

The Bovine Tuberculosis Eradication Programme in Northern Ireland

Proposals from the Tuberculosis Strategic Partnership Group (TBSPG)

Scientific peer review

Final report

Simon J. More
University College Dublin
Dublin, Ireland

31 August 2016

Minor revisions
13 September 2016

Table of Contents

SUMMARY	5
INTRODUCTION	9
THE RECOMMENDED APPROACH: IMPLEMENT THE TBSPG RECOMMENDATIONS IN FULL	10
AN INTEGRATED APPROACH	10
PROGRAMME ELEMENTS BASED ON BEST AVAILABLE SCIENTIFIC KNOWLEDGE.....	11
ONGOING PROGRAMME REVIEW.....	12
<i>Measures of progress, including indicators and targets</i>	12
Evaluating the likelihood of eradication success.....	12
Specific measures/indicators of programme progress.....	12
Programme targets.....	14
<i>International benchmarking</i>	15
SCIENTIFIC SUPPORT	16
<i>Rationale</i>	16
<i>Key values</i>	16
Scientific independence.....	16
Scientific quality.....	17
Policy relevance.....	17
<i>Resource requirements</i>	18
<i>Expected output</i>	19
<i>Initial research questions</i>	19
INTERNATIONAL VACCINATION SYMPOSIUM	22
THE 'TEST AND VACCINATE OR REMOVE' (TVR) STUDY	22
ADDITIONAL AREAS FOR CONSIDERATION	23
<i>Non-technical issues</i>	23
<i>Defining eradication</i>	23
<i>Estimating time to eradication</i>	24
THE ALTERNATIVES	25
DO NOTHING OPTION	25
STATUS QUO OPTION	25
TOOLS AND PROCESSES (ANNEX B)	26
INTRODUCTION.....	26
IMPROVED SURVEILLANCE	26
Abattoir surveillance [proposed measure 10].....	26
A new contract for the provision of bTB testing services [proposed measure 5].....	27
IMPROVED MANAGEMENT OF INFECTED HERDS.....	28
<i>Increasing the likelihood that infected animals will be detected</i>	29
Severe interpretation of the skin test [proposed measure 1].....	29
Increased usage of interferon γ [proposed measure 2].....	29
<i>Increasing the probability that infected animals will be eliminated</i>	29
Full and partial depopulations [proposed measure 6].....	29
<i>Reducing the infection risk posed to other herds</i>	30
Introducing an additional 6 month test for derestricted herds [proposed measure 14].....	30
Reducing the number of NVL reactor animals required for a herd to be considered OTW [proposed measure 9].....	31
Chronic herds [proposed measure 7].....	33
Requiring a herd test prior to restocking after a bTB breakdown [proposed measure 8].....	33
<i>Removing restrictions where not epidemiologically justified</i>	34
Fattening herds operating under alternative conditions [proposed measure 11].....	34
ADDITIONAL CONTROL STRATEGIES.....	34
Genetic susceptibility of bovines [proposed measure 13].....	34
PROGRAMME INTEGRITY	34

DNA tagging [proposed measure 3]	34
TB reactor – quality assurance checks [proposed measure 15]	34
ADDITIONAL RESOURCES TO SUPPORT DECISION-MAKING AND SCIENTIFIC SUPPORT	35
<i>Geographic information systems</i>	35
Geographic information system (GIS) [proposed measure 12]	35
<i>M. bovis genotyping</i>	35
Genotyping of Mycobacterium bovis [proposed measure 4]	35
WILDLIFE AND VACCINATION (ANNEX C)	37
NORTHERN IRELAND’S BADGER ROAD TRAFFIC ACCIDENT (RTA)/FOUND DEAD SURVEY	37
<i>Relevant epidemiological concepts</i>	37
Study validity	37
Study precision	38
<i>Comments relevant to the Review</i>	38
In general.....	38
Relating to study validity and precision	39
Related work.....	40
WILDLIFE VACCINATION.....	40
<i>Wildlife and the epidemiology of bTB in cattle</i>	40
The role of badgers	40
A role for other wildlife species?	41
<i>Key epidemiological concepts</i>	42
<i>A critique of the proposed approach</i>	44
The Review proposal.....	44
Is badger intervention necessary?	45
What options are available, to limit badger-to-cattle transmission?	45
Will widespread badger vaccination work?.....	45
Is it reasonable to move to oral vaccination, once available?.....	46
Is badger removal necessary before implementing a widespread vaccination policy?	47
The scientific rationale	47
Field concerns.....	47
Evidence from modelling.....	48
<i>Further critique of the proposed badger intervention programme</i>	49
<i>Additional comments</i>	52
FARM PRACTICE AND BIOSECURITY (ANNEX D).....	53
GENERAL COMMENTS.....	53
IMPROVED BIOSECURITY CONTRIBUTES TO DISEASE PREVENTION	54
<i>Risk mitigation options</i>	54
<i>Risk mitigation responsibilities</i>	56
<i>Policy considerations</i>	57
IMPROVEMENT NOTICES	59
INFORMED PURCHASING	59
FARM FRAGMENTATION	61
REFERENCES.....	62

Summary

1. International experience has shown that the eradication of bovine tuberculosis (bTB) will only be achieved by simultaneously addressing all factors that meaningfully contribute to the persistence and spread of *Mycobacterium bovis* in all infected animal populations. Eradication success in Northern Ireland will only be possible with an integrated approach to *M. bovis* eradication.
2. Throughout the programme, programme managers (and other policy-makers) will require ongoing access to objective science-based information, to inform policy decision-making. In this review, detailed information is presented with respect to the rationale, key values, resource requirements and expected output of scientific support. The independence, quality, relevance and timeliness of this support will be critical to eradication success.
3. Concerning tools and processes
 - a. The 15 proposed measures can be grouped within the following key strategies relevant to bTB eradication: improved surveillance, improvement management of known infected herds, additional control strategies, programme integrity, and additional information to support decision-making and scientific knowledge.
 - b. These measures are each important components of an integrated approach to national bTB eradication. Nonetheless, some of the measures are likely to be more effective than others in reducing the infection risk to other herds. The effectiveness of these measures should be evaluated through ongoing review and relevant supporting scientific research.
4. Concerning wildlife and vaccination
 - a. Northern Ireland's badger road traffic accident (RTA)/found dead survey will provide valuable insights into the impact on badger populations of the national bTB eradication programme. As acknowledged in the Review, it is critical that the survey is designed and conducted so as to maximise both the validity and precision of the study.

- b. Badgers are an important maintenance host¹ for *M. bovis*, acting as a reservoir of infection² with spillover of infection to cattle, on the island of Ireland. The presence of an infected wildlife reservoir is a key constraint to bTB control or eradication. As highlighted previously, bTB eradication will only be achieved by simultaneously addressing all factors that meaningfully contribute to the persistence and spread of *M. bovis* in all infected animal populations. Therefore, intervention to limit badger-to-cattle transmission is necessary, as part of an integrated national approach to eradication.
- c. Options to limit transmission from badgers to cattle are limited, either to reducing the adequacy of contact (through badger culling or improved biosecurity) or reducing the proportion of the population susceptible (through badger or cattle vaccination). Based on current scientific knowledge, badger culling and badger vaccination each have the potential to contribute to national bTB eradication.
- d. The Review proposes a long-term strategy of widespread badger vaccination throughout Northern Ireland, in time through oral vaccination.
 - i. Based on available knowledge, it is reasonable to expect badger vaccination to reduce *M. bovis* prevalence in badgers, and in cattle in high bTB prevalence areas, over time. However, no data are yet publicly available to assess the magnitude and timing of these effects. The results from the Kilkenny badger vaccine trial will become available shortly.
 - ii. The concept of oral vaccination is attractive, particularly in terms of ease (and potentially, cost) of delivery. However, a number of issues need to be addressed, relating to both the safety and efficacy of the oral vaccine.
- e. In areas of increased bTB risk, the Review recommends that badger removal precede vaccination. In these areas, a badger intervention programme is

¹ A single host population capable of maintaining a pathogen over the long term (Viana et al., 2014).

² One or more epidemiologically connected populations or environments in which a pathogen can be permanently maintained and from which infection is transmitted to the target population (Viana et al., 2014).

proposed, including a 'ring vaccination' area surrounding a control (removal) area.

- i. Consistent with current knowledge, culling will be required in areas of high bTB risk prior to mass vaccination, specifically to reduce the prevalence of *M. bovis* infection in the re-emergent badger population.
- ii. The proposed badger intervention programme is seeking to balance two competing objectives, namely the requirement for a low prevalence population in which to introduce a badger vaccination programme, and concerns that a perturbation effect may occur following badger removal. On balance, the proposed approach seems both reasonable and prudent:
 - Using this approach, an area suitable for vaccination will be achieved, whilst also reasonably mitigating against a potential adverse effect.
 - Ring vaccination will also have the effect of facilitating the immigration of vaccinated badgers from border areas.
- iii. Research should be conducted in Northern Ireland, as part of the badger intervention programme, to clarify whether the perturbation effect occurs following badger removal.

5. Concerning farm practice and biosecurity

- a. Biosecurity is a critical aspect of good farming practice, protecting a herd (or industry) from the spread of a broad range of infectious diseases. There are numerous reports, from a range of countries, of problems with the widespread adoption of effective biosecurity on farms.
- b. With bTB, there are two key biosecurity-related risks, including contact with infected cattle and contact with infected wildlife.
 - i. Risk mitigation measures to limit cattle-related biosecurity risks are robust and generally well understood.
 - ii. With respect to wildlife-associated biosecurity risks, there are important gaps in knowledge. Further, there is as yet no empirical

evidence linking improved biosecurity with reduced wildlife-related bTB risks.

- c. The Review highlights several strategies to increase awareness of biosecurity on farms in Northern Ireland, including the development of a checklist to guide biosecurity assessment and the provision of farm-specific biosecurity advice. At this stage, farmer effort should primarily focus on cattle-related biosecurity risks. Several supporting research projects are proposed.
- d. Improvement notices may be helpful, but there is a need to first review progress and available evidence concerning the impact of improved farm-level biosecurity on future bTB risk in Northern Ireland.
- e. The Review proposes the introduction of informed purchasing, to allow farmers to make purchasing decisions informed by knowledge of past testing history, of the animal and herd. The underpinning principle is sound, however, limited progress has been made towards the development of predictive tools, to allow accurate prediction of future bTB risk. There is a need for ongoing research to critically evaluate the value of informed purchasing with respect to infection control benefit to the national bTB eradication programme.
- f. There is currently little understanding of the risks posed by farm fragmentation to national bTB control and eradication. The proposed strategy, including research to quantify the impact of farm fragmentation on future infection risk, is welcome.

Introduction

1. This document presents a review of the scientific elements of the proposals from the Tuberculosis Strategic Partnership Group (TBSPG) for a bovine tuberculosis (bTB) eradication programme for Northern Ireland. These were made available on 26 July 2016 in relevant TBSPG documents. A meeting was held with the TBSPG on 01 August 2016 to provide further context and other relevant information.
2. This work was conducted as a narrative review, based on detailed reference to relevant international peer-reviewed scientific literature.

The recommended approach: implement the TBSPG recommendations in full

3. This reviewer agrees that this is the only feasible option, and that the TBSPG recommendations should be implemented as described. Several points of caution, relating to specific elements of the proposal, are outlined later in this document.
4. A number of factors are critical to the success of the recommended approach, as outlined below.

An integrated approach

5. A number of models have been developed to represent *Mycobacterium bovis* infection in animal populations (including Hardstaff et al., 2013; Abdou et al., 2016). Invariably, they highlight the complexity of the overall system, particularly where more than one maintenance host is involved. In complex systems such as this, infection control (and more particularly eradication) can be very challenging.
6. International experience has shown that bTB eradication will only be achieved through an integrated approach, to simultaneously address all factors that meaningfully contribute to the persistence and spread of *M. bovis* in all infected animal populations. To illustrate:
 - a. In Australia, effective control was primarily achieved through cattle controls, using a range of strategies to limit the potential for damage from undetected, residually infected animals. Feral animal reservoir hosts (water buffalo (*Bubalus bubalis*), feral pigs (*Sus scrofa*)) were also removed during the eradication programme (Radunz, 2006; More et al., 2015).
 - b. In New Zealand, control efforts have greatly reduced the bTB burden in cattle, from 11% of mature cattle in 1905 to <0.003% in 2012/13 (Livingstone et al., 2015). Until 1995, control was based on established cattle-based methods of test and slaughter, and movement controls. Wildlife involvement was suspected following unexplained regional control failures and serious disease outbreaks, with the Australian brushtail possum (*Trichosurus vulpecula*; a feral species in New Zealand) subsequently identified as a true maintenance host of *M. bovis* infection. In recent years, New Zealand has implemented a multifaceted approach to bTB eradication, including test and slaughter programmes and risk-based movement

controls (in both cattle and farmed deer) as well as extensive possum control and wildlife surveillance (Livingstone et al., 2015).

- c. In the USA, *M. bovis* infection is problematic in several areas, with different epidemiological drivers. In recent years, bTB has been detected in a number of very large dairies in California. The root cause(s) of these breakdowns is believed to be multifactorial, linked with the importation of animals of greater risk of previous bTB exposure, and complex interactions of herd management practices (McCluskey et al., 2014). In Michigan, bTB in cattle is associated with a self-sustaining reservoir of infection in free-ranging white-tailed deer (*Odocoileus virginianus*) (O'Brien et al., 2002). Progress towards control, leading to a reduction in apparent prevalence in deer in the core (affected) area of greater than 60%, was primarily achieved through a reduction in deer densities through hunting, and restrictions on public feeding and baiting of deer (O'Brien et al., 2011).
7. Efforts to eradicate bTB from Northern Ireland will likely also only be achieved through an integrated programme that addresses all factors that contribute to the persistence and spread of *M. bovis* in all infected animal populations.

Programme elements based on best available scientific knowledge

8. As suggested in the Review, it is generally not possible to quantify the relative contribution of different interventions within a successful integrated approach. Nor is it currently possible, with any degree of certainty, to determine the number of interventions that would be required to achieve eradication. International experience clearly suggests that eradication is only possible if robust and comprehensive measures are applied to simultaneously address all factors that meaningfully contribute to the persistence and spread of *M. bovis* in all infected animal populations.
9. At programme start, programme elements should be both identified and designed based on, or consistent with, best available scientific knowledge. A detailed critique of each of the elements in the science-related elements of the integrated programme (Annex A: Tools and processes; Annex B: Wildlife and vaccination; Annex C: Farm practices and biosecurity) is presented later in this document.
10. Once the programme has started, there should be ongoing review, as outlined below.

Ongoing programme review

11. As indicated in the Review, an ongoing process will be needed ‘to review targets, review progress, identify reasons for variance and adapt the programme accordingly’. This ongoing process is strongly supported.

Measures of progress, including indicators and targets*Evaluating the likelihood of eradication success*

12. In theoretical terms, the eradication of infection will be achievable once the reproductive number, R^3 , of the overall system can be sustainably reduced below 1. In a multi-host system, the R of the system is influenced by both within- and between-species transmission (Dobson et al., 2004), as highlighted later. With bTB, therefore, the R of the system will be influenced by infection transmission between all relevant hosts (including cattle-cattle [animal, herd], badger-cattle, badger-badger etc).
13. In Ireland, the R of the overall system is currently being estimated, after accounting for both within- and between-species transmission (Aznar et al., 2014). Further, this information is guiding the development of the national strategy for badger vaccination, specifically to determine whether (and how⁴) badger vaccination, in addition to all existing bTB controls, has the potential to sustainably reduce the R of the system below 1. These calculations are drawing on available national data, including *M. bovis* prevalence in cattle herds and in badgers, and vaccine efficacy for susceptibility and infectiousness, derived from the Kilkenny badger vaccine trial.

Specific measures/indicators of programme progress

14. Measures of programme progress should provide insights into overall programme progress. Where possible, measures should also provide insights into the biological processes contributing to the persistence and spread of *M. bovis* in all infected animal populations.

³ The expected number of secondary cases caused by each infectious individual in a partially immune population.

⁴ For example, different levels of vaccine coverage.

- a. Overall programme progress
 - i. Measures of overall programme progress are well recognised, including herd prevalence and incidence, and animal prevalence.
- b. Assessing defined biological processes. In recent years, there has been increased focus on methods to quantify the relative importance of factors contributing to persistence and spread of *M. bovis*, primarily in cattle populations. The following points highlights progress that had been made, as well as points of caution, when seeking to quantify the relative importance of different infection sources for cattle, and to attribute programme impact:
 - i. With most cattle-based measures of programme progress, it is not possible to distinguish different infection sources, such as cattle (*cattle-to-cattle transmission*) or badgers (*badger-to-cattle transmission*). Further, cattle-to-cattle transmission can be indirectly attributable to badgers, for example, primary infection from a badger leading to residual infection within a herd. Therefore, a rise in cattle incidence (including herd incidence, animal prevalence) will occur following an increase in either cattle-to-cattle or (either direct or indirect) badger-to-cattle transmission.
 - ii. Increased attention is being paid on methods that would allow routine assessment of different infection source, and of associated programme impact:
 - Introduced cattle. Several authors have used methods to estimate breakdowns attributable to introduced cattle, thereby quantifying the role of introduced animals during different time-periods of a national programme (Clegg et al., 2008, 2015; Good et al., 2011).
 - Residual infection. In a recent review, More and Good (2015) outline the evidence in support of residual (persistent but undetected) infection in cattle as an increase concern in the Irish eradication programme. Breakdowns during the period following derestriction are mainly (but not entirely) attributable to residual infection (from cattle), rather than new

infection (either from cattle or badgers). Survival curves post-eradication have been used in several countries (Gallagher et al., 2013; Dawson et al., 2014), again during different time-periods, to evaluate programme progress in addressing this issue.

Programme targets

15. Drawing on international experience, there has been very limited success in developing targets that realistically reflect programme 'checkpoints' into the future. In large part, this reflects the complexity of infection in a multi-host system, including the multiple (often interacting) factors (some unknown, some poorly quantified) that influence the spread and persistence of *M. bovis*. In other words, future predictions need to be interpreted with considerable care. That said, the following points may be of assistance:
 - a. The Australian experience provides an example of the trajectory that might be expected in a successful programme (but noting the limited role of wildlife in that situation) (Radunz, 2006; More et al., 2015).
 - b. As highlighted previously, the R of the overall system in Ireland and of component parts is currently being estimated. Once the proposed eradication strategy is in place in Northern Ireland, it will be possible using this methodology to periodically evaluate the likelihood of eradication success, and to identify areas of potential weakness. This methodology could be used to estimate R at different geographical scales, including within the wildlife intervention areas. This approach may provide insights to complement (and perhaps refine) formal programme targets.
 - c. Computer modelling is increasingly used as a means to represent complex systems of infectious diseases. Models are only a simple representation of reality, but can prove very useful in predicting programme progress given current actions. For example, individual-based, spatially explicit ecological-epidemiological models have been developed for several animal diseases, including African swine fever (Lange et al., 2014) and classical swine fever (Lange et al., 2012), specifically to assist with national (and EU-wide) policy decision-making. A similar approach has been taken in Ireland, with the development of a computer model to represent bovine viral diarrhoea (BVD) in the national herd (Thulke et al., in preparation). The model has

contributed greatly to decision-support in the national eradication programme, estimating 'time to eradication' given a range of different control scenarios.

International benchmarking

16. International benchmarking offers a further opportunity to identify programme strengths and weaknesses.
 - a. To date, the approach to benchmarking has relied on the development of agreed summary measures of performance, separate country-level calculation of these measures, then the sharing of measures to allow country-level comparisons:
 - i. Abernethy et al. (2013) describe spatial and temporal trends in England, Ireland, Northern Ireland, Scotland and Wales during 1995 and 2010, focusing on measures of cattle demographics, testing, herd-level bTB statistics (annual herd prevalence, annual herd incidence), animal-level bTB statistics (apparent animal prevalence), abattoir surveillance and post-outbreak surveillance.
 - ii. In further work covering the period from 2003 to 2015 in these five countries, currently nearing completion, more detailed performance measures are being compared, relating to:
 - Episode recurrence (number of recurrences, inter-episode length),
 - Episode duration (number of days restricted, restriction length), and
 - Episode severity (number of standard and non-standard reactors at the start of the restriction, total number of infected animals during the restriction).
 - b. In the future, more detailed analyses are possible to facilitate sophisticated country-level comparison of programme progress. However, this would require the sharing of raw data across borders.

Scientific support

17. Throughout the programme, programme managers (and other policy-makers) will require ongoing access to objective science-based information, to inform policy decision-making. This is only considered briefly in the Review, and much greater detail would be helpful. The independence, quality, relevance and timeliness of this scientific information will be critical to the success of the bTB eradication programme in Northern Ireland. Further detail about the rationale, key values, resource requirements and expected output is outlined below.

Rationale

18. Programme success – *the successful eradication of bTB from Northern Ireland* - will be contingent on the generation, throughout the programme, of independent, high quality, relevant and timely scientific information in support of policy decision-making.

Key values

Scientific independence

19. It is critical that policy-makers and the general public have access to independent scientific advice on issues relevant to the national programme. The following context highlights this importance:
 - a. Public confidence in food safety was severely shaken during several food-related, public health crises in the 1980s and 1990s, including the emergence of BSE, first detected in 1985 in the UK (Bradley and Wilesmith, 1993), and the dioxin crisis in Belgium in 1999 (van Larebeke et al., 2001). At the time and subsequently, questions have been asked about the relationship between scientific advice and regulatory policy (Frewer and Salter, 2002), and in particular whether scientific advice was genuinely independent. Concern was also raised about the effectiveness of scientific communications and the potential for undue influence, for example from lobbying.
 - b. In response, fundamental changes have occurred throughout Europe, leading to clear lines of separation between science (often referred to as risk assessment) and policy (risk management).

- c. The European Food Safety Authority (EFSA) was established in 2002 to provide independent scientific advice, primarily to the Commission, with key values including scientific excellence, independence and openness⁵. EFSA is a European agency separate from the Commission. A similar separation between science and policy has occurred in many Member States. As one example, the Centre of Veterinary Epidemiology and Risk Analysis (CVERA, www.ucd.ie/cvera) was established in University College Dublin in large part to facilitate the independence of scientific animal health and welfare research conducted in support of national decision-making in Ireland.

Scientific quality

20. The quality of the science is paramount, to provide assurance to all stakeholders in Northern Ireland about the robustness of the information underpinning key policy decisions.
21. This is best protected through publication of the results of scientific research in international peer reviewed journals, to assure the quality of science conducted in Northern Ireland. It also facilitates the sharing of this information to interested stakeholders and the international scientific community. International scientific collaboration is also critical, to facilitate the sharing of knowledge and methodologies.

Policy relevance

22. It is critical that the scientific support is policy-relevant. This is best achieved through close collaboration and ongoing interaction between the scientists and policy-makers throughout the project lifecycle, from inception through to finalisation.

⁵ EFSA Strategy 2020. Trusted science for safe food. Protecting consumers' health with independent scientific advice on the food chain.
<http://www.efsa.europa.eu/en/corporate/pub/strategy2020>

23. Throughout this collaborative process, scientists and policy-makers have important, but differing, roles:
- a. The role of scientists is to generate information to support policy decision-making. It is critical, as highlighted previously, that this information is independent, of high quality, relevant and timely.
 - b. The role of policy-makers is to maximise returns from scientific effort, by prioritising research needs, helping scientists to clearly understand the research context, by aligning research objectives with these needs, by facilitating collaboration and communications, etc.

Resource requirements

24. As highlighted above, programme success will be contingent on timely, objective and independent scientific information. Therefore, resourcing will be critical, with respect to funding, scientific expertise and data access. Although each is critically important, only data access will be considered further here.
- a. It is likely that the research support will primarily use epidemiological (and ecological) research methodologies using observational study designs (but noting that modelling may also be important). In large part, this work will be conducted using routinely collected data (for example, animal identification and movement, bTB test data for herds and animals), or additional data collected as part of programme-related activities (for example, the proposed badger intervention programme). This research cannot be undertaken without ready and ongoing access to these and all other relevant national databases. This will include national spatial (GIS) data such as the national land parcel information system, noting the importance of a spatial perspective in many epidemiological and ecological studies.
 - b. Relevant to data access for epidemiological (and ecological) research, questions are often asked about data protection, and the impact of relevant legislation on epidemiological and ecological research.
 - i. The Data Protection Act 1998 defines UK law on the processing of data on identifiable living people. Consistent with EU legislation, there is a requirement for the protection of personal data.

- ii. There is a clear tension between the national good achieved through policy-relevant research and the need to protect individual's personally identifiable information. This is particularly relevant to epidemiological and ecological research given the central role played by identifier data fields such as *'farm ID'* and *'vet ID'* in joining databases, identifying land parcels, etc. Anonymised databases (for example, by removing *farm ID*) are generally of little to no epidemiological research value. Without *farm ID*, it would not be possible, for example, to attribute test results to a specific land parcel.
- iii. In Ireland, CVERA conducts policy-relevant epidemiological research whilst remaining compliant with the relevant national data protection legislation⁶. Under this legislation, CVERA is designated a data processor, and the national Department of Agriculture, Food and the Marine (DAFM) a data controller. To ensure legislative compliance, only summarized data are represented in CVERA's scientific outputs (as would be the norm for any epidemiological study), so that natural persons can never be identified. Further, a number of methods have been introduced (including point jittering, the use of grids etc) when representing the spatial results of CVERA's work, to ensure that individual farms cannot be identified.

Expected output

- 25. It is likely that there will be two main types of scientific output, each important for policy decision-making, including:
 - a. Scientific research: clear answers to defined policy-relevant questions.
 - b. Scientific support: ongoing assessment of progress, for example through the generation of agreed measures of programme performance.

Initial research questions

- 26. In this document, a number of research questions have been identified based on current perceived knowledge gaps. These are included here, to provide an

⁶ Data Protection Act 1988 and the Data Protection (Amendment) Act 2003. The latter legislation brought Irish law into line with the EU Data Protection Directive 95/46/EC.

overview of current questions where knowledge is limited. They might reflect current priorities with respect to scientific support, however, it is certain that additional questions will arise throughout the programme. Relevant to the three themes considered in this Review, these initial questions include:

- a. Tools and processes
 - i. Reasons for slaughterhouse-level differences in submission and confirmation rates among cattle at routine slaughter.
 - ii. The future infection risk in herds following full/partial depopulation.
 - iii. The practicality and effectiveness of measures to prevent reintroduction of bTB, following full/partial depopulation, both through restocked cattle or as a result of bTB persistence in the locality.
 - iv. The utility of additional measures to limit the spread of infection between herds, including an additional 6 month test for derestricted herds following all or higher risk breakdowns, pre-movement testing, herd testing delayed according to date of reactor removal, restrictions to the movement of inconclusive reactors, and contiguous testing.
 - v. The impact of animal introductions during a bTB restriction on future herd risk.
 - vi. Review the appropriateness of existing threshold for OTW and OTS breakdowns in Northern Ireland, in terms of future herd bTB risk.
 - vii. Whether fattening herds, operating as outlined in the Review, pose an increased risk of local bTB persistence in comparison with non-infected herds.
 - viii. The impact of different measures to limit fraud in the national programme.

b. Wildlife and vaccination

i. Relating to the badger found dead survey

- A critical evaluation of the RTA/badger found dead survey results in comparison to more detailed studies conducted under the auspices of the localized TVR study.
- The use of modelling to investigate biases associated with the RTA/badger found dead survey in Northern Ireland, and potential practical sampling alternatives. This approach would also allow sample size calculations to be refined.

ii. An assessment of the role of deer in the epidemiology of bTB in cattle in Northern Ireland, and whether intervention is warranted.

iii. Relating to the proposed badger intervention programme

- Consideration of cost, effectiveness and safety with respect to the deployment of an oral vaccine in Northern Ireland.
- An improved understanding of badger ecology and epidemiology, based on data collected during the proposed intervention studies.
- Revision of existing models on *M. bovis* infection in badgers and cattle in Northern Ireland, based on relevant new data that emerges.
- The likelihood and magnitude of the perturbation effect following badger removal in Northern Ireland.

c. Farm practice and biosecurity

i. A clearer understanding of the relative importance of the two key biosecurity-related risks in Northern Ireland, including:

- Contact with infected cattle, and
- Contact with infected wildlife.

- ii. The impact of improved farm biosecurity on future bTB risk, including the relative contribution of different biosecurity practices.
- iii. An improved understanding of motivators for, and constraints to, improved on-farm biosecurity on farms in Northern Ireland.
- iv. The impact of informed purchasing, with respect to infection control benefit to the national bTB eradication programme.
- v. The impact of farm fragmentation on future infection risk following a bTB breakdown.

International Vaccination Symposium

27. An International Vaccination Symposium was held in Belfast during 14-16 May 2012, with the objective *'to identify and evaluate factors that may determine the effectiveness of a TB vaccination strategy in wildlife, particularly the Eurasian badger, which will result in a reduction in bovine TB incidence'*. Many experts attended, representing the diversity of bTB policy and science internationally. Detailed meeting notes are available, but have yet to be distilled into a single summary document for publication in the international peer reviewed literature. There is much to be gained from such a summary publication:

- a. This was a unique event, and reflections from this meeting are of considerable international interest.
 - b. Subsequent international progress has been relatively limited; therefore, this information is not yet dated.
 - c. This information is highly relevant to the current Review.
28. Further issues, relevant to wildlife vaccination in Northern Ireland, were discussed during a follow-on meeting, hosted by APHA in May 2016 (the 'TB in European wildlife and control measures based on vaccines' workshop).

The 'test and vaccinate or remove' (TVR) study

29. The TVR study commenced in Northern Ireland in May 2014, informed by modelling (Smith et al., 2013), with the aim *'to describe the effects of implementing*

*a 'test and vaccinate or remove' intervention on badgers in an area of high badger and cattle density and with high levels of bTB in cattle*⁷. Field aspects of the study will be completed by late 2018. It is critical that the results of this study are published in the international scientific peer reviewed literature, as a means of quality assurance and also to share knowledge about aspects of badger ecology and bTB epidemiology, including the dynamics of bTB in badgers in Northern Ireland, and of TVR, including impact and logistics. This information is relevant to the national bTB eradication programme.

Additional areas for consideration

Non-technical issues

30. Based on international experiences, including from Australia (More et al., 2015) and New Zealand (Livingstone et al., 2015), a number of non-technical issues are critical to programme success, including programme governance, cost and responsibility sharing, etc. These have been addressed by the TBSPG in their overall report, but are not considered further in this scientific peer review.

Defining eradication

31. In the document, eradication is defined in terms of legislative targets (that is, infection below a defined level, as defined by relevant EU legislation) rather than true biological freedom (the absence of the causative organism, either entirely or from defined animal populations). However, legislative targets for freedom, once reached, are likely harder to maintain in the presence, compared to the absence, of an infected wildlife reservoir. In New Zealand in recent years, there has been a rapid fall in the number of infected herds (of cattle and deer), to very low levels (annual infected herd prevalence < 0.2%). Although cattle controls continue, in recent years the primary focus of the national eradication programme has shifted towards the eradication of infection from possums and other wildlife (Livingstone et al., 2015).

⁷ The test and vaccinate or remove (TVR) wildlife intervention research project. Year 2 report – 2015. Department of Agriculture, Environment and Rural Affairs. <https://www.daera-ni.gov.uk/sites/default/files/publications/daera/tvr-wildlife-intervention-research-project-year-2-report-2015.pdf>.

Estimating time to eradication

32. The estimates for time to eradication are reasonable, but only if there are tools available to reasonably control infection in all animal populations that are epidemiologically linked. In Australia, for example, eradication was achieved after 27 years, in the absence of any meaningful wildlife reservoir (More et al., 2015). The New Zealand programme will take longer, relying on similar cattle controls to those applied in Australia plus additional controls specifically to address *M. bovis* infection in infected wildlife.
33. Figures 4 & 7 present important messages, in particular an ongoing exponential decline in national measures of infection (herd breakdowns, herd incidence etc) in both New Zealand and Ireland. However, these figures should be interpreted with care, given the differing challenges (for which there may not be ready solutions) that may emerge towards the end of an eradication programme.

The alternatives

34. The TBSPG briefly described two alternative approaches, as required in their terms of reference, including the 'do nothing' and '*status quo*' options. The TBSPG has rejected both of these options, for reasons that are scientifically sound.

Do nothing option

35. The overall scientific assessment is sound.
36. It is not possible to predict the 'post-Brexit' future with respect to the trade in cattle and associated commodities. Nonetheless, there is little doubt that trade of animals and animal products into the EU will continue on the basis of equivalence, that is compliance with current EU legislation, in particular Council Directive 64/432⁸.

Status quo option

37. The overall scientific assessment is sound.
38. The argument is predicated on the assumption of existing costing arrangements, namely the taxpayer covering all costs and most disease risks. However, it should be noted that other cost-sharing models are available, including the exacerbator-pays approach to cost allocation, as applied in New Zealand⁹. Therefore, the *status quo* could continue, but under a different cost-sharing model, as acknowledged in the Review.

⁸ Council Directive 64/432/EEC of 26 June 1964 on animal health problems affecting intra-Community trade in bovine animals and swine.

⁹ An illustration of the use of this approach in the context of bTB eradication in New Zealand is presented in Bovine TB Strategy, Review of Costs, 2010.
<http://www.biosecurity.govt.nz/files/pests/bovine-tb/tb-strategy-cost-review.pdf>.

Tools and processes (Annex B)

Introduction

39. The 15 proposed measures have been evaluated after grouping them under the following 5 key strategies relevant to bTB eradication:
 - a. improved surveillance (to first detect infection).
 - b. improvement management of known infected herds.
 - c. additional control strategies.
 - d. programme integrity.
 - e. additional information to support decision-making and scientific knowledge.
40. These 5 strategies are important components of an integrated approach to national bTB eradication.

Improved surveillance

41. The Review proposes several measures to increase the likelihood of detecting infection in herds that are currently free to trade, including those relating to:
 - a. Abattoir surveillance (*routine post mortem examination*).
 - b. Field surveillance (*bTB testing services*).
42. The 2 relevant measures [*proposed measures 5 and 10*] will each contribute to improved surveillance in the national programme.

Abattoir surveillance [proposed measure 10]

43. Variations between abattoirs with respect to submission and confirmation rates have previously been noted in Ireland (Frankena et al., 2007; Olea-Popelka et al., 2012) and England (Shittu et al., 2013). In Ireland, a reduction in between-abattoir variation between 2003-04 (Frankena et al., 2007) and 2005-07 (Olea-Popelka et al., 2012) was observed, once this concern had been highlighted to relevant staff.

44. As suggested in the Review, objective measures of performance are needed, to enable ongoing critical review.
- a. In Australia, abattoir submission targets were integral to the National Granuloma Submission Program (NGSP) (More et al., 2015), noting the background incidence of granulomas that would be expected (in Australia, at least one granuloma for every 1000 cattle slaughtered with two or more permanent teeth) in the absence of bTB. Throughout this Program, meat inspectors were encouraged to submit all granulomas, noting that it is not possible to accurately distinguish bTB and non-bTB lesions by gross examination alone (Liebana et al., 2008). There was ongoing laboratory support throughout this Program, to clarify the aetiology of submitted granulomas.
 - b. The suggested minimum submission target of 1 per 1,000 cattle slaughtered is a reasonable starting point for Northern Ireland. This may need subsequent adjustment because the background incidence of non-bTB granulomas (due to infection other than *M. bovis*) is certain to vary in different geographical regions.

A new contract for the provision of bTB testing services [proposed measure 5]

45. In several jurisdictions, including Northern Ireland, private veterinary practitioners (PVPs) play a key role in field surveillance, but have very little input otherwise. The Review proposes a broadened role for PVPs in the national programme, including the provision of advice on issues relating to the management of infected herds. This move is welcomed, but care will be needed to ensure that farmers continue to receive a robust and consistent message, underpinned by the best available science.
46. The Review only briefly considers quality control as it relates to bTB testing by PVPs. Until recently, quality control of bTB testing has mainly focused on key inputs, such as personnel training, standard operating procedures, equipment and test reagents. As a consequence of IT advances, objective measurement and benchmarking of outputs is now relatively straightforward. In Ireland, for example, Duignan et al. (2012) describe the introduction of a special report (the so-called ER13A) providing an objective summary of:

- a. Administrative performance, including advanced itinerary submission, test report submission for clear tests and for positive tests, private test approvals and test amendments.
- b. Field performance, including reactor detection rate, inconclusive reactor detection rate, back-traced reactors, and bTB confirmed lesions in attested cattle.

Improved management of infected herds

47. Difficulties in clearing infected herds, leading to herd recurrence, has been identified as a key challenge to bTB eradication, both in Ireland (More and Good, 2015) and New Zealand (Dawson et al., 2014). The fundamental problem is that it is not always possible, with current diagnostic tools, to identify all of the infected animals within infected herds. These animals are likely at differing stages of infection. Either currently or into the future, they can pose an infection risk to the index or neighbouring herds, or to herds to which the animal subsequently moves. The primary effect (noting that there may be multiple) of the measures proposed in the Review includes:

- a. Increasing the likelihood that infected animals will be detected (*severe interpretation of the skin test, increased usage of interferon γ testing*).
- b. Increasing the probability that infected animals (and infection within the herd) will be eliminated (*full and partial depopulations*).
- c. Reducing the infection risk posed to other herds including neighbours and herds to which animals are traded (*reducing the number of NVL reactor animals required for a herd to be considered OTW, requiring a herd test prior to restocking after a TB breakdown, introducing an additional 6 month test for derestricted herds, chronic herds*).
- d. Removing restrictions where not epidemiologically justified (*fattening herds operating under alternative arrangements*).

Increasing the likelihood that infected animals will be detected*Severe interpretation of the skin test [proposed measure 1]**Increased usage of interferon γ [proposed measure 2]*

48. Each of these measures [*proposed measures 1 and 2*] will increase the likelihood that infected animals will be detected.
49. There are now a number of studies highlighting the future infection risk posed by animals in known infected herds that are positive under severe interpretation (referred to in the Review) or inconclusive under standard interpretation (including Clegg et al., 2011a,b). Animals positive to the interferon γ test are also at increased future infection risk, in circumstances where the positive predictive value of this test is likely to be high (Gormley et al., 2013; Lahuerta-Marín et al., 2015) (see below).
50. As test sensitivity is increased, it is inevitable that some non-infected animals will be incorrectly identified (false positive reactions).
51. As acknowledged in the Review, there are two very different contexts in which interferon γ testing is conducted, including:
- a. To increase the likelihood that infected animals will be detected. As outlined in the Review, it is critical that interferon γ testing is limited to high-risk groups in known infected herds. Due to the imperfect specificity of the interferon γ test (including Álvarez et al., 2012), the positive predictive value of this test will be low unless the test is used in situations where the prevalence of infection is likely to be high.
 - b. To provide assurance that bTB skin reactors are indeed infected (this is discussed later).

Increasing the probability that infected animals will be eliminated*Full and partial depopulations [proposed measure 6]*

52. There is limited published information about the long-term infection risk following full/part herd depopulation. This strategy was shown to be effective in Irish herds depopulated with bTB during 2003-05 (Good et al., 2011). However, as also suggested in the Review, associated measures to prevent reintroduction were

critical. In the UK after the FMD epidemic, future TB risk on restocked farms was associated with both the restocked cattle (from high risk areas) and bTB persistence on the restocked farm (Carrique-Mas et al., 2008).

53. Research will be needed to consider key elements relating to this measure, including:
 - a. The future infection risk in herds following full/partial depopulation.
 - b. The practicality and effectiveness of measures to prevent reintroduction of bTB, following full/partial depopulation, both through restocked cattle or as a result of bTB persistence in the locality.

Reducing the infection risk posed to other herds

54. Some of the proposed measures [*proposed measures 7, 8, 9 and 14*] are more effective than others in reducing the infection risk to other herds.

Introducing an additional 6 month test for derestricted herds [proposed measure 14]

55. There is compelling evidence of continuing infection risk, both in herds and individual animals, for an extended period after derestriction, as a consequence of both newly introduced and residual infection. Multiple studies from a range of countries have highlighted the contribution of residual infection to bTB persistence in a herd or locality (including Karolemeas et al., 2011; Dawson et al., 2014; More and Good, 2015; More et al., 2015), and a recent study has noted that higher-than-baseline risk persists for many years subsequent to high-risk breakdowns (Clegg et al., 2015).
56. Under Council Directive 64/432, restricted herds are free to trade (and considered at no greater risk than non-infected herds) once two consecutive clear full-herd SICTT tests are achieved. In contrast, in the Australian programme, all animals present during a breakdown were considered at risk for the rest of their life, and infected herds took a minimum of 8 years to attain the lowest herd risk status (More et al., 2015). The EU legislation is at odds with current scientific thinking.
57. In each of the bTB-affected countries in Europe, there are difficulties with respect to feasible options that can be applied to manage herd risk during the post-derestriction period. In particular, compromises have been sought to limit

disruptions to the normal business of farming whilst also managing the heightened herd risk that can persist for an extended period following infection.

58. The Review proposes an additional 6 month test for derestricted herds (in other words, full-herd tests at 6 and 12 months following derestriction). This proposal is beneficial, as it will provide an additional opportunity, as acknowledged in the Review, to evaluate the infection status of herds known to be at higher-than-baseline risk. However, it will not impact on the free movement of animals once herd derestriction has occurred, with the potential for ongoing spread of infection through animal movement.
59. The Review proposes a range of options, including an additional 6 month test for derestricted herds following all or higher risk breakdowns¹⁰, pre-movement testing, herd testing delayed according to date of reactor removal, restrictions to the movement of inconclusive reactors¹¹, and contiguous testing. Each of these options has epidemiological merit in reducing the infection risk posed to other herds, and, as acknowledged in the Review, many are currently applied in other countries. Further research would be beneficial to further evaluate these options.

Reducing the number of NVL reactor animals required for a herd to be considered OTW [proposed measure 9]

60. As outlined in the Review, the management of herds with OTW (Officially TB free status Withdrawn) and OTS (Officially TB free status Suspended) breakdowns is fundamentally different. Specifically, disease control is much more rigorous and continues for a longer period following OTW compared to OTS breakdowns.
61. The distinction between OTW and OTS breakdowns varies by country, as described by Abernethy et al. (2013):
- a. In Great Britain, OTF status is suspended if infection is not confirmed, and only one further negative herd test is required.
 - b. The policy in Northern Ireland is similar to GB, except that outbreaks with six or more unconfirmed reactors are treated as OTW.

¹⁰ Noting that breakdown size is an important predictor of future risk (Olea-Popelka et al., 2004; Karolemeas et al., 2011)

¹¹ As currently occurs in Ireland based on the work of Clegg et al. (2011a,b)

- c. In Ireland, almost all outbreaks are considered OTW, except a small proportion (those classified under the so-called ‘singleton policy’) released early following an epidemiological risk assessment and laboratory analysis (Good and Duignan 2011; Murray et al., 2012).
62. The OTW/OTS threshold has very important implications for national disease control. In particular, there is a need to maximise control efforts in infected herds, and to minimise the problems of residual infection (the persistence of infection in a herd and locality) and infection risk associated with animal movement (More and Good, 2015). As highlighted previously, bTB risk persists in many infected herds despite current legislative requirements for OTW herds (two consecutive clear tests prior to derestriction) (Clegg et al., 2015). Therefore, it is particularly important that requirements are not inadvertently weakened by incorrectly classifying an infected herd as OTS.
63. The appropriate threshold between OTW and OTS is best determined through epidemiological research. There are many relevant insights from existing research including:
 - a. Future herd bTB risk is not generally associated with either detection of lesions at slaughter (Olea-Popelka et al., 2004) or lesion confirmation (Karolemeas et al., 2011; Doyle et al., 2014).
 - b. Future bTB herd risk is associated with the size of the breakdown (the number of skin reactors identified) (Olea-Popelka et al., 2004; Karolemeas et al., 2011; Doyle et al., 2014).
64. Research was recently completed in Ireland specifically to address questions relating to the classification of H-herds¹² for the purposes of future bTB risk (Clegg et al., 2016). This study confirmed the key role of past bTB history in determining the future risk of Irish herds, with the odds related to both the severity of and time since the previous restriction. It also illustrates the difficulty in clearly defining H-herds, noting that risk persists for extended periods following a bTB restriction,

¹² In Ireland, herds are considered at higher risk (so-called H-herds) if they experience a bTB episode with at least 2 standard reactors or at least 1 standard reactor and an animal with a TB lesion found at slaughter, and where infection was acquired within the herd. H-herds are deemed OTW, but additional control measures are applied, including badger removal if the infection source was attributed to badgers and three SICTTs (single intradermal comparative bTB tests) at 6 month intervals following de-restriction before returning to annual testing.

regardless of breakdown severity. The study concluded that there is a need for robust controls on H-herds for an extended period post de-restriction.

65. In the Review, the proposed designation of OTW breakdowns is generally consistent with the policy in Ireland, as recently confirmed by the work of Clegg et al. (2015). It would be valuable, in time, to review the proposed policy for Northern Ireland, as further relevant data become available.

Chronic herds [proposed measure 7]

66. As highlighted in the Review, there is currently imperfect knowledge about chronic herds, including measures to resolve or minimise their impact. Given this, the proposal for critical review and further research is entirely reasonable, to address knowledge gaps.
67. Many of the individual measures as proposed in this Review should contribute to reducing the number and impact of these herds. The recent publication by Doyle et al. (2016) is particularly welcome, highlighting risk factors associated with duration and recurrence of chronic herd breakdowns in Northern Ireland.

Requiring a herd test prior to restocking after a bTB breakdown [proposed measure 8]

68. As highlighted in the Review, this requirement provides some disease control benefit, at least in Ireland (Clegg et al., 2013), but is also driven by EU legislation.
69. Legislative requirements are certain to influence future policy decision-making. Therefore, this is a need for clarity post-Brexit concerning the ongoing relevance of EU legislation to Northern Ireland. In addition, it would be valuable to conduct research, similar to that completed in Ireland (Clegg et al., 2013):
- a. To provide an overview of movement events associated with each bTB episode.
 - b. To determine whether introduction of animals during a bTB episode is associated with increased future bTB risk.
 - c. To identify practices relating to the introduction of animals that are the most risky.

Removing restrictions where not epidemiologically justified*Fattening herds operating under alternative conditions [proposed measure 11]*

70. Similar measures for fattening herds are also applied in other jurisdictions.
71. Based on the measures as described, there is no infection risk directly associated with the movement of cattle.
72. The risk posed by fattening herds, compliant with each of the suggested conditions, would only be negligible (ie acceptable in the context of national bTB eradication) if there were no contribution from fattening herds to bTB persistence in the locality. The robustness of this measure needs to be formally tested, in time, by determining whether fattening herds pose an increased risk of local bTB persistence in comparison with non-infected herds.

Additional control strategies*Genetic susceptibility of bovines [proposed measure 13]*

73. It is now clear that genetics can make a significant contribution to animal health, including resistance to bTB and other infectious diseases. Further, tools for simultaneous selection on these traits and other performance traits are available (Berry et al., 2011). Advances in this area are likely relevant to animal breeding in Northern Ireland.

Programme integrity*DNA tagging [proposed measure 3]**TB reactor – quality assurance checks [proposed measure 15]*

74. The Review proposes several measures relating to programme integrity [*proposed measures 3 and 15*], including DNA tagging and quality assurance checks on bTB reactors. Any steps to limit errors, fraudulent or otherwise, should be supported.
75. There is evidence from several countries, primarily anecdotal, of substantial improvements following the introduction of measures to limit fraud. Ongoing discussions will be helpful, to share experiences with international colleagues.
76. Further, supporting research should be conducted to determine the impact of different measures to limit fraud in the national programme. Field trials are

suggested in the Review, however, observational studies should also be considered.

77. Additional comments with respect to the quality control of field surveillance are included previously, concerning the new contract for the provision of bTB testing services [*proposed measure 5*].

Additional resources to support decision-making and scientific support

78. In the Review, additional measures [*proposed measures 4 and 12*] are proposed to support decision-making and scientific support, including improvements to existing geographic information systems (GIS) and an expansion of genotyping of *M. bovis*. These proposals are very strongly supported.

Geographic information systems

Geographic information system (GIS) [proposed measure 12]

79. GIS is a fundamental tool for many relevant aspects of science, including epidemiology and ecology. It is critical that GIS is able to seamlessly link with other relevant national databases. The data requirements for epidemiological and ecological research are considered in greater detail previously.

M. bovis genotyping

Genotyping of Mycobacterium bovis [proposed measure 4]

80. Molecular epidemiology is a well-established discipline, providing insights into the dynamics of infection in animal and human populations. Until recently, however, the use of molecular epidemiology in bTB science has been relatively limited, in large part due to the absence of suitable molecular tools. As outlined in the Review, such tools are now emerging (spoligotyping and VNTR for some years, whole genome sequencing currently), offering an extraordinary opportunity to address a broad range of questions, many with direct implications for the national bTB eradication programme. Of particular interest are questions relating to the maintenance and transmission of bTB, at a range of different scales, both spatial (on a farm, in a locality [potentially between different animal species], nationally) and temporal.

81. Northern Ireland is a world leader in the development and application of these technologies.
82. The genotyping work is a critical national resource, and should be supported, both from Northern Ireland and elsewhere (eg UK competitive funding). It is important that genotyping-related research is conducted in close collaboration with national policy-makers and field veterinarians, to ensure that priority is given to those questions, once answered, that are most likely to contribute to bTB eradication. There is also a need to move rapidly from concept to practical application.

Wildlife and vaccination (Annex C)

Northern Ireland's badger road traffic accident (RTA)/found dead survey

Relevant epidemiological concepts

83. Several important epidemiological concepts are relevant to the badger RTA/found dead survey, relating to the validity and precision of survey results. These issues are considered in detail in the STROBE statement (von Elm et al., 2007; Vandembroucke et al., 2007).

Study validity

84. Internal validity characterizes the quality of conclusions relative to the population under study (Toma et al., 1999).
- a. Minimising confounding, namely those variables that could lead to a distortion in the effect estimate between another variable and an outcome (Toma et al., 1999). In related work from Wales, a range of potential confounders were considered during multivariable analyses, including age, sex and season (Goodchild et al., 2012). Carcass weight and stage of decomposition are further potential confounders.
 - b. Minimising information (measurement) bias¹³. The Review outlines research conducted in support of an agreed uniform protocol for sample collection and laboratory procedures (organs to be collected, diagnostic methodology to be used etc). The protocol used for culled badgers in Ireland is described elsewhere (Byrne et al., 2015).
85. External validity relates to the possibility of extrapolating the study conclusions (assuming that the internal validity has been confirmed) to other populations (other times and places) (Toma et al., 1999).
- a. A number of inherent weaknesses with badger RTA/found dead surveys have previously been identified, in Northern Ireland and elsewhere, each relating to the representativeness of the study population (Abernethy et al., 2003, 2011; Nusser et al., 2008).

¹³ Bias is defined as systematic error that leads to incorrect quantitative findings (Toma et al., 1999)

- i. A non-random sub-population of badgers, collected through a convenience sample. Badgers found dead are a non-random sample, and are unlikely to be typical of all badgers with respect to age, sex or severity of bTB infection.
 - ii. Geographical bias. The sample population is limited to areas surrounding roads. Therefore, regions with a lower road density or with roads with fewer cars may be under-represented.
 - iii. Temporal bias. Temporal trends may be missed, if RTAs are not collected over a sufficiently short period of time.
 - iv. Reporting bias is a particularly concern given the passive nature of the surveillance effort. Motivation to report among farmers, for example, may be linked to the bTB status of their herds.
 - v. Collection bias. There may be fewer RTAs collected from highways, both because carcasses may be too damaged to retrieve and also because retrieval from these roads is particularly dangerous.
- b. A number of strategies have previously been implemented in Northern Ireland to at least partly address these concerns, including strict controls on reporting (solely by defined public officials), the setting of regional quotas, and proactive searching for badger carcasses (Abernethy et al., 2011).

Study precision

86. Study precision corresponds to a reduction in random error (Rothman et al., 2008).

Comments relevant to the Review

In general

87. The Review highlights some of the key questions to be addressed, including temporal and spatial trends in infection prevalence in the badger, and the interrelationship, in terms of *M. bovis* infection, between cattle and badger populations. Similar objectives have been in place since 1999 (Abernethy et al., 2003, 2011). As argued in the Review, the survey will provide valuable insights into the impact on badger populations of the national bTB eradication programme.

Relating to study validity and precision

88. As clearly acknowledged in the Review, it is critically important that the badger RTA/found dead survey is designed and conducted so as to maximise both the validity and precision of the study.
- a. With respect to study validity:
 - i. As reflected above, a range of strategies have previously been used in Northern Ireland to maximise study validity, both internal and external. Concerns with study validity are clearly acknowledged in the Review and in supporting documentation¹⁴, and a range of strategies to maximise study validity are outlined. A critical evaluation of these results with those using different methods (the TVR results, the interventions areas), as outlined below, is recommended.
 - b. With respect to study precision:
 - i. The sample size calculations are presented in the Review and in supporting documentation¹⁴. This work highlights both:
 - The current study limitations, with respect to study precision: *'the current sampling strategy provides weak [statistical] power to detect significant inter-annual changes in prevalence at a national Northern Ireland level, with the exception of very large changes in prevalence ...'*¹⁴
 - The study precision required: sufficient to allow *'moderate variations in infection prevalence across time (inter-annually) and space (between counties)'*¹⁴ to be detected.
 - ii. The Review presents a robust case for an increase in sample size.
 - iii. The sample calculations should be considered a guide to the numbers required. As a point of caution, these calculations are based on the assumptions of probability sampling and random distribution of infection/disease, which do not strictly apply to this survey

¹⁴ Byrne, A.W., 2016, unpublished. Power study guidelines for RTA badger surveillance: the power to detect true inter-annual change

methodology. No alternative methodology is immediately available, although a suggestion for further research is outlined below.

89. Given the challenges faced with respect to study validity and precision, it is important that the study results are published in the international scientific literature, as an important means of ongoing scientific quality control. This is acknowledged in the Review.

Related work

90. The TVR study and the work proposed in intervention area (as outlined later) will each provide insights into both the prevalence and incidence of *M. bovis* in badgers, but in quite localised areas of Northern Ireland. As acknowledged in the Review, a comparison of results from these studies and from the badger RTA/found dead survey in equivalent areas will provide valuable insights into potential biases affecting the badger RTA/found dead survey.
91. Nusser et al. (2008) have outlined some of the challenges of disease surveillance in wildlife populations using convenience sampling. These are further elaborated by several authors, including Rees et al. (2011) and Leslie et al. (2014), primarily in the context of early detection and case finding. Consideration should be given to the use of modelling to investigate biases associated with the RTA/badger found dead survey in Northern Ireland, and potential practical sampling alternatives. This approach would also allow sample size calculations to be refined.

Wildlife vaccination

Wildlife and the epidemiology of bTB in cattle

The role of badgers

92. The Review correctly summarises current scientific knowledge with respect to the role played by badgers in the epidemiology of bTB cattle (Corner et al., 2011; Ní Bhuachalla et al., 2015). Badgers are an important maintenance host¹ for *M. bovis*, acting as a reservoir of infection² with spillover of infection to cattle, on the island of Ireland (More, 2009), in Great Britain (Godfray et al., 2015), and likely in parts of mainland Europe (Payne et al., 2013, Hardstaff et al., 2014).

93. The Review is also correct in suggesting that eradication will only be possible if all infection sources, including badgers, are addressed. There are several supporting evidence sources:
- a. As suggested in the Review, several large-scale culling projects have demonstrated substantial and sustained reduction in the bTB risk of associated cattle herds. The measured impact was greater in Ireland (Griffin et al., 2005; Kelly et al., 2008) than in England and Wales (Donnelly et al., 2007; Jenkins et al., 2008).
 - b. As stated previously, in complex systems such as *M. bovis* infection in animal populations, it is extremely unlikely that eradication is achievable unless all factors contributing to persistence and spread are addressed. This has been a key lesson from the successful Australian bTB eradication programme (More et al., 2015), and from experiences in several other countries, including New Zealand (Livingstone et al., 2015) and the USA (O'Brien et al., 2011).
 - c. The presence of an infected wildlife reservoir is recognised as a key constraint to bTB control or eradication in many countries, both in Europe (Godfray et al., 2013; Gortázar et al., 2014, 2015) and elsewhere (O'Brien et al., 2011; Miller and Sweeney, 2013; Gortázar et al., 2015; Warburton and Livingstone, 2015).

A role for other wildlife species?

94. Currently, there is limited knowledge of the role of other wildlife species in the epidemiology of bTB on the island of Ireland. In this context, concern has been raised about the considerable expansion in several deer species in Ireland, including red, sika and fallow deer (Carden et al., 2011). As outlined in the Review, research is needed to assess the role of deer in the epidemiology of bTB in cattle in Northern Ireland, and whether intervention is warranted.

Key epidemiological concepts

95. The following concepts from basic epidemic theory are relevant (see Halloran, 1998; Viana et al., 2014):
- a. The basic reproductive number, R_0 , is the expected number of secondary cases caused by a single infectious individual in a fully susceptible population.
 - i. R_0 is a composite of the number of contacts per unit time, the duration of infectiousness and the transmission potential per potentially infective contact (collectively known as '*the adequacy of contact*').
 - b. R , the effective reproductive number, is the expected number of secondary cases caused by each infectious individual in a partially immune population.
 - i. $R = R_0x$, where x is the proportion of contacts that are susceptible.
 - c. In a single host system, infection would be eradicated if R could sustainably be reduced to <1 . In a multi-host system, the R of the system is influenced by both within- and between-species transmission (Dobson et al., 2004).
 - d. It is not necessary to immunise every individual in order to stop transmission of an infectious agent through a population. Herd immunity refers to the reduction of infection or disease in the unimmunised segment as a result of immunising a proportion of the population (John and Samuel, 2000).
96. Consistent with basic epidemic theory, options to limit transmission from badgers to cattle are reliant on either a reduction in either the adequacy of contact or the proportion of the population susceptible.
- a. Options to limit adequacy of contact have been restricted to efforts to reduce the number of contacts per unit time, noting that practical options are currently not available to limit the duration of infectiousness in badgers or the transmission potential from infectious badger to cattle given contact. Strategies to reduce the number of contacts per unit time between badgers and cattle has been undertaken either through:
 - i. Badger culling. This strategy is currently being used extensively throughout Ireland, in areas with cattle bTB breakdowns that cannot

be attributed to cattle movement (for further detail, see Byrne et al., 2015).

- ii. Improved biosecurity (relevant to badger-to-cattle transmission), to limit contact between badgers and cattle. As discussed later in this document, a number of risk mitigation strategies are proposed, and strategies to successfully exclude badgers from housing have been demonstrated (Judge et al., 2011). As yet, however, there is as yet no empirical evidence linking improved biosecurity with reduced wildlife-related risks (More, 2009; O'Hagan et al., 2016).
- b. Options to reduce the proportion of the population susceptible. Two strategies are being considered, including:
- i. Badger vaccination. As outlined in the Review and by Robinson et al. (2012), considerable progress has been made towards a bTB vaccine for badgers, primarily as a collaborative effort between Ireland and the UK. In pen-based trials, in vaccinated compared to control badgers, there was a significant decrease in the number and severity of gross lesions, lower bacterial load in the lungs, and fewer sites of infection (Corner et al., 2010; Chambers et al., 2011; Murphy et al., 2014). Several field trials have been conducted, in the UK (Carter et al., 2012) and Ireland (Aznar et al., 2011, 2013, 2014), and final results from the Irish trials will become available shortly. A further trial is underway in six counties of Ireland, scheduled to end in December 2017, to determine whether vaccination is not inferior to area-wide targeted badger culling in maintaining a herd-level risk of bTB in cattle (O'Keeffe et al., 2016).
 - ii. Cattle vaccination. Under current EU legislation, bTB vaccination in cattle is prohibited, because it may interfere with current bTB diagnostic methods. Detailed consideration of the research required in support the use of cattle vaccination in the UK is outlined by the European Food Safety Authority (EFSA Panel on Animal Health and Welfare (AHAW), 2013), focusing on evaluation of both vaccine efficacy and the performance of a test to '*Detect Infected among Vaccinated Animals*' (DIVA). Chambers et al. (2014) provide further detail.

97. Of these four options, only two are considered in detail in the section immediately following, namely badger culling and badger vaccination. The other two options (improved biosecurity (relevant to badger-to-cattle transmission), cattle vaccination) may prove useful into the future. At this point, however, there is insufficient scientific knowledge to support the inclusion of either improved biosecurity (relevant to badger-to-cattle transmission) or cattle vaccination in a national bTB eradication programme.
- a. Improved biosecurity (relevant to badger-to-cattle transmission). This issue is considered in further detail under 'Farm practice and biosecurity'. *At this point in time*, there is no empirical evidence linking improved biosecurity with reduced wildlife-related risks.
 - b. Cattle vaccination. A substantial number of technical and non-technical issues will need to be resolved before cattle vaccination could be considered for use in Northern Ireland. A summary of current knowledge is presented elsewhere (EFSA Panel on Animal Health and Welfare (AHAW), 2013; Chambers et al., 2014).

A critique of the proposed approach

The Review proposal

98. The TBSPG recommends a long-term strategy of widespread badger vaccination throughout Northern Ireland, using the only available licensed bTB vaccine bacille Calmette–Guérin (BCG), specifically to limit badger to cattle transmission, as an integral part of a national disease control strategy. The TBSPG indicate that injectable vaccine will be used initially, moving to oral vaccination once available. Oral vaccine will be deployed via an effective bait.
99. In areas of increased bTB risk, the TBSPG recommends that badger removal precede vaccination, using an intervention area design. This includes a control (removal) area surrounded by a ring vaccination area. This is considered in detail later.

Is badger intervention necessary?

100. As highlighted elsewhere, there is now conclusive evidence, both from Ireland and the UK, that badgers are an important contributor to bTB epidemiology in cattle.
101. International experience has shown that bTB eradication will only be achieved through an integrated approach, by simultaneously addressing all factors that meaningfully contribute to the persistence and spread of *M. bovis* in all infected animal populations.
102. The rationale for badger intervention is consistent with current knowledge.

What options are available, to limit badger-to-cattle transmission?

103. Based on the material presented under 'Key epidemiological concepts', only two options are current feasible for further consideration, including:
- a. Limiting the adequacy of contact
 - i. Badger culling. As outlined by Abdou et al. (2016), culling strategies can be either selective (based on the infection status of the animal, as determined by an animal-side test) or not.
 - b. Reducing the proportion of the population susceptible
 - i. Badger vaccination. As outlined by Abdou et al. (2016), options for vaccination include either oral (dependent on bait uptake rates) or parenteral (dependent on trapping efficacy) administration.

Will widespread badger vaccination work?

104. This question has been the focus of intensive research for some years, both in Ireland and the UK. The results of pen trials have been very encouraging (Corner et al., 2010; Chambers et al., 2011), with the main protective effect being a reduction in the severity and progression of disease following *M. bovis* challenge. Field trials have been completed in both the UK and Ireland, and again early results from the UK have been encouraging (Carter et al., 2012). It is reasonable to expect vaccination to reduce *M. bovis* prevalence in badgers, and in cattle in high bTB prevalence areas, over time. However, no data are yet publicly available to assess the magnitude and timing of these effects (Godfray et al., 2013). The final

results from the Irish field trial (the Kilkenny vaccine trial; Aznar et al., 2011, 2013, 2014) will be available shortly.

105. A number of modelling studies have been undertaken (including Smith et al., 2001; Hardstaff et al., 2013; Abdou et al., 2016), generally highlighting the value of long-term vaccination in reducing bTB incidence in badger populations. Further detail is given below.
106. The proposal is scientifically sound, but will need to be reviewed as further information becomes available, in particular the results of the Kilkenny badger vaccine trial. It is important to note that widespread vaccine deployment, in areas previously subject to badger culling, is currently being conducted in six Irish counties. This study is seeking to determine whether vaccination is not inferior to area-wide targeted badger culling in maintaining a herd-level risk of bTB in cattle (O’Keeffe et al., 2016). The results of this work, to be completed at the end of 2017, will also be of relevance to the proposed national strategy in Northern Ireland.

Is it reasonable to move to oral vaccination, once available?

107. Only two practical vaccination routes are available: parenteral (subcutaneous, intramuscular) and oral (Robinson et al., 2012). Both routes have been used extensively, but only in pen and field trials. Parenteral vaccine is currently licensed for use in badgers in the UK, whereas oral vaccine is not.
108. The concept of oral vaccination is attractive, particularly in terms of ease (and potentially, cost) of delivery. However, a number of issues need to be addressed, relating to both vaccine safety and efficacy. Several authors highlight some of the issues under consideration, including the potential for BCG exposure by non-target species, including cattle (Robinson et al., 2012, 2015).
109. The proposal is scientifically sound, but will need to be reviewed as further information becomes available. This is clearly acknowledged in the Review. In the proposed trial of an oral vaccine in Northern Ireland, key questions to be considered would need to include the cost, effectiveness and safety of this method of vaccine deployment (Chambers et al., 2014)

Is badger removal necessary before implementing a widespread vaccination policy?

The scientific rationale

110. In humans, the BCG vaccine is included as part of the childhood vaccination programme in many countries. However, the efficacy¹⁵ of BCG in preventing pulmonary tuberculosis (TB; caused by infection with *M. tuberculosis*) is known to vary greatly in different circumstances. In a recent meta-analysis of randomized clinical trials¹⁶, it was demonstrated that BCG confers protection against pulmonary TB when administered both in infancy and at school age, provided children were not already infected with *M. tuberculosis* or sensitized to other mycobacteria (Mangtani et al., 2014).
111. As with people (Andersen and Doherty, 2005), there is no evidence of either a beneficial or detrimental effect of BCG in infected badgers (Chambers et al., 2014). Because it is ineffective in infected animals (Robertson et al., 2012; Chambers et al., 2014), vaccination has the potential to provide benefit only to those animals that are not infected (or otherwise sensitized to mycobacteria) at the time of vaccination¹⁷.
112. Pseudo-vertical transmission¹⁸ is believed to be an important feature of *M. bovis* infection in badgers, and may be a key factor in maintaining infection within local populations (Ní Bhuachalla et al., 2015). In infected setts, therefore, it is plausible that cubs may become infected with *M. bovis*, or exposed to other (environmental) mycobacteria, whilst young. Logically, the force of infection¹⁹ will be greater in high compared with lower prevalence badger populations.

Field concerns

113. Collectively, these issues have raised concerns as to whether badger vaccination alone will be sufficient to limit transmission, initially between badgers, and subsequently to cattle, given the current force of infection in the badger population. In Ireland, *M. bovis* infection in badgers can be very high (reaching

¹⁵ The percentage of vaccinated individuals that are protected.

¹⁶ Randomised clinical trials are recognised as the most rigorous way to determine whether a case-effect relationship exists between treatment and outcome (Sibbald and Roland, 1998).

¹⁷ There may be additional benefit subsequently, if herd immunity is established, to non-infected animals.

¹⁸ Via the respiratory route during the rearing phase rather than *in utero*.

¹⁹ Hazard rate of infection from a defined source to susceptible host individuals in a defined population (Viana et al., 2014).

43.2% in hot-spot areas in Ireland; Corner et al., 2012). Further, in undisturbed badger populations, infected animals are long-lived (Ní Bhuachalla et al., 2015), and therefore an ongoing source of infection.

114. Differing results have been observed in Ireland and GB with respect to the impact of culling on the prevalence of *M. bovis* in the emergent badger population.
 - a. In Ireland, a significant reduction in the prevalence of *M. bovis* infection over time has been observed in areas of proactive culling, both in the four area trial (among the emergent badger population during 1997-2002; Corner et al., 2008) and throughout the country during 2007-13 as part of the national programme of targeted badger removal (Byrne et al., 2015). In the national programme, repeated culling is conducted once an area has been recruited.
 - b. In GB, in contrast, an increase in *M. bovis* infection in badgers was observed with successive proactive culls in the Random Badger Culling Trial (RBCT), especially where landscape features allow badgers from neighbouring land to recolonize culled areas (Woodroffe et al., 2006).
115. Given this background, and based on the evidence presented, the current thinking in Ireland is that culling will be required in areas of high bTB risk prior to mass vaccination, specifically to reduce the prevalence of *M. bovis* infection in the re-emergent badger population. Practical steps have been taken in this direction, as described by O’Keeffe et al. (2016), with the establishment of widespread vaccine deployment, in areas previously subject to badger culling, in six Irish counties. As indicated previously, this study has been designed as a non-inferiority trial, to determine whether vaccination is not inferior to area-wide targeted badger culling in maintaining a herd-level risk of bTB in cattle. By default, the study is also evaluating the transition from focused culling to a national badger vaccination programme.

Evidence from modelling

116. Modelling studies have provided further insights into this question.
 - a. Using a spatial simulation model, Abdou et al. (2016) have highlighted the limited impact of vaccination alone on bTB infection in badger populations, and the substantial improvement when vaccination was preceded by 5 years of culling.

- b. The results of Smith et al. (2001) and Wilkinson et al. (2004) are broadly similar. Smith et al. (2001) found that cattle herd breakdowns and bTB prevalence were most effectively reduced by first introducing a proactive element (such as proactive culling) followed by vaccination or culling. If this proactive element was avoided, the impact of vaccination alone on bTB prevalence in badgers took many more years to achieve (Wilkinson et al., 2004).
 - c. Hardstaff et al. (2013) found that vaccination alone could be an effective disease control strategy for bTB in higher-density badger populations, but only with annual deployment of a vaccine with an efficacy of around 80%. The effectiveness of this strategy was reduced by the presence of external sources of infection.
117. The proposal is consistent with current scientific knowledge, apart from the above-mentioned RBCT results. Ongoing research in Northern Ireland will be critical, in particular questions relating to badger ecology and epidemiology during the proposed intervention studies. The modelling studies should be updated as further information becomes available.

Further critique of the proposed badger intervention programme

118. The Review proposes a badger intervention programme, to allow for the strategic removal of badgers from areas of high bTB prevalence in cattle. The proposed programme would be implemented in bTB problem areas, where bTB incidence in cattle is high, recurrent and/or persistent. A 'ring vaccination' area would surround the control (removal) area, with intervention in each area (either culling or vaccination, as relevant) continuing over a four-year period. An appropriate sample strategy will be implemented to establish the prevalence (and where possible, also the incidence) of infection in the target intervention area.
- a. The control (removal) area
 - i. The stated rationale is to reduce the level of bTB in a wildlife reservoir directly associated with bTB breakdowns in associated cattle herds.
 - ii. Relevant scientific issues are considered in detail in the section above.
 - iii. The Review presents the expected consequences of removal versus TVR, given a series of simplistic assumptions. If these assumptions are

correct, the fall in *M. bovis* prevalence in badgers will be much more rapid following badger removal compared with TVR.

b. The ring vaccination area

- i. The stated rationale is linked to events in both the control area (facilitating the immigration of vaccinated badgers from border areas) and bordering areas (mitigating against perturbation).
- ii. The ring vaccination area has been introduced directly in response to concerns that the perturbation effect will occur as a consequence of badger removal in the control (removal) area. As outlined in the Review, the perturbation effect (Godfray et al., 2013) is a hypothesized chain of consecutive effects triggered by badger culling, including substantial changes to the spatial and social organisation and territorial behaviour of badger populations (social perturbation), increased contact and transmission of *M. bovis* infection between badgers, increased contact between cattle and the disturbed badger population, and increased infection risk in associated cattle (More et al., 2007).
- iii. During the RBCT in the southwest of GB, there was evidence in support of the perturbation effect, including social perturbation and associated increases in *M. bovis* prevalence in both badgers (Woodroffe et al., 2006) and cattle (Donnelly et al., 2007).
- iv. In Ireland, social perturbation is well described (O’Corry-Crowe et al., 1996). However, there has been no evidence of associated increases in *M. bovis* prevalence following badger removal, either in badgers (Corner et al., 2008; Byrne et al., 2015) or cattle (Griffin et al., 2005; Kelly et al., 2008; Olea-Popelka et al., 2009). In recent years, there has been a long-term trend of falling herd bTB prevalence in Ireland (Abernethy et al., 2013). This has occurred coincident with a long history of targeted badger removal (for example, 7,284 badgers removed in 2008 (Sheridan, 2011) from an estimated national population of approximately 84,000 (Sleeman et al., 2009)) in response to herd bTB breakdowns where badgers are implicated (Byrne et al., 2015).

- v. Reasons for the observed differences between southwest GB and Ireland are uncertain. O'Connor et al. (2012) has suggested a range of possibilities, relating to cattle, badgers and badger controls.
 - vi. It is also uncertain whether the perturbation effect might occur in Northern Ireland. The Irish experience may be more applicable than the experience from southwest GB as influencing factors, such as bTB epidemiology and badger ecology, are likely more similar across the island of Ireland than between Northern Ireland and southwest England. Byrne et al. (2012) presents a detailed review of badger ecology in Ireland, but with considerable reference to Northern Ireland.
119. The proposed badger intervention programme is seeking to balance two competing objectives, namely the requirement for a low prevalence population in which to introduce a badger vaccination programme, and concerns that a perturbation effect may occur following badger removal. Based on current knowledge, it seems very likely that a mass vaccination programme will be largely ineffective in a high prevalence population. It is much less certain, however, whether badger removal will result in a perturbation effect in Northern Ireland. On balance, and noting the critical need to address badger-to-cattle transmission within an integrated national approach to bTB eradication, the approach as proposed in the Review seems both reasonable and prudent. Using this approach, an area suitable for vaccination will be achieved, whilst also reasonably mitigating against a potential adverse effect. Ring vaccination will also have the effect, as described in the Review, of facilitating the immigration of vaccinated badgers from border areas.
120. Smith et al. (2012) has previously modeled the impact of ring vaccination around areas of badger culling, but using parameter estimates from the RBCT in southwest GB. These authors found that culling plus ring vaccination did mitigate some, but not all, of the adverse effects of the perturbation effect. It would be valuable to rerun this model using parameter estimates from Northern Ireland, once these are available.
121. It is critical that research is conducted in Northern Ireland, as part of the badger intervention programme, to clarify whether the perturbation effect occurs following badger removal.

Additional comments

122. Several authors highlight issues to be considered when designing, implementing and evaluating a badger vaccination programme, including Robinson et al. (2012), Chambers et al. (2014) and Ní Bhuachalla et al. (2015). These include:

- a. Clearly defined objectives and criteria for success
- b. Area of coverage, underpinned by knowledge of bTB prevalence (in time, in space, between social groups)
- c. Strategies to maximise vaccine coverage
- d. Revaccination, taking account of duration of immunity, protection following revaccination, vaccine coverage, population recruitment (births, immigration)
- e. Monitoring strategy, relating to coverage (uptake if oral vaccine), vaccine effectiveness and the epidemiological consequences of vaccination
- f. Length of programme

Farm practice and biosecurity (Annex D)

General comments

123. Biosecurity has been defined as a strategy of management practices to prevent the introduction of diseases and pathogens to an operation and to control spread within the operation (Wells, 2000). In the literature, there are varying uses of terms relating to biosecurity, some different from those used in the Review. For example, in Mee et al. (2012):
- a. Bioexclusion relates to preventive measures (risk reduction strategies) designed to avoid the introduction of pathogenic infections (hazards), and
 - b. Biocontainment relates to measures to limit within-farm transmission of infectious hazards and onward spread to other farms.
124. Biosecurity is a critical aspect of good farming practice, protecting a herd (or industry) from the spread of a broad range of infectious diseases. Further, appropriate risk mitigation measures are well described (including Mee et al., 2012).
125. The implementation of bioexclusion plans on beef and dairy farms (covering both the entry and exit of infectious agents from a farm) is voluntary in most countries (Mee et al., 2012). One exception is larger dairy farms in Denmark, where farmers are required to introduce measures to reduce the risk of introducing animal diseases into the dairy herd and minimize the impact of outbreaks, should they occur (Kristensen and Jakobsen, 2011).
126. There are numerous reports of problems with the widespread adoption of effective biosecurity on farms (for example, Nöremark et al., 2010; Sayers et al., 2013). A number of barriers to adoption have been identified, including:
- a. A lack of consensus regarding effective biosecurity protocols
 - b. The efficacy of such protocols, and
 - c. Their cost-effectiveness.
127. The Review highlights problems of biosecurity on farms in Northern Ireland (also O'Hagan et al., 2016). Farm biosecurity is relatively poor throughout the island of Ireland, in part due to land fragmentation and animal movement. These were

important considerations leading to the decision for national eradication (as opposed to farm-by-farm control) of BVD from Ireland (Barrett et al., 2011).

Improved biosecurity contributes to disease prevention

Risk mitigation options

128. Risk factors for bTB are increasingly understood, including many that relate to biosecurity risks. In broad terms, there are two key biosecurity-related risks, including:
- a. Contact with infected cattle, either through animal movement or contiguous contact.
 - b. Contact with infected wildlife (including an infected environment).
 - c. [The role of fomites, visitors, biological material in the spread of *M. bovis* is likely minor]
129. The relative importance of each of these risks is uncertain, and will likely vary in different countries (differing epidemiology, ecology, programme management etc) and over time (Broughan et al., 2016). Therefore, generalization is difficult. In Ireland, White et al. (2013) focused on the relative importance of ‘neighbourhood’, specifically farm-to-farm spread and spread from wildlife, on bTB persistence. Among the study farms, they attributed 15% of bTB episodes in the study to residual infection, between 0% and 20% to contiguous spread, and between 19% and 39% to wildlife (More and Good, 2015).
130. With respect to the biosecurity risk ‘*contact with infected cattle*’, risk mitigation measures are robust and generally well understood.
- a. As outlined by Mee et al. (2012), strategies relevant to bTB include:
 - i. Concerning animal movement – maintaining a closed herd, minimizing the number of cattle purchased and the number of source herds, purchasing from herds with likely low disease (infection) prevalence, obtaining cattle disease history, testing cattle before movement.
 - ii. Concerning contiguous spread – attention to boundary fencing to prevent nose-to-nose contact.

- b. Considerable empirical evidence is available highlighting the impact of these measures on infection risk, both for bTB and other directly transmissible diseases of cattle. For many bovine infections, including BVD and Johne's disease, farm-to-farm spread mainly occurs through the movement of infected animals. Therefore, efforts to prevent such movement, through the methods listed above, are critical to success in national control programmes (Lindberg and Alenius, 1999; Geraghty et al., 2014). The infection benefit of closed herds, with no introductions or contact with cattle in neighbouring herds, is well recognised (van Schaik et al., 2002). With bTB, many studies have highlighted the disease risk associated with cattle movement (including Gilbert et al., 2005; Doyle et al., 2016). Further, movement controls are central to relevant EU bTB control legislation⁸, and to progress in successful national bTB eradication programmes (More et al., 2015).

131. With respect to the biosecurity risk '*contact with infected wildlife or an infected environment*', the issues are more problematic. To illustrate:

- a. There are important gaps in knowledge about aspects of *M. bovis* epidemiology in badgers (see reviews by Corner et al., 2011; Ní Bhuachalla et al., 2015; Broughan et al., 2016).
- b. There is uncertainty about how *M. bovis* is transmitted between badgers and cattle (Godfray et al., 2013). Available evidence would suggest that transmission between badgers, and by extrapolation from badgers to cattle, is primarily via aerosol during direct contact. The high prevalence of pulmonary infection strongly supports the lungs as the principal site of primary infection in badgers, with inhalation of infectious aerosol particles ('droplet nuclei', with an aerodynamic diameter of 0.7-7 μm) the principal mode of transmission. Droplet nuclei are formed during normal respiratory air movements, as well as during coughing and sneezing. The conditions required for aerosol transmission and establishment of infection are exacting and principally involve the aerodynamic diameter of the aerosol particle. The most vulnerable sites for primary infection of *M. bovis* are the alveoli, alveolar sacs or alveolar ducts, which are not reached by particles in the respiratory tract of $>5\mu\text{m}$ (Corner et al., 2011).

- c. The relative importance of different locations (housing, pasture) with respect to badger-to-cattle transmission remains uncertain. Contacts between badgers and cattle have been reported both in housing (including Ward et al., 2010) and at pasture (including Payne et al., 2015). A recent study has found that direct contact between badgers and cattle is very infrequent, irrespective of whether cattle were housed or at pasture (Woodroffe et al., 2016).
- d. A number of risk mitigation strategies have been proposed, including cattle grazing regimes, habitat manipulation, management of latrines, and protection of farm buildings (Ward et al., 2010), addressing risks at pasture and in housing. Further, strategies to successfully exclude badgers from housing have been demonstrated (Judge et al., 2011).
- e. The costs of these strategies are likely to vary by farm, but could be substantial. Given the nature of some of these strategies, their effectiveness may be greatly affected by farmer diligence (Judge et al., 2011).
- f. To this point, relevant research has primarily focused on the effectiveness with which these strategies might reduce or prevent contact. No work has yet been undertaken (for example, using controlled field trials) to critically evaluate the impact of one or more of these strategies on herd bTB risk.
- g. In conclusion, there is as yet no empirical evidence linking improved biosecurity with reduced wildlife-related bTB risks (More, 2009; O'Hagan et al., 2016). Further, the relative effectiveness of different risk mitigation strategies in terms of bTB risk from infected wildlife is also unknown.

Risk mitigation responsibilities

132. It may be useful to view biosecurity from the perspectives of '*who can meaningfully control/who's responsible*'? This has some similarities to the exacerbator-pays approach to cost allocation in the New Zealand bTB eradication programme⁹. Using this approach:

- a. Farmers have responsibility for cattle-related biosecurity measures on their farms, noting that farmers can meaningfully control each of the above-mentioned risk mitigation measures to limit contact with infected cattle.

- b. The government has responsibility for wildlife-related measures. Protected wildlife can be considered a public good²⁰, and are therefore the responsibility of government rather than individual farmers. Further, as argued above, there is currently no empirical evidence demonstrating that individual farmers can reduce wildlife-related bTB risks by implementing biosecurity measures on their farms.

Policy considerations

133. The Review highlights several strategies to increase awareness of biosecurity on farms in Northern Ireland, including the development of a checklist to guide biosecurity assessment and the provision of farm-specific biosecurity advice.

- a. The proposed strategies are supported, but with caution.
- i. At this point in time, proven options are available to effectively limit cattle-, but not wildlife-, -related bTB risks. It is important that resources and advice for farmers reflect the best-available science, to ensure that farmers focus their efforts on those biosecurity strategies with proven effectiveness in reducing future farm-level bTB risk. In this context, greater emphasis should be placed on conclusions from prospective (cohort [observational], field trials [experimental]) studies in comparison to cross-sectional or retrospective (case-control [observational]) studies, given the need to distinguish association and causation.
 - ii. Contact with infected cattle is one of a range of factors that influence future farm-level bTB risk. Further, the effectiveness of cattle-related biosecurity is greatly influenced by ongoing farmer diligence. For these reasons, improved on-farm biosecurity will not always lead to reduced bTB risk. As a consequence, the impact of the proposed measures will be variable, and on many farms may be minimal.

²⁰ Public goods and services (for example, national institutions for law and order, public roads, education, hospitals etc.) are generally funded through compulsory taxation and are, therefore, available to all. In economic terms, public and private goods are distinguished using the principles of excludability and rivalry. Purely public goods are those goods from which it is not possible to exclude one consumer without excluding all (non-excludability) and of which the consumption by one person does not reduce its availability for consumption by others (non-rivalry) (Ahuja, 2004).

134. It may also be possible to consider improved farm biosecurity within a broader context, in particular efforts currently being made by Animal Health and Welfare Northern Ireland (AHWNI) with respect to BVD eradication and, shortly, JD control. Of note:
- a. Concerns with respect to biosecurity adoption have also been noted on Irish farms (Sayers et al., 2013), despite many years of biosecurity messages as part of the national bTB control programme. Over the last 3 or so years, however, there appears to have been a substantial increase in farmer understanding and awareness of farm biosecurity. This has been attributed to the national BVD eradication programme, which has been compulsory since 2013. Under this programme, all farmers are required to consider farm-level biosecurity decisions to prevent or control BVD on their farms. The national discussion on the issue of PI (animals persistently infected with BVD virus) retention has been particularly useful in this regard, including research quantifying the future risk posed by PI retention to both the index herd (Graham et al., 2015) and its neighbours (Graham et al., 2016).
 - b. AHI have produced a series of information leaflets for Irish farmers, being practical evidence-based biosecurity measures suited to Irish farms²¹. A scientific paper was also produced, to provide a robust foundation for recommendations (Mee et al., 2012). The leaflets address the following topics:
 - i. Understanding infectious diseases
 - ii. Bioexclusion: keeping infectious diseases out of your herd
 - iii. Purchasing stock: reducing disease risks
 - iv. Preventing disease spread within your farm - biocontainment

²¹ See http://animalhealthireland.ie/?page_id=395

135. It would be valuable to complement the proposed strategies with supporting research:
- a. There is a need for a clearer understanding of the relative importance of the two key biosecurity-related bTB risks in Northern Ireland, including contact with infected cattle and contact with infected wildlife.
 - b. In time, there is a need for a critical evaluation of the impact of improved farm biosecurity on future bTB risk, including the relative contribution of different biosecurity practices.
 - c. There is a building body of international literature on motivators for, and constraints to, improved on-farm biosecurity (including Kristensen and Jakobsen, 2011; Frössling and Nöremark, 2016; Shortall et al., 2016). It would be valuable to consider similar studies from Northern Ireland, to identify strategies to encourage livestock farmers to best achieve improved biosecurity.

Improvement notices

136. As highlighted in the Review, improved on-farm biosecurity has the potential to substantially improve animal health on individual farms. However, as highlighted above, improved on-farm biosecurity will not always lead to reduced bTB risk. Given this context, there is a need to review progress and available evidence concerning the impact of improved farm-level biosecurity on future bTB risk in Northern Ireland. This is foreshadowed in the Review.

Informed purchasing

137. As outlined previously, a number of strategies are available to limit the infection risk posed through animal movement, as outlined previously. Each is underpinned by the same principle – seeking to limit the probability that an introduced animal is infected with *M. bovis*. These strategies (and strategy combinations) vary with respect to their failure rate (the % infected among all animals traded). Maintaining a closed herd has a zero failure rate whereas the failure rate for pre-movement testing may be relatively high (given the imperfect operating characteristics of the tests used).

138. In the context of bTB eradication, there are substantial differences between countries with respect to the implementation of these strategies. Using two examples:
- a. During the bTB eradication programme in Australia, the movement of cattle between herds was determined on the basis of herd and area risk, to limit the potential for spread of infection to lower risk herds and areas (More et al., 2015).
 - b. In England and Wales, there has been considerable discussion about a potential role for a risk-based trading scheme, based on the probability of bTB in a herd (Adkin et al., 2016a,b).
139. 'Informed purchasing', as outlined in this Review, is consistent with the above-mentioned principle, seeking to limit the probability that an introduced animal is infected with *M. bovis*. Using this strategy, purchasing decisions are informed by knowledge of past testing history, of the animal and herd. As highlighted in the Review, it is critical that an informed purchasing scheme is practical, transparent and based on accurate and available data.
140. Substantial progress has been made in the identification of animal- and herd-level risk factors for bTB (so-called explanatory models; see reviews by Skuce et al., 2012; Broughan et al., 2016) As yet, however, there has been limited success in translating this knowledge into tools that allow accurate prediction of future bTB risk (so-called predictive models; Karolemeas et al., 2010; Wolfe et al., 2010). For this reason, while past testing history will be useful, future prediction of bTB risk is certain to be imperfect. For this reason, the failure rate from informed purchasing will be non-negligible, and may be relatively high.
141. There is a need for ongoing research to critically evaluate the value of informed purchasing, with respect to infection control benefit to the national bTB eradication programme.
142. From an epidemiological perspective, care will be needed in identifying the data that will most accurately assist an interested buyer. Animal-level test results may be of limited value, or worse, without a clear understanding of the current and past bTB risk of the herd(s).

Farm fragmentation

143. Farm fragmentation is a feature of farming in several bTB-affected countries, including Northern Ireland. As yet, however, this issue has never been comprehensively addressed, in terms of the risk posed to national bTB control and eradication. Therefore, the proposed review is welcomed. As part of this review, epidemiological research will be needed to quantify the impact of farm fragmentation on future infection risk following a bTB breakdown.
144. The proposed interim measure (a notice to limit the risk of spread associated with fragmentation) is epidemiologically sound, but should be revisited once the above-mentioned review has been completed.

References

- Abernethy, D.A., Denny, G.O.D., Pfeiffer, D.U., Wrigley, K., 2003. Survey for *Mycobacterium bovis* infection in road-traffic-accident badgers in Northern Ireland. Proceedings of the 10th International Symposium on Veterinary Epidemiology and Economics, Viña del Mar, Chile.
- Abernethy, D.A., Upton, P., Higgins, I.M., McGrath, G., Goodchild, A.V., Rolfe, S.J., Broughan, J.M., Downs, S.H., Clifton-Hadley, R.S., Menzies, F.D., la Rua-Domenech, de, R., Blissitt, M.J., Duignan, A., More, S.J., 2013. Bovine tuberculosis trends in the UK and the Republic of Ireland, 1995-2010. *Vet Rec* 172, 312–312.
- Abernethy, D.A., Walton, E., Menzies, F., Courcier, E., Robinson, P., 2011. *Mycobacterium bovis* surveillance in European badgers (*Meles meles*) killed by vehicles in Northern Ireland: an epidemiological evaluation. *Epidémiol et santé anim* 59-60, 216-218.
- Adkin, A., Brouwer, A., Downs, S.H., Kelly, L., 2016a. Assessing the impact of a cattle risk-based trading scheme on the movement of bovine tuberculosis infected animals in England and Wales. *Prev Vet Med* 123, 23–31.
- Adkin, A., Brouwer, A., Simons, R.R.L., Smith, R.P., Arnold, M.E., Broughan, J., Kosmider, R., Downs, S.H., 2016b. Development of risk-based trading farm scoring system to assist with the control of bovine tuberculosis in cattle in England and Wales. *Prev Vet Med* 123, 32–38.
- Ahuja, V., 2004. The economic rationale of public and private sector roles in the provision of animal health services. *Rev sci tech Off int Epiz* 23, 33-45.
- Álvarez, J., Pérez, A., Bezos, J., Marqués, S., Grau, A., Saez, J.L., Mínguez, O., de Juan, L., Domínguez, L., 2012. Evaluation of the sensitivity and specificity of bovine tuberculosis diagnostic tests in naturally infected cattle herds using a Bayesian approach. *Vet Microbiol* 155, 38–43.
- Andersen, P., Doherty, T.M., 2005. The success and failure of BCG - implications for a novel tuberculosis vaccine. *Nat Rev Microbiol* 3, 656–662.
- Aznar, I., Frankena, K., Byrne, A.W., More, S.J., De Jong, M.C.M., 2014. Infection dynamics and effective control options of tuberculosis in cattle and badgers. VI International *M. bovis* Conference, 16-19 June 2014, Cardiff, Wales. p37.
- Aznar, I., McGrath, G., Frankena, K., More, S.J., Martin, S.W., Martin, W., O'Keeffe, J., De Jong, M.C.M., 2011. Trial design to estimate the effect of vaccination on tuberculosis incidence in badgers. *Vet Microbiol* 151, 104–111.
- Aznar, I., More, S.J., Frankena, K., De Jong, M.C.M., 2013. Estimating the power of a *Mycobacterium bovis* vaccine trial in Irish badgers. *Prev Vet Med* 111, 297–303.
- Aznar, I., Frankena, K., More, S.J., Whelan, C., Martin, S.W., Martin, W., Gormley, E., Corner, L.A.L., Murphy, D., De Jong, M.C.M., 2014. Optimising and evaluating the characteristics of a multiple antigen ELISA for detection of *Mycobacterium bovis* infection in a badger vaccine field trial. *PLoS One* 9, e100139.

- Barrett, D., More, S.J., Graham, D., O'Flaherty, J., Doherty, M.L., Gunn, H.M., 2011. Considerations on BVD eradication for the Irish livestock industry. *Ir Vet J* 64, 12.
- Berry, D.P., Bermingham, M.L., Good, M., More, S.J., 2011. Genetics of animal health and disease in cattle. *Ir Vet J* 64, 5.
- Bradley, R., Wilesmith, J.W., 1993. Epidemiology and control of bovine spongiform encephalopathy (BSE). *Br Med Bull* 49, 932–959.
- Broughan, J.M., Judge, J., Ely, E., Delahay, R.J., Wilson, G., Clifton-Hadley, R.S., Goodchild, A.V., Bishop, H., Parry, J.E., Downs, S.H., 2016. A review of risk factors for bovine tuberculosis infection in cattle in the UK and Ireland. *Epidemiol Infect* 1–28.
- Byrne, A.W., Kenny, K., Fogarty, U., O'Keeffe, J.J., More, S.J., McGrath, G., Teeling, M., Martin, S.W., Dohoo, I.R., 2015. Spatial and temporal analyses of metrics of tuberculosis infection in badgers (*Meles meles*) from the Republic of Ireland: Trends in apparent prevalence. *Prev Vet Med* 122, 345–354.
- Byrne, A.W., O'Keeffe, J.J., Sleeman, D.P., Davenport, J., 2012. The ecology of the European badger (*Meles meles*) in Ireland: a review 112B, 105–132
- Carden, R.F., Carlin, C.M., Marnell, F., Mcelholm, D., Hetherington, J., Gammell, M.P., 2011. Distribution and range expansion of deer in Ireland. *Mammal Rev* 41, 313–325.
- Carrique-Mas, J.J., Medley, G.F., Green, L.E., 2008. Risks for bovine tuberculosis in British cattle farms restocked after the foot and mouth disease epidemic of 2001. *Prev Vet Med* 84, 85–93.
- Carter, S.P., Chambers, M.A., Rushton, S.P., Shirley, M.D.F., Schuchert, P., Pietravalle, S., Murray, A., Rogers, F., Gettinby, G., Smith, G.C., Delahay, R.J., Hewinson, R.G., McDonald, R.A., 2012. BCG vaccination reduces risk of tuberculosis infection in vaccinated badgers and unvaccinated badger cubs. *PLoS One* 7, e49833.
- Chambers, M.A., Carter, S.P., Wilson, G.J., Jones, G., Brown, E., Hewinson, R.G., Vordermeier, M., 2014. Vaccination against tuberculosis in badgers and cattle: an overview of the challenges, developments and current research priorities in Great Britain. *Vet Rec* 175, 90–96.
- Chambers, M.A., Rogers, F., Delahay, R.J., Lesellier, S., Ashford, R., Dalley, D., Gowtage, S., Davé, D., Palmer, S., Brewer, J., Crawshaw, T., Clifton-Hadley, R.S., Carter, S., Cheeseman, C., Hanks, C., Murray, A., Palphramand, K., Pietravalle, S., Smith, G.C., Tomlinson, A., Walker, N.J., Wilson, G.J., Corner, L.A.L., Rushton, S.P., Shirley, M.D.F., Gettinby, G., McDonald, R.A., Hewinson, R.G., 2011. Bacillus Calmette-Guérin vaccination reduces the severity and progression of tuberculosis in badgers. *Proc Biol Sci B* 278, 1913–1920.
- Clegg, T.A., Blake, M., Healy, R., Good, M., Higgins, I.M., More, S.J., 2013. The impact of animal introductions during herd restrictions on future herd-level bovine tuberculosis risk. *Prev Vet Med* 109, 246–257.
- Clegg, T.A., Good, M., Duignan, A., Doyle, R., Blake, M., More, S.J., 2011a. Longer-term risk of *Mycobacterium bovis* in Irish cattle following an inconclusive diagnosis to the single intradermal comparative tuberculin test. *Prev Vet Med* 100, 147–154.

- Clegg, T.A., Good, M., Duignan, A., Doyle, R., More, S.J., 2011b. Shorter-term risk of *Mycobacterium bovis* in Irish cattle following an inconclusive diagnosis to the single intradermal comparative tuberculin test. *Prev Vet Med* 102, 255–264.
- Clegg, T.A., Good, M., More, S.J., 2015. Future risk of bovine tuberculosis recurrence among higher risk herds in Ireland. *Prev Vet Med* 118, 71–79.
- Clegg, T.A., More, S.J., Higgins, I.M., Good, M., Blake, M., Williams, D.H., 2008. Potential infection-control benefit for Ireland from pre-movement testing of cattle for tuberculosis. *Prev Vet Med* 84, 94–111.
- Corner, L.A.L., More, S.J., Williams, D.H., O'Boyle, I., Costello, E., Sleeman, D.P., Griffin, J.M., 2008. The effect of varying levels of population control on the prevalence of tuberculosis in badgers in Ireland. *Res Vet Sci* 85, 238–249.
- Corner, L.A.L., Costello, E., O'Meara, D., Lesellier, S., Aldwell, F.E., Singh, M., Hewinson, R.G., Chambers, M.A., Gormley, E., 2010. Oral vaccination of badgers (*Meles meles*) with BCG and protective immunity against endobronchial challenge with *Mycobacterium bovis*. *Vaccine* 28, 6265–6272.
- Corner, L.A.L., Murphy, D., Gormley, E., 2011. *Mycobacterium bovis* infection in the Eurasian badger (*Meles meles*): the disease, pathogenesis, epidemiology and control. *J Comp Pathol* 144, 1–24.
- Corner, L.A.L., O'Meara, D., Costello, E., Lesellier, S., Gormley, E., 2012. The distribution of *Mycobacterium bovis* infection in naturally infected badgers. *Vet J* 194, 166–172.
- Dawson, K.L., Stevenson, M., Sinclair, J.A., Bosson, M.A., 2014. Recurrent bovine tuberculosis in New Zealand cattle and deer herds, 2006–2010. *Epidemiol Infect* 142, 2065–2074.
- Dobson, A., 2004. Population Dynamics of Pathogens with Multiple Host Species. *Am Nat* 164, S64–S78.
- Donnelly, C.A., Wei, G., Johnston, W.T., Cox, D.R., Woodroffe, R., Bourne, F.J., Cheeseman, C.L., Clifton-Hadley, R.S., Gettinby, G., Gilks, P., Jenkins, H.E., Le Fevre, A.M., McInerney, J.P., McInerney, J.P., Morrison, W.I., 2007. Impacts of widespread badger culling on cattle tuberculosis: concluding analyses from a large-scale field trial. *Int J Infect Dis* 11, 300–308.
- Doyle, L.P., Courcier, E.A., Gordon, A.W., O'Hagan, M.J.H., Menzies, F.D., 2016. Bovine tuberculosis in Northern Ireland: Risk factors associated with duration and recurrence of chronic herd breakdowns. *Prev Vet Med* 131, 1–7.
- Doyle, L.P., Gordon, A.W., Abernethy, D.A., Stevens, K., 2014. Bovine tuberculosis in Northern Ireland: Risk factors associated with time from post-outbreak test to subsequent herd breakdown. *Prev Vet Med* 116, 47–55.
- Duignan, A., Good, M., More, S.J., 2012. A review of quality control in the national bovine tuberculosis control programme in Ireland. *Rev sci tech Off int Epiz* 31, 845–860.
- EFSA Panel on Animal Health and Welfare (AHAW), 2013. Scientific opinion on field trials for bovine tuberculosis vaccination. *EFSA J* 11, 3475.

Frankena, K., White, P., O'Keeffe, J., Costello, E., Martin, S.W., Martin, W., van Grevenhof, I., More, S.J., 2007. Quantification of the relative efficiency of factory surveillance in the disclosure of tuberculosis lesions in attested Irish cattle. *Vet Rec* 161, 679–684.

Frewer, L., Salter, B., 2002. Public attitudes, scientific advice and the politics of regulatory policy: The case of BSE. *Sci Publ Policy* 29, 137–145.

Frössling, J., Nöremark, M., 2016. Differing perceptions – Swedish farmers' views of infectious disease control. *Vet Med Sci* 2, 54–68.

Gallagher, M.J., Higgins, I.M., Clegg, T.A., Williams, D.H., More, S.J., 2013. Comparison of bovine tuberculosis recurrence in Irish herds between 1998 and 2008. *Prev Vet Med* 111, 237–244.

Geraghty, T., Graham, D., Mullowney, P., More, S.J., 2014. A review of bovine Johne's disease control activities in 6 endemically infected countries. *Prev Vet Med* 116, 1–11.

Gilbert, M., Mitchell, A., Bourn, D., Mawdsley, J., Clifton-Hadley, R.S., Wint, W., 2005. Cattle movements and bovine tuberculosis in Great Britain. *Nature* 435, 491–496.

Godfray, H.C.J., Donnelly, C.A., Kao, R.R., Macdonald, D.W., McDonald, R.A., Petrokofsky, G., Wood, J.L., Woodroffe, R., Young, D.B., McLean, A.R., 2013. A restatement of the natural science evidence base relevant to the control of bovine tuberculosis in Great Britain. *Proc Biol Sci B* 280, 20131634.

Good, M., Duignan, A., 2011. An evaluation of the Irish Single Reactor Breakdown Protocol for 2005-2008 inclusive and its potential application as a monitor of tuberculin test performance. *Vet Microbiol* 151, 85–90.

Good, M., Clegg, T.A., Duignan, A., More, S.J., 2011. Impact of the national full herd depopulation policy on the recurrence of bovine tuberculosis in Irish herds, 2003 to 2005. *Vet Rec* 169, 581.

Goodchild, A.V., Watkins, G.H., Sayers, A.R., Jones, J.R., Clifton-Hadley, R.S., 2012. Geographical association between the genotype of bovine tuberculosis in found dead badgers and in cattle herds. *Vet Rec* 170, 259–259.

Gormley, E., Doyle, M., Duignan, A., Good, M., More, S.J., Clegg, T.A., 2013. Identification of risk factors associated with disclosure of false positive bovine tuberculosis reactors using the gamma-interferon (IFN γ) assay. *Vet Res* 44, 117.

Gortázar, C., Che Amat, A., O'Brien, D.J., 2015. Open questions and recent advances in the control of a multi-host infectious disease: animal tuberculosis. *Mammal Rev* 45, 160–175.

Gortázar, C., Diez-Delgado, I., Barasona, J.A., Vicente, J., La Fuente, De, J., Boadella, M., 2015. The wild side of disease control at the wildlife-livestock-human interface: A Review. *Front Vet Sci* 1, 443.

Graham, D., Clegg, T.A., O'Sullivan, P., More, S.J., 2015. Influence of the retention of PI calves identified in 2012 during the voluntary phase of the Irish national bovine viral diarrhoea virus (BVDV) eradication programme on herd-level outcomes in 2013. *Prev Vet Med* 120, 298–305.

- Graham, D., Clegg, T.A., Thulke, H.H., O'Sullivan, P., McGrath, G., More, S.J., 2016. Quantifying the risk of spread of bovine viral diarrhoea virus (BVDV) between contiguous herds in Ireland. *Prev Vet Med* 126, 30–38.
- Griffin, J.M., Williams, D.H., Kelly, G.E., Clegg, T.A., O'Boyle, I., Collins, J.D., More, S.J., 2005. The impact of badger removal on the control of tuberculosis in cattle herds in Ireland. *Prev Vet Med* 67, 237–266.
- Halloran, M.E., 1998. Concepts of infectious disease epidemiology. In: Rothman, J.J., Greenland, S. (Eds.), *Modern Epidemiology*, 2nd edition. Lippincott-Raven, Philadelphia, pp. 529–554.
- Hardstaff, J.L., Bulling, M.T., Marion, G., Hutchings, M.R., White, P.C.L., 2013. Modelling the impact of vaccination on tuberculosis in badgers. *Epidemiol Infect* 141, 1417–1427.
- Hardstaff, J.L., Marion, G., Hutchings, M.R., White, P.C.L., 2014. Evaluating the tuberculosis hazard posed to cattle from wildlife across Europe. *Res Vet Sci* 97 Suppl, S86–93.
- Jenkins, H.E., Woodroffe, R., Donnelly, C.A., 2008. The effects of annual widespread badger culls on cattle tuberculosis following the cessation of culling. *Int J Infect Dis* 12, 457–465.
- John, T.J., Samuel, R., 2000. Herd immunity and herd effect: new insights and definitions. *Eur J Epidemiol* 16, 601–606.
- Judge, J., McDonald, R.A., Walker, N., Delahay, R.J., 2011. Effectiveness of biosecurity measures in preventing badger visits to farm buildings. *PLoS One* 6, e28941.
- Karolemeas, K., McKinley, T.J., Clifton-Hadley, R.S., Goodchild, A.V., Mitchell, A., Johnston, W.T., Conlan, A.J.K., Donnelly, C.A., Wood, J.L.N., 2010. Predicting prolonged bovine tuberculosis breakdowns in Great Britain as an aid to control. *Prev Vet Med* 97, 183–190.
- Karolemeas, K., McKinley, T.J., Clifton-Hadley, R.S., Goodchild, A.V., Mitchell, A., Johnston, W.T., Conlan, A.J.K., Donnelly, C.A., Wood, J.L.N., 2011. Recurrence of bovine tuberculosis breakdowns in Great Britain: risk factors and prediction. *Prev Vet Med* 102, 22–29.
- Kelly, G.E., Condon, J., More, S.J., Dolan, L.A., Higgins, I.M., Eves, J., 2008. A long-term observational study of the impact of badger removal on herd restrictions due to bovine TB in the Irish midlands during 1989–2004. *Epidemiol Infect* 136, 1362–1373.
- Kristensen, E., Jakobsen, E.B., 2011. Danish farmers' perception of biosecurity. *Prev Vet Med* 99, 122–129.
- Lahuerta-Marin, A., Gallagher, M., McBride, S., Skuce, R.A., Menzies, F., McNair, J., McDowell, S.W.J., Byrne, A.W., 2015. Should they stay, or should they go? Relative future risk of bovine tuberculosis for interferon-gamma test-positive cattle left on farms. *Vet Rec* 46, 1.
- Lange, M., Kramer-Schadt, S., Thulke, H.-H., 2012. Efficiency of spatio-temporal vaccination regimes in wildlife populations under different viral constraints. *Vet Rec* 43, 37.

- Lange, M., Siemen, H., Blome, S., Thulke, H.H., 2014. Analysis of spatio-temporal patterns of African swine fever cases in Russian wild boar does not reveal an endemic situation. *Prev Vet Med* 117, 317–325.
- Leslie, E., Cowled, B., Garner, G.M., Toribio, J.-A.L.M.L., Ward, M.P., 2014. Effective surveillance strategies following a potential classical swine fever incursion in a remote wild pig population in north-western Australia. *Transbound Emerg Dis* 61, 432–442.
- Liebana, E., Johnson, L., Gough, J., Durr, P., Jahans, K., Clifton-Hadley, R.S., Spencer, Y., Hewinson, R.G., Downs, S.H., 2008. Pathology of naturally occurring bovine tuberculosis in England and Wales. *Vet J* 176, 354–360.
- Lindberg, A., Alenius, S., 1999. Principles for eradication of bovine viral diarrhoea virus (BVDV) infections in cattle populations. *Vet Microbiol* 64, 197–222.
- Livingstone, P., Hancox, N., Nugent, G., de Lisle, G.W., 2015. Toward eradication: the effect of *Mycobacterium bovis* infection in wildlife on the evolution and future direction of bovine tuberculosis management in New Zealand. *NZ Vet J* 63 Suppl 1, 4–18.
- Mangtani, P., Abubakar, I., Ariti, C., Beynon, R., Pimpin, L., Fine, P.E.M., Rodrigues, L.C., Smith, P.G., Lipman, M., Whiting, P.F., Sterne, J.A., 2014. Protection by BCG vaccine against tuberculosis: a systematic review of randomized controlled trials. *Clin Infect Dis* 58, 470–480.
- McCluskey, B., Lombard, J., Strunk, S., Nelson, D., Robbe-Austerman, S., Naugle, A., Edmondson, A., 2014. *Mycobacterium bovis* in California dairies: A case series of 2002–2013 outbreaks. *Prev Vet Med* 115, 205–216.
- Mee, J.F., Geraghty, T., O'Neill, R., More, S.J., 2012. Bioexclusion of diseases from dairy and beef farms: risks of introducing infectious agents and risk reduction strategies. *Vet J* 194, 143–150
- Miller, R.S., Sweeney, S.J., 2013. *Mycobacterium bovis* (bovine tuberculosis) infection in North American wildlife: current status and opportunities for mitigation of risks of further infection in wildlife populations. *Epidemiol Infect* 141, 1357–1370.
- More, S.J., 2009. What is needed to eradicate bovine tuberculosis successfully: An Irish perspective. *Vet J* 180, 275–278.
- More, S.J., Clegg, T.A., McGrath, G., Collins, J.D., Corner, L.A.L., Gormley, E., 2007. Does reactive badger culling lead to an increase in tuberculosis in cattle? *Vet Rec* 161, 208–209.
- More, S.J., Good, M., 2015. Understanding and managing bTB risk: perspectives from Ireland. *Vet Microbiol* 176, 209–218.
- More, S.J., Radunz, B., Glanville, R.J., 2015. Lessons learned during the successful eradication of bovine tuberculosis from Australia. *Vet Rec* 177, 224–232.
- Murray, D., More, S.J., 2012. Evaluation of single reactor bovine tuberculosis breakdowns based on analysis of reactors slaughtered at an Irish export meat plant. *Vet Rec* 170, 516–516.

Murphy, D., Costello, E., Aldwell, F.E., Lesellier, S., Chambers, M.A., Fitzsimons, T., Corner, L.A.L., Gormley, E., 2014. Oral vaccination of badgers (*Meles meles*) against tuberculosis: Comparison of the protection generated by BCG vaccine strains Pasteur and Danish. *Vet J* 200, 362–367.

Ní Bhuachalla, D., Corner, L.A.L., More, S.J., Gormley, E., 2015. The role of badgers in the epidemiology of *Mycobacterium bovis* infection (tuberculosis) in cattle in the United Kingdom and the Republic of Ireland: current perspectives on control strategies. *Vet Med Res Rep* 6, 27–38.

Nöremark, M., Frössling, J., Lewerin, S.S., 2010. Application of routines that contribute to on-farm biosecurity as reported by Swedish livestock farmers. *Transbound Emerg Dis* 57, 225–236.

Nusser, S.M., Clark, W.R., Otis, D.L., Huang, L., 2008. Sampling Considerations for Disease Surveillance in Wildlife Populations. *J Wildl Manage* 72, 52–60.

O'Brien, D.J., Schmitt, S.M., Fierke, J.S., Hogle, S.A., Winterstein, S.R., Cooley, T.M., Moritz, W.E., Diegel, K.L., Fitzgerald, S.D., Berry, D.E., Kaneene, J.B., 2002. Epidemiology of *Mycobacterium bovis* in free-ranging white-tailed deer, Michigan, USA, 1995–2000. *Prev Vet Med* 54, 47–63.

O'Brien, D.J., Schmitt, S.M., Fitzgerald, S.D., Berry, D.E., 2011. Management of bovine tuberculosis in Michigan wildlife: Current status and near term prospects. *Vet Microbiol* 151, 179–187.

O'Connor, C.M., Haydon, D.T., Kao, R., 2012. An ecological and comparative perspective on the control of bovine tuberculosis in Great Britain and the Republic of Ireland. *Prev Vet Med* 104, 185–197.

O'Corry-Crowe, G., Hammond, R.F., Eves, J., Hayden, T.J., 1996. The effect of reduction in badger density on the spatial organisation and activity of badgers *Meles meles* L. in relation to farms in central Ireland. *Biol Environ* 96B, 147–158.

O'Hagan, M.J.H., Matthews, D.I., Laird, C., McDowell, S.W.J., 2016. Herd-level risk factors for bovine tuberculosis and adoption of related biosecurity measures in Northern Ireland: A case-control study. *Vet J* 213, 26–32.

O'Keeffe, J.J., Martin, S.W., Byrne, A., White, P., Doyle, R., Fitzgerald, M., Hayes, M., Lynch, D., Monaghan, P., O'Brien, M., 2016. Replacing reactive culling of badgers by badger vaccination: part of the end-game of the eradication of bovine tuberculosis in the Republic of Ireland. *Proceedings of the World Buiatrics Congress 2016, Dublin*, p231.

Olea-Popelka, F.J., Fitzgerald, P., White, P., McGrath, G., Collins, J.D., O'Keeffe, J., Kelton, D.F., Berke, O., More, S.J., Martin, W., Martin, S.W., 2009. Targeted badger removal and the subsequent risk of bovine tuberculosis in cattle herds in county Laois, Ireland. *Prev Vet Med* 88, 178–184

Olea-Popelka, F.J., Freeman, Z., White, P., Costello, E., O'Keeffe, J., Frankena, K., Martin, S.W., Martin, W., More, S.J., 2012. Relative effectiveness of Irish factories in the surveillance of slaughtered cattle for visible lesions of tuberculosis, 2005-2007. *Ir Vet J* 65, 2.

- Olea-Popelka, F.J., White, P., Collins, J.D., O'Keeffe, J., Kelton, D.F., Martin, S.W., 2004. Breakdown severity during a bovine tuberculosis episode as a predictor of future herd breakdowns in Ireland. *Prev Vet Med* 63, 163–172.
- Payne, A., Moyon, J.L., Boschioli, M.L., Gueneau, E., Rambaud, T., Dufour, B., Gilot-Fromont, E., Hars, J., 2013. Bovine tuberculosis in “Eurasian” badgers (*Meles meles*) in France. *Eur J Wildl Res* 59, 331–339.
- Payne, A., Chappa, S., Hars, J., Dufour, B., Gilot-Fromont, E., 2015. Wildlife visits to farm facilities assessed by camera traps in a bovine tuberculosis-infected area in France. *Eur J Wildl Res* 62, 33–42.
- Radunz, B., 2006. Surveillance and risk management during the latter stages of eradication: experiences from Australia. *Vet Microbiol* 112, 283–290.
- Rees, E.E., Bélanger, D., Lelièvre, F., Coté, N., Lambert, L., 2011. Targeted surveillance of raccoon rabies in Québec, Canada. *J Wildl Manage* 75, 1406–1416.
- Robertson, A., Chambers, M.A., Delahay, R.J., McDonald, R.A., Palphramand, K.L., Rogers, F., Carter, S.P., 2015. Exposure of nontarget wildlife to candidate TB vaccine baits deployed for European badgers. *Eur J Wildl Res* 1–7.
- Robinson, P.A., Corner, L.A.L., Courcier, E.A., McNair, J., Artois, M., Menzies, F.D., Abernethy, D.A., 2012. BCG vaccination against tuberculosis in European badgers (*Meles meles*): A review. *Comp Immunol Microbiol Infect Dis* 35, 277–287.
- Rothman, K.J., Greenland, S., Lash, T.L., 2008. *Modern Epidemiology*, 3rd edition. Lippincott Williams & Wilkins, Philadelphia.
- Sayers, R.G., Sayers, G.P., Mee, J.F., Good, M., Bermingham, M.L., Grant, J., Dillon, P.G., 2013. Implementing biosecurity measures on dairy farms in Ireland. *Vet J* 197, 259–267.
- Sheridan, M., 2011. Progress in tuberculosis eradication in Ireland. *Vet Microbiol* 151, 160–169.
- Shittu, A., Clifton-Hadley, R.S., Ely, E.R., Upton, P.U., Downs, S.H., 2013. Factors associated with bovine tuberculosis confirmation rates in suspect lesions found in cattle at routine slaughter in Great Britain, 2003-2008. *Prev Vet Med* 110, 395–404.
- Shortall, O., Ruston, A., Green, M.J., Brennan, M., Wapenaar, W., Kaler, J., 2016. Broken biosecurity? Veterinarians’ framing of biosecurity on dairy farms in England. *Prev Vet Med*. doi:10.1016/j.prevetmed.2016.06.001
- Sibbald, B., Roland, M., 1998. Understanding controlled trials: Why are randomised controlled trials important? *BMJ* 316, 201.
- Skuce, R.A., Allen, A.R., McDowell, S., 2012. Herd-level risk factors for bovine tuberculosis: a literature review. *Vet Med Int* 2012, 621210.
- Sleeman, D.P., Davenport, J., More, S.J., Clegg, T.A., Collins, J.D., Martin, S.W., Williams, D.H., Griffin, J.M., O'Boyle, I., 2009. How many Eurasian badgers *Meles meles* L. are there in the Republic of Ireland? *Eur J Wildl Res* 55, 333–344.

Smith, G.C., Budgey, R., Delahay, R.J., 2013. A simulation model to support a study of test and vaccinate or remove (TVR) in Northern Ireland. <https://www.daera-ni.gov.uk/sites/default/files/publications/dard/fera-tvr-modelling-report.pdf>

Smith, G.C., Cheeseman, C.L., Clifton-Hadley, R.S., Wilkinson, D., 2001. A model of bovine tuberculosis in the badger *Meles meles*: an evaluation of control strategies. *J Appl Ecol* 38, 509–519.

Smith, G.C., McDonald, R.A., Wilkinson, D., 2012. Comparing badger (*Meles meles*) management strategies for reducing tuberculosis incidence in cattle. *PLoS One* 7, e39250.

Toma, B., Vaillancourt, J.-P., Dufour, B., Eloit, M., Moutou, F., Marsh, W., Bénét, J.J., Sanaa, M., Michel, P. (eds), 1999. *Dictionary of Veterinary Epidemiology*. Iowa State University Press, Ames.

Warburton, B., Livingstone, P., 2015. Managing and eradicating wildlife tuberculosis in New Zealand. *NZ Vet J* 63 Suppl 1, 77–88.

Vandenbroucke, J.P., von Elm, E., Altman, D.G., Gøtzsche, P.C., Mulrow, C.D., Pocock, S.J., Poole, C., Schlesselman, J.J., Egger, M., STROBE Initiative, 2007. Strengthening the Reporting of Observational Studies in Epidemiology (STROBE): explanation and elaboration. *Epidemiology* 18, 805–835.

van Larebeke, N., Hens, L., Schepens, P., Covaci, A., Baeyens, J., Everaert, K., Bernheim, J.L., Vlietinck, R., De Poorter, G., 2001. The Belgian PCB and dioxin incident of January–June 1999: exposure data and potential impact on health. *Environ Health Perspect* 109, 265–273.

van Schaik, G., Schukken, Y.H., Nielen, M., Dijkhuizen, A.A., Barkema, H.W., Benedictus, G., 2002. Probability of and risk factors for introduction of infectious diseases into Dutch SPF dairy farms: a cohort study. *Prev Vet Med* 54, 279–289.

Viana, M., Mancy, R., Biek, R., Cleaveland, S., Cross, P.C., Lloyd-Smith, J.O., Haydon, D.T., 2014. Assembling evidence for identifying reservoirs of infection. *Trends Ecol Evol* 29, 270–279.

von Elm, E., Altman, D.G., Egger, M., Pocock, S.J., Gøtzsche, P.C., Vandenbroucke, J.P., 2007. The Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) statement: guidelines for reporting observational studies. *Lancet* 370, 1453–1457.

Ward, A.I., Judge, J., Delahay, R.J., 2010. Farm husbandry and badger behaviour: Opportunities to manage badger to cattle transmission of *Mycobacterium bovis*? *Prev Vet Med* 93, 2–10.

Wells, S.J., 2000. Biosecurity on dairy operations: Hazards and risks. *J Dairy Sci* 83, 2380–2386.

White, P., Martin, S.W., Martin, W., De Jong, M.C.M., O’Keeffe, J.J., More, S.J., Frankena, K., 2013. The importance of “neighbourhood” in the persistence of bovine tuberculosis in Irish cattle herds. *Prev Vet Med* 110, 346–355.

Wilkinson, D., Smith, G.C., Delahay, R.J., Cheeseman, C.L., 2004. A model of bovine tuberculosis in the badger *Meles meles*: an evaluation of different vaccination strategies. *J Appl Ecol* 41, 492–501.

Wolfe, D.M., Berke, O., Kelton, D.F., White, P., More, S.J., O'Keeffe, J., Martin, W., Martin, S.W., 2010. From explanation to prediction: a model for recurrent bovine tuberculosis in Irish cattle herds. *Prev Vet Med* 94, 170–177.

Woodroffe, R., Donnelly, C.A., Ham, C., Jackson, S.Y.B., Moyes, K., Chapman, K., Stratton, N.G., Cartwright, S.J., 2016. Badgers prefer cattle pasture but avoid cattle: implications for bovine tuberculosis control. *Ecol Lett* doi:10.1111/ele.12654

Woodroffe, R., Donnelly, C.A., Jenkins, H.E., Johnston, W.T., Cox, D.R., Bourne, F.J., Cheeseman, C.L., Delahay, R.J., Clifton-Hadley, R.S., Gettinby, G., Gilks, P., Hewinson, R.G., McInerney, J.P., Morrison, W.I., 2006. Culling and cattle controls influence tuberculosis risk for badgers. *PNAS* 103, 14713–14717.