before (emergence) or that had been eradicated in the past (re-emergence). Noteworthy, examples of (re-)emerging animal diseases are: the foot-and-mouth disease (FMD) epidemic in the United Kingdom in 2001 (Knowles et al., 2001) and in Japan in 2010 (Muroga et al., 2012) and the continuing outbreaks of highly pathogenic avian influenza (HPAI) since 2003-2004 around the world (Elbers et al., 2004), the Bluetongue epidemic in Western Europe (Carpenter et al., 2009; Wilson & Mellor, 2009) and the newly identified Schmallenberg disease in Germany in 2011, which has further spread to other parts of Europe, like The Netherlands, Belgium and Northern Ireland (Afonso et al., 2014; Anonimous, 2013). Also, in 2016, cases of highly pathogenic avian influenza were reported to the OIE from different European member states including Belgium (World Organisation for Animal Health, 2018). Another very important recent emerging livestock disease reported specifically in Belgium at the end of 2018 was African swine fever. Although cases so far have been reported only in wild boars (Linden et al., 2019). Its emergence is of great concern for the pig industry of the region and being a disease, which until now has been exotic for Belgium. It shows how diseases may re-emerge unexpectedly with most likely origin attributable to a human activity (Saegerman, 2018).

The (re-)emergence of diseases shift in relation to several underlying set of factors inherent to modern society, i.e. the so-called "drivers". The joint presence of these drivers can create an environment in which infectious disease can (re-)emerge and be maintained in animal and/or human compartments (King, 2004). Many drivers have been identified, such as climate change, global travel, immigration patterns, increase of the human population, environmental degradation and others (Altizer et al., 2013; King, 2004; Daszak et al., 2000).

The threat of (re-)emergence is more likely to increase and past experience has shown that no country, however economically well-developed it may be, is capable of ensuring 100% security of its borders, even by imposing measures such as quarantine protocols or import bans on animals and animal products (Ben Jebara, 2004). In Belgium, the monitoring and reporting of livestock diseases are subjected mostly on self-reporting of suspected clinical cases by the farmers to the Federal Agency for the Safety of the Food Chain (FASFC), with an established list of mandatory notifiable diseases for livestock and other species (aquatic, exotic) (Federal Agency for the Safety of the Food Chain, 2019). Each suspicion is then confirmed by laboratory analysis (Federal Agency for the Safety of the Safety of the Food Chain, 2019). Thus, a rational priority setting approach is needed to assist decision makers in identifying and prioritize diseases that are more likely to (re-)emerge and as such allocating the right resources tailored to a particular disease threat. One such

approach used is disease prioritization, which has as main objectives: to optimise financial and human resources for the surveillance, prevention, control, and eradication of infectious disease and to target surveillance for early detection of any emerging diseases (Humblet et al., 2012).

Some studies identified key characteristics of potential emerging infectious diseases and prioritized infectious diseases according to their risk of (re-)emergence or impact in some countries (Cox et al., 2013; Humblet et al., 2012; Havelaar et al., 2010; Cardoen et al., 2009). Hence, these focused on human or zoonotic diseases and the impact they would have in certain countries. In this study, the focus is livestock epidemic diseases and the aim was to identify (re-)emergence drivers' criteria and with it use expert elicitation to prioritize livestock epidemic diseases that may emerge in Belgium.

A multi-criteria decision analysis (MCDA) method was chosen because it provides a systematic way to integrate information from a range of sources (Cox et al., 2013) and it aims to improve transparency and repeatability (European Centre for Disease Prevention and Control, 2015). MCDA requires identifying criteria and scoring criteria according to the pathogen/disease. By weighting each criterion and calculating weighted scores from the criteria, an overall score per pathogen/disease was calculated (Humblet et al., 2012; European Centre for Disease Prevention and Control, 2015).

This is the first study to prioritize livestock epidemic disease using drivers as criteria. This prioritization list could be an aid to decision makers to make an informed decision on course of actions to be taken and use the correct resources when there is a threat of a disease (re-)emerging in Belgium.

MATERIALS AND METHODS

Selection of diseases

We compiled a list of livestock-associated infectious diseases (**Figure 1**) using a systematic approach. This was done by collating in a single database notifiable terrestrial animal diseases from different governmental official lists from Belgium (Federal Agency for the Safetuy of the Food Chain, 2015) and neighbouring countries (Luxembourg was excluded because of high similarity), i.e. Germany (Federal Ministry of Food and Agriculture of Germany, 2015), France (Légifrance, 2015a and 2015b), The Netherlands (Ministerie van Landbouw, 2015) and Great Britain (Scottish Government, 2015). In order to broaden the spectrum, diseases included in two

other lists of official international organisations, i.e. the World Organisation for Animal Health (OIE) (World Organisation for Animal Health, 2015) and the European Union (European Commission, 2012), were also added to the database. Only diseases that affect cattle, sheep, goats, swine and poultry (livestock) were selected from the official lists and included in database.

After completion of the database, diseases were excluded if: a) they were not of the epidemic type; b) by the time the list was compiled (January 2015) no cases were reported in Belgium over the past year (i.e. during the year 2014). The disease duplicates were removed. Four diseases that were not in any of the official lists were added to the list of diseases for prioritization: Schmallenberg, Aino, Akabane and novel swine enteric coronavirus. Schmallenberg virus is a novel pathogen detected in 2011 in three adjoining countries: Germany, the Netherlands and Belgium, which eventually caused an outbreak in Northern Europe from 2011 to 2013 (Lievaart-Peterson et al., 2012). Aino and Akabane viruses were added because both viruses belong to the same Simbu serogroup of the genus Orthobunyavirus of the Bunyaviridae family as Schmallenberg virus. Additionally, a number of publications have highlighted that viruses from the Simbu group circulate within the Mediterranean basin (Chaintoutis et al., 2014; Yilmaz et al., 2014; Azkur et al., 2013; Lievaart-Peterson et al., 2012). Thus, the risk of any of these viruses to (re-)emerge may be present, which further prompted the necessity of adding these three viruses to the list of diseases to be prioritized. The appearance of the novel swine enteric coronavirus disease, first in the United States in February 2014 and later in March 2014 in Ontario Canada (European Food Safety Authority, 2014), raised concerns in European Members States, as this emerging diseases could affect the health status of pig holding in Europe and their production. For this reason, we decided to include it in the final list of epidemic livestock diseases.

Questionnaire Design

The main objective was to prioritize the diseases according to drivers of (re-)emergence. A driver was defined as a factor, which has the potential to directly or indirectly precipitate ('drive') or lead to the (re-)emergence of a livestock infectious disease. We identified different criteria considered as drivers through scientific literature and previous disease prioritization exercises, and discussion with experts from academia, government agencies and international bodies.

A total of 50 criteria were identified and classified under 8 different domains (**Table 1**): (A) pathogen/disease characteristics (N 9 criteria); (B) distance to Belgium (N =3 criteria); (C) ability to monitor, treat and control the disease (N =7 criteria); (D) farm/production characteristics (N = 7 criteria); (E) changes in climate conditions (N = 3) criteria; (F) wildlife interface (N = 6 criteria) (G); human activity (N = 6 criteria); and (H) economic and trade activity (N = 9 criteria). The questionnaire was formatted in Excel® (Microsoft, Redmond, WA, USA, 2013) file with one spreadsheet per domain including corresponding criteria with an addition of a last spreadsheet, with the eight listed domains (N=8 Domains).

Each criterion had a definition of the coefficient, which ranged from 0 to 4 accordingly (**Appendix** 1).

Scoring and weighting system

Each domain spreadsheet had a number of criteria. For each criterion, coefficients were clearly defined for a good comprehension and standardisation. Coefficients were from scores of 0 to 4 or from 1 to 4 (a number of criteria could not be scored with a zero; e.g. current species specificity of the disease causing agent). Each spreadsheet included two columns. Experts had to fill both of them. The first one corresponding to the coefficient for the choice for the criterion, and the second one for weighting they gave to the criterion (intra-domain weighting). Regarding the weighting system, a Las Vegas method was applied (Gore, 1987). The number of points to be distributed was proportional to the number of criteria per category multiplied by ten. Indeed, the criterion with the most points allocated is considered the one that weighs the most in the category. If, on the other hand, all the criteria have the same weight in the category, the distribution is equitable, with 10 points for each criterion. For example, 90 points were to be distributed between the 9 criteria of the "pathogen characteristics" domain. Indeed the criterion with the most points allocated is considered the one that weighs the most in the pathogen characteristics. Such process illustrated the experts' opinion on the relative importance of criteria within one domain.

The last spreadsheet was dedicated to the inter-domain weighting. Experts were asked to distribute a total of 80 points (N = 8 domains) among the domains to classify the domains according to their opinion.

Expert elicitation

Two rounds of expert elicitation were implemented. The first round consisted in the questionnaire assessment; experts were asked to verify if the questions were in relation with the drivers and if the scoring systems were correctly defined and identified. The questionnaire and related instructions were sent to 14 experts (**Appendix 2**) by e-mail. The experts were asked to complete questionnaire by scoring and additionally to assess and give comments on the criteria and coefficient definitions. The questionnaire was then refined according to experts' comments and suggestions.

For the second round, 62 experts were identified (**Appendix 3**) via internet searching and recommendations from the project partners and recruited participants. These experts were asked to answer the questionnaire in order to rank the diseases. Thus, they had to choose the defined coefficient for each criterion (i.e. criterion scoring), then distribute the points for within each domain (i.e. the intra-domain weighting), and lastly distribute the points within the domains (i.e. inter-domain weighting).

They were invited to participate via a project summary e-mail and were sent the reviewed questionnaire via e-mail if they agreed to participate. Experts were recruited until a minimum of 4 experts per disease was obtained with a maximum of 5 experts. In some cases, one expert could answer several questionnaires (one per disease), if the diseases were within is area of expertise.

Calculation of total scores for each disease

To obtain the overall score for the ranking an aggregation method that combined the 2 types of weighting (i.e. the intra and inter-domain) was used. First the criterion score (coefficients attributed by experts) had to be standardised. Indeed, some criteria were allocated coefficients from 0 to 4 and others from 1 to 4. This standardised score was then multiplied by the intradomain weight as given by the expert. These results were summed to obtain a domain score.

In this formula, DSj = domain score, crit= criterion, SCj= standardized score of the criterion, and WdWj = intra-domain weight for each criterion.

Each domain score was then multiplied by the inter-domain weight. These results were summed and an overall weighted score calculated, per expert and per disease.

OWS=
$$\sum cat (DSj \times IdWj)$$
 [Equation 2]

In this formula, OWS = overall weighting score of each expert for a specific disease, cat = category, DSj = domain score, and IdWj = inter-domain weight.

Each disease had 4 or 5 OWS (since there were 4 or 5 experts per disease), thus for each disease, the final score was the average of all disease experts' OWS. The final score was then used to rank the diseases, based on drivers, from the highest score to the lowest. The highest score corresponded to the disease with the highest risk of (re-)emerging according to the drivers. In addition, the median and range among the scores of all the disease experts were also obtained. With the median, a ranking was done to observe if there was any significant difference with the ranking obtained using the mean. The range was used to note which diseases had the highest and lowest level of variation/uncertainty among the final experts' average score.

Ranking of the perceived drivers (domains)

In order to determine which driver(s) was/were considered as the most influential for the (re-) emergence of diseases, the domains were ranked. Domain ranking was performed using the interdomain scores (weights). The sum of each domain-weight (∑IdWj) per disease and per domain given by each expert was ranked from the high to the low, i.e. 1 to 8. Then for each domain, the frequency of their rank was used to display in graph.

Cluster analysis

A cluster analysis was implemented using regression tree analysis (Salford Predictive Modeler®, Version 8.2, Salford Systems, San Diego, California, USA). The normalized disease score are a continuous variable and the aim was to obtain groups in qualitative categories of importance (e.g. very high, high, moderate and low) with minimal within-group variance.

Sensitivity Analysis

Two sensitivity analysis were assessed, i.e. on expert elicitation and influence of a domain. This was achieved by repeating the disease ranking with a "reduced" version of the model and comparing the new ranking to the complete model.

The experts' sensitivity analysis consisted in dividing them into 4 groups. Scores were then recalculated by deleting a group of experts. Each reduced ranking model was compared to the full complete model by using the Spearman's rank test to establish if the ranking was correlated between the complete and the reduced models. The sensitivity analysis on the domains was done by deleting one domain and re-calculating the mean scores to rank the diseases. This "reduced" ranking was then compared with the complete model and the Spearman's rank test was applied. If the ranking position changed to less than three places, then the final score was considered as robust. If it changed to more than two places, then it was considered as a domain of drivers influencing greatly disease (re-)emergence.

RESULTS

Disease selection

We compiled a list of 29 diseases (**Table 2**) after applying inclusion and exclusion criteria. Nearly all of them were viral with the exception of three bacterial diseases: contagious bovine pleuropneumonia (CBPP), contagious caprine pleuropneumonia (CCPP) and haemorrhagic septicaemia. Out of the 29 diseases, 13 were caused by arboviruses. Six diseases, i.e. eastern equine encephalitis (EEE), western equine encephalitis (WEE), Venezuelan equine encephalitis (VEE), Japanese encephalitis, West Nile fever and Nipah disease were zoonotic.

Questionnaire survey

All 14 experts contacted for the first phase (questionnaire assessment) answered positively (**Appendix 2**). There was a general agreement on which criteria and coefficients were clear or not. No criterion, nor coefficient were deleted but only amended according to experts' suggestions.

For the second phase of expert elicitation, a total of 62 experts agreed to participate and answered the questionnaires (**Appendix 3**). The objective of minimum 4 experts per disease was reached and the maximum of 5 experts was reached for 8 diseases.

Ranking of diseases

The final disease ranking based on the average final scores are shown in **Figure 2**. The higher the mean score, the higher the ranking, which means the disease is most likely to (re-)emerge in Belgium.

The top 5 diseases in decreasing order were: porcine epidemic diarrhoea (PED), FMD, low pathogenic avian influenza (LPAI), African horse sickness (AHS) and HPAI (**Table 3**). On the

other end, the diseases with the lowest mean scores were: haemorrhagic septicaemia, Japanese encephalitis, WNF, *peste des petits ruminants* (PPR) and Nipah disease.

When comparing the ranking obtained using the average of the scores of the experts and the ranking obtained with the median of the experts' score, the Spearman's test, a Rho of 0.8044 was obtained (p-value 0.05), showing that there was a significant correlation in both rankings (Appendix 4). However, important change in the ranking for 4 diseases (CBPP, CCPP, Bluetongue and Newcastle) was noted (Appendix 4). The range obtained showed that the 4 highest range values (i.e. the diseases which experts had a high disagreement on their (re)emergence in Belgium) were: CBPP, CCPP, Vesicular stomatitis and Nipah virus (Figure 2). On the other end the 5 smallest range values were: Novel Swine Enteric Coronavirus Disease, HPAI, Haemorrhagic Septicaemia, CSF and Schmallenberg (Figure 2).

Cluster analysis

The regression tree analysis determined 4 clusters (**Figure 2**). The clusters distinguished five, eleven, nine and four diseases, and were classified, respectively as of 'low importance', 'moderate importance', 'high importance', and 'very high importance' (i.e. highly influenced by drivers). The diseases belonging to the node 'highest importance' were PED, FMD, LPAI and AHS. The node of the lowest importance included haemorrhagic septicaemia, Japanese encephalitis, PPR, Nipah disease and WNF.

Drivers influence

The relative importance of the 8 domains varied depending on the disease. However, when considering all domains for all 29 diseases, 'economy and trade activities' obtained the highest number of points, being ranked first 15 times and zero times last ranked (8th). The opposite can be said about 'characteristics of farm/production system', as it was never ranked 1st nor 2nd (**Figure 3**).

Sensitivity Analysis

The sensitivity analysis done on the groups of experts showed that the ranking of diseases was not affected in the reduced models. Indeed, the Spearman's rank-order correlation indicated a strong

positive association of ranks when using different groups of experts for different reduced models, showing that there was a consistency among the scoring of the experts

As for the domain sensitivity analysis, **Table 3** displays the mean scores and ranking of the diseases without the scores. The domain which showed the strongest influence on the ranking of a disease (changing the ranking of a disease for more than 3 spots) was 'economic and trade activity'. When discarding that domain, 22 diseases moved three places up or down in the ranking. The Spearman rank correlation test for comparing the base model with the reduced model without the 'economic and trade activity' showed a 0.42-Rho (p < 0.05).

Figure 4 illustrates the movements of the top 5 diseases after performing the sensitivity analysis. When discarding the domain (A) (pathogen characteristics), FMD moved from the 2nd to the 6th place in the ranking, thus highlighting the strong influence of the domain (A) on that specific disease. The ranking of AHS changed notoriously without 'economy and trade activities' (domain H), moving from the 4th to the 25th place. LPAI was also strongly influenced, lowering from the 4th to the 23rd place, in the model without the wildlife interface domain.

DISCUSSION

The MCDA approach allowed the selection of 29 livestock diseases exotic to Belgium and their prioritization based on drivers. Whilst such approach was used in previous disease prioritization exercises, this is one of the first to consider livestock epidemic diseases only and to use criteria related to drivers of (re-)emergence. Only diseases exotic to Belgium were prioritized.

The diseases that fitted the eligibility criteria were all of viral origin, except haemorrhagic septicaemia ($Pasteurella\ multocida$, serotypes 6:B, 6:E), CCPP and CBPP. Few zoonoses were included in the list (n = 6) as the prioritization exercise focused on livestock epidemic diseases. Therefore, several zoonoses included in other prioritization processes were excluded.

Regarding prioritization, PED ranked top of the list. Although currently not reportable neither in the EU (except in the UK) nor to the OIE, it ranked high in all models (high mean score), possibly due to its highly transmissible character and the difficulty to control it; furthermore, the disease mainly concerns intensive production. Cases have already been reported in EU Member States: e.g. in May 2014, an outbreak of diarrhoea occurred in fattening pigs on German farms. An

outbreak of diarrhoea occurred on a Belgian fattening pig farm at the end of January 2015; this was the first confirmed PED case in Belgium in decades (Theuns et al., 2015). When the list of diseases was compiled the outbreak had not occurred yet, but when the expert's answered the questionnaire it had and therefore this was most likely the reason why it ranked at the top of the prioritized list.

LPAI ranked slightly higher than HPAI in this multi-criteria analysis on the risk of (re)-emergence (LPAI ranked 3rd whilst HPAI ranked 5th). However, by the time this paper was written, no cases of LPAI were registered on the OIE WAHIS interface for Belgium (World Organisation for Animal Health, 2018), whereas HPAI was detected in Hungary in October 2016 and later in 19 other Member States: Austria, Belgium, Bulgaria, Croatia, Czech Republic, Denmark, France, Germany, Greece, Hungary, Italy, Luxembourg, the Netherlands, Poland, Slovakia, Spain, Sweden, Romania and the United Kingdom (European Comission, 2018). LPAI shows less signs and symptoms than the HPAI and the vast majority of LPAI viruses are maintained in asymptomatic wild birds (Center for Food Security and Public Health, 2015), thus an incursion of LPAI in an area free of the virus is more likely to happen and go undetected. Additionally, the HPAI viruses can evolve directly from low-pathogenic (LPAI) virus precursors following introduction into domestic poultry (Monne et al., 2014). Hence, these characteristics of the virus give in this prioritization LPAI a higher score than HPAI, but HPAI is more likely to be detected and notified.

AHS surprisingly ranked 4th, although its last know incursion in Europe (Portugal and Spain) was in 1987 and its eradication dates back to 1990. Such high position in the ranking could be related to its vector borne transmission, i.e. by *Culicoides* biting midges. These vectors are often highly abundant, across most of Africa, the Middle East, Europe, and southern Asia (Carpenter et al., 2017). Additionally, the recent changes in the epidemiology of bluetongue and its latest epidemic in Europe) and the emergence of Schmallenberg disease (Carpenter et al., 2009; Wison & Mellor, 2009; Alfonso et al., 2014; Anonimous, 2013), highlight the uncertainty about the variables controlling the spread and persistence of *Culicoides*-borne arboviruses. These different factors have raised concerns that AHS may also amount similar incursions, hence explaining such high mean final score in the prioritization process.

In this prioritization most of the diseases were in cluster 2 (high importance, N=9) and 3 (moderate importance, N=11). Cluster 2 of high importance includes the diseases HPAI, CSF,

LSD, sheep and goat pox and CBPP, all of which have been well described in the past, have had epidemics and still have important outbreaks worldwide. The new swine enteric coronavirus disease, which was added on interest basis, also belongs to this cluster. The three diseases which were added to the prioritization although not present in any of the official list of notifiable diseases; Aino, Akabane and Schmallenberg were categorized in cluster 3, even though only Schmallenberg has had outbreaks in Europe. It is therefore considered that the Simbu serogroup could be of moderate importance in (re-)emerging.

From the complete list of the livestock diseases prioritized it is important to highlight ASF. In this prioritisation, ASF did not obtain the highest ranking score at 16th place and placed in the group of moderate importance of the regression tree analysis. However, ASF has become more prevalent in the Caucasus regions since its spread from eastern Africa to Georgia in 2007 and the virus reached the European Union member states of Estonia, Latvia, Lithuania and Poland; in 2016, Moldova; in 2017, the Czech Republic and Romania (Chenais et al., 2018) and in 2018, the Hungary and Bulgaria. It emerged in Belgium in September of 2018 when authorities in Belgium reported that ASF had been confirmed in 2 wild boars (Linden et al., 2019). The detection of ASF in Belgium was unexpected as ASF appears to have jumped a considerable distance from previously affected countries: ~500 km from the border with the Czech Republic, 800 km from Hungary, and 1,200 km from the border with Romania (Garigliany et al., 2019) and how it was introduced in the wild boar population until the writing this article is unknown (presumably related to illegal human activities) (Saegerman, 2018). The ASF score (ranked 16th place and was in group of moderate importance of the regression tree analysis) may be explained that although there was an awareness of the risk of ASF spreading to EU member states, when the questionnaire was answered by the experts (year 2016) the risk that ASF would become endemic in domestic pigs in Ukraine and Belarus was considered to be moderate and the risk to further spread into unaffected areas was also considerate moderate (European Food Safety Authority, 2014). Furthermore, the score reflected the geographical position of where ASF had been reported and it was unexpected that ASF skipped neighbouring countries and directly entered Belgium (Garigliany et al., 2019). In addition, any ranking cannot include unforeseen circumstances such as the human factors; the vigilance should be always implemented for new introduction.

This score can only be compared with the prioritization work done by Humblet and collaborators (Humblet et al., 2012) as other prioritization works using the MCDA method, such as those by Cardoen et al., 2009 and Havelaar et al., 2010, only included zoonoses. Indeed, in regression tree analysis of prioritized diseases of food-producing animals and zoonoses, ASF also fell in the 3th group of importance out of the 4 (Humblet et al., 2012), just like in this prioritization work. Another study, which may be used for comparison as it used MCDA approach and had swine diseases, done by Brookes et al. 2014, ASF ranked higher, but in this study only exotic diseases for the pig industry in Australia were ranked using criteria related to impact and the experts were pig producers which changes the importance in the scores, giving ASF a higher ranking.

The livestock diseases at the bottom of the list were Nipah disease, PPR, WNF, Japanese encephalitis and haemorrhagic septicaemia. In other prioritization exercises, Nipah, Japanese encephalitis and WNF were ranked in a higher category (Humblet et al., 2012; Cox et al., 2013; Havelaar et al., 2010). The prioritization model presented here was based on criteria reflecting only drivers; no criteria linked to societal or economic impacts were considered, which affects the weights given to the different domains. Therefore, diseases that otherwise would have scored high in the ranking, were in the lower end ('low importance' group in the regression tree analysis). Moreover, until recently only WNF had been reported in Europe (Sambria et al., 2013). However, when writing the results of this article, in June 2018, Bulgaria reported the first outbreak in the European Union of PPR, in farms close to the border with Turkey (Altan et al., 2018). Thus, although PPR here is in the low importance group this unexpected introduction would make this disease become suddenly a priority.

Drivers are a complex set of factors and their convergence can cause the (re-)emergence of a disease. Several drivers have a stronger impact on diseases compared to others, as shown in the results section. PED ranked at the top in all models, except in the reduced models of production system characteristics. PED affects mainly intensive production systems, thus, the driver category 'production system characteristics' logically influences a lot. When using the reduced model, the mean score decreases and the disease moved from the 1st place to the 8th place. In comparison, FMD ranked high in the prioritization process (2nd), but lowered to the 12th place in the reduced model, which excluded disease pathogen characteristics. For FMD, the strongest driver was the 'pathogens characteristics'. The virus is highly contagious, spreads via airborne and direct contact and affects different livestock species, giving this driver category a strong weight.

All experts considered that 'economy and trade activities' was the most important driver (high weight). It was ranked first more often than others. In the reduced model (without the 'economy and trade activities' domain), all diseases with the exception of 7, moved up or down in the ranking by more than 3 places. This is of no surprise, as economic and trade activity has priority in the age of globalisation; increased movement of live animals and animal products crossing oceans and international boundaries increase the risk of spread for animal and zoonotic diseases (Domenech et al., 2006). On the other side of the scale, the domain defined as 'characteristics of farm/production system' was given the least weight, therefore with the least influence. Although this true for many diseases within the EU Member States, it is important to consider that for some other diseases in certain cases this domain could be a strong influence. One example is farms, which may have backyard pigs, with no biosecurity set in place and not always under the full control of veterinary services. This type of farming could well explain the dissemination of diseases such as ASF, thus making characteristics of farm/production an importance driver.

As only diseases exotic to Belgium were considered, the results presented here are specific to the country. Nevertheless, a similar prioritization exercise could be applied to other countries, in particular EU Member States, because their animal sanitary status, regulations and controls are similar. Indeed, the focus of the questionnaire was to prioritize diseases according to their drivers and not to the impact on the country nor other criteria country-specific. Furthermore, the sensitivity analysis of experts also showed a high correlation among the ranking of models, which confirms that experts were in agreement in regards to the scores.

Overall, the importance of validating each generated model is highlighted. Two types of validations can be used. This involves testing the internal validity of the model (e.g., by performing a sensitivity analysis on the domains of criteria and/or testing the effect of deleting groups of experts on the results) and the external validity of the model (e.g., comparing results of each model with other driver-based prioritization exercises if they exist).

The tool provided here clearly defines each criterion and its coefficients in order to ensure standardisation of answers. Although this study cannot account for the complexity of drivers in the (re-)emergence of a disease, it can provide, through a quick assessment, a general picture of what drivers can influence the (re-)emergence of a disease. Furthermore, this MCDA tool, which could be made available to third parties upon request to the main authors, can be used with a subset of criteria and/or impact criteria or public health aspects can be easily added, and it could be applied

to a broader set of diseases. The resulting scores could be translated into practical recommendations tailored to the needs of a specific country's national public or governmental agencies.

ACKNOWLEDGMENTS

The authors would like to thank the colleagues who participated to expert elicitation (all cited in **Appendix 1 and Appendix 2**) and shared their experiences and expertise. This study was supported by the Federal Public Service of Health, Food Chain and Environment as part of the EPIDIACAP research project RT13/3 implemented by Liege University and Sciensano. The authors declare no conflict of interest.

ETHICAL APPROVAL

Ethical statement is not applicable to this study as the data were gathered through questionnaire survey without any animal experimentation.

REFERENCES

Afonso, A., Abrahantes, J.C., Conraths, F., Veldhuis, A., Elbers, A., et al. (2014). The Schmallenberg virus epidemic in Europe-2011-2013. Prev Vet Med., 116(4), 391-403. doi: 10.1016/j.prevetmed.2014.02.012.

Altan, E., Parida, S., Mahapatra, M., Turan, N. & Yilmaz, H. (2018). Molecular characterization of Peste des petits ruminants viruses in the Marmara Region of Turkey. Transbound Emerg Dis., 66(2), 865-872. doi: 10.1111/tbed.13095.

Altizer, S., Ostfeld, R.S., Johnson, P.T., Kutz, S. & Harvell, C.D. (2013). Climate change and infectious diseases: from evidence to a predictive framework. Science, 341(6145), 514-519. doi: 10.1126/science.1239401.

Anonimous. (2013). Schmallenberg virus continues to spread across Europe. Vet Rec., 172(21), 543. doi: 10.1136/vr.f3270.

Azkur, A.K., Albayrak, H., Risvanli, A., Pestil, Z., Ozan, E., et al. (2013). Antibodies to Schmallenberg virus in domestic livestock in Turkey. Trop Anim Health Prod., 45(8), 1825-1828. doi: 10.1007/s11250-013-0415-2.

Brookes, V.J., Hernandez-Jover, M., Cowled, B., Holyoake, P.K., Ward M.P. Building a picture: Prioritisation of exotic diseases for the pig industry in Australia using multi-criteria decision analysis. Prev Vet Med., 113(1), 103-107. doi: 10.1016/j.prevetmed.2013.10.014.

Cardoen, S., Van Huffel, X., Berkvens, D., Quoilin, S., Ducoffre, G., et al. (2009). Evidence-based semiquantitative methodology for prioritization of foodborne zoonoses. Foodborne Pathog Dis., 6(9), 1083-1096. doi: 10.1089/fpd.2009.0291.

Carpenter, S., Mellor, P.S., Fall, A.G., Garros, C. & Venter, G.J. (2017). African Horse Sickness Virus: History, Transmission, and Current Status. Annu Rev Entomol., 62, 343-358. doi: 10.1146/annurev-ento-031616-035010.

Carpenter, S., Wilson, A. & Mellor, P.S. (2009). Culicoides and the emergence of bluetongue virus in northern Europe. Trends Microbiol., 17(4), 172-178. doi: 10.1016/j.tim.2009.01.001.

Center for Food Security and Public Health (2015). Iowa State University. Avian Influenza. http://www.cfsph.iastate.edu/Factsheets/pdfs/highly_pathogenic_avian_influenza-citations.pdf

Chaintoutis, S.C., Kiossis, E., Giadinis, N.D., Brozos, C.N., Sailleau, C. et al. (2014). Evidence of Schmallenberg virus circulation in ruminants in Greece. Trop Anim Health Prod., 46(1), 251-255. doi: 10.1007/s11250-013-0449-5.

Chenais, E., Ståhl, K.& Guberti, V.K.D. (2018). Identification of Wild Boar–Habitat Epidemiologic Cycle in African Swine Fever Epizootic. Emerg Infect Dis., 24(4), 810-812. doi: 10.3201/eid2404.172127.

Cox, R., Sanchez, J. & Revie, C.W. (2013). Multi-criteria decision analysis tools for prioritising emerging or re-emerging infectious diseases associated with climate change in Canada. PLoS One., 8(8), e68338. doi: 10.1371/journal.pone.0068338.

Daszak, P., Cunningham, A.A. & Hyatt, A.D. (2000). Emerging infectious diseases of wildlifethreats to biodiversity and human health. Science, 287(5452), 443-449. DOI: 10.1126/science.287.5452.443.

Domenech, J., Lubroth, J., Eddi, C., Martin, V., Roger, F. (2006). Regional and international approaches on prevention and control of animal transboundary and emerging diseases. Ann N Y Acad Sci., 1081, 90-107. doi.org/10.1196/annals.1373.010.

Elbers, A.R., Fabri, T.H., de Vries, T.S., de Wit, J.J., Pijpers, A. & Koch, G. (2004). The highly pathogenic avian influenza A (H7N7) virus epidemic in The Netherlands in 2003-lessons learned from the first five outbreaks. Avian Dis., 48(3), 691-705. doi.org/10.1637/7149.

European Centre for Disease Prevention and Control (2015). Best practices in ranking emerging infectious disease threats. A literature review. Stockholm: European Centre for Disease Prevention and Control. https://ecdc.europa.eu/sites/portal/files/media/en/publications/Publications/emerging-infectious-disease-threats-best-practices-ranking.pdf

European Comission (2018). Avian Influenza. https://ec.europa.eu/food/animals/animal-diseases/control-measures/avian-influenza en

European Commission (2012). Decision 2012/737/EU of 27 November 2012 amending Annexes I and II to Council Directive 82/894/EEC on the notification of animal diseases within the Community. Official Journal of the European Union, L 329, 19-22. Available from: https://eurlex.europa.eu/legal-content/EN/TXT/PDF/?uri=CELEX:32012D0737& from=FR

European Food Safety Authority (2014). Scientific Opinion on African swine fever. EFSA Journal, 12(4), 3628. https://www.efsa.europa.eu/fr/efsajournal/pub/3628

European Food Safety Authority (2014). Scientific Opinion on porcine epidemic diarrhoea and emerging porcine deltacoronavirus. EFSA Journal, 12(10), 3877, 68 pp. doi:10.2903/j.efsa.2014.3877.

Federal Agency for the Safetuy of the Food Chain (2015). Situation zoosanitaire et maladies à déclaration obligatoire en Belgique [in French]. [cited 2015 January]. http://www.afsca.be/santeanimale/zoosanitaire-belgique/

Federal Agency for the Safetuy of the Food Chain (2019). Notification Obligatoire [in French]. http://www.afsca.be/professionnels/notificationobligatoire/

Federal Ministry of Food and Agriculture of Germany [in German] (2015). Anzeigepflichtige Tierseuchen. https://www.bmel.de/DE/Tier/Tiergesundheit/Tierseuchen_texte/ AnzeigepflichtigeTierseuchen.html

Food and Agricultural Organization (2018). Animal Production Health. [cited 2018 September 15th]; Available from: http://www.fao.org/ag/againfo/programmes/en/empres/ diseases.asp

Garigliany, M., Desmecht, D., Tignon, M., Cassart, D., Lesenfant, C. et al. (2019). Phylogeographic Analysis of African Swine Fever Virus, Western Europe, 2018. Emerg Infect Dis., 25(1), 184-186. doi: 10.3201/eid2501.181535.

Gore, S.M. (1987). Biostatistics and the Medical Research Council. MRC News, 35, 19-20.

Havelaar, A.H., van Rosse, F., Bucura, C., Toetenel, M.A., Haagsma, J.A., et al. (2010). Prioritizing emerging zoonoses in the Netherlands. PLoS One, 5(11), e13965. doi: 10.1371/journal.pone.0013965.

Humblet, M.F., Vandeputte, S., Albert, A., Gosset, C., Kirschvink, N., et al. (2012). Multidisciplinary and evidence-based method for prioritizing diseases of food-producing animals and zoonoses. Emerg Infect Dis., 18(4), doi: 10.3201/eid1804.111151.

Ben Jebara, K.B. (2004). Surveillance, detection and response: managing emerging diseases at national and international levels. Rev Sci Tech., 23(2), 709-715. http://web.oie.int/boutique/extrait/709716benjebara.pdf.

King L. (2004). Emerging and re-emerging zoonotic diseases: challenges and opportunities. 72nd General Session, 23-28 May 2004, World Organisation for animal health, Paris, France, 9 p. https://pdfs.semanticscholar.org/17c9/7352041fa26c8a0e21e5a094d22e67834ad7.pdf.

Knowles, N.J., Samuel, A.R., Davies, P.R., Kitching, R.P. & Donaldson, A.I. (2001). Outbreak of foot-and-mouth disease virus serotype O in the UK caused by a pandemic strain. Vet Rec., 148(9), 258-259. PMID: 11292084.

Légifrance (2015a). Le service public de l'accès au droit [in French]. https://www.legifrance.gouv.fr/affichCodeArticle.do;jsessionid= 1900B905EE29F3195D79F324211FA5BE.tpdjo07v_2?idArticle=LEGIARTI000006588118&cid Texte=LEGITEXT000006071367&dateTexte=20080505

Légifrance (2015b). Le service public de l'accès au droit [in French]. https://www.legifrance.gouv.fr/affichCodeArticle.do?cidTexte=LEGITEXT000006071367&idArt icle=LEGIARTI000006588115&dateTexte=&categorieLien=cid

Lievaart-Peterson, K., Luttikholt, S.J.M., Van den Brom, R. & Vellema, P. (2012). Schmallenberg virus infection in small ruminants – First review of the situation and prospects in Northern Europe. Small Rumin Res., 106(2-3, 71-76. https://doi.org/10.1016/j.smallrumres.2012.03.006.

Linden, A., Licoppe, A., Volpe, R., Paternostre, J., Lesenfants, C., et al. (2019). Summer 2018: African swine fever virus hits north-western Europe. Transbound Emerg Dis., 66(1), 54-55. doi: 10.1111/tbed.13047.

Ministerie van Landbouw (2015). Nederlandse Voedsel- en Warenautoriteit [in Dutch]. https://www.nvwa.nl/onderwerpen/dierziekten/lijst-aangifteplichtige-dierziekten/aangifteplichtige-dierziekten-bij-vee

Monne, I., Fusaro, A., Nelson, M.I., Bonfanti, L., Mulatti, P. et al. (2014). Emergence of a highly pathogenic avian influenza virus from a low-pathogenic progenitor. J Virol., 88(8), 4375-4388. doi: 10.1128/JVI.03181-13.

Muroga, N., Hayama, Y., Yamamoto, T., Kurogi, A., Tsuda, T. & Tsutsui, T. (2012). The 2010 foot-and-mouth disease epidemic in Japan. J Vet Med Sci., 74(4), 399-404. doi.org/10.1292/jvms.11-0271.

Saegerman, C. (2018). Unexpected discovery of African swine fever in Belgium [in French]. Épidémiol. et santé anim., 73, 147-164 [in French].

Sambria, V., Capobianchi, M., Charrelde, R., Fyodorova, M., Gaibanig, P. et al. (2013). West Nile virus in Europe: emergence, epidemiology, diagnosis, treatment, and prevention. Clin Microbiol Infect., 19(8), 699 -704. doi: 10.1111/1469-0691.12211.

Scottish Government (2015). List of Notifiable Diseases Great Britain. https://www.gov.scot/Topics/farmingrural/Agriculture/animal-welfare/Diseases/disease/notifiable

Theuns, S., Conceicao-Neto, N., Christiaens, I., Zeller, M., Desmarets, L.M., et al. (2015). Complete genome sequence of a porcine epidemic diarrhea virus from a novel outbreak in Belgium, January 2015. Genome Announc, 3(3), e00506-15. doi: 10.1128/genomeA.00506-15.

Wilson, A.J. & Mellor, P.S. (2009). Bluetongue in Europe: past, present and future. Philos Trans R Soc Lond B Biol Sci., 364(1530), 2669-2681. doi: 10.1098/rstb.2009.0091.

World Organisation for Animal Health (2015). OIE-Listed diseases, infections and infestations in force in 2018. http://www.oie.int/animal-health-in-the-world/oie-listed-diseases-2018/

World Organisation for Animal Health (2018). WAHIS Interface. http://www.oie.int/wahis_2/public/wahid.php/Countryinformation/countryhome

Yilmaz, H., Hoffmann, B., Turan, N., Cizmecigil, U.Y., Richt, J.A. & Van der Poel, W.H. (2014).

Detection and partial sequencing of Schmallenberg virus in cattle and sheep in Turkey. Vector Borne Zoonotic Dis., 14(3), 223-225. doi: 10.1089/vbz.2013.1451.

Figures captions

Figure 1. Systematic process for selecting the livestock diseases

Legend: * Livestock diseases were those which affected cattle, sheep, goats, swine and poultry.

Figure 2. (Re-)emerging livestock diseases prioritized. Mean scores and standard deviations are mentioned. Four clusters were identified by regression tree analysis marked by brackets

Figure 3. Frequency of rank (from 1 to 8) for each domain

Legend: (A) Disease/pathogen characteristics; (B) Distance to Belgium; (C) Ability to monitor, treat and control the disease; (D) Farm/production system characteristics; (E) Changes in climatic conditions; (F) Wildlife interface; (G) Human activity; and (H) Economic and trade activity. Colour of each bar: white (ranked 1st) until black (ranked 8th).

Figure 4. Sensitivity analysis for the five diseases with highest mean scores; the graph illustrates their up or down movements in the ranking.

Legend: * ranking changed by more than 3 positions. (A) Disease/pathogen characteristics; (B) Distance from Belgium; (C) Ability to monitor, treat and control the disease; (D) Farm/production system characteristics; (E) Changes in climate change; (F) Wildlife interface; (G) Human activity; and (H) Economic and trade activity. AHS, African horse sickness; FMD, Foot and mouth disease; HPAI, high pathogenic avian influenza; LPAI, low pathogenic avian influenza; PED, porcine epidemic diarrhoea.

DOMAIN A. DISEASE / PATHOGEN CHARACTERISTICS

| A1 | Current l | knowledge of the pathogen. |
|-----------|------------|--|
| | Score 0 | |
| | Score 1 | Very high: deep scientific knowledge on the pathogen, extensive scientific literature available on its biology (transmission mode, knowledge on vector(s), infectivity, etc.) |
| | Score 2 | High: detailed scientific knowledge on the pathogen but conflicting scientific results; some elements of the pathogen's biology are still not elucidated |
| | Score 3 | Moderate : limited scientific knowledge on the pathogen agent because it is still under characterization; pathogen recently discovered/isolated but belonging to a well-known and studied family of pathogens; the pathogen is characterized by multiple variants not characterized yet |
| | Score 4 | Low: lack of scientific knowledge on the pathogen (multiplication, infectivity, incubation period, transmission mode, etc.); pathogen agent recently discovered and emerging |
| A2 | The curr | ent species specificity of the causing agent of the disease |
| | Score 0 | |
| | Score 1 | Low. Only one host is involved belonging to the same family. e.g. only bovines, only equines, only avian, only porcines |
| | Score 2 | Medium: two species involved |
| | Score 3 | High: three species involved |
| | Score 4 | Very high: affects more than 3 types of families |
| А3 | Genetic v | rariability of the infectious agent |
| | Score 0 | Negligible. The infectious agent is genetically stable |
| | Score 1 | Low. The genetic variability is low therefore it has a low effect in the (re)emergence of the pathogen |
| | Score 2 | Medium The pathogen can be considered with a medium genetic variability. |
| | Score 3 | High. The pathogen is considered with a high genetic variability |
| | Score 4 | Very high. Very high genetic instability (e.g. high mutation rate, re-assortment and recombination). Potentially the three phenomena can characterise the pathogen's evolution |
| A4 | Transmis | ssion of the agent in relation of the possible spread of the epidemic (i.e. ease/speed of spread) |
| | Score 0 | |
| | Score 1 | Low: Low and slow transmission within farms. Between farms only if an infected animal is introduced, close contact |
| | Score 2 | Medium: Medium ease/speed transmission within the farm. Between farms medium |
| | Score 3 | High. Fast transmission within a farm. In a short period of time all animals of the farm are infected. Adjacent farms become infected fast |
| | Score 4 | Very High. Very fast and high transmission within the farms and between farms. A complete area is infected in a very short period of time. |
| A5 | Risk of sl | nowing no clinical signs and silent spread during infection and post infection |
| | Score 0 | Null: Silent spread is not part of the pathogen's characteristics |
| | Score 1 | Low: Very short incubation period and signs of infections easily detected/recognised. |
| | Score 2 | Moderate: Very short incubation period and signs of infection are <u>NOT</u> easily detected/recognised |
| | Score 3 | Medium: Long incubation period, clinical signs are not characteristics and therefore specific diagnosis is necessary to detect infection. |

| | Score 4 | Very high. Long incubation period. Disease/infection shows not clinical symptoms during the infectious period. Chronic shedder |
|-----------|-----------|--|
| 4.6 | | reservoir and potential spread from it |
| A6 | | |
| | Score 0 | Null: no known wildlife reservoir. Disease has never been reported in wildlife species |
| | Score 1 | Low: few clinical cases have been reported in wildlife and no transmission to livestock has ever been documented. |
| | Score 2 | Moderate: wildlife is a reservoir of the disease but only accidental spill overs to livestock have been reported. |
| | Score 3 | High: wildlife is a reservoir for the pathogen/disease but certain environmental conditions (e.g. floods, farms crossing the farmland-bush division, etc.) have to occur for the pathogen/disease to (re)emerge in livestock. |
| | Score 4 | Very high: Disease establishes itself in wildlife as a reservoir and very hard to eradicate it from wildlife. Livestock easily gets infected with the contact with wildlife. |
| Α7 | Existence | of vectors (vertebrate and invertebrate, e.g. mosquitoes, bats, rodents, ticks, midges, culicoids) and potential spread. |
| | Score 0 | Null: No known vector |
| | Score 1 | Low: only one type of vector is present in the country but it's role in the transmission is presumed low (has not been assessed to date). |
| | Score 2 | Moderate: only one type of vector exists in the country and has only been suspected as source and spread of disease |
| | Score 3 | High: only one competent vector is present and can carry and spread the disease |
| | Score 4 | Very high: more than one type of vector can carry and spread the disease and are found spread in most of the territory |
| A8 | Transmis | sion of the pathogen. |
| | Score 0 | |
| | Score 1 | Low: Animals only get infected by <u>direct</u> close contact with other infected animals and vertical transmission. |
| | Score 2 | Moderate: transmission by direct and indirect contact only (e.g. through vehicles, clothes, instruments) or non flying vector (e.g., ticks). |
| | Score 3 | High: Exclusively vector transmission by flying vectors (e.g. culicoides, mosquitoes) |
| | Score 4 | Very high: more than three modes of transmission and/or airborne transmission |
| A9 | Environn | nental persistence |
| | Score 0 | Null: pathogen does not survive in the environment |
| | Score 1 | Low: only anecdotal isolation of the pathogen from the environment has been recorded |
| | Score 2 | Moderate: The survival of the agent in the environment is limited (only temporary) and it's dependent on certain environmental conditions such as humidity, temperature, rainfall, etc. |
| | Score 3 | High: The survival of the agent in the environment is limited (only temporary) and <u>NOT</u> dependent on certain environmental conditions such as humidity, temperature, rainfall, etc. |
| | Score 4 | Very high: agent naturally surviving in the environment (soil, water) and organic materials were it has a long term-survival. |

Number of Criteria = 9, hence 90 points to be distributed within this domain for the intra-domain weighing

DOMAIN B. DISTANCE TO BELGIUM

| B1 | Current | incidence (cases)/prevalence of the disease in the world |
|-----------|---------|--|
| | Score 0 | |
| | Score 1 | Pathogen has been reported only in the countries of the Australasia (Australia, New Zealand, New Guinea and Neighbouring Pacific Islands) region |
| | Score 2 | Disease was reported in countries of the Americas, Caribbean and Asia (excluding the Russian Federation) |
| | Score 3 | Disease was reported/present in the African continent |
| | Score 4 | Disease was reported in countries of the Mediterranean Basin, Middle East and the Russian Federation |
| B2 | Europea | n geographic proximity of the pathogen/disease to Belgium |
| | Score 0 | |
| | Score 1 | Disease has never been present in Europe |
| | Score 2 | Disease has been reported in Europe in the past but is currently exotic. |
| | Score 3 | Disease is currently present in at least one European country which is NOT bordering Belgium |
| | Score 4 | Diseases is currently present in at least one of the countries bordering Belgium |
| В3 | To your | knowledge when was the disease last reported in Europe |
| | Score 0 | More than 20 years ago |
| | Score 1 | More than 10 years ago |
| | Score 2 | More than 5 years ago |
| | Score 3 | More than 1 year ago |
| | Score 4 | Currently present in Europe |
| N.T. | 1 60 | torio = 2, hanno 20 nointe to ha distributed within this domain for the intro domain weighing |

Number of Criteria = 3, hence 30 points to be distributed within this domain for the intra-domain weighing

DOMAIN C. ABILITY TO MONITOR, TREAT AND CONTROL THE DISEASE

| C | Ability of preventive/control measures to stop the disease from entering the country or spreading (containment of the epidemic), EXCLUDING treatment, vaccination and | | |
|---|---|--|--|
| | vector(s) | vector(s)/reservoir(s) control | |
| | Score 0 | | |
| | Score 1 | Very High Sanitary certificate; effective traceability of animals and by-products; effective disinfection measures; no contact between domestic and wild animals; effective | |
| | Score 1 | biosecurity measures | |
| | Score 2 | High No sanitary certificate; effective traceability of animals and by-products; effective disinfection measures; limited or incomplete possibilities to restrict contacts between | |
| | Score 2 | domestic and wild animals; effective biosecurity measures | |
| | Score 3 | Low No sanitary certificate; incomplete traceability of animals and by-products; ineffective disinfection measures; incomplete restriction of contacts between domestic and wild | |
| | Score 3 | animals; ineffective biosecurity measures | |
| | | Very low No sanitary certificate; no traceability of animals and by-products; ineffective disinfection measures; impossibility to restrict contact between farms or between domestic and wild animals; biosecurity measures totally ineffective | |
| | Score 4 | and wild animals; biosecurity measures totally ineffective | |
| C | Vaccine availability | | |
| F | | | |
| | Score 0 | | |

| Score 1 Very high Commercialized vaccine available at a regional/national scale and/or for a targeted species (not systematically available for a global fight plan) Score 3 Low Experimental vaccine, not commercialized to date; severe adverse reaction when applied; limited protector effect Score 4 Very low Absence; no vaccine available on the market for a use in the species considered in the study, no experimental vaccine either C3 Control of reservoir(s) and/or vector(s) Score 0 Null No vector-borne transmission and/or no reservoir(s) known to date Score 1 Very high Effective. Limited reservoir(s) with limited geographical repartition, easy-to-identify; high scientific knowledge on vector(s)/reservoir(s); effective fighting measures bn/OT applicable at a large scale; limited fighting measures Score 2 NoT applicable at a large scale; limited fighting measures Low Numerous reservoirs vectors identified with limited geographical repartition; hard to identify. Lack of scientific knowledge on vector(s)/reservoir(s)/reservoir(s). Fighting measures poorly effective - resistances and/or negative impact on environment; Very low Numerous Vector(s)/reservoir(s)/identified with wide geographic distribution; hard to identify, absence of scientific knowledge on vector(s)/reservoir(s). NO effecting fighting measures applied) Score 0 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 1 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 2 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 1 Very high: Generalized surveillance implemented by ALL EU Member States and worldwide surveillance (i.e. OIE reported) | re |
|--|---------------|
| Score 3 Low Experimental vaccine, not commercialized to date; severe adverse reaction when applied; limited protector effect | re |
| C3 Control of reservoir(s) and/or vector(s) Score 0 Null No vector-borne transmission and/or no reservoir(s) known to date Score 1 Very high Effective. Limited reservoir(s) with limited geographical repartition, easy-to-identify; high scientific knowledge on vector(s)/reservoir(s); effective fighting measures Score 2 High Limited reservoir(s)/vector(s) with limited geographical repartition; easy-to-identify, high scientific knowledge on vector(s)/reservoir(s); effective fighting measures by NOT applicable at a large scale; limited fighting measures Score 3 Low Numerous reservoirs vectors identified with limited geographical repartition; hard to identify. Lack of scientific knowledge on vector(s)/reservoir(s). Fighting measures poorly effective - resistances and/or negative impact on environment; Very low Numerous Vector(s)/reservoir(s)/identified with wide geographic distribution; hard to identify, absence of scientific knowledge on vector(s)/reservoir(s); NO effect fighting measure against vector(s) (no active molecule, resistance to measures applied) C4 Availability and quality of diagnostic tools in Belgium Score 0 Score 1 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 2 High Tests used in local/regional laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | re |
| Score 1 Very high Effective. Limited reservoir(s) with limited geographical repartition, easy-to-identify; high scientific knowledge on vector(s)/reservoir(s); effective fighting measures Score 2 Score 3 Score 4 Score 4 Score 4 Score 5 Score 5 Score 5 Score 6 Score 6 Score 7 Score 8 Score 8 Score 9 Score 9 Score 9 Score 9 Score 9 Score 1 Score 9 Score 1 Score 1 Score 1 Score 9 Score 1 Score 1 Score 1 Score 1 Score 1 Score 1 Score 2 Score 1 Score 3 Score 1 Score 3 Score 1 Score 3 Score 4 Score 9 Score 1 Score 9 Score 9 Score 1 Score 9 Score 9 Score 1 Score 9 S | re |
| Score 1 Very high Effective. Limited reservoir(s) with limited geographical repartition, easy-to-identify; high scientific knowledge on vector(s)/reservoir(s); effective fighting measures by NOT applicable at a large scale; limited fighting measure | re |
| Score 2 High Limited reservoir(s)/vector(s) with limited geographical repartition; easy-to-identify, high scientific knowledge on vector(s)/reservoir(s); effective fighting measures by NOT applicable at a large scale; limited fighting measures Score 3 Low Numerous reservoirs vectors identified with limited geographical repartition; hard to identify. Lack of scientific knowledge on vector(s)/reservoir(s). Fighting measures poorly effective - resistances and/or negative impact on environment; Score 4 Very low Numerous Vector(s)/reservoir(s) identified with wide geographic distribution; hard to identify, absence of scientific knowledge on vector(s)/reservoir(s); NO effect fighting measure against vector(s) (no active molecule, resistance to measures applied) C4 Availability and quality of diagnostic tools in Belgium Score 0 Score 1 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 2 High Tests used in local/regional laboratories by not in the field Score 3 Low tests only used in specialized laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | re |
| Score 3 Score 4 NOT applicable at a large scale; limited fighting measures Low Numerous reservoirs vectors identified with limited geographical repartition; hard to identify. Lack of scientific knowledge on vector(s)/reservoir(s). Fighting measures poorly effective - resistances and/or negative impact on environment; Very low Numerous Vector(s)/reservoir(s)identified with wide geographic distribution; hard to identify, absence of scientific knowledge on vector(s)/reservoir(s); NO effect fighting measure against vector(s) (no active molecule, resistance to measures applied) Score 0 Score 1 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 2 High Tests used in local/regional laboratories by not in the field Score 3 Low tests only used in specialized laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | ·e |
| Score 3 Low Numerous reservoirs vectors identified with limited geographical repartition; hard to identify. Lack of scientific knowledge on vector(s)/reservoir(s). Fighting measures a poorly effective - resistances and/or negative impact on environment; Score 4 Very low Numerous Vector(s)/reservoir(s) identified with wide geographic distribution; hard to identify, absence of scientific knowledge on vector(s)/reservoir(s); NO effecting fighting measure against vector(s) (no active molecule, resistance to measures applied) Score 0 Score 1 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 2 High Tests used in local/regional laboratories by not in the field Score 3 Low tests only used in specialized laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 Score 0 Score 0 Score 1 Score 1 Score 2 Score 3 Score 3 Score 3 Score 4 Score 3 Score 4 Score 5 Score 6 Score 6 Score 6 Score 6 Score 7 Score 8 Score 9 | |
| fighting measure against vector(s) (no active molecule, resistance to measures applied) C4 Availability and quality of diagnostic tools in Belgium Score 0 Score 1 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 2 High Tests used in local/regional laboratories by not in the field Score 3 Low tests only used in specialized laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | ve |
| Score 1 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 2 High Tests used in local/regional laboratories by not in the field Score 3 Low tests only used in specialized laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | |
| Score 1 Very High Field test(s) available and easy to use, with highly discriminating sensitivity and specificity Score 2 High Tests used in local/regional laboratories by not in the field Score 3 Low tests only used in specialized laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | |
| Score 2 High Tests used in local/regional laboratories by not in the field Score 3 Low tests only used in specialized laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | |
| Score 3 Low tests only used in <i>specialized</i> laboratories/national reference laboratory Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | |
| Score 4 Very Low no diagnostic tools available to date C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | |
| C5 Disease is currently under surveillance overseas (OIE, EU) Score 0 | |
| Score 0 | |
| TO BE A STATE OF THE STATE OF T | |
| Score 1 Very high: Generalized surveillance implemented by ALL EU Member States and worldwide surveillance (i.e. OIE reported) | |
| | |
| Score 2 High Surveillance of the pathogen only EU member states | |
| Score 3 Low Surveillance only in some EU member states (because they had cases of the disease) and only in some NON-EU countries (not a disease reported in any international organisations) | |
| Score 4 Very low Absence of surveillance of the pathogen in ALL EU member countries AND world wide | |
| C6 Eradication experience in other countries and/or Belgium | |
| Score 0 | |
| Score 1 Very high Previous experience on eradication has been applied, fast and successfully | |
| Score 2 High Previous experience on eradicating the disease but with some setbacks in the process | |
| Score 3 Low Knowledge on eradication procedures but have never had to implement an eradication program in Belgium | |
| Score 4 Very low It is a novel disease, first time countries are faced with a new disease to eradicate | |
| 7 Detection of emergence - e.g. difficulties for the farmer/veterinarian to declare the disease or clinical signs not so evident. | $\overline{}$ |
| Score 0 | |

| Score 1 | Very high Disease is easily detected with clinically signs and farmers are aware of the disease and willing to notify it as soon as possible it |
|---------|---|
| Score 2 | High Disease is easily detected by the clinical signs but farmers don't have sufficient knowledge/awareness nor interest to notify it |
| Score 3 | Moderate Disease is not as easily detect by the clinical signs and farmers don't have sufficient knowledge/awareness nor interest to notify. |
| Score 4 | Low The infected animal does not show any pathognomonic clinical sign(s); farmer is reluctant to declare/notify any abnormality. |

Number of Criteria = 7, hence 70 points to be distributed within this domain for the intra-domain weighing

DOMAIN D. FARM/PRODUCTION SYSTEM CHARACTERISTICS Mana species forms. One single formed unimal (e.g. only begins)

| D1 | | ecies farms - One single farmed animal (e.g. only bovines) or multi species farms (farms with more than one species e.g. goats and bovines in the same |
|----|----------|---|
| | farm/lan | d/premises). |
| | Score 0 | |
| | Score 1 | Negligible: the type of farm does not influence in any form (re)emergence of the disease among the livestock population. |
| | Score 2 | Low: mono or multi species farm has a low effect on the risk of disease to emerge or re-emerge. |
| | Score 3 | Moderate: the type or types of farmed animals has a moderate effect on the emergence of the disease in Belgium. |
| | Score 4 | High: the type of farmed animals has a high influence for the disease to emerge and spread in Belgium. |
| D2 | | mography/management: such as type of dairy or beef (cattle) production. For pigs - reproduction, fattening, finishing farm or both . Chickens- only laying eggs chickens finishing broilers |
| | Score 0 | |
| | Score 1 | Negligible: population demography does not influence in any form the (re)emergence of the disease among the livestock population. |
| | Score 2 | Low: the demographic population of the farm is a low influencing factor for disease (re)emergence. E.g. Disease only clinically affects only one age strata (i.e.) newborns, therefore adults are immune to it. |
| | Score 3 | Moderate: the demographic of the population has a moderate effect on the (re)emergence of the disease, as it can (re)emerge in more than one type of demography but other conditioning factors have to occur in conjunction. |
| | Score 4 | High: the type of demographic of the farm has a high effect on the (re)emergence of the disease as it can (re)emerge in different types of farmed animals and all types of age groups |
| D3 | Animal d | lensity of farms. Extensive (small holders with a few animals) v/s intensive farming |
| | Score 0 | |
| | Score 1 | Negligible: animal farm density is not a risk factor for the disease to emerge in belgium |
| | Score 2 | Low: farm density (extensive or intensive) of animals has a <u>low effect</u> on the pathogen's/ disease (re)emergence |
| | Score 3 | Moderate: farm density of animals in the farm (extensive v/s intensive) has a moderate effect on the emergence of pathogen/disease |
| | Score 4 | High : farm density of animals has a high effect on the (re)emergence of pathogen/disease. |
| D4 | Feeding | practices of farms |
| | Score 0 | |
| | Score 1 | Negligible : Feeding practices have a negligible effect on the (re)emergence of the pathogen/disease |

| | Score 2 | Low: Feeding practices have a low effect on the (re)emergence of the pathogen/disease |
|----|---------|---|
| | Score 3 | Moderate: Feeding practices have a moderate effect on the (re)emergence of the pathogen/disease |
| | Score 4 | High: Feeding practices have a high effect on the (re)emergence of the pathogen/disease |
| D5 | Human | novements among premises - Veterinarians or farm staff. |
| | Score 0 | |
| | Score 1 | Negligible: disease is spread by other means |
| | Score 2 | Low: movement of human staff has a low effect on the introduction or spread of the disease |
| | Score 3 | Moderate: movement of human staff has a moderate effect on the introduction or spread of the disease |
| | Score 4 | High: movement of human staff has a high effect on the introduction or spread of the disease |
| D6 | | y of livestock farm to wildlife and wildlife reservoirs of disease e.g. contact with wild or feral birds and animals which have been scavenging on landfill sites that contain nated animal products |
| | Score 0 | • |
| | Score 1 | Negligible: Disease (re)emergence from wildlife and wildlife reservoir never reported. |
| | Score 2 | Low: Disease (re)emergence from wildlife and wildlife reservoir rarely reported. |
| | Score 3 | Moderate: Disease (re)emergence from wildlife and wildlife reservoir is documented regularly. |
| | Score 4 | High: wildlife is a reservoir for the disease and the main source of infection for livestock. |
| D7 | Changes | of land use, e.g. field fragmentation, creation of barriers, landfill sites. |
| | Score 0 | |
| | Score 1 | Negligible: Changes in land use have a negligible effect on the (re)emergence of pathogen/disease. |
| | Score 2 | Low: changes in land use have a low effect on the (re)emergence of the disease/pathogen but need other factors (e.g. land use changes combined with higher winter temperatures) |
| | Score 3 | Moderate: land use changes increases the availability of vectors or increases the pathogen's survival. Also empty land can create a suitable environment for certain wildlife carrying the disease (e.g. migratory birds) |
| | Score 4 | High: land use changes are one of the main drivers for pathogen or its vectors |

Number of Criteria = 7, hence 70 points to be distributed within this domain for the intra-domain weighing

DOMAIN E. CHANGES IN CLIMATIC CONDITIONS

| E1 | Influenc | e of annual <u>rainfall</u> in the survival and transmission of the pathogen/disease |
|----|----------|--|
| | Score 0 | |
| | Score 1 | Negligible: Pathogen survival and mode of transmission of the disease are not influenced by increased rainfall |
| | Score 2 | Low: pathogen survival and mode of transmission fo the disease are slightly influenced by increased rainfall |
| | Score 3 | Moderate: pathogen survival and mode of transmission of the disease are moderatly influenced by increased rainfall |
| | Score 4 | High: pathogen survival and mode of transmission of the disease are highly influenced by increased rainfall |

| E2 | Influenc | e of annual <u>humidity</u> in the survival and transmission of the pathogen/disease |
|-----------|----------|---|
| | Score 0 | |
| | Score 1 | Negligible: Pathogen survival and mode of transmission of the disease are not influenced by increased humidity |
| | Score 2 | Low: pathogen survival and mode of transmission of the disease are slightly influenced by increased humidity |
| | Score 3 | Moderate: pathogen survival and mode of transmission of the disease are moderatly influenced by increased humidity |
| | Score 4 | High: pathogen survival and mode of transmission of the disease are highly influenced by increased humidity |
| E3 | Influenc | e of annual <u>temperature</u> in the survival and transmission of the pathogen/disease |
| | Score 0 | |
| | Score 1 | Negligible: Pathogen survival and mode of transmission of the disease are not influenced by increased temperature |
| | Score 2 | Low: pathogen survival and mode of transmission fo the disease are slightly influenced by increased temperature |
| | Score 3 | Moderate: pathogen survival and mode of transmission of the disease are moderatly influenced by increased temperature |
| | Score 4 | High: pathogen survival and mode of transmission of the disease are highly influenced by increased temperature |

Number of Criteria = 3, hence 30 points to be distributed within this domain for the intra-domain weighing

DOMAIN F. WILDLIFE INTERFACE

| F1 | Potential r | oles of zoo's in the (re)emergence of the pathogen |
|----|-------------|---|
| | Score 0 | |
| | Score 1 | Negligible: The disease can be present in zoo animals but it is not known to have been transmitted from zoo animals to livestock. |
| | Score 2 | Low: The disease can enter a zoo (e.g. with introduction of an infected exotic animal) but only accidental transmissions of the disease from zoo animals to livestock have been reported. Hence, zoos have a low effect on the (re)emergence of the disease in Belgium's livestock |
| | Score 3 | Moderate: The disease can enter a zoo and be present in zoo animals but it needs a vector (biological/mechanical) for its transmission into livestock. Therefore, zoos have a moderate effect on the (re)emergence of the disease in Belgium. |
| | Score 4 | High: Disease can be introduced to a zoo via an infected imported animal, zoo animals can carry the disease that can easily jump to livestock animals |
| F2 | The rural(| farm)-wildlife interface |
| | Score 0 | |
| | Score 1 | Negligible: the disease has never (re)emerged from the narrowing of the farm-wild interface |
| | Score 2 | Low: the disease has a low probability to (re)emerge via the livestock farm-forest interface. The disease has been known to (re)emerge from the wild bush but very rarely |
| | Score 3 | Moderate: the disease has a moderate probability of (re)emergence via the farm/wildlife interface. Barriers (natural or artificial) are needed to keep the disease/pathogen (re)emerging in livestock |
| | Score 4 | High: there is a high probability for the disease to (re)emerge via the farm/forest interface. Barriers (natural or artificial) separating farms from natural forests are ineffective |
| F3 | Increase o | f autochthons (indigenous animal) wild mammals in Belgium and neighbouring countries |
| | Score 0 | Null: disease has not been reported in wildlife |
| | Score 1 | Negligible: the increase the autochthonous mammals population does not affect the risk of the diseases to (re)emergence |

| | Score 2 | Low: The slight increase of autochthonous mammals can slightly increase the probably of the disease emerging |
|----|------------------|--|
| | Score 3 | Moderate : The increase of wild mammals has been associated with the re-emergence of the disease |
| | Score 4 | High: The increase of wild mammals <u>IS the only factor</u> associated with outbreaks of the disease in livestock |
| F4 | Increase in | n endemic/migrating populations of wild birds. |
| | Score 0 | Null: Wild/migrating birds are not a reservoir of the disease |
| | Score 1 | Negligible: there is a negligible probability of disease (re)emerging in livestock because of an increase in populations of endemic/migrating wild birds. |
| | Score 2 | Low : there is a low probability of the disease (re)emerging and spreading through increased populations of endemic/migrating wild birds. Disease has spread from the endemic/migrating wild birds but only accidentally or under exceptional circumstances |
| | Score 3 | Moderate : there is a moderate probability of disease being introduced and spread through increased populations of endemic/migrating wild birds. They are hosts and in close contact with domestic livestock (i.e. poultry farms) may spread the disease |
| | Score 4 | High: there is a high probability for a disease to (re)emerge through increased populations of wild/migrating birds. These are hosts or reservoirs of the disease |
| F5 | Hunting A | ctivities: hunted animals can be brought back to where livestock is present |
| | Score 0 | |
| | Score 1 | Negligible: The risk of the disease/pathogen of (re)emerging in livestock due to hunting activities is practically null |
| | Score 2 | Low: disease is present in hunted wildlife and birds and only accidental cases have been reported in livestock that have (re)emerged because of hunting. The risk of the disease/pathogen of (re)emerging in livestock due to hunting activities is practically null |
| | Score 3 | Moderate: disease is present in hunted wildlife and birds but a certain control is established by the hunter |
| | Score 4 | High: disease is present in hunted wildlife and birds and hunting is one of the main modes of transmission of the disease to livestock |
| F6 | Transbou | ndary movements of terrestrial wildlife from other countries |
| | Score 0 | Null: Disease is not carried by terrestrial wildlife |
| | Score 1 | Negligible: (re)emergence of the disease by terrestrial movements of wildlife has only been suspected but never confirmed. |
| | Score 2 | Low: There is a low probability for the disease to (re)emerge and spread through transboundary movements of terrestrial wildlife |
| | Score 3 | Moderate: There is a moderate probability for the disease to (re)emerge and spread through transboundary movements of terrestrial wildlife |
| | Score 4 | High : There is a high probability for the disease to (re)emerge and spread through transboundary movements of terrestrial wildlife. These are host and may spread/carry the disease along. |

Number of Criteria = 6, hence 60 points to be distributed within this domain for the intra-domain weighing

DOMAIN G. HUMAN ACTIVITIES

| G1 | In- and out- people movements linked to tourism | | | | | | |
|--------------------------------|--|--|--|--|--|--|--|
| | Score 0 | | | | | | |
| | Score 1 | 77 78 77 77 77 77 77 77 77 77 77 77 77 7 | | | | | |
| | Score 2 | | | | | | |
| | Score 3 | | | | | | |
| | Score 4 | Highs towers may amont is a high driver on the (re) amorganes of a disease. Towers are highly likely to being the disease into Delegium in their belongings and highesqueity. | | | | | |
| G2 | Human Imi | n Immigration | | | | | |
| | Score 0 | | | | | | |
| | Score 1 | Negligible: the immigration movements are a negligible driver of the disease (re)emergence in Belgium | | | | | |
| | Score 2 | Low: the immigration movements are a low driver of the disease (re)emergence in Belgium | | | | | |
| | Score 3 Moderate: the disease is currently present in countries where more immigrants come from and pathogen highly likely to enter through, clothes, shoes and or pathogen by the current biosecurity measures in place are able to prevent the emergence of the disease in Belgium | | | | | | |
| | Score 4 | High: the immigration movement has a high effect as a driver on the emergence or re-emergence of disease in Belgium. Disease is highly likely to emerge using this route as biosecurity measures are not enough to avoid emergence of the disease | | | | | |
| G3 | Transport n | ort movements: more specifically commercial flights, commercial transport by ships, cars or military (EXCLUDING TRANSPORT VEHICLES OF LIVE ANIMALS). | | | | | |
| | Score 0 | | | | | | |
| | Score 1 | Negligible: the role of commercial movements as a driver on the (re)emergence of the disease in Belgium is negligible. | | | | | |
| | Score 2 | Low: the role of commercial movements as a driver on the (re)emergence of the disease in Belgium is low. It is easily preventable by implementing biosecurity measures | | | | | |
| | Score 3 | Moderate: the role of commercial movements as a driver on the (re)emergence of a disease in Belgium is moderate. Disease can be prevented if biosecurity measures are tightened. | | | | | |
| | Score 4 | High: the role of commercial movements as a driver on the (re)emergence of a disease in Belgium is high. Disease is hard to control via the current biosecurity measures. | | | | | |
| G4 | Transport v | sport vehicles of live animals | | | | | |
| | Score 0 | | | | | | |
| | Score 1 | Negligible: the role of transport vehicles of live animals as a driver for the (re)emergence of the disease in Belgium is negligible | | | | | |
| Score 2 Low: the role of trans | | Low: the role of transport vehicles of live animals as a driver for the (re)emergence of the disease in Belgium is low. | | | | | |
| 35015 5 | | Moderate : the role of transport vehicles of live animals as a driver for (re)emergence of the disease in Belgium is moderate. | | | | | |
| | Score 4 High: the role of transport vehicles of live animals as a driver for (re)emergence of the disease in Belgium is high | | | | | | |
| G5 | Bioterrorisi | ism potential | | | | | |
| | Score 0 | . 0 | | | | | |
| | Score 1 | Negligible : the role of bioterrorism as a driver for a disease to (re)emerge is negligible: agent is available but difficult to handle or has a low potential of spread or generates few economic consequences | | | | | |
| | Score 2 Low: the role of bioterrorism as a driver for a disease to (re)emerge is low: agent is available and easy to handle by professionals and labs but has a low spread | | | | | | |
| | Score 3 Moderate: the role of bioterrorism as a driver for a disease to (re)emerge is moderate: agent available and easy to handle by professionals and labs and rapidly spreads | | | | | | |
| | Score 4 High: the role of bioterrorism as a driver for a disease to (re)emerge is high: Agent is available and easy to handle by individuals and rapidly spreads | | | | | | |

| G6 | Inadvertent | Inadvertent release of an exotic infectious agent from a containment facility e.g. Laboratory | | | | |
|----|---|--|--|--|--|--|
| | Score 0 | | | | | |
| | Score 1 Negligible: the pathogen is not currently present in any laboratory | | | | | |
| | Score 2 | Score 2 Low: the pathogen is present in a containment facility but its release is very unlikely as it is very easily contained | | | | |
| | Score 3 Moderate: the pathogen is present in a containment facility and its release can occur as not easily contained | | | | | |
| | Score 4 | High: pathogen is handled in a risk 3 or 4 laboratory (BSL3 or BSL4) in the country. It can leave the facility if the correct biosecurity measures are not implemented correctly and easily spread to livestock | | | | |

Number of Criteria = 6, hence 60 points to be distributed within this domain for the intra-domain weighing

DOMAIN H. ECONOMIC AND TRADE ACTIVITIES

| Н1 | Decrease of | se of resources allocated to the disease surveillance | | | | | | |
|----|-------------|---|--|--|--|--|--|--|
| | Score 0 | | | | | | | |
| | Score 1 | Negligible: resources allocated to the disease surveillance have no effect on the (re)emergence of the disease in Belgium. Disease has never been under surveillance | | | | | | |
| | Score 2 | Low: resources allocated to the disease surveillance have a low effect on the (re)emergence of the disease in Belgium. Disease has been under surveillance in the past and no change has happened after surveillance has been stopped. | | | | | | |
| | Score 3 | Medium: resources allocated to the disease surveillance have a moderate effect on the (re)emergence of the disease in Belgium. Disease is under passive surveillance (reported only when observed) but with no need to further increase its surveillance | | | | | | |
| | Score 4 | High: resources allocated to the disease surveillance have a high effect on the (re)emergence of the disease in Belgium. Disease needs to be under active and passive surveillance as its (re)emergence can easily occur, therefore if its surveillance decreases it's highly likely to (re)emerge | | | | | | |
| Н2 | Modificatio | ion of the disease status (i.e. reportable disease becoming <u>not</u> reportable) or change in screening frequency due to a reduced national budget. | | | | | | |
| | Score 0 | | | | | | | |
| | Score 1 | Negligible: modification of the disease status due to a reduced national budget has a negligible effect on the (re) emergence of the disease in Belgium | | | | | | |
| | Score 2 | Low: modification of the disease status due to a reduced national budget has a low effect on the (re) emergence of the disease in Belgium | | | | | | |
| | Score 3 | Moderate: modification of the disease status due to a reduced national budget has a moderate effect on the (re) emergence of the disease in Belgium | | | | | | |
| | Score 4 | High: modification of the disease status due to a reduced national budget has a high effect on the (re) emergence of the disease in Belgium | | | | | | |
| Н3 | Decrease of | of resources allocated to the implementation of biosecurity measures at border controls (e.g. harbors or airports). | | | | | | |
| | Score 0 | | | | | | | |
| | | Negligible: decreasing the resources allocated to the implementation of biosecurity measures has a negligible effect on the (re)emergence of the disease in Belgium. Disease has never been detected in the past in a harbor or airport | | | | | | |
| | | Low: decreasing the resources allocated to the implementation of biosecurity measures has a low effect on the (re)emergence of the disease in Belgium. The disease has been suspected to have entered other countries because of deficient biosecurity at border controls. | | | | | | |
| | Score 3 | Medium: decreasing the resources allocated to the implementation of biosecurity measures has a moderate effect on the (re)emergence of the disease in Belgium. The disease has been introduced in other countries because of deficient biosecurity at border controls | | | | | | |
| | Score 4 | High: decreasing the resources allocated to the implementation of biosecurity measures highly increases the risk of (re)emergence of the disease in Belgium. In the past, the | | | | | | |

| Н4 | Most likely i Belgium. | Most likely influence of (il)legal movements of live animals (livestock, pets, horses etc) from neighbouring/European Union member states (MS) for the disease to (re)emerge in selgium. | | | | | | |
|----|---|--|--|--|--|--|--|--|
| | Score 0 | | | | | | | |
| | Score 1 | Negligible: (il)legal movements of live animals (livestock, pets, horses etc) from neighbouring/European Union MS have a <u>negligible influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 2 | Low: (il)legal movements (livestock, pets, horses etc) from neighbouring/European Union MS have a <u>low influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 3 | Moderate: (il)legal movements (livestock, pets, horses etc) from neighbouring/European Union MS have a moderate influence on the pathogen/disease (re)emergence Belgium. | | | | | | |
| | Score 4 | High: (il)legal movements (livestock, pets, horses etc.) from neighbouring/European Union MS have a high influence on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| Н5 | Influence of | increased (il)legal imports of animal subproducts such as skin, meat and edible products from EU member states for the disease/pathogen to (re)emerge in Belgium | | | | | | |
| | Score 0 | | | | | | | |
| | Score 1 | Negligible: increased (il)legal imports of animal subproducts such as skin, meat and edible products from EU member states have a <u>negligible influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 2 | Low: increased (il)legal imports of animal subproducts such as skin, meat and edible products from EU member states have a <u>low influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 3 | Moderate: increased (il)legal imports of animal subproducts such as skin, meat and edible products from EU member states have a <u>moderate influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 4 | High: increased (il)legal imports of animal subproducts such as skin, meat and edible products from EU member states have a <u>high influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| Н6 | Most likely i Belgium. | influence of increased (il)legal imports of NON-animal products such as tires, wood, furniture from EU member states for the disease/pathogen to (re)emerge in | | | | | | |
| | Score 0 | | | | | | | |
| | Score 1 | Negligible: increased (il)legal imports of NON-animal products such as tires, wood, furniture from EU member states have a <u>negligible influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 2 | Low: increased (il)legal imports of NON-animal products such as tires, wood, furniture from EU member states have a <u>low influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 3 | Moderate: increased (il)legal imports of NON-animal products such as tires, wood, furniture from EU member states have a <u>moderate influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | High: increased (il)legal imports of NON-animal products such as tires, wood, furniture from EU member states have a <u>high influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | | |
| Н7 | Most likely i | nfluence of (il)legal movements of live animals (livestock, pets, horses etc) from Third countries for the disease to (re)emerge in Belgium. | | | | | | |
| | Score 0 | | | | | | | |
| | Score 1 | Negligible: (il) legal movements of live animals (livestock, pets, horses etc) from Third countries have a <u>negligible influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 2 | Low: (il)legal movements of live animals (livestock, pets, horses etc) from Third countries have a <u>low influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 3 | Moderate: (il)legal movements of live animals (livestock, pets, horses etc) from Third countries have a <u>moderate influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 4 | High: (il)legal movements of live animals (livestock, pets, horses etc) from Third countries have a <u>high influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| Н8 | Most likely i | influence of increased imports of animal subproducts such as skin, meat and edible products from Third countries, for the disease to (re)emerge in Belgium. | | | | | | |
| | Score 0 | | | | | | | |
| | Score 1 | Negligible: Increased imports of animal subproducts such as skin, meat and edible products from Third countries have a <u>negligible influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |

| | Score 2 | Low: Increased imports of animal subproducts such as skin, meat and edible products from Third countries have a <u>low influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
|--|--|--|--|--|--|--|--|--|
| | Score 3 Moderate: Increased imports of animal subproducts such as skin, meat and edible products from Third countries have a moderate influence on the pathogen/of (re)emergence in Belgium. | | | | | | | |
| | Score 4 | High: Increased imports of animal subproducts such as skin, meat and edible products from Third countries have a high influence on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| Н9 | Most likely influence of increased (il)legal imports of NON-animal products such as tires, wood, furniture from Third countries, for the disease to (re)emerge in Belgium. | | | | | | | |
| | Score 0 | | | | | | | |
| Score 1 Negligible: increased (il)legal imports of NON-animal products such as tires, wood, furniture from Third countries have a <u>negligible influence</u> on the pathoge (re)emergence in Belgium. | | | | | | | | |
| Score 2 Low: increased (il)legal imports of NON-animal products such as tires, wood, furnished Belgium. | | Low: increased (il)legal imports of NON-animal products such as tires, wood, furniture from Third countries have a <u>low influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| Score 3 Moderate: increased (il)legal imports of NON-anima (re)emergence in Belgium. | | Moderate: increased (il)legal imports of NON-animal products such as tires, wood, furniture from Third countries have a <u>moderate influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |
| | Score 4 | High: increased (il)legal imports of NON-animal products such as tires, wood, furniture from Third countries have a <u>high influence</u> on the pathogen/disease (re)emergence in Belgium. | | | | | | |

Legend: Number of Criteria = 9, hence 90 points to be distributed within this domain for the intra-domain weighing.

Appendix 2. List of experts enrolled (N= 14) in the phase I (questionnaire assessment) with their gender, affiliation, country and field of expertise

| Expert | Gender | Institution | Background | Country | Field of expertise | Keywords |
|-------------------------------|--------|----------------------------|--|---------|--|---|
| Kris de Clercq | M | Sciensano | DVM, MSc, PhD, Head of Unit, Sciensano | Belgium | Exotic viruses and transmissible spongiform | Exotic diseases |
| Philippe Leonard | M | University Hospital Center | Medical doctor | Belgium | encephalopathies Infectious diseases | Travel medicine |
| Dirk Berkvens | M | University | Ir, PhD, Institute of Tropical Medicine , Antwerp | Belgium | Epidemiology and quantitative risk analysis | Veterinary |
| Etienne Thiry | M | University | DVM, PhD, Dipl. ECVPH, Professor, Liege University | Belgium | Virology and viral diseases | Veterinary |
| Nathalie Kirschvink | F | University | | Belgium | Animal physiology | Arboviruses |
| Thierry van den Berg | M | Sciensano | DVM, MSc, PhD, Operational Director Viral diseases at Sciensano | Belgium | Viral diseases, Avian influenza, Newcastle , Schmallenberg | Avian viruses, viral diseases |
| Christian Gortazar Schmidt | M | University | DVM, PhD, Professor at the University of Castilla - La Mancha, Spain. Head of SaBio (Sanidad y | Spain | Diseases and | Population dynamics, Epidemiology, Ecology, animal health |

Biotechnologia) of

IREC

| Hendriks Pascal | M | Anses | DVM, PhD, France | Animal health, | Surveillance |
|-----------------|-----|--------|------------------------|----------------------|------------------|
| Hendriks Fascar | IVI | Allses | | | |
| | | | Scientific director of | surveillance, | systems |
| | | | epidemiology and | veterinary | |
| | | | surveillance | epidemiology | |
| Fabiana Dal | F | AMCRA | DVM, MSc, PhD, Belgium | Viral diseases, | Viral diseases, |
| Pozzo | | | Scientific | Bluetongue, | arboviruses, |
| | | | Coordinator at | laboratory | Antibiotic |
| | | | AMCRA | diagnostics, Q | resistance |
| | | | | fever | |
| Morgane | F | OIE | DMV, PhD, OIE France | Epidemiology, | Veterinary |
| Dominguez | | | project officer | Risk analysis in | epidemiology, |
| | | | | veterinary sciences | biosecurity |
| Boelaert Frank | M | EFSA | DVM, MSc, PhD, Italy | Zoonoses, public | Surveillance, EU |
| | | | Dipl. ECVPH, | health, surveillance | surveillance |
| | | | Senior Scientific | of zoonoses and | |
| | | | Office at the | food-borne | |
| | | | Biological hazards | outbreaks | |
| | | | and contaminants | | |
| | | | Unit of EFSA | | |
| Vanholme Luc | M | FASFC | DVM, Federal Belgium | Veterinary | Animal diseases |
| | | | agency for the Safety | medicine, Animal | |
| | | | of the Food Chain, | diseases, Control | |
| | | | General Direction of | policy | |
| | | | Control policy | | |
| | | | • | | |

| Laetitia | F | University | DVM, PhD | , Dipl. | Belgium | Parasitology, | Tick-borne |
|------------------|---|------------|---------------|-----------|---------|-------------------|-----------------|
| Lempereur | | | EVPC, | Assistant | | Vector-borne | animal diseases |
| | | | Professor | of | | diseases | |
| | | | parasitology, | Liege | | | |
| | | | University | | | | |
| Depoorter Pieter | M | FASFC | DMV, | Federal | Belgium | Veterinary | Animal diseases |
| | | | Agency fo | or the | | medicine, Animal | |
| | | | Safety of th | e Food | | diseases, Control | |
| | | | Chain, | General | | policy | |
| | | | Direction of | Control | | | |
| | | | Policy, | Risk | | | |
| | | | Direction | | | | |
| | | | | | | | |

Appendix 3. List of experts enrolled (N=62) in phase II (disease prioritization) with their gender, affiliation, country, field of expertise, and disease they answered for

| Expert | Condon | Institution | Background | Country | Field of | Keywords | Disease expert |
|-------------|--------|-------------|----------------|---------|-------------------|---------------|------------------|
| Expert | Genuer | msutution | Dackground | Country | expertise | Keywords | answered |
| Agnes Waret | F | University | DVM, | France | Epidemiology of | Animal health | Peste des petits |
| | | | MSc,PhD, | | animal infectious | | Ruminants |
| | | | Assistant | | diseases in | | |
| | | | Lecturer, | | southern | | |
| | | | Swine | | countries, animal | | |
| | | | production and | | health economy | | |
| | | | pathology, | | | | |
| | | | University of | | | | |
| | | | Toulouse, | | | | |
| | | | France | | | | |
| Alexandre | M | CIRAD | DVM, PhD, | France | Disease ecology | Disease | Peste des petits |
| Caron | | | CIRAD-UPR | | at the | ecology | Ruminants |
| | | | AGIRs | | wildlife/domestic | | |
| | | | | | interface in | | |
| | | | | | border | | |
| | | | | | conservation | | |
| | | | | | areas, thinking | | |
| | | | | | sustainable and | | |
| | | | | | resilient socio- | | |
| | | | | | ecosystems in | | |
| | | | | | borders of | | |
| | | | | | conservation | | |
| | | | | | areas | | |
| Ana Alba | F | CReSA | DVM, PhD, | Spain | Data Mining and | West Nile | West Nile Fever |
| Casals | | | Epidemiology | | knowledge | Fever | |
| | | | Unit, CReSA | | discovery | | |
| | | | om, Cresa | | uiscovery | | |

| Ana de la | F | University | DMV, PhD, | Portugal | Virology and | African horse | African horse |
|---------------|---|------------|-----------------|----------|----------------|----------------|-----------------|
| Grandière | | | Department of | | viral diseases | sickness | sickness |
| | | | infectious and | | | | |
| | | | parasitic | | | | |
| | | | diseases, Liege | | | | |
| | | | University | | | | |
| Ana Sofia | F | University | DMV, MSC, | Germany | Infectious | Infectious | Contagious |
| Ramirez | | | Heidelberg | | Diseases, | diseases | Bovine |
| | | | University, | | Epidemiology, | | Pleuropneumonia |
| | | | Germany | | Ventilation, | | Contagious |
| | | | | | Tuberculosis, | | Caprine |
| | | | | | Airway | | pleuropneumonia |
| | | | | | obstruction | | |
| Andrea | M | CIRAD | M.A., Physics, | France | Modelling of | Computational | Contagious |
| Apolloni | | | Phd, | | infectious | epidemiology | Bovine |
| | | | Researcher at | | diseases | | Pleuropneumonia |
| | | | CIRAD | | | | Contagious |
| | | | | | | | Caprine |
| | | | | | | | pleuropneumonia |
| Anette Botner | F | DTU VET | DMV, PhD, | Denmark | Veterinary | Viral diseases | Porcine |
| | | | Division of | | virology | | Epidemic |
| | | | Diagnostics & | | | | Diarrohea |
| | | | Scientific | | | | |
| | | | Advice - | | | | |
| | | | Virology, | | | | |
| | | | National | | | | |
| | | | Veterinary | | | | |
| | | | Institute | | | | |

| Ann Brigitte | F | Sciensano | Bio Engineer, | Belgium | Molecular | Horse diseases | West Nile fever |
|--------------|---|-------------|-----------------|---------|-------------------|----------------|-------------------|
| Cay | | | PhD, Head of | | Biology, | | |
| | | | Unit Enzootic | | Molecular | | |
| | | | and Re- | | Cloning, Cell | | |
| | | | emerging viral | | Biology, | | |
| | | | diseases, | | Infection | | |
| | | | Sciensano | | | | |
| Annelise | F | CIRAD | PhD, Animal et | France | Spatial Analysis, | Arboviruses | Rift Valley fever |
| Tran | | | Gestion | | Remote Sensing, | | |
| | | | Intégrée des | | Geographic | | |
| | | | Risques | | Information | | |
| | | | (AGIRS), | | System, | | |
| | | | CIRAD | | Environmental | | |
| | | | | | science | | |
| Axel Mauroy | M | University | DVM, PhD, | Belgium | Virology, Viral | Arboviruses | Aino, akabane, |
| | | | Assistant | | diseases | | Low pathogenic |
| | | | Professor of | | | | avian influenza, |
| | | | Veterinary | | | | High pathogenic |
| | | | Virology at the | | | | avian influenza, |
| | | | University of | | | | Porcine epidemic |
| | | | Liege | | | | diarrhoea, |
| | | | | | | | Schmallenberg, |
| | | | | | | | Vesicular |
| | | | | | | | stomatitis |
| Bart Pardon | M | Ghent | DVM, PhD, | Belgium | Internal | Respiratory | Haemorragic |
| | | University, | Dip ECBHM, | | Medicine, | Diseases, | Septicaemia |
| | | Assistant | Ghent | | Infectious | Internal | |
| | | | University, | | Diseases | Medicine, | |
| | | | Doctor | | | Infectious | |
| | | | Assistant of | | | Diseases | |

| | | | : | | | | |
|-------------|---|------------|------------------|---------|------------------|---------------|----------------|
| | | | internal | | | | |
| | | | medicine of | | | | |
| | | | large animals at | | | | |
| | | | Ghent | | | | |
| | | | University. | | | | |
| Bénédicte | M | Sciensano | DVM, PhD, | Belgium | Avian virology | Newcastle | Newcastle |
| Lambrecht | | | Head of | | and immunology | disease | disease |
| | | | Scientific | | | | |
| | | | Service Avian | | | | |
| | | | virology and | | | | |
| | | | immunology, | | | | |
| | | | Sciensano | | | | |
| Benoît | M | ANSES | DVM, MSc, | France | Epidemiology | Animal | Western Equine |
| Durand | | | PhD, | | unit | diseases, | Encephalitis, |
| | | | Epidemiology | | | modelling | Eastern Equine |
| | | | unit, ANSES | | | | Encephalitis, |
| | | | | | | | Venezuelan |
| | | | | | | | Equine |
| | | | | | | | Encephalitis, |
| | | | | | | | Foot and mouth |
| | | | | | | | disease |
| Benoit | M | University | DVM, PhD, | Belgium | Virology (herpes | Arboviruses | Akabane |
| Muylkens | | | Professor at the | | virus, | | |
| | | | University of | | vaccination) | | |
| | | | Namur | | control of viral | | |
| | | | | | genetics | | |
| | | | | | expression | | |
| Cecile Beck | F | ANSES | DVM, PhD, | France | Virology | Antibodies, | Venezuelan |
| | | | Laboratory of | | | ELISA, Virus, | equine |
| | | | | | | Vaccination | encephalitis |
| | | | | | | | |

| | | | animal health, | | | | |
|------------|---|------------|-----------------|----------|------------------|-----------------|------------------|
| | | | ANSES | | | | |
| Chris Oura | M | University | DVM, PhD, | Trinidad | Virology, One- | Exotic diseases | African Swine |
| | | | Senior lecturer | and | Health, Zoonotic | | fever |
| | | | in Veterinary | Tobago | and animal | | |
| | | | Virology, | | pathogens, | | |
| | | | University of | | Emerging | | |
| | | | the West | | infectious | | |
| | | | Indies, | | diseases | | |
| | | | Trinidad and | | | | |
| | | | Tobago | | | | |
| Christelle | F | ANSES | DEA, Biology | France | Epidemiologist, | Epidemiology, | Novel swine |
| Fablet | | | and production | | Animal | One health | enteric |
| | | | animals, PhD, | | Productions, | initiative. | coronavirus |
| | | | Epidemiologist | | Respiratory | | |
| | | | at ANSES | | Diseases, Swine | | |
| Dirk | M | University | Ig., MSc, PhD, | Belgium | Epidemiology | Epidemiology, | Bluetongue, Rift |
| Berkvens | | | Institute of | | and quantitative | modelling | Valley fever |
| | | | Tropical | | risk analysis | | |
| | | | Medicine, | | | | |
| | | | Antwerp | | | | |
| Ducatez | F | University | DVM, PhD, | France | PCR, | Influenza | Low pathogenic |
| Mariette | | | Host-pathogen | | Genotyping, | viruses | avian influenza, |
| | | | interaction, | | Emerging | | High pathogenic |
| | | | University of | | Infectious | | avian influenza |
| | | | Toulouse | | Diseases, Viral | | |
| | | | | | infection | | |
| Ethienne | M | University | DVM, PhD, | Belgium | Virology | Virus, Animal, | Aino, Akabane, |
| Thiry | | | University | | | emerging | Vesicular |
| | | | Professor, Unit | | | | stomatitis |
| | | | | | | | |

| | | | of Virology | | | diseases, | |
|-------------|---|-------|-----------------|---------|--------------------|------------------|------------------|
| | | | and Viral | | | genetics | |
| | | | Diseases, | | | | |
| | | | University of | | | | |
| | | | Liège | | | | |
| Emmanuel | M | ANSES | DMV, PhD, | France | PCR, Cell | Arboviruses | Bluetongue, |
| Bread | | | Laboratory for | | culture, | | Epizootic |
| | | | Animal Health, | | Infection, | | haemorragic |
| | | | ANSES | | Immunology of | | disease, |
| | | | | | infectious | | Schmallenberg |
| | | | | | diseases | | |
| Fabiana Dal | F | AMCRA | DVM, MSc, | Belgium | Viral diseases, | Viral diseases, | African horse |
| Pozzo | | | PhD, Scientific | | bacterial diseases | poxviruses, | sickness, |
| | | | Coordinator at | | | arboviruses, | Bluetongue, |
| | | | AMCRA | | | antibiotics | Epizootic |
| | | | | | | resistance | haemorragic |
| | | | | | | | diseases, Sheep |
| | | | | | | | and goat pox |
| Francois | M | CIRAD | DVM, MSc, | France | Epidemiology, | One Health | Peste des petits |
| Roger | | | PhD, Animals, | | Infectious | | Ruminant |
| | | | Health, | | diseases, | | |
| | | | Territories, | | Biostatistics | | |
| | | | Risks and | | | | |
| | | | Ecosystems | | | | |
| | | | Unit, CIRAD | | | | |
| Francois | M | CIRAD | DVM, PhD, | France | Animal Science, | Animal | Contagious |
| Thiaucourt | | | Researcher at | | Cattle, Vaccine | Science, Cattle, | Bovine |
| | | | CIRAD | | Development | Diagnostics, | Pleuropneumonia |
| | | | | | | Molecular | , Contagious |

| | | | | | | Biological | Caprine |
|--------------|-----|------------|-----------------|-----------|------------------|-----------------|-------------------|
| | | | | | | Techniques | pleuropneumonia |
| Frank | M | Sciensano | DVM, PhD, | Belgium | Surveillance, | Classical Swine | African Swine |
| Koenen | 171 | Sciensano | One Health | Deigium | Swine diseases | | |
| Koenen | | | | | Swille diseases | Fever, African | Fever, Classical |
| | | | Unit, Sciensano | | | Swine Fever | swine fever |
| Gaby Van | F | University | DVM, MSc, | Australia | Equine medicine | Internal | African horse |
| Galen | | | PhD, DES, | | | Medicine and | sickness, Eastern |
| | | | Dipl. ECEIM, | | | Surgery | equine |
| | | | Dipl ECVECC, | | | | encephalitis, |
| | | | Associate | | | | Western equine |
| | | | Professor, | | | | encephalitis, |
| | | | University of | | | | Japanese |
| | | | Sidney | | | | encephalitis |
| Gilles Meyer | M | University | DMV, PhD, | France | Veterinary | Veterinary | Aino, |
| | | | ECBHM, | | Virology, Viral, | virology, | Schmallenberg |
| | | | University of | | Ruminant | vector-borne | |
| | | | Toulouse, | | Pathology | diseases | |
| | | | Professor | | | | |
| Grasland | F | ANSES | PhD, ANSES | France | Swine virology | Virology, | Novel swine |
| Beatrice | | | | | and diseases | Nomenclature, | enteric |
| | | | | | | Swine | coronavirus |
| | | | | | | Diseases, | |
| | | | | | | PRRS | |
| Guy | M | ARSIA | DVM, MSc, | Belgium | Laboratory | Animal | Foot and mouth |
| Czaplicki | | | Head of a | | diagnosis | serology, | disease, swine |
| | | | veterinary | | | bovine | vesicular |
| | | | diagnostic | | | pathology, | diseases, |
| | | | laboratory | | | swine | vesicular |
| | | | , | | | pathology, | stomatitis |
| | | | | | | epidemiology, | |
| | | | | | | epidennology, | |

| | | | | | | animal | |
|---------------|---|------------|------------------|---------|-------------------|-----------------|-----------------|
| | | | | | | infectiology | |
| | | | | | | | |
| Guy-Pierre | M | University | DVM, PhD, | France | Medicine and | Pig production | Novel swine |
| Martineau | | | Diplomate of | | porcine | | enteric |
| | | | ECPHM, | | production | | coronavirus, |
| | | | Professor at the | | | | Swine vesicular |
| | | | National | | | | disease |
| | | | Veterinary | | | | |
| | | | School of | | | | |
| | | | Toulouse | | | | |
| Ignacio | M | Universty | DVM, PhD, | Spain | Animal health, | Wildlife | West Nile Fever |
| Garcia | | | Dip. ECZM, | | wildlife | population | |
| Bocanegra | | | Professor of | | population health | health | |
| | | | animal Health | | | | |
| | | | at the | | | | |
| | | | University of | | | | |
| | | | Cordoba, Spain | | | | |
| James Wood | M | University | DVM, MSc, | United | Epidemiology, | Horse diseases, | African horse |
| | | | PhD, Dipl. | Kingdom | infection | Bat ecology | Sickness, Nipah |
| | | | ECVPH, | | dynamic, control | | virus |
| | | | Professor, | | of diseases in | | |
| | | | Department of | | Africa and | | |
| | | | Veterinary | | globally | | |
| | | | Medicine, | | | | |
| | | | University of | | | | |
| | | | Cambridge | | | | |
| Jaques Mainil | M | University | DVM, PhD, | Belgium | Bacteriology | Bacteriology, | Haemorragic |

Professor,

Bacteriology

Septicaemia

pathogeny,

genetics

| | | | and | | | (prokaryotes), | |
|----------------|---|------------|-------------------|---------|-------------------|----------------|--------------------|
| | | | Bacteriologic | | | molecular | |
| | | | Diseases, | | | epidemiology, | |
| | | | University of | | | plasmidology | |
| | | | Liège | | | | |
| Jean Guillotin | M | Departeman | DMV, | France | Diagnosis of | Swine diseases | Classical swine |
| | | tal | Departmental | | animal diseases | | fever |
| | | laboratory | laboratory | | | | |
| Jean Pierre | M | University | DMV, PhD, | France | Mandatory | Animal | Peste des petits |
| Ganière | | | Oniris | | diseases | diseases | Ruminants |
| Jean-Pierre | M | University | DVM, MSc, | Canada | Epidemiology of | Public health, | Newcastle |
| Vaillancourt | | | PhD, Professor | | zoonosis and | biosecurity | disease |
| | | | titulaire, | | public health, | | |
| | | | University of | | Infectious | | |
| | | | Montreal | | diseases of swine | | |
| | | | | | and poultry | | |
| Jordi Casal | M | University | DVM, | Spain | Animal Health | Animal | Foot and mouth |
| | | | University | | | epidemiology, | disease, lumpy |
| | | | Professor, | | | zoonoses, | skin disease, Rift |
| | | | Universidad | | | biosecurity | valley fever, |
| | | | Autonoma de | | | | vesicular |
| | | | Barcelona | | | | stomatitis |
| Joseph | M | FASFC | DVM, MSc, | Belgium | Animal diseases, | Epidemic | African swine |
| Hooyberghs | | | Federal agency | | virology | diseases | fever, classical |
| | | | for safety of the | | | | swine fever, |
| | | | food chain, | | | | porcine epidemic |
| | | | General | | | | diarrhoea |
| | | | Direction of | | | | |
| | | | Control Policy | | | | |

| Julien | M | CIRAD | DVM, PhD, | France | Wildlife ecology | Ecology, | Nipah virus |
|------------|---|------------|----------------|-----------|--------------------|-----------------|-----------------|
| Cappelle | | | Health | | | epidemiology, | |
| | | | Ecologist, | | | Wildlife | |
| | | | CIRAD | | | | |
| Kris De | M | Sciensano | DVM, MSc, | Belgium | Exotic viruses | Exotic diseases | Foot and mouth |
| Clercq | | | EU Reference | | and transmissible | | disease, lumpy |
| | | | Laboratory for | | spongiform | | skin disease, |
| | | | FMD viruses, | | encephalopathies | | sheep and goat |
| | | | Sciensano | | | | pox |
| Labib | M | ANSES | DVM, PhD, | France | Virology, | Laboratory, | Foot and mouth |
| Bakkali | | | Head of FAO | | immunology, | Foot and mouth | disease |
| Kassimi | | | reference | | molecular | disease | |
| | | | centre and OIE | | biology | | |
| | | | reference | | | | |
| | | | laboratory for | | | | |
| | | | FMD at | | | | |
| | | | ANSES | | | | |
| Lecoq | F | University | DVM, DES, | Belgium | Equine medicine | Horse diseases | Japanese |
| Laureline | | | MSc, Dipl. | | | | encephalitis |
| | | | ACVIM | | | | |
| Louis | M | University | DMV, MSc, | Australia | Management of | Animal | Contagious |
| Lignereux | | | Liege | | wildlife diseases, | diseases | caprine |
| | | | University | | Animal diseases | | Pleuropneumonia |
| Ludovic | M | University | DVM, MSc, | Belgium | Epidemiology, | Pathogenesis, | Aino, Akabane, |
| Martinelle | | | PhD, Head of | | patogenesis of | Bluetongoue, | Epizootic |
| | | | the | | Bluetongue and | Schmallenberg | haemorragic |
| | | | Experimental | | Shmallenberg | | disease |
| | | | Station | | | | |
| | | | (CARE-FePex) | | | | |
| | | | | | | | |

| ot | Liogo |
|----|-------|
| aı | Liege |

| | | | 2 | | | | |
|--------------|---|------------|------------------|---------|------------------|---------------|-------------------|
| | | | University | | | | |
| Marie-France | F | University | DVM, MSC, | Belgium | Biosecurity, | Biosecurity, | Japanese |
| Humblet | | | PhD, | | epidemiology | Hygiene, | encephalitis, |
| | | | Department of | | | Epidemiology | Newcastle |
| | | | Occupational | | | | disease, |
| | | | Protection and | | | | Venezuelan |
| | | | Hygiene, | | | | equine |
| | | | Biosafety and | | | | encephalitis, |
| | | | Biosecurity | | | | West Nile fever |
| | | | section, Liege | | | | |
| | | | University | | | | |
| Marilena | F | Sciensano | DVM, PhD, | Belgium | Veterinary | Disease | Rift Valley fever |
| Filippitzi | | | Dipl. ECVPH, | | epidemiology, | surveillance, | |
| | | | Veterinary | | Risk assessment, | Antimicrobial | |
| | | | epidemiology, | | Antimicrobial | resitance | |
| | | | Sciensano | | resistance, | | |
| | | | | | Biosecurity | | |
| Marius | M | University | Applied | Belgium | Spatial | Ecology, | Low pathogenic |
| Gilbert | | | Biological | | epidemiology of | population | avian influenza, |
| | | | Sciences, PhD, | | animal diseases | biology, | High pathogenic |
| | | | Head of spatial | | | | avian influenza |
| | | | epidemiology | | | | |
| | | | Lab, FNRS | | | | |
| | | | Research | | | | |
| | | | Associate at the | | | | |
| | | | Universite | | | | |
| | | | Libre de | | | | |
| | | | Bruxelles. | | | | |
| | | | | | | | |

| Marylene | F | Sciensano | Lic., MSc, | Belgium | Veterinary | Diagnosis | African horse |
|--------------|---|------------|---------------|---------|-------------------|----------------|------------------|
| Tignon | | | PhD, Virology | | virology, | | sickness |
| | | | Department, | | Porcine, bovine | | |
| | | | Sciensano | | and horse viral | | |
| | | | | | diseases | | |
| Mutien-Marie | M | University | DVM, PhD, | Belgium | Pathologist of | Influenza, | Bluetongue, |
| Garigliany | | | Dipl. ECVP, | | infectious | Pathology | Epizootic |
| | | | General | | disease, avian | | haemorragic |
| | | | pathology, | | influenza | | disease, |
| | | | Liege | | | | Schmallenberg |
| | | | University | | | | |
| Nick De | M | Sciensano | DMV, PhD, | Belgium | Infectious animal | Vector-borne | Western Equine |
| Regge | | | Virology | | diseases, | diseases, | Encephalitis, |
| | | | Department, | | Enzootic and | Arthropod | Eastern Equine |
| | | | Sciensano | | vector-borne | vectors | Encephalitis, |
| | | | | | diseases. | | Venezuelan |
| | | | | | | | Equine |
| | | | | | | | Encephalitis, |
| | | | | | | | Swine vesicular |
| | | | | | | | diseases, |
| | | | | | | | vesicular |
| | | | | | | | stomatitis |
| Nicolas Rose | M | ANSES | DVM, PhD, | France | Swine | Epidemiology, | African swine |
| | | | Swine | | epidemiology | Animal welfare | fever, Classical |
| | | | Epidemiology | | | | swine fever, |
| | | | and Welfare | | | | Novel swine |
| | | | Unit, ANSES | | | | enteric |
| | | | | | | | coronavirus, |
| | | | | | | | Porcine epidemic |
| | | | | | | | diarrhoea |

| Patrick | M | University | DVM, PhD, | Belgium | Microbiology | Microbiology, | Heamorragic |
|---------------|---|------------|-----------------|---------|-----------------|----------------|-----------------|
| Butaye | | | School of | | | Antimicrobial | septicaemia |
| | | | Veterinary | | | resistance | |
| | | | Medicine, Ross | | | | |
| | | | University | | | | |
| Paul Kitching | M | The | DMV, PhD, | United | Virology | Poxviruses | Lumpy skin |
| | | Pirbright | The Pirbright | Kingdom | | | disease, sheep |
| | | Institute | Institute | | | | and goat pox |
| Philippe | M | CIRAD | DVM, PhD, | France | Virology, | Poxviruses | Lumpy skin |
| Caufour | | | Department | | Immune response | | disease, sheep |
| | | | BIOS, CIRAD | | | | and goat pox |
| Ruben | M | University | DMV, PhD, | Spain | Veterinary | Infectious | Contagious |
| Rosales | | | Universidad de | | science, | diseases | Bovine |
| | | | Las Palmas de | | Veterinary | | Pleuropneumonia |
| | | | Gran Canaria | | diagnostics, | | , Contagious |
| | | | | | Veterinary | | Caprine |
| | | | | | infectious | | pleuropneumonia |
| | | | | | diseases, | | |
| | | | | | Veterinary | | |
| | | | | | epidemiology | | |
| Stephan | M | Anses | DVM, MSc, | France | Virology | Foot-and | Bluetongue, |
| Zientara | | | PhD, Head of | | | mouth disease, | Epizootic |
| | | | Virology and | | | Bluetongue | haemorragic |
| | | | of the National | | | West Nile | disease |
| | | | Reference | | | Fever, Equine | |
| | | | Laboratory for | | | viral diseases | |
| | | | Foot-and | | | | |
| | | | Mouth Disease, | | | | |
| | | | Bluetongue, | | | | |
| | | | West Nile and | | | | |

African Horse

Sickness

| Steven Van Gutch | M | Sciensano | DVM, MSC, PhD, Head of Viral Diseases, Sciensano | Belgium | Virology | Bat diseases | Nipah virus |
|------------------|---|------------|--|-----------|------------------|----------------|------------------|
| Sylvie | F | ANSES | DVM, PhD, | France | PCR, Infection, | Viruses, | Western equine |
| Lecollinet | | | Laboratory for | | ELISA, Viral | Equine | encephalitis, |
| | | | Animal health, | | Infection | Medicine | Eastern equine |
| | | | ANSES | | | | encephalitis, |
| | | | | | | | Japanese |
| | | | | | | | encephalitis, |
| | | | | | | | Venezuelan |
| | | | | | | | equine |
| | | | | | | | encephalitis, |
| | | | | | | | West nile fever |
| Thierry van | M | Sciensano | DMV, PhD, | Belgium | Viral diseases, | Avian viruses, | Low pathogenic |
| den Berg | | | MSc, | | Avian influenza, | viral disease | avian influenza, |
| | | | Operational | | Newcastle | | High pathogenic |
| | | | Director Viral | | | | avian influenza, |
| | | | diseases at | | | | Newcastle |
| | | | Sciensano | | | | |
| Thomas | M | University | DMV, PhD, | The | Biology, | Swine diseases | Swine vesicular |
| Hagennarts | | | Bacteriology | Netherlan | Ecology, | | disease |
| | | | and | ds | Epidemiology, | | |
| | | | Epidemiology, | | Mathematics, | | |
| | | | | | | | |

| | | | University of | | Veterinary | | |
|-------------|---|-----------|-----------------|----------|-----------------|------------------|-------------|
| | | | Wageningen | | science | | |
| Pierre | M | Sciensano | Bachelor | Belgium | Laboratory | Laboratory | Heamorragic |
| Wattiau | | | degree in | | techniques, | Microbiology | septicaemia |
| | | | industrial | | Bacterial | | |
| | | | Chemistry, | | isolation and | | |
| | | | MSc, PhD, | | identification, | | |
| | | | Veterinary | | Antibiotic | | |
| | | | bacteriology | | susceptibility | | |
| | | | Department, | | testing, | | |
| | | | Sciensano | | Molecular | | |
| | | | | | detection | | |
| Weerapong | M | Ministry | DVM, PhD, | Thailand | Animal Health, | Spatial analysis | Nipah virus |
| Thanapongth | | | Senior | | livestock | | |
| arm | | | Veterinary | | development | | |
| | | | Office at | | | | |
| | | | Ministry of | | | | |
| | | | Agriculture and | | | | |
| | | | Cooperatives, | | | | |
| | | | Thailand | | | | |

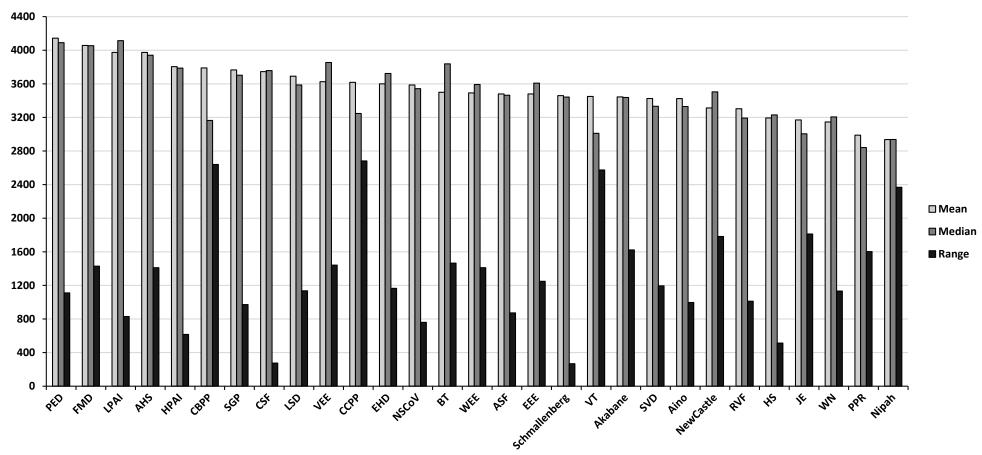
Appendix 4

Appendix 4, Table 1. Means, Standard deviation, Median and Range of the scores of the diseases. Ranking of the diseases according to the mean score and to the median score are also shown

| Disease | Mean (SD ^a) | Rank Mean ^b | Median | Rank Median ^c | Ranged |
|---|-------------------------|---------------------------|----------|-----------------------------|---------|
| Porcine Epidemic Diarrhoea | 4143.38 (469.88) | 1 | 4090 | 2 | 1111 |
| Foot and Mouth Disease | 4057.36 (546.83) | 2 | 4053.75 | 3 | 1428.75 |
| Low Pathogenic Avian Influenza | 3974.13 (376.09) | 3 | 4114.5 | 1 | 830 |
| African Horse Sickness | 3974.1 (527.52) | 4 | 3940.75 | 4 | 1411 |
| Highly Pathogenic Avian Influenza | 3804.5 (327.9) | 5 | 3787.375 | 7 | 616.75 |
| Contagious Bovine Pleuropneumonia | 3789.35 (1297.83) | 6 | 3164 | 25 | 2640.6 |
| Sheep and Goat Pox | 3765.06(434.19) | 7 | 3702.125 | 10 | 972 |
| Classical Swine Fever | 3745.33 (117.13) | 8 | 3758.15 | 8 | 275 |
| Lumpy Skin Disease | 3691.29 (488.16) | 9 | 3586.75 | 13 | 1135.85 |
| Venezuelan Equine Encephalitis | 3625.75 (671.92) | 10 | 3853.75 | 5 | 1441.25 |
| Contagious Caprine Pleuropneumonia | 3617.45 (1099.65) | 11 | 3247.25 | 21 | 2681.75 |
| Epizootic Haemorrhagic Disease | 3599.63 (532.13) | 12 | 3723.75 | 9 | 1165.65 |
| Novel Swine Enteric Coronavirus Disease | 3586 (322.33) | 13 | 3542.125 | 14 | 760.25 |
| Bluetongue | 3499.22 (652.21) | 14 | 3837.5 | 6 | 1465 |
| Western Equine Encephalitis | 3491.81 (647.42) | 15 | 3591.875 | 12 | 1411 |
| African Swine Fever | 3479.96 (411.22) | 16 | 3464.375 | 16 | 872.6 |
| Eastern Equine Encephalitis | 3479.38 (590.71) | 17 | 3608.125 | 11 | 1248.75 |
| Schmallenberg | 3459.19 (113.93) | 18 | 3442.125 | 17 | 267.5 |
| Vesicular Stomatitis | 3450.4 (1043.85) | 19 | 3011.25 | 26 | 2574.25 |
| Akabane Disease | 3444.55 (814.42) | 20 | 3437.6 | 18 | 1623 |
| Swine Vesicular Disease | 3425.25 (512.82) | 21 | 3333 | 19 | 1195 |
| Aino Disease | 3424.75 (455.24) | 22 | 3330.375 | 20 | 996.75 |
| NewCastle | 3312.75 (770.34) | 23 | 3504 | 15 | 1783 |
| Rift Valley Fever | 3303.6 (433.98) | 24 | 3192 | 24 | 1011.6 |
| Haemorrhagic Septicaemia | 3193.44 (218.2) | 25 | 3230 | 22 | 513.75 |
| Japanese Encephalitis | 3169.56 (763.67) | 26 | 3005 | 27 | 1811.75 |
| West Nile Fever | 3146.47 (419.96) | 27 | 3206.25 | 23 | 1132.5 |
| Peste des Petits Ruminants | 2989.31 (698.7) | 28 | 2841.25 | 29 | 1602.75 |
| Nipah Virus | 2936.56 (1038.14) | 29 | 2937.125 | 28 | 2369 |

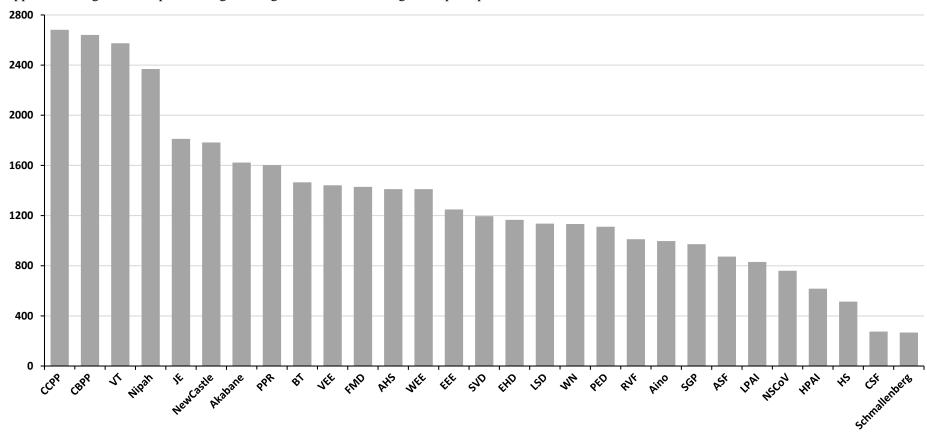
Legend: ^a SD = Standard Deviation , ^b Rank Mean= The ranking of the disease obtained with the mean scores, ^c Rank Median = The ranking of the disease obtained with the median, ^d Range = The range of the scores obtained from the expert's scores.

Appendix 4, Figure 1. Graph showing the mean, median scores and the range of the scores among the experts per disease



Legend: PED =Porcine Epidemic Diarrhoea, FMD= Foot and Mouth Disease, LPAI= Low Pathogenic Avian Influenza, AHS= African Horse Sickness, HPAI= Highly Pathogenic Avian Influenza, CBPP= Contagious Bovine Pleuropneumonia, SGP= Sheep and Goat Pox, CSF= Classical Swine Fever, LSD= Lumpy Skin Disease, VEE= Venezuelan Equine Encephalitis, CCPP= Contagious Caprine Pleuropneumonia, EHS= Epizootic Haemorrhagic Disease, NSCoV= Novel Swine Enteric Coronavirus Disease, BT=Bluetongue, WEE= Western Equine Encephalitis, ASF= African Swine Fever, EEE=Eastern Equine Encephalitis, VT= Vesicular Stomatitis, SVD = Swine Vesicular Disease, RVF= Rift Valley Fever, HS= Haemorrhagic Septicaemia, JE= Japanese Encephalitis, WN= West Nile Fever, PPR= Peste des Petits Ruminants.

Appendix 4 Figure 2: Graph showing the range of the scores among the experts per disease



Legend: CCPP= Contagious Caprine Pleuropneumonia; CBPP= Contagious Bovine Pleuropneumonia; VT= Vesicular Stomatitis; Nipah= Nipah virus; JE= Japanese Encephalitis; PPR= Peste des Petits Ruminants; BT=Bluetongue; VEE= Venezuelan Equine Encephalitis; FMD= Foot and Mouth Disease; AHS= African Horse Sickness; WEE= Western Equine Encephalitis; EEE=Eastern Equine Encephalitis; SVD = Swine Vesicular Disease; EHD= Epizootic Haemorrhagic Disease; LSD= Lumpy Skin Disease; WN= West Nile Fever; PED = Porcine Epidemic Diarrhoea; RVF= Rift Valley Fever; SGP= Sheep and Goat Pox; ASF= African Swine Fever; LPAI= Low Pathogenic Avian Influenza; NSCoV= Novel Swine Enteric Coronavirus Disease; HPAI= Highly Pathogenic Avian Influenza; HS= Haemorrhagic Septicaemia; CSF= Classical Swine Fever.

Table 1. List of criteria used to prioritise (re)emerging infectious diseases, according to their likelihood of (re)emergence in Belgium in response to different categories of drivers

A. DISEASE / PATHOGEN CHARACTERISTICS

- A.1 Current knowledge on the pathogen
- A.2 Current species specificity of the disease causing agent
- A.3 Genetic variability of the infectious agent
- A.4 Transmission of the pathogen in relation with the possible spread of the epidemic
- A.5 Risk of showing no clinical signs and silent spread during infection and post infection
- A.6 Wild reservoir and potential spread from it
- A.7 Existence of vectors (vertebrates and invertebrates, e.g. mosquitoes, bats, rodents, ticks, culicoid biting midges, etc.) and potential spread
- A.8 Transmission of the pathogen
- A.9 Environmental persistence

B. DISTANCE TO BELGIUM

- B.1 Current incidence (cases)/prevalence of the disease in the world
- B.2 European geographic proximity of the pathogen/disease to Belgium
- B.3 To your knowledge, when was the disease last reported in Europe

C. ABILITY TO MONITOR, TREAT AND CONTROL THE DISEASE

- C.1 Ability of preventive/control measures to stop the disease from entering the country or spreading (containment of the epidemic). Excluding treatment, vaccination and vector(s)/reservoir(s) control
- C.2 Vaccine availability
- C.3 Control of reservoir(s) and/or vector(s)
- C.4 Availability and quality of diagnostic tool(s) in Belgium
- C.5 Disease is currently under surveillance overseas (OIE, EU)
- C.6 Eradication experience in other countries and/or Belgium
- C.7 Detection of emergence, e.g. difficulties for the farmer/veterinarian to declare the disease or clinical signs not so evident

D. FARM/PRODUCTION SYSTEM CHARACTERISTICS

- D.1 Mono-species farms (one single farmed animal species, e.g. only cattle) or multispecies farms (more than one species e.g. goats and cattle, are raised in the same farm/land/premises).
- D.2 Farm demography/management: such as type of dairy or beef (cattle) production. For pigs reproduction, fattening, finishing farm or both. Chickens only laying eggs chickens or solely finishing broilers
- D.3 Animal density of farms. Extensive (small holders with a few animals) v/s intensive farming
- D.4 Feeding practices of farms
- D.5 Human movements among premises veterinarians or farm staff
- D.6 Proximity of livestock farm to wildlife and wildlife reservoirs of disease e.g. contact with wild or feral birds and animals which have been scavenging on landfill sites that contain contaminated animal products
- D.7 Changes of land use, e.g. field fragmentation, creation of barriers, landfill sites

E. CHANGES IN CLIMATIC CONDITIONS

- E.1 Influence of annual rainfall on the survival and transmission of the pathogen/disease
- E.2 Influence of annual humidity on the survival and transmission of the pathogen/disease
- E.3 Influence of annual temperature on the survival and transmission of the pathogen/disease

F. WILDLIFE INTERFACE F.1 Potential roles of zoo's in

- F.1 Potential roles of zoo's in the (re)emergence of the pathogen
- F.2 The rural(farm)-wildlife interface
- F.3 Increase of indigenous wild mammals in Belgium and neighbouring countries
- F.4 Increase in endemic/migrating populations of wild birds
- F.5 Hunting activities: hunted animals can be brought back to where livestock is present
- F.6 Transboundary movements of terrestrial wildlife from other countries

G. HUMAN ACTIVITIES

- G.1 In- and out- people movements linked to tourism
- G.2 Human immigration
- G.3 Transport movements: more specifically commercial flights, commercial transport by ships, cars or military (excluding transport vehicles of live animals)
- G.4 Transport vehicles of live animals
- G.5 Bioterrorism potential
- G.6 Inadvertent release of an exotic infectious agent from a containment facility e.g. laboratory

H. ECONOMIC AND TRADE ACTIVITIES

- H.1 Decrease of resources allocated to the disease surveillance
- H.2 Modification of the disease status (i.e. reportable disease becoming not reportable) or change in screening frequency due to a reduced national budget
- H.3 Decrease of resources allocated to the implementation of biosecurity measures at border controls (e.g. harbours or airports)
- H.4 Most likely influence of (il)legal movements of live animals (livestock, pets, horses, etc.) from neighbouring/MSs for the on the disease (re)emergence in Belgium
- H.5 Influence of increased (il)legal imports of animal products such as skin, meat and edible products from MSs on the disease (re)emergence in Belgium
- H.6 Most likely influence of increased (il)legal imports of NON-animal products such as tires, wood, furniture from MSs on the disease (re)emergence in Belgium.
- H.7 Most likely influence of (il) legal movements of live animals (livestock, pets, horses etc.) from Third countries on the disease (re)emergence in Belgium.
- H.8 Most likely influence of increased imports of animal products such as skin, meat and edible products from Third countries on the disease (re)emergence in Belgium
- H.9 Most likely influence of increased (il)legal imports of NON-animal products such as tires, wood, furniture from Third countries on the disease (re)emergence in Belgium

Legend: MS, European Union Member State.

Table 2. List of 29 diseases selected for prioritization, including the family and genus it belongs to and species it affects

| Name of disease | Family | Species affected |
|------------------------------------|---------------------|--|
| Eastern equine encephalitis | F: Togaviridae | Wild birds, horses, humans |
| | G: Alphavirus | |
| Western equine encephalitis | F: Togaviridae | Wild birds, horses, humans |
| | G: Alphavirus | |
| Venezuelan equine encephalitis | F: Togaviridae | Wild birds, horses, humans |
| | G: Alphavirus | |
| Japanese Encephalitis | F: Flaviviridae | Equids, wild birds, humans, swine |
| | G: Flavivirus | |
| West Nile fever | F: Flaviviridae | Wild birds, equids, humans |
| | G: Flavivirus | |
| Aino disease | F: Bunyaviridae | Bovines, cervids, sheep |
| 1 | G: Orthobunyavirus | |
| Akabane disease | F: Bunyaviridae | Bovines, goats, sheep |
| | G: Orthobunyavirus | |
| Schmallenberg disease | F: Bunyaviridae | Bovines, sheep, goats |
| | G: Orthobunyavirus | |
| Rift Valley fever | F: Bunyaviridae | Sheep, bovines and goats. |
| | G: Phlebovirus | |
| African horse sickness | F: Reoviridae | Equids |
| | G: Orbivirus | |
| Bluetongue | F: Reoviridae | Bovines, sheep, goats and wild ruminants |
| | G: Orbivirus | |
| Epizootic haemorrhagic disease | F: Reoviridae | Bovines and wild ruminants |
| | G: Orbivirus | |
| African swine fever | F: Asfivirus | Pigs and wild boar |
| | G: Asfivirus | |
| High pathogenic avian influenza | F: Orthomyxoviridae | Poultry, wild birds |
| | G: Influenzavirus A | |
| Low pathogenic avian influenza | F: Orthomyxoviridae | Poultry, wild birds |
| | G: Influenzavirus A | |
| Contagious bovine pleuropneumonia | Mycoplasma | Bovines |
| | Mycoides | |
| Contagious caprine pleuropneumonia | Mycoplasma | Goats |
| | capricolum | |
| Classic swine fever | F: Flaviviridae | Pigs and wild boar |
| | G: Pestivirus | |

| Foot and mouth disease | F: Picornaviridae | All cloven-hoofed animals |
|---|-----------------------|---------------------------|
| | G: Aphthovirus | |
| Haemorrhagic septicaemia | Pasteurella multocida | Bovines |
| | (Serotypes 6:B, 6:E) | |
| Lumpy skin disease | F: Poxviridae | Cattle |
| | G: Capripoxvirus | |
| Newcastle disease | F: Paramyxoviridae | Poultry |
| | G: Avulavirus | |
| Nipah virus encephalitis | F: Paramyxoviridae | Pigs |
| | G: Henipavirus | |
| Novel swine enteric coronavirus disease | F: Coronaviridae | Pigs |
| | G: Deltacorona Virus | |
| Peste des petits ruminants | F; Paramyxoviridae | Sheep and goats |
| | G: Morbillivirus | |
| Porcine epidemic diarrhoea | F: Coronavirus | Pigs |
| | G: Alphacoronavirus | |
| Sheep and goat pox | F: Poxviridae | Sheep and goats |
| | G: Capripoxvirus | |
| Swine vesicular disease | F: Picornaviridae | Pigs |
| | G: Enterovirus | |
| Vesicular stomatitis | F: Rhabdoviridae | Equids, cattle and goats |
| | G:Vesiculovirus | |

Legend: F, Family; G, Genus.

Table 3. Ranking and mean scores grouped by regression tree analysis of the 29 diseases according to the base model and the other "reduced" models

| | | | Deleted Domain | | | | | | | |
|----------------------------|--|----------------|--|------------------------|---|---|--------------------------------|-----------------------|---------------------|------------------------------|
| Disease | Regression tree Cluster ^a | 0 ^b | Disease Pathogen Characteristics | Distance to Belgium | Monitoring, treatment and control of the disease | Production system characteristics | Changes in climatic conditions | Wildlife interface | Human activities | Economy and trade activities |
| | | (Rank) | (Rank) | (Rank) | (Rank) | (Rank) | (Rank) | (Rank) | (Rank) | (Rank) |
| | | Mean Score | Mean Score | Mean Score | Mean Score | Mean Score | Mean Score | Mean Score | Mean Score | Mean Score |
| Porcine Epidemic Diarrhoea | 1 | (1) | (1) | (3) | (3) | (8)* | (1) | (1) | (3) | (5)* |
| 7 | | 4143.38 | 3454.63 | 3839.38 | 3572.13 | 3461.81 | 4124.63 | 4129.94 | 3599 | 2822.13 |
| Foot and Mouth Disease | 1 | (2) | (12)* | (2) | (1) | (2) | (2) | (2) | (6)* | (8)* |
| | | 4057.36 | 2938.61 | 3841.11 | 3731.11 | 3773.01 | 4007.26 | 3954 | 3390.86 | 2765.56 |
| Low Pathogenic Avian | 1 | (3) | (8)* | (1) | (5) | (6)* | (3) | (23)* | (2) | (1) |
| Influenza | | 3974.13 | 3019.5 | 3881.13 | 3386.88 | 3467.88 | 3851.938 | 3017.06 | 3609.438 | 3585.06 |
| African Horse Sickness | 1 | (4) | (2) | (4) | (2) | (1)* | (10)* | (3) | (1)* | (25)* |
| | | 3974.1 | 3370.8 | 3797.35 | 3578.85 | 3882.1 | 3501.1 | 3837.8 | 3639.1 | 2211.6 |
| Highly Pathogenic Avian | 2 | (5) | (6) | (9)* | (6) | (10)* | (6) | (17)* | (7) | (2)* |
| Influenza | | 3804.5 | 3053.86 | 3507.75 | 3357.63 | 3377.94 | 3684.19 | 3153.31 | 3381.375 | 3115.44 |
| Contagious Bovine | 2 | (6) | (5) | (5) | (23)* | (3)* | (4) | (6) | (8) | (11)* |
| Pleuropneumonia | | 3789.35 | 3071.25 | 3650.54 | 2824.66 | 3615.98 | 3761.23 | 3614.66 | 3350.6 | 2636.54 |
| Sheep and Goat Pox | 2 | (7) | (7) | (8) | (9) | (4)* | (7) | (4)* | (17)* | (16)* |
| | | 3765.06 | 3045.89 | 3514.49 | 3186.19 | 3485.31 | 3678.81 | 3736.06 | 3211.94 | 2496.75 |
| Classical Swine Fever | 2 | (8) | (3)* | (11)* | (4)* | (15)* | (5)* | (20)* | (19)* | (6) |
| | | 3745.33 | 3280.125 | 3402.83 | 3550.01 | 3235.01 | 3732.2 | 3045.83 | 3174.39 | 2796.89 |
| Lumpy Skin Disease | 2 | (9) | (11) | (14)* | (8) | (9) | (9) | (5)* | (11) | (19)* |
| | | 3691.29 | 2946.05 | 3347.29 | 3193.24 | 3455.41 | 3523.79 | 3627.79 | 3326.79 | 2418.66 |
| Venezuelan Equine | 2 | (10) | (4)* | (6)* | (7)* | (7)* | (20)* | (13)* | (20)* | (24)* |

| Encephalitis | | 3625.75 | 3168.5 | 3582.5 | 3353.25 | 3465.75 | 3093.25 | 3271.75 | 3119.5 | 2325.75 |
|-----------------------------|---|---------|---------|---------|---------|---------|---------|----------|----------|---------|
| Contagious Caprine | 2 | (11) | (10) | (7)* | (19)* | (13) | (8)* | (19)* | (10) | (10) |
| Pleuropneumonia | | 3617.45 | 2952.3 | 3516.6 | 2920.45 | 3275.7 | 3587.45 | 3049.5 | 3328.7 | 2691.45 |
| Epizootic Haemorrhagic | 2 | (12) | (15)* | (13) | (14) | (5)* | (14) | (12) | (4)* | (20)* |
| Disease | | 3599.63 | 2880.52 | 3360.03 | 3056.33 | 3484.88 | 3319.63 | 3273.96 | 3429.13 | 2392.93 |
| New Swine Enteric | 2 | (13) | (22)* | (18)* | (15) | (27)* | (11) | (7)* | (5)* | (4)* |
| Coronavirus disease | | 3586 | 2639.25 | 3263.88 | 3056.31 | 2870.69 | 3499.13 | 3532.625 | 3391.625 | 2848.5 |
| Bluetongue | 3 | (14) | (14) | (22)* | (16) | (11)* | (16) | (14) | (15) | (23)* |
| | | 3499.22 | 2885.64 | 3112.02 | 3028.04 | 3368.72 | 3255.22 | 3260.21 | 3223.97 | 2360.72 |
| Western Equine Encephalitis | 3 | (15) | (13) | (10)* | (10)* | (12)* | (18)* | (25)* | (16) | (21)* |
| | | 3491.81 | 2909.38 | 3404.31 | 3110.25 | 3276.19 | 3241.81 | 2892.13 | 3223.06 | 2385.56 |
| African Swine Fever | 3 | (16) | (9)* | (19)* | (11)* | (20)* | (12)* | (24)* | (22)* | (12)* |
| | | 3479.96 | 2963.81 | 3181.34 | 3072.46 | 3090.59 | 3456.71 | 2933.65 | 3079.03 | 2582.15 |
| Eastern Equine Encephalitis | 3 | (17) | (23)* | (12)* | (13)* | (14)* | (19) | (18) | (13)* | (15) |
| | | 3479.38 | 2600 | 3391.88 | 3056.88 | 3263.75 | 3152.81 | 3075.313 | 3280.94 | 2534.06 |
| Schmallenberg disease | 3 | (18) | (26)* | (23)* | (24)* | (16) | (21)* | (11)* | (9)* | (3)* |
| | | 3459.19 | 2532.94 | 3108.56 | 2788.44 | 3231.38 | 3071.06 | 3279 | 3336.06 | 2866.88 |
| Vesicular stomatitis | 3 | (19) | (21) | (15)* | (18) | (17) | (15)* | (10)* | (12)* | (26)* |
| | | 3450.4 | 2667.5 | 3342.9 | 2953.4 | 3127.9 | 3297.4 | 3310.4 | 3287.9 | 2165.4 |
| Akabane disease | 3 | (20) | (20) | (16)* | (17)* | (18) | (22) | (15)* | (18) | (14)* |
| | | 3444.55 | 2681.93 | 3332.05 | 3013.19 | 3108.94 | 2978.61 | 3244.55 | 3211.175 | 2541.43 |
| Swine Vesicular Disease | 3 | (21) | (18)* | (21) | (20) | (26)* | (13)* | (8)* | (21) | (17)* |
| | | 3425.25 | 2704.94 | 3131.56 | 2896.5 | 2906.5 | 3400.88 | 3360.875 | 3100.25 | 2475.25 |
| Aino disease | 3 | (22) | (16)* | (17)* | (21) | (19)* | (23) | (9)* | (14)* | (22) |
| | | 3424.75 | 2784.18 | 3306.94 | 2853.26 | 3107.19 | 2965.38 | 3313.25 | 3266.81 | 2376.25 |
| NewCastle | 3 | (23) | (17)* | (24) | (12)* | (25) | (17)* | (21) | (29)* | (18)* |
| | | 3312.75 | 2722.88 | 3107.06 | 3059 | 2934 | 3242.13 | 3028.063 | 2647.75 | 2448.38 |
| Rift Valley Fever | 3 | (24) | (28)* | (20)* | (22) | (23) | (24) | (16)* | (25) | (13)* |
| | | 3303.6 | 2483.38 | 3134.79 | 2851.79 | 3005.85 | 2954.23 | 3211.1 | 2925.48 | 2558.6 |

| Haemorrhagic Septicaemia | 4 | (25) | (19)* | (26) | (25) | (21)* | (27) | (22)* | (23) | (28)* |
|----------------------------|---|---------|---------|---------|---------|---------|---------|----------|---------|---------|
| | | 3193.44 | 2683.75 | 2973.44 | 2759.69 | 3052.81 | 2859.06 | 3019.688 | 2993.44 | 2012.19 |
| Japanese Encephalitis | 4 | (26) | (29)* | (25) | (29)* | (22)* | (28) | (27) | (26) | (9)* |
| | | 3169.56 | 2480.31 | 3069.56 | 2344.56 | 3010.19 | 2847.69 | 2828.313 | 2860.19 | 2746.13 |
| West Nile fever | 4 | (27) | (25) | (29) | (26) | (24)* | (29) | (28) | (24)* | (7)* |
| | | 3146.47 | 2577.93 | 2756.78 | 2640.74 | 2941.07 | 2738.17 | 2631.66 | 2954.97 | 2783.97 |
| Peste des Petits Ruminants | 4 | (28) | (24)* | (27) | (27) | (28) | (25)* | (26) | (28) | (29) |
| | | 2989.31 | 2585 | 2812.75 | 2523.06 | 2684 | 2953.38 | 2883.688 | 2748.38 | 1734.94 |
| Nipah Virus | 4 | (29)) | (27) | (28) | (28) | (29) | (26)* | (29) | (27) | (27) |
| | | 2936.56 | 2486.19 | 2795.31 | 2498.94 | 2514.69 | 2919.69 | 2500.813 | 2796.88 | 2043.44 |
| | | | | | | | | | | |

Legend: Highlighted numbers represent an up or down movement of more than 3 steps in the ranking; a Regression tree analysis clusters group: 1= very high importance; 2 = high importance; 3 = moderate importance; and 4 = low importance; b Base model of the ranking; * denotes more than three changes in the ranking.

31 Notifiable 21 Notifiable 36 Notifiable 28 Notifiable 53 Notifiable 59 Notifiable 78 Notifiable livestock livestock* livestock livestock livestock livestock livestock diseases diseases diseases diseases diseases diseases diseases European U.K The Netherlands Belgium France OIE Germany Union 306 animal diseases identified 161 animal diseases excluded (not epidemic type) and no cases over the last year Removal of duplicates (n=120) 25 animal diseases identified 4 animal diseases identified of interest 29 animal diseases identified

Means

