Mapping of poverty and likely zoonoses hotspots

Zoonoses Project 4

Report to Department for International Development, UK
The objective of this report is to present data and expert knowledge on poverty and zoonoses hotspots to inform prioritisation of study areas on the transmission of disease in emerging livestock systems in the developing world, where prevention of zoonotic disease might bring greatest benefit to poor people.

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**Introduction**

Mapping and measuring the burden of zoonoses, the density and number of poor livestock keepers and emerging markets for livestock products can help identify the ‘hotspots’ where zoonoses not only impose significant burdens but where zoonoses management is likely to be pro-poor (targeted at poor livestock keepers and poor consumers of livestock products) and have most impact on helping small farmers reach emerging markets.

All zoonoses are not equal and a first step of the study was to categorise zoonoses according to epidemiology and impact. We considered three groups of zoonoses:

- **Endemic zoonoses** are present in many places and affect many people and animals.
- **Outbreak or epidemic zoonoses** are sporadic in temporal and spatial distribution.
- **Emerging zoonoses** newly appear in a population or have existed previously but are rapidly increasing in incidence or geographical range. Many occur as outbreaks.

The first chapter reviews the substantial literature on prioritising disease and identifies prioritisation criteria relevant to this study, namely: burden of human disease; impacts on livestock production and productivity; amenability to agricultural intervention; and, concern because of emergence or severity. This allowed us to identify 24 zoonoses of high importance to poor people, 13 of which we investigated in depth. Our identified priorities were broadly similar to comparable exercises.

The next chapter reviews current evidence on poverty and livestock, on livestock systems and their dynamics, and on zoonoses and how they are currently mapped. We update the map of poor livestock keepers of Thornton et al. (2002) and present an additional map based on sub-national data. Maps of livestock systems that are changing most rapidly in response to emerging markets are taken from Herrero et al. 2009 and Notenbaert et al. 2009), and vulnerability to climate change from Ericksen et al. 2011). The strengths and weaknesses of different maps are noted and quantitative examples provided on the massive under-reporting of zoonoses and animal diseases in poor countries.

The next chapter presents evidence from a systematic review of over 1,000 studies on the prevalence of the 13 priority zoonoses in people and animals. It focuses on the endemic zoonoses that impose greatest burden and a ‘top 20’ list is given of geographical hotspots. Data on zoonoses are also extracted from the WHO Global Burden of Disease and the ‘top 20’ countries identified. We include a case study that compares our systematic review with an ‘in-country review’ focusing on grey literature and literature in a language other than English. Finally, we discuss some of the challenges of the study and caution in interpreting the results. Maps are presented.

The next chapter updates the map of emerging disease events of Jones et al. (2008). For the first time, we map emerging zoonoses as distinct from other emerging disease events. A ‘top 20’ of geographical hotspots is given. Maps are presented. The last chapter provides maps of regional agro-ecosystems and summarises numbers of livestock, people and poor livestock keepers by system as well as the zoonoses context. It also draws some global conclusions from the study.

Annexes provide references for the papers in the systematic review of endemic zoonoses, the in-country review, and the systematic review of emerging zoonotic events. They provide information on the long list of zoonoses and the selection of the 13 most important to poor people in terms of burden and economic impacts.
**Key points**

There is a strong association between poverty, hunger, livestock keeping, and zoonoses.  
*Strength of evidence: strong*

Zoonotic disease has many aspects and existing disease reporting systems do not adequately capture the impact of zoonoses or identify investment opportunities. There is much unpublished information in grey literature of developing countries.  
*Strength of evidence: strong*

Across a range of zoonoses burden, poverty burden, and reliance on livestock, the hotspots for poverty, emerging livestock systems and zoonoses are (in decreasing order of importance both by region and country; countries in red appear in multiple listings):

- **South Asia**: India > Bangladesh > Pakistan
  - *Is higher than:* East and Central Africa: Ethiopia > Nigeria > Congo DR > Tanzania > Sudan
  - *Is higher than:* South East Asia: China > Indonesia > Myanmar > Vietnam
  - *Is higher than:* West Africa: Burkina Faso > Mali > Ghana

*Strength of evidence: moderate*

We updated maps of poor livestock keepers (table 0.1). Around 70% of the rural poor and 10% of the urban poor are dependent on livestock. The last decade has seen declines in density of poor livestock keepers in South America and South East Asia and lesser declines in parts of West Africa and South Asia. High density of poor livestock keepers is focal: around 6 hotspots and 14 countries bear the brunt. **Four** countries (India > Nigeria > Ethiopia > Bangladesh) have 44% of the world’s poor livestock keepers (table 0.1 and chapter 4.1).  
*Strength of evidence: moderate*

Areas with both high livestock populations and strong rising demand for livestock products offer highest opportunities for livestock to be a pathway out of poverty. Demand is largely driven by urbanisation, demographic growth and increasing wealth. Monogastric (poultry and pig) production responds more to increased demand because of their high reproduction rates and ease of intensification. Hence total number of monogastrics and magnitude of change in monogastric population are proxies for identifying emerging livestock systems (table 0.1 and chapter 4.3).  
Countries with both high numbers and large change include: India > Myanmar > Pakistan > Bangladesh = China  
*Strength of evidence: moderate*

The study distinguishes between three categories of zoonoses:

- **Endemic zoonoses**, present in many places and affecting many people and animals are responsible for the great majority of human cases of illness (we estimate 99.9%) and deaths (we estimate 96%) as well as the greatest reduction in livestock production. Examples are: brucellosis, leptospirosis, and salmonellosis. *Endemic zoonoses are of most concern where the objective is lowering the burden of human disease and increasing the productivity and profitability of livestock for poor people.*

- **Outbreak or epidemic zoonoses** are zoonoses that typically occur as outbreaks. Examples are: anthrax, rabies, Rift Valley fever, and, leishmaniasis. They are much more sporadic in temporal and spatial distribution than endemic zoonoses but may be more feared because of their unpredictability and in some cases, severity. They are often present in neglected populations with poor health services and infrastructure. *Outbreak zoonoses are of concern when there is an objective of reducing vulnerability of neglected populations.*
• Emerging zoonoses newly appear in a population or have existed previously but are rapidly increasing in incidence or geographical range. Many occur as outbreaks. They are relatively rare, around 300 events in the last 70 years. Most are of minimal impact, but historically, emerging diseases have been responsible for massive impacts (e.g. HIV AIDS). Emerging zoonoses are of concern when the object is foresight, and understanding disease emergence in order to try and avert pandemics of major impact.

**Strength of evidence: strong**

The study assessed 56 zoonoses, together responsible for around 2.5 billion cases of human illness and 2.7 million human deaths a year. We identified the 13 zoonoses most important to poor livestock keepers because of their impacts on human health, livestock sector, amenability to agriculture-based control, and other criteria (chapter 2). These were, in descending order: zoonotic gastrointestinal disease; leptospirosis; cysticercosis; zoonotic tuberculosis; rabies; leishmaniasis; brucellosis; echinococcosis; toxoplasmosis; Q fever; zoonotic trypanosomosis, hepatitis E; and anthrax.

**Strength of evidence: moderate**

The study searched for papers on zoonotic disease emergence events since 2004. Out of 43 new or newly identified events, most are viral and originate in wild animal hosts. Although the mappable zoonotic new events (n = 30) are globally spread across every continent, there may be clusters in northeast US, South America, Europe and South East Asia. These trends may reflect surveillance differences. There is a possible trend to more events in developing countries in recent years, which may reflect increased attention over this period. Combined with existing data on zoonotic EID events from 1940-2004 (n = 202), the clearest potential hotspots are USA and Western Europe, (this may also reflect historical surveillance differences). Countries with most events are USA, UK, Australia, and France (table 0.1).

**Strength of evidence: weak-moderate**

Massive under-reporting constrains our ability to understand and prevent disease. In sub Saharan Africa, 99.9% of livestock losses do not appear in official reports. At least 50% of these losses are probably due to notifiable diseases (farmers and experts rank many notifiable diseases as major causes of mortality including Newcastle disease, African swine fever, classical swine fever, trypanosomosis, East Coast fever, peste de petits ruminants).

**Strength of evidence: moderate to strong**

The study accessed information around 1,000 surveys on prevalence of endemic zoonoses, covering over 16 million subjects. A qualitative and semi-quantitative analysis suggests a strongly spatial distribution, with a few countries bearing most of the human and animal disease burden (chapter 3). The study also assessed the burden of zoonoses in the Global Burden of Disease (GBD) extracting data on 7 important zoonoses. This also shows a highly skewed distribution of human disease burden: 19 countries are responsible for 75% of the total burden in the GBD. Hotspots are: Nigeria, Ethiopia, Tanzania, Togo, and India.

**Strength of evidence: moderate**

**Countries appearing multiple times at the top of multiple metrics are (in descending order of importance):** India, China, Bangladesh, Ethiopia, Nigeria, Pakistan, Congo DR, Indonesia, Myanmar, and Tanzania.

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11 However, this analysis does not consider trends in reducing the burden of zoonoses and the numbers of poor livestock keepers. If this were to be factored in, then China, Brazil and perhaps Indonesia would have a lower rank.
Table 0.1: The top 20 countries at the interface of poverty, emerging livestock systems and zoonoses according to different metrics (in descending order of importance)

<table>
<thead>
<tr>
<th>POVERTY INTERFACE</th>
<th>EMERGING MARKET INTERFACE</th>
<th>ZOONOSES INTERFACE</th>
</tr>
</thead>
<tbody>
<tr>
<td>Poor livestock keepers</td>
<td>Protein energy malnutrition&lt;sup&gt;a&lt;/sup&gt;</td>
<td>Monogastrics (TLU) 2010</td>
</tr>
<tr>
<td>India</td>
<td>India</td>
<td>China</td>
</tr>
<tr>
<td>Nigeria</td>
<td>Ethiopia</td>
<td>Brazil</td>
</tr>
<tr>
<td>Ethiopia</td>
<td>Nigeria</td>
<td>Indonesia</td>
</tr>
<tr>
<td>Bangladesh</td>
<td>China</td>
<td>India</td>
</tr>
<tr>
<td>Congo DR</td>
<td>Congo DR</td>
<td>Viet Nam</td>
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<tr>
<td>Pakistan</td>
<td>Bangladesh</td>
<td>Iran</td>
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<tr>
<td>Kenya</td>
<td>Pakistan</td>
<td>Philippines</td>
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<tr>
<td>Sudan</td>
<td>Indonesia</td>
<td>Thailand</td>
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<tr>
<td>China</td>
<td>Angola</td>
<td>Nigeria</td>
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<tr>
<td>Tanzania</td>
<td>Afghanistan</td>
<td>Ukraine</td>
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<tr>
<td>Indonesia</td>
<td>Tanzania</td>
<td>Pakistan</td>
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<tr>
<td>Madagascar</td>
<td>Brazil</td>
<td>Myanmar</td>
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<tr>
<td>Niger</td>
<td>Philippines</td>
<td>Bangladesh</td>
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<tr>
<td>Uganda</td>
<td>Uganda</td>
<td>Peru</td>
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<tr>
<td>Turkey</td>
<td>Mali</td>
<td>Colombia</td>
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<td>Philippines</td>
<td>Sudan</td>
<td>Ecuador</td>
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<tr>
<td>Afghanistan</td>
<td>Mozambique</td>
<td>Morocco</td>
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<tr>
<td>Egypt</td>
<td>Malawi</td>
<td>South Africa</td>
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<tr>
<td>Mozambique</td>
<td>South Africa</td>
<td>Bolivia</td>
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<tr>
<td>Burkina</td>
<td>Viet Nam</td>
<td>Egypt</td>
</tr>
<tr>
<td>Pakistan</td>
<td>Uganda</td>
<td>Zimbabwe</td>
</tr>
</tbody>
</table>

<sup>a</sup> Protein-energy malnutrition is the term used in the WHO GBD. WHO defines it as a nutritional deficiency resulting from either inadequate energy (caloric) or protein intake and manifesting in either marasmus or kwashiorkor.

*More than 20 countries because of tied ranks

Data sources for table 0.1. Poor livestock keepers, this study; Protein energy malnutrition: extracted from WHO GBD, 2009; Monogastrics, number of poultry and pigs in developing countries converted to tropical livestock units (TLU), FAOSTAT, 2012; Rapid change monogastrics: % increase in pigs and poultry in developing countries from 2000 to 2030, this study (based on IMPACT model); Zoonoses burden GBD: Burden of zoonoses extracted from WHO GBD, 2004 using the assumptions set out in chapter 3; Endemic zoonoses prevalence, this study; Emerging zoonoses events, this study,
Key maps

Density of poor livestock keepers (update of Thornton et al., 2002 by Kruska, this study)

This map shows the density of poor livestock keepers (number of poor livestock keepers per km
square). Countries with most poor livestock keepers are ranked in table 0.1 and estimates of the
absolute number of poor livestock and density of livestock keepers per country are given in chapter 4.

Hotspots for poverty & livestock keeping
- Absolute numbers: India > Nigeria > Bangladesh > Congo > Pakistan > Kenya > Sudan
- High density: South Asia, East Africa highlands, great lakes, Nigeria, west Africa, littoral
  South-East Asia

Key points
- Around 1 billion poor people (<$2 a day) depend on livestock
- Around two thirds of the rural poor and one third of the urban poor depend on livestock
- Livestock provide one fifth to one half of household income for the poor
- In poor countries, livestock provide from 6 to 36% of protein intake

More details on poor livestock keepers are provided in chapter 2.
This map shows the percentage change in poultry numbers from 2000 to 2030 based on projections by Herrero et al. (2009) using the International Model for Policy Analysis of Agricultural Commodities and Trade (IMPACT) model for the ‘business as usual’ scenario used in the International Assessment of Agriculture Science and technology for Development (IAASTD). Increase in poultry production is one possible proxy for the demand-driven increase in livestock production called the ‘livestock revolution’. According to that study:

**Hotspots** for livestock sector growth in developing countries are:
- Poultry in South and East Asia > bovines in South and East Asia > poultry in sub Saharan Africa = pigs in sub Saharan Africa

**Key points**
- Livestock production is increasing rapidly in response to growth in population, growth in income, urbanisation and changing diets: the so-called livestock revolution
- Herrero et al. (2009), projects that, over the next 40 years, absolute growth in consumption will be greatest in South Asia and South East Asia and relative growth greatest in sub-Saharan Africa
- On the supply side, growth will be greatest in the poultry sector followed by bovine then small ruminants then pigs
- Emerging livestock systems offer opportunities for smallholders if they can access the inputs needed to reach emerging markets

More details on poor livestock keepers are provided in chapter 2.
This map shows the surveys on endemic zoonoses reviewed in this report which reported zoonoses affecting >1% of humans or animal. Over 1,000 surveys were accessed covering over 14 million animals, humans and livestock products.

**Hotspots for high prevalence of endemic disease confirmed by multiple surveys**
Nigeria, Ethiopia, Tanzania, Togo, India, Mali, Vietnam, Sudan, Bangladesh

**Key points**
In poor countries as a whole:
- 12% of animals have recent or current infections with brucellosis, reducing production by 8%
- 10% of livestock in Africa are infected with trypanosomosis, reducing their production by 15%
- 7% of livestock are currently infection with tuberculosis (TB), reducing their production by 6%
- and from 3-10% of human TB cases may be caused by zoonotic TB
- 17% of smallholder pigs show signs of current infection with cysticercosis, reducing their value and creating the enormous burden of human cysticercosis
- 27% of livestock show signs of current or past infection with bacterial food-borne disease, a major source of food contamination and illness in people
- 26% of livestock show signs of current or past infection with leptospirosis reducing production and acting as a reservoir for infection
- 25% of livestock show signs of current or past infection with Q fever, and are a major source of infection of farmers and consumers
This map shows locations of zoonotic emerging disease events between 2004 and 2011.

**Geographical hotspots**

Combined with existing data on zoonotic EID events from 1940-2004 (n = 202), the clearest potential hotspots are USA, South America, South East Asia and Western Europe, which may reflect historical surveillance differences.
Chapter 1: Zoonoses of most relevance to poor people in emerging livestock systems

Summary
This chapter identifies three categories of zoonoses important for different reasons: endemic zoonoses, outbreak zoonoses, emerging zoonoses, and, old zoonoses. We discuss previous work to prioritise zoonoses and some of the challenges in identifying zoonoses of importance to the poor. We identified 56 zoonoses that appeared in multiple listings and selected criteria to prioritise them. Together, the 56 zoonoses are responsible for an estimated 2.7 human million deaths and around 2.5 billion cases of human illness a year. For the top 13 zoonoses, the figures were 2.2 million human deaths and 2.4 billion cases of illness. Our prioritisation is broadly compatible with other exercises.

1.1 Introduction
Zoonoses are diseases transmissible between animals (domestic and wildlife) and humans. Around 60% of all human diseases and around 75% of emerging infectious diseases are zoonotic (Taylor et al., 2001; Woolhouse et al., 2005). In aggregate, they have high impacts on human health, livelihoods, animals and ecosystems. In the first global syntheses of the impact (partial) of zoonotic diseases, Grace et al. (2011a) estimated that, in least developed countries, 20% of human sickness and death was due to zoonoses or diseases recently jumped species from animals to people.

1.2 The rationale for prioritising zoonoses
What cannot be measured cannot be managed and the first recommendation of a high-level WHO-convened group was to assess the societal burden of disease attributable to zoonoses (Molyneux et al., 2011). Assessing, including mapping, of zoonoses is key to helping decision-makers and implementers plan and manage disease control.

Zoonoses can threaten human health in different ways:

- **Endemic zoonoses** are continually present to a greater or lesser degree in certain populations. Examples are cysticercosis, brucellosis, bovine tuberculosis, leptospirosis and food-borne zoonoses. They are common in poor populations and are responsible around a billion illnesses and millions of deaths every year (table 2.1). However, endemic zoonoses have been neglected by the international donor, standard setting, and research communities. Maps exist for human health burden of individual zoonoses, usually at country level but there are no available maps of endemic zoonoses as a group, and few maps for the impact of zoonoses on livestock.

- **Outbreak or epidemic zoonoses** typically occur intermittently. Examples are anthrax, rabies, Rift Valley fever, and leishmaniasis. Endemic zoonoses may occur as outbreaks in naïve populations or when triggered by events such as climate changes, flooding, waning immunity or concomittent hunger or disease. They typically have high temporal and spatial variability. Their overall impact in terms of morbidity, mortality and production loss is much less than endemic zoonoses but because they can ‘shock’ systems they are often of high priority to farmers and decision makers. They can also cause important economic losses, which are often related to reaction to the disease rather than the disease itself (Butler and Grace, forthcoming). Some diseases which now occur in endemic foci have in the past resulted in major outbreaks or epidemics.

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2 Some of these also occur as outbreaks but are differentiated from the outbreak zoonoses in that community surveys will generally show that the disease is present in communities, although it may only get attention when there is an outbreak involving multiple cases.
Emerging zoonoses newly appear in a population or have existed previously but are now rapidly increasing in incidence or geographical range. They are relatively rare, around 300 events in the last 70 years (Jones et al., 2008). Most are of minimal impact. Diseases novel in one place may be endemic in other places and burdens are not necessarily linked to site of emergence, so mapping the point of emergence may not correlate to impact on poor people at that point. Donors and decision-makers are often concerned about emerging diseases, whose impacts on poor farmers are orders of magnitude less than the impacts of endemic zoonoses. However, the potential impact (e.g. a new HIV AIDS) is at least of similar magnitude to endemic zoonoses. Good maps exist but may not be useful for informing research aimed at identifying poor at risk from zoonoses.

‘Old zoonoses’ were originally zoonotic but are now spread mainly or entirely by human-to-human transmission (some with remaining zoonotic reservoirs) (Grace and McDermott, 2011). These include HIV-AIDs, influenza, malaria, measles and dengue. These diseases have jumped species in many places and their burden is not linked to site of emergence. Their current order of magnitude is about similar to that of the endemic zoonoses (almost all due to HIV-AIDs). The ability of the livestock sector to predict, prevent and control these diseases is small and maps not likely to be useful for directing research activities so these will not be further discussed.

1.3 Review of zoonoses prioritisation exercises

In order to map zoonoses and poverty, information is needed on which zoonoses pose risk to the poor. One of the earliest attempts to prioritise zoonoses was conducted by ILRI (Perry et al., 2002) with support from DfID. Recent years have seen several other prioritisation exercises for zoonoses and animal health. These have been reviewed by the ENHanCE group (http://www.liv.ac.uk/enhance/) (Enhance, undated). Most use experts, criteria setting, and weighting to come up with lists. However, when evidence is both highly scarce and highly scattered (as is the case for zoonoses) then expert opinion is less useful; this is illustrated by major discrepancies between different systems of prioritising (Perry et al., 2009).

Prioritisation exercises use various criteria of importance including: livestock pathogens with a high actual human disease burden; rare zoonotic pathogens with severe disease manifestations in people; arthropod-borne and wildlife associated pathogens which may pose a severe risk in future (Havelaar et al., 2010; health consequences in animals, economic consequences in animals (Dufour et al., 2006); public health (severity and occurrence in humans), animal health (severity of disease coupled with economic consequences and occurrence in animals), and food (occurrence in food) (Cardoen et al., 2009).

Most of these prioritisation exercises were done in rich countries. Other challenges in identifying the zoonoses that matter most to the poor include:

- Capturing multiple impacts: Many endemic zoonoses and some emerging zoonoses have impacts on livestock causing death and reduced productivity as well as costs for control but prioritisation may focus only on some impacts.
- Lack of evidence on the adverse impacts caused by disease: Many zoonoses are not notifiable so are not recorded in official statistics. Even for notifiable diseases, many national reports are highly unreliable. As described in this report, under-reporting is a serious problem both in animal and human populations.
- Variability: zoonoses are often focal and some vary from year to year (predictably or not). For example, Rift Valley fever may be absent for decades before causing severe problems. Human trypanosomosis currently affects only thousands of people, but historically there have been major epidemics affecting millions of people.
1.4 Selection of zoonoses for prioritisation

In order to select ‘important’ zoonoses for further study, we used information from five listings of priority zoonoses or priority diseases that included zoonoses and were relevant to developing countries:

1) The World Health Organisation Global Burden of Disease
2) The World Animal Health Organisation list of notifiable zoonoses
3) Zoonoses important to poor people identified by expert consultation (Perry et al., 2002)
4) The Rosetta listing of infectious causes of death
5) A systematic review of zoonoses commissioned by DFID, which identified 373 zoonoses as important (Grace et al., 2011).

Zoonoses that appeared in more than one list were considered (n=56). We ranked these 56 zoonoses according to criteria considered important by the authors of this study. We selected the following criteria (Table 2.1):

- Human mortality (>1,000 deaths per year)
- Human morbidity (>1 million people affected)
- High impact on livestock sector
- Amenability to agriculture-based control
- Emergence or severity of disease in people

The major difference between our criteria and criteria used in previous studies was the inclusion of ‘amenability to agricultural intervention’ as a criterion. The rationale was that the ability to do something about a problem was an important criterion for prioritisation for donor agencies and decision makers. The complete table and weighting used is given in annex 1. By these criteria, 13 zoonoses were defined as most important (Table 1.3) to poor people. These 13 were selected for in-depth systematic literature review and mapping.

Together, the 56 zoonoses are responsible for an estimated 2.7 human million deaths and around 2.5 billion cases of human illness a year. For the top 13 zoonoses, the figures were 2.2 million human deaths and 2.4 billion cases of illness. Nine of the 13 top-ranked zoonoses were considered to have high impact on livestock, all have a wildlife interface, and all are amenable to agriculture-based interventions.
Table 2.1 The most important zoonoses in terms of human health impact, livestock impact, amenability to agricultural interventions, severity of disease and emergence (data from WHO and authoritative literature: when several authoritative estimates the mid point is given)

<table>
<thead>
<tr>
<th>Disease</th>
<th>Wildlife interface</th>
<th>Deaths human annual</th>
<th>Affected humans annual</th>
<th>Death &gt;1000 people</th>
<th>Affected&gt;1 million people</th>
<th>Animal impacts high</th>
<th>Farm intervention</th>
<th>Other (score =1)</th>
<th>Total score</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gastrointestinal (zoonotic)</td>
<td>Important</td>
<td>1,500,000</td>
<td>2,333,000,000</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Leptospirosis</td>
<td>Very important</td>
<td>123,000</td>
<td>1,700,000</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Cysticercosis</td>
<td>Some importance</td>
<td>50,000</td>
<td>50,000,000</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
</tr>
<tr>
<td>Tuberculosis (zoonotic)</td>
<td>Some importance</td>
<td>100,000</td>
<td>554,500</td>
<td>2</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Rabies</td>
<td>Important</td>
<td>70,000</td>
<td>70,000</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Leishmaniasis</td>
<td>Important</td>
<td>47,000</td>
<td>2,000,000</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Brucellosis</td>
<td>Some importance</td>
<td>25,000</td>
<td>500,000</td>
<td>2</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Echinococcosis</td>
<td>Important</td>
<td>18,000</td>
<td>300,000</td>
<td>2</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Toxoplasmosis</td>
<td>Important</td>
<td>10,000</td>
<td>2,000,000</td>
<td>2</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Q fever</td>
<td>Important</td>
<td>3,000</td>
<td>3,500,000</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Trypanosomosis (zoonotic)</td>
<td>Important</td>
<td>2,500</td>
<td>15,000</td>
<td>2</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Anthrax</td>
<td>Some importance</td>
<td>1,250</td>
<td>11,000</td>
<td>2</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Hepatitis E *</td>
<td>Some importance</td>
<td>300,000</td>
<td>14,000,000</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Chagas</td>
<td>Important</td>
<td>10,000</td>
<td>8,000,000</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>3</td>
</tr>
<tr>
<td>Chikungunya</td>
<td>Very important</td>
<td>12,500</td>
<td>500,000</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>Emerge 3</td>
</tr>
<tr>
<td>Clostridium difficile disease</td>
<td>Possible importance</td>
<td>3,000</td>
<td>300,000</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>Emerge 3</td>
</tr>
<tr>
<td>Dengue fever</td>
<td>Minor</td>
<td>20,000</td>
<td>50,000,000</td>
<td>2</td>
<td>1</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>3</td>
</tr>
<tr>
<td>Ebola</td>
<td>Very important</td>
<td>500</td>
<td>800</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>Severe 3</td>
</tr>
<tr>
<td>Hanta disease</td>
<td>Very important</td>
<td>1,750</td>
<td>175,000</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>Emerge 3</td>
</tr>
<tr>
<td>Avian influenza</td>
<td>Important</td>
<td>77</td>
<td>145</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>Emerge 3</td>
</tr>
<tr>
<td>Bov. Spongiform Encephalopathy^</td>
<td>Some importance</td>
<td>182</td>
<td>188</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>Severe 3</td>
</tr>
<tr>
<td>Psittacosis</td>
<td>Important</td>
<td>2,250</td>
<td>22,000</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>3</td>
</tr>
<tr>
<td>Japanese encephalitis</td>
<td>Possibly, bats</td>
<td>11,000</td>
<td>40,000</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>0</td>
<td>3</td>
</tr>
<tr>
<td>Buffalo pox</td>
<td>Not important</td>
<td>Negligible</td>
<td>Common</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>3</td>
</tr>
<tr>
<td>Rift Valley fever</td>
<td>Important</td>
<td>45</td>
<td>150</td>
<td>0</td>
<td>0</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td>Emerge 3</td>
</tr>
</tbody>
</table>

Note: high human mortality gets a double weight of as the most important criterion for many stakeholders. Total score = (human death x 2) + (humans affected) + (high livestock impacts) + (farm intervention possible) + (other concerns: severe or emerging disease). The maximum possible score is therefore 6 and the minimum 0.

* Importance of zoonotic transmission not fully known  ^ Not a problem in poor countries
1.6 Comparing with other assessments
ENhanCE (undated) reviewed 12 methods of disease prioritisation. Two were global (FAO/OIE and WHO), one focused on Rajasthan in India, while the rest focused on developed countries. A variety of methods were used: risk assessment approach, multi-criteria decision tools, and qualitative methods. Together the studies reviewed covered animal diseases, human diseases, and zoonoses. Of the 99 diseases appearing in the rankings reviewed, 33 were zoonoses.

Zoonoses appearing in multiple listings according to the ENhanCE review, in declining order of number of appearances, were:
- Salmonellosis
- Leptospirosis = rabies
- Campylobacteriosis = tuberculosis = West Nile virus = toxoplasmosis
- Listeriosis = anthrax = echinococcosis = E. coli infection = BSE = botulism
- Cryptosporidiosis = Japanese encephalitis = Q fever = Rift Valley fever = tetanus

Out of the top 18 ranked zoonoses across the 12 studies, 17 appeared in our top 25 listing. The exception was West Nile, which did not appear in our review. This has been most problematic in the Americas. Among all the 33 zoonoses listed in the review, 24 appeared in our list of top 25 zoonoses, suggesting reasonable similarity given the different criteria and focus.

A notable characteristic of recent and expert-driven prioritisations is the high ranking given to common, food-borne diseases (salmonellosis, campylobacteriosis, toxoplasmosis, listeriosis, toxigenic E. coli, and cryptosporidiosis). Decision makers and implementers using unstructured prioritisation often focus on classical zoonoses they have studied at university or emerging diseases with high media and donor attention rather than food-borne zoonoses which cause a greater burden of disease (Grace et al., 2010).

1.7 Mapping zoonoses: strengths and weaknesses
There are three types of existing zoonoses maps: emerging disease event maps, disease report maps and research-derived prevalence maps. A summary along with strengths and weaknesses are presented in table 1.2.

1. Emerging disease event maps. Jones et al. (2008) have identified emerging disease events as “the first temporal emergence of a pathogen in a human population which was related to the increase in distribution, increase in incidence or increase in virulence or other factor which led to that pathogen being classed as an emerging disease”. They identified 335 events between 1940 and 2004: 60% of which are zoonotic. An updated map, which shows only zoonotic emerging disease events, is presented in Chapter 4.

2. Disease report maps. There are several systems for reporting disease outbreaks: these are summarised in table 1.2. The most authoritative is the World Animal Health Organisation (OIE) World Animal Health Information Database (WAHID) maps. HealthMap (www.healthmap.org) aggregates all the major disease reporting systems and information sources. Members of the OIE (currently 178 countries) have a legal obligation to report certain diseases (currently 115). Maps are generated from the information provided.

3. Prevalence maps. These are based on studies assessing the prevalence of zoonoses in livestock, livestock products and people. Global prevalence maps exist for some individual zoonoses but data is often at country level. Some zoonoses have been mapped using geo-spatial data – notably trypanosomosis. The World Health Organisation (WHO) Global Burden of Disease (GBD) has also been mapped at country level.
Table 1.2 Zoonoses disease and disease outbreak reporting systems

<table>
<thead>
<tr>
<th>Source of data</th>
<th>Strengths</th>
<th>Weaknesses</th>
</tr>
</thead>
<tbody>
<tr>
<td>Reports from state veterinary services.</td>
<td>Notifiable, all 172 OIE member states obliged to report.</td>
<td>Little reporting of endemic disease by developing countries.</td>
</tr>
<tr>
<td>Reports from state veterinary services.</td>
<td>Supported by FAO. Resolution to village level possible</td>
<td>Not widely used. Access to data limited to users.</td>
</tr>
<tr>
<td>All reports verified by qualified moderators.</td>
<td>Quality, comprehensiveness and timeliness dependent on quality of surveillance.</td>
<td></td>
</tr>
<tr>
<td>Media sources and then verified by GPHIN officials.</td>
<td>Real time reports in 7 languages.</td>
<td>Developing countries under-reported. Not universally accessible and costs associated with use.</td>
</tr>
<tr>
<td>Combines alert mechanisms of OIE, FAO, WHO.</td>
<td>Official reports.</td>
<td>Developing countries under-reported.</td>
</tr>
<tr>
<td>Aggregates data from many sources.</td>
<td>Real time, most comprehensive.</td>
<td>Reflects weaknesses in the source data.</td>
</tr>
</tbody>
</table>

1.8 Challenges in reporting systems for zoonoses in developing countries

The challenges of mapping the multiple burdens of zoonoses include:

- Reporting systems cover only few of the important zoonoses. There are over 600 zoonoses and around 100 of these are of some importance (Grace et al., 2011). However, WHO GBD and OIE only cover 11 and 33 zoonoses respectively.
- The GBD does not distinguish between zoonotic and non-zoonotic causes of disease and for several diseases (including tuberculosis, schistosomiasis, gastro-intestinal disease) the proportion of disease attributable to zoonoses is not accurately known.
- Zoonoses are often confused with other diseases (e.g. malaria and typhoid) and this misdiagnosis leads to systematic under-reported in the human health system.
- OIE reporting grossly underestimates the importance of endemic zoonoses – see next section.
- Emerging disease databases give little information on actual burden on poor people or which diseases are likely to be problematic in the future.
- Meta-disease reporting systems (summarised in HealthMap) are only as good as the data they aggregate.

An important conclusion of our study is that massive under-reporting of zoonotic (and other diseases) in developing countries is a major impediment to understanding prevalence and impacts of disease and developing appropriate control. We illustrate this with the examples of brucellosis in poor countries and Q fever in Africa and also compare official reports of notifiable diseases with probable mortality of livestock in Africa.

a) The case of brucellosis – a well-known, widespread, notifiable zoonosis
Brucellosis is an important disease of cattle, sheep, goats and pigs. It is also important zoonosis and is notifiable to the OIE. In cattle, it can be suspected on clinical signs, as it causes late abortion with characteristic lesions on the placenta. It also causes carpal hygromas, a very specific indicator of brucellosis. Diagnostic tests are widely available and relatively inexpensive. Brucellosis is both important and easy to detect.

Commonly used tests for brucellosis detect antibodies produced in response to infection. Antibodies tested for persist for several months (IGG) or several years (IGM). Positive tests (to both antibodies) indicate the animal is currently sick, is chronically infected or has been infected in the last year or so. Hence positive tests are roughly equivalent to annual cases.

Our review captured information from 241 community surveys (that is, surveys from the general livestock community and not targeting high risk animals) of bovine, sheep and goat populations, representing 475,968 samples. The prevalence for different regions is shown in Table 1.2. From the number of ruminants, the prevalence of seropositive cases, and the relation between sero-positivity and disease we can predict the number of cases of brucellosis a year. The discrepancy between the number reported and the number predicted is several orders of magnitude. For example, for every 1 million cases in East Africa less than one case is reported to OIE. The situation is similar for other diseases reported to OIE. When there are 999,999 missed cases for every one report, surveillance is not fulfilling its purpose.

<table>
<thead>
<tr>
<th>Livestock prevalence %</th>
<th>Number of ruminants</th>
<th>Predicted cases a year</th>
<th>Cases reported 2010</th>
</tr>
</thead>
<tbody>
<tr>
<td>East Africa</td>
<td>8.2</td>
<td>257,377,760</td>
<td>21,104,976</td>
</tr>
<tr>
<td>West Africa</td>
<td>15.5</td>
<td>197,716,517</td>
<td>30,646,060</td>
</tr>
<tr>
<td>South Africa</td>
<td>14.2</td>
<td>59,806,724</td>
<td>8,492,555</td>
</tr>
<tr>
<td>North Africa</td>
<td>13.8</td>
<td>57,629,367</td>
<td>7,952,853</td>
</tr>
<tr>
<td>South Asia</td>
<td>16.0</td>
<td>683,181,040</td>
<td>109,308,966</td>
</tr>
<tr>
<td>South East Asia</td>
<td>2.9</td>
<td>21,247,586</td>
<td>616,180</td>
</tr>
</tbody>
</table>

b) The case of Q fever – a less well-known, difficult to diagnose, notifiable zoonosis

Q fever is an infectious disease of animals and humans caused by a species of bacteria (Coxiella burnetii). The main reservoirs are sheep, goats and cattle. It is highly contagious to humans and typically causes influenza-like illness, although some infections are asymptomatic and in rare cases fatal complications can ensue.

Q fever is a notifiable disease and appeared in the top 13 zoonoses in terms of impact on human health, livestock sector and other criteria in our listing (table 2.1).

Most tests for Q fever detect antibodies. Antibodies may persist for several years. Most of the surveys in our review were community based. For these, a positive result indicates current infection, chronic infection or infection in the last few years.

In our review the average sero-prevalence from community surveys in Africa was 26% suggesting half a billion animals infected each year.
We reviewed cases of Q fever reported to OIE between 2006 and 2010, retrieving 742 reports from 54 African countries. Only one report had numbers of animals affected, no report had population at risk. The reports were:

- Disease outbreak report: 1 report, 8 animals affected, 1 death
- Confirmed infection without clinical signs: 10 reports
- Disease present but without quantitative data: 16 reports
- Diseases suspected but not confirmed: 10 reports
- Disease absent: 226 reports
- No information available: 479 reports

Given that surveys carried out in the field suggests millions of cases occur in livestock in Africa each year, and that all surveys conducted in Africa found evidence of sero-positive animals indicating infection is present. It is obvious that official reporting seriously under-estimates the occurrence of this important notifiable zoonosis.

c) Comparing probable livestock mortality with notifiable disease reports

The World Bank and OIE produced a very useful atlas summarising animal disease reports between 2006 and 2009 (World Bank, 2011). This also allows us to assess under-reporting for developing countries. We do this by estimating number of livestock in Africa from FAOSTAT, annual mortality from systematic reviews, proportion of mortality likely to be due to notifiable diseases from expert opinion, and we compare these with official reports to OIE.

Number of livestock in Africa

FAO estimate that globally there are 24 billion livestock in 2010 (FAOSTAT, 2012), corresponding to 2.4 billion livestock standard units (using the OIE definition given in the aforementioned Atlas) (World Bank, 2011). Sub-Saharan Africa has two billion livestock corresponding to 253 million standard livestock units.

Number lost each year

Numerous studies on African livestock indicate annual mortality is high. Otte and Chilona (2002) reviewed production parameters of ruminants in traditional and non-traditional production systems reported in published and grey literature between 1973 and 2000 (table 1.3). Depending on species and age category, mortality ranged from 6-28% with three quarters of the species-age categories having a mortality of 10% or more.

<table>
<thead>
<tr>
<th></th>
<th>Young animals</th>
<th>Growing female</th>
<th>Growing male</th>
<th>Adult female</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cattle</td>
<td>22%</td>
<td>7%</td>
<td>9%</td>
<td>6%</td>
</tr>
<tr>
<td>Sheep</td>
<td>27</td>
<td>10</td>
<td>11</td>
<td>11</td>
</tr>
<tr>
<td>Goats</td>
<td>28</td>
<td>13</td>
<td>14</td>
<td>12</td>
</tr>
</tbody>
</table>

From Otte & Chilona, 2002

Otte and Chilona did not include poultry in the review but production parameters and characteristics of family poultry production have been compiled and published for eleven African countries (IAEA 2002). These give a range of annual mortalities from around 30% to 80% depending on the age category and country. Rege and Gibson (2009) estimate mortality in backyard poultry in Africa at 70% per year. There is little comprehensive information on mortality among smallholder pigs, but mortality is often high among pre-weaned piglets in smallholder systems (around one fifth) and very high losses occur during outbreaks of African swine fever and other epidemics (Wabacha et al., 2004).
Proportion of losses due to notifiable disease
Some of the annual livestock losses are due to non-infectious causes (mainly accidents, poisoning, predation and malnutrition). Other losses will be due to non-notifiable diseases (such as endoparasites) but farmers and experts agree that the 87 notifiable diseases are among the most important causes of mortality for livestock in Africa (not surprising when notifiable diseases include such high impact diseases as Newcastle disease, trypanosomosis, classical swine fever, East Coast fever, contagious bovine pleuropneumonia, and peste des petits ruminants). The authors of the report consider at least 50% of mortality is attributable to notifiable diseases

Combining these assumptions indicates a major discrepancy between the probable losses from notifiable disease (around 10 million) and the losses reported to the OIE (around 100,000) that can only be explained by under-reporting of several orders of magnitude. The following example makes this clear:

- Livestock in Africa = 253, 00, 000 standard livestock units (FAOSTAT, 2011)
- Livestock death, slaughter, or destruction reported to OIE = 82,319 units (World Bank, 2011)
- Livestock annual estimated losses = 25,300,000 TLUs (literature: 10% as a conservative estimate\(^3\))
- Estimated losses due to notifiable diseases = 12, 800,000 TLUs (expert opinion: 50% of losses)
- Losses (notifiable) reported to OIE = 0.2% of total units (less than one fifth of one percent)
- Losses (notifiable) probably not reported to OIE = 99.8% of losses
- Losses (notifiable & non notifiable) not officially reported = 99.9%

\(^3\) Mortality is lowest in cattle which contribute the most to tropical livestock units so we chose a low estimate of mortality
Chapter 2: Mapping of poverty, livestock, zoonoses and vulnerability to climate change

Summary
Information of reasonable quality is available on number of people in poverty by country and the number of livestock by country and by farming system. Literature provides estimates on the number and proportion of poor people keeping livestock. Emerging zoonotic disease events have been mapped, but because of the nature of the data, maps may not be informative about the impacts on poor people. The World Health Organisation information on the Global Burden of Disease provides information by country on the human health impact of around 11 important zoonoses and also on protein-energy malnutrition which is indirectly linked to livestock product availability and hence zoonoses. The World Animal Health Organisation collates information on 33 zoonoses but data from developing countries is prone to under-reporting.

2.1 Poverty
Poverty can be defined as a pronounced deprivation in wellbeing. No single indicator exists to measure all dimensions of poverty simultaneously, however, internationally comparable metrics, such as the US 1$ a day ($1.25), are useful for spatial and temporal comparisons. Estimates of poverty are probably reasonably accurate. The proportion of people living in poverty (<$1.25 per day\(^4\)) dropped by half between 1990 and 2010, but 1.3 billion people still live on less than $1.25 a day and around 2.5 billion on less than two dollars a day (World Bank, 2012).

In the past 3 decades, dramatic drops in poverty are mainly due to development in China: in Africa and South Asia numbers of people in poverty or stable or increasing. In terms of numbers, more than 75% of the people living in poverty live in 9 countries and 80% of poor people in 12 countries. In terms of intensity of poverty, 17 countries have more than 50% of the population living on less than $1.25 per day. Whereas in 1990, nine tenths of the poor lived in poor countries, presently three quarters live in middle-income countries (mainly India, China and Brazil).

2.2 Livestock
How many livestock are kept and where are they?
In 2012, the human population reached 7 billion and the production animal population around 24 billion (FAOSTAT, 2012). Global livestock systems have been recently re-mapped (Robinson et al., 2011). Poultry and pigs increasingly dominate in terms of number of animals kept (although in terms of tropical livestock units, ruminants are more important): 85% of all domestic animals alive are now pigs or poultry. As disease transmission is dependent on numbers and contact rates, and monogastrics are kept in higher numbers and more intensive systems, monogastrics may become more important in disease emergence.

Livestock density maps
Livestock density reflects the number of livestock and the level of intensification. Human population density is a major determinant of livestock density. High density is also an important factor in the transmission of disease through increasing the probability and number of contacts. However, density may also be associated with better biosecurity and control systems, which reduce risk. High livestock density, especially of monogastrics, often reflects intensification and tends to be inversely correlated with poverty. In our study, livestock density seems more correlated with zoonotic disease event emergence than burden of zoonotic disease. Figures 1.1 to 1.3 show global cattle density from the FAO gridded livestock maps (FAO, 2007).

\(^4\) 2005 international prices
Figure 2.1 Global poultry density (Robinson et al., 2011)

Figure 2.2 Global pig density (Robinson et al., 2011)
2.3 Poverty and livestock

How many poor people depend on livestock? Where are they?

Livestock-keeping had been variously regarded as a symptom of being poor, an important pathway out of poverty, and a transitional stage as burgeoning developing world populations shift from agriculture to urban livelihoods (Perry et al., 2010). Recent estimates suggest nearly 1 billion people living on less than two dollars a day are dependent to some extent on livestock (Staal et al., 2009). Over 600 million are found in South Asia, mostly in India. Sub-Saharan Africa has over 300 million poor livestock keepers, concentrated in East and West Africa, with fewer in southern and central Africa. A breakdown by region is provided in chapter 4.

What proportion of the poor depends on livestock?

Earlier estimates were around 70% of the rural poor depended on livestock (LID, 1999). Others have estimated that 40-50% of those living in poverty ($1.25 threshold) are at least partially dependent on livestock (Thomas & Rangnekar, 2004; IFAD, 2004). A more recent 12-country study supports this, finding that on average, around 68% of rural households in the bottom 40% as regards expenditure kept some farm animal compared to 65-58% of those in the top 40%; in urban areas 22-26% of the poor kept livestock, and 8-12% of the well-off (Pica-Ciamarra et al., 2011).

To what extent do poor people depend on livestock?

Staal et al. (2009) analysed 92 case studies from the developing world and found that livestock contributions made up on average 38% of household incomes (33% of the income in mixed crop-livestock systems, and 55% of total income in pastoral systems). The 12-country study of Pica-Ciamarra et al. found livestock contributed on average 12% to household income, with no statistical differences between contribution in rich and poor countries. (This study over-represented emerging countries, which may explain the lower contribution of livestock compared to the study of Staal et al., 2009). There is strong evidence that poor people depend on livestock, but more research is needed on the extent and nature of this dependence.
Livestock provide many benefits besides income. These include traction, manure, food, social status as well as economic services such as insurance and guarantees. Several studies show that manure, while seldom marketed is highly valued in smallholder systems (ranking higher than milk in West Africa (Grace et al., 2009). A study in Kenya, found that non-marketed values comprised approximately 20% of the animals total perceived value (Ouma et al., 2003).

**How much do livestock products contribute to nutrition in developing countries?**
Across a range of developing countries, livestock products contribute 6-36% of protein and 2-12% of total calories (Nzuma & Randolph, 2008). In South Asia and East Africa dairy products account for most livestock product consumption; in the rest of Africa, dairy, poultry, beef and shoats are balanced, while in South East Asia poultry and pork predominate. Countries with low livestock consumption (e.g. Bangladesh) may offset this with high fish consumption.

**What livestock do the poor keep?**
Poorer households are more likely to keep small ruminants and richer to keep large ruminants. Poultry keeping tends to be evenly distributed across wealth groups. However, species ownership is system and country specific.

**Which livestock systems contribute most value in poor countries?**
Around half the value of livestock production in sub-Saharan Africa is derived from cattle (55%), followed by poultry (25%) and small ruminants (20%). In South Asia these proportions were 61%, 21% and 18%, respectively (FAOSTAT, 2012). In both regions the arid/semi-arid zone contributed most to value of production. However, the trends are towards more value from poultry and pigs and more production from intensified systems. A recent comprehensive rural poverty mapping is the CGIAR Geographic Domain Analysis (2009). The most recent public domain maps on global poor livestock-keepers are those produced by ILRI in 2002 (Thornton et al., 2002) and subsequently updated.

**2.4 Mapping poverty and livestock systems**
Understanding the spatial distribution of livestock keepers can guide the allocation of resources as a first step in reaching the poor; identify areas of opportunity for livestock as a catalyst to growth; and, target hotspots of potential livestock-associated disease and environmental degradation. However, our knowledge of the location, characteristics and trends of change among poor livestock keeping populations is very patchy, both spatially and temporally. Here we outline a rapid broad-brush global assessment of spatial distribution of poor livestock keepers, and describe parallel activities in high-resolution poverty mapping for countries in East Africa using sophisticated econometric techniques pioneered at the World Bank.

In 2001 the UK Government’s Department for International Development (DFID) commissioned a study to produce sets of maps locating the significant populations of poor livestock keepers in the world, and to assess in very broad terms how these populations are likely to change over the next three to five decades. These data are reported by Thornton et al. (2002). The map presented in this document is updated using more current poverty estimates from World Bank’s 2011 WDI with the majority of the information being from 2005-2010 yet with a few countries with estimates still from the 1990s. Several countries within each region also have no estimates and have to be extrapolated by region.

The Thornton et al. (2002) study made use of existing data and spatial data layers, together with information from the literature and expert opinion. The central element of the analysis is a global livestock classification based on that of Seré and Steinfeld (1996), who present a typology based on
mixed crop-livestock systems, livestock-only rangeland-based systems, and landless production systems. We defined the classification primarily in terms of landuse/cover and climate-based length of growing period (LGP), supplemented by existing global coverages of human population, irrigated lands, and urban areas. Human population scenarios to 2050 were developed for Africa, Latin America and Asia.

The study also provided a breakdown of poverty information by country and livestock production system that was available for most of the countries only at the national level including: World Bank rural and national rates and two internationally comparable poverty lines: less than 1$/day and 2$/day. But this information did not include any information on how many of these poor were livestock keepers. So one additional layer was created by assigning differential poverty rates by broad livestock systems (mixed, pastoral and other) within each country providing at least some further sub-national distribution of the poor with livestock.

2.5 Maps of poor livestock keepers
The updated map of density of “poor livestock keepers” 2010 based on the methodology of Thornton et al. (2002) is shown in Figure 2.4. There are many assumptions and extrapolations involved in map development; however, despite caveats various conclusions can be drawn. In terms of numbers of livestock keepers, the critical regions remain South Asia and sub-Saharan Africa. The mixed farming systems (crop and livestock) contain large numbers of poor (over 1 billion), and numbers of poor people dependent to some extent on livestock in these systems are considerable. Mixed rainfed systems have more poor livestock keepers than mixed irrigated systems. Rangeland systems have least absolute numbers of the poor but the poor in this system have highest dependency on livestock. Almost half of the poor in rangeland systems are located in sub-Saharan Africa.

Figure 2.4 Density of poor livestock keepers in developing countries based on national data (updated March 2012)
Change in number of poor livestock keepers

Because of different methods in developing the maps, the map of 2000 is not directly comparable with updates. Some changes are evident since the map of 2002. There has been a marked decrease in the density of poor livestock keepers in South America and South East Asia. There has been some improvement, but to a lesser degree in parts of francophone West Africa and South Asia. However, these improvements have been more than offset by increases in Africa, most of South Asia, the middle East and Central Asia. Overall, the number of poor livestock keepers is estimated to have by 56 million in the eight years from 2000 to 2008 (FAO, 2011).

Figure 2.5 Density of poor livestock keepers as mapped in 2002

The International Food Policy Research Institute IFPRI (Wood et al., 2009) have released their sub-national rural poverty rates that cover most of sub Saharan Africa, but not yet the rest of the developing world. We used these to prepare Poor Livestock Keeper (PLK) maps for the < $1.25/day and $2/day poverty lines (Figure 2.6 and 2.7).

The two methods (national and sub-national) have broadly comparable results. In both maps, Ethiopia, Nigeria, the Great Lakes region, parts of West Africa and Malawi have the highest density of poor livestock keepers. However, the sub-national data reveals differences within countries. Because endemic zoonoses and emerging zoonotic events are geo-located, maps derived from sub-national data are more useful in exploring associations between zoonoses and poor livestock keepers (Figure 2.6 and Figure 2.7).
Figure 2.6 Density of poor livestock keepers (<$1.25 a day) in sub-Saharan based on sub-national data (update May 2012)

Figure 2.7 Density of poor livestock keepers (<$2 a day) in sub-Saharan based on sub-national data (update May 2012)
2.6 Maps of livestock system change
Livestock systems are changing rapidly in response to various drivers. Figure 2.8 shows estimates of livestock systems in 2000 and 2030 (Kruska et al., 2003, Hererro et al., 2008).

Figure 2.8 Farming systems in 2000 and 2030
Herrero et al. 2009 modelled growth in different livestock systems using the IMPACT model (Fig 2.9).

Figure 2.9 Changes in monogastric populations 2000-2030 (Herrero et al 2009)
Figure 2.9 shows percentage changes in pig and poultry densities between 2000 and 2030 (Herrero et al 2009). These are used as a proxy for emerging livestock systems. These use estimates from the ‘baseline scenario’, that is the most probable development of the sectors.

Under the reference, or ‘business as usual’ scenario, considered the most plausible, the hotspots in terms of rapid growth are in descending order, poultry in South and East Asia > poultry in South America > bovines in South and East Asia > poultry in sub Saharan Africa = pigs in sub Saharan Africa

Does changing livestock systems change the risk of zoonotic disease emergence?
The maps in Figure 2.9 show some of the geographical areas where change is most rapid; countries which are in the top 20 for both high numbers of monogastrics and rapid change are found in South Asia and South East Asia: Myanmar, India, Pakistan, Bangladesh, Thailand and China.

Change in livestock systems extends beyond intensification of poultry and pigs. Rapid change and growth is often associated with erosion of the natural resource base. Globally, anthropogenic changes are driving climate change with implication for livestock keeping and zoonoses. These three change processes are discussed in this section: intensification, interface with wildlife and climate change.

Agricultural intensification and zoonoses
A systematic review of zoonoses at the livestock/wildlife interface recently commissioned by DFID examined evidence for links between livestock intensification and disease emergence. Evidence is as yet insufficient for definitive conclusions, and in some cases intensification is associated with less disease, but overall it seems that intensification is linked with disease emergence and spread.

There are several contributing factors. Increased livestock numbers are themselves a risk factor for increased disease transmission. Moreover, selection, breeding and management for increased productivity in livestock create host populations conducive to pathogen evolution and persistence (through lack of genetic diversity, high numbers and contact opportunities, stress-induced immunosuppression and other factors). This provides opportunity for “wild” microorganisms to invade and amplify or for livestock pathogens to evolve to new and more pathogenic forms. In addition, corollaries of intensification such as high livestock and pest densities, extensive transportation networks, sale of live animals for food and pets, landscape modification, poor waste management, and juxtaposition of agriculture or recreation with wildlife all contribute to “emergence” and shifting virulence of diseases (Grace et al., 2011). Furthermore, much of the current intensification is driven by rising demand in developing countries, where the systems for disease control and reporting are relatively weak.

The literature review conducted for this survey found that zoonotic food-borne pathogens were markedly higher in poultry and pigs than in small ruminants and cattle. This suggests that as monogastric systems expand, so may food-borne disease.

Agricultural intensification is likely to have different impacts on the key zoonoses depending on their epidemiology. Probable impacts are discussed in chapter 3 and summarised here. Of the priority zoonoses, 9 are likely to become more of a problem with intensification, 4 are likely to decrease and for the remaining there is no clear link. Whether disease increases or decreases, also depends on the type of intensification and other factors. Where intensification occurs in close association with wildlife, risks for disease spillover are higher. If intensification is accompanied by improvements in biosecurity and a disease control programme, then diseases such as bovine tuberculosis may decrease. Other diseases tend to increase in the early stages of intensification but may decrease after.
Table 2.1 Probable impact of intensification on priority zoonoses

<table>
<thead>
<tr>
<th>Zoonosis</th>
<th>Likely impacts of agricultural intensification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gastrointestinal (zoonotic)</td>
<td>Most gastro-intestinal zoonoses are food-borne and likely to increase with intensification and associated lengthening and branching of food supply chains. Many gastro-intestinal zoonoses cause little visible signs in animals reducing farmer incentives for control.</td>
</tr>
<tr>
<td>Leptospirosis</td>
<td>Leptospirosis is associated with smaller farms, and pasture-grazing especially where there is stagnant water. Intensification may reduce prevalence.</td>
</tr>
<tr>
<td>Cysticercosis</td>
<td>Associated with free-range, scavenging pigs. Intensification will reduce prevalence.</td>
</tr>
<tr>
<td>Tuberculosis (zoonotic)</td>
<td>Associated with larger farms and confined systems. Intensification likely to increase.</td>
</tr>
<tr>
<td>Rabies</td>
<td>No clear link. Most human transmission from dog bites or wildlife.</td>
</tr>
<tr>
<td>Leishmaniasis</td>
<td>No clear link. Transmitted by sandflies. Domestic dogs are the most important reservoir.</td>
</tr>
<tr>
<td>Brucellosis</td>
<td>Associated with larger farms and confined systems. Intensification will increase. However, artificial insemination, often associated with intensification, will decrease.</td>
</tr>
<tr>
<td>Echinococcosis</td>
<td>Associated with feeding offal to dogs. More common in extensive systems.</td>
</tr>
<tr>
<td>Toxoplasmosis</td>
<td>Some evidence this is more common in extensive systems. Associated with rodents.</td>
</tr>
<tr>
<td>Q fever</td>
<td>No clear link.</td>
</tr>
<tr>
<td>Trypanosomosis (zoonotic)</td>
<td>Intensification reduces risk by removing tsetse habitat and wildlife hosts.</td>
</tr>
<tr>
<td>Anthrax</td>
<td>No clear link.</td>
</tr>
<tr>
<td>Hepatitis E</td>
<td>Extent of transmission from pigs not clear.</td>
</tr>
<tr>
<td>Chagas</td>
<td>Most associated with extensive systems</td>
</tr>
<tr>
<td>Chickungunya</td>
<td>Associated with incursion into forest areas.</td>
</tr>
<tr>
<td>Clostridium difficile disease</td>
<td>No clear relation. Present in farm animals but role in transmission not clear</td>
</tr>
<tr>
<td>Dengue fever</td>
<td>Most transmission anthroponotic: livestock systems no clear role</td>
</tr>
<tr>
<td>Ebola</td>
<td>Intensification around bats is a risk</td>
</tr>
<tr>
<td>Hanta disease</td>
<td>Spread by rodents. Not farm associated</td>
</tr>
<tr>
<td>Avian influenza</td>
<td>Associated high poultry density – link with intensification not clear</td>
</tr>
<tr>
<td>Bov. Spongiform Encephalopathy</td>
<td>Associated intensive systems</td>
</tr>
<tr>
<td>Psittacosis</td>
<td>No clear link.</td>
</tr>
<tr>
<td>Japanese encephalitis</td>
<td>Associated with intensive rice systems</td>
</tr>
<tr>
<td>Buffalo pox</td>
<td>No clear link.</td>
</tr>
<tr>
<td>Rift Valley fever</td>
<td>May increase with intensification and irrigation</td>
</tr>
</tbody>
</table>

Zoonoses with a wildlife interface

Fifteen of the ‘top 25’ zoonoses have important wildlife (including synanthropic wildlife) reservoirs across many regions, including 9 of the ‘top 13’ zoonoses, namely: gastro-intestinal zoonoses, leptospirosis, rabies, leishmaniasis, toxoplasmosis, echinococcosis, Q fever, trypanosomosis and anthrax. For some of the other zoonoses, wildlife may play an important role in some epidemiological circumstances. For example, tuberculosis associated with conservation areas in Tanzania and South Africa, brucellosis associated with buffaloes, and hepatitis E and cysticercosis with wild pigs. Where a
wildlife interface exists, zoonoses control is much more complex (Grace et al., 2011). Other zoonoses in the ‘top 25’ but not ‘top 13’ with an important wildlife interface are: Chagas, Chickungunya, Ebola, Hanta disease, avian influenza, psittacosis and Rift Valley fever.

2.7 Climate change and zoonoses
There are several metrics for vulnerability to climate change (Cutter et al., 2009; Fussel et al., 2009). Tropical African countries and Asian coastal countries are usually among the countries considered most vulnerable to climate change. The CGIAR Research Program on Climate Change, Agriculture and Food Security (CCAFS) commissioned ILRI/ CCAFS to conduct a rapid assessment across the global tropics of the vulnerability of food security to climate change (Ericksen et al., 2011). The goal was to identify ‘hotspot’ locations where climate change impacts are projected to become increasingly severe by 2050 and food insecurity is currently a concern, using a range of indicators. The maps mainly focused on change with implications for crop growth but some of these changes also have implications for livestock production and disease.

Figure 2.10 and 2.11 predict areas where rainfall and flooding will increase. This is expected to increase the risk associated with vector-borne zoonoses including tick-borne and mosquito-borne diseases. It will also increase risk of bacterial pathogens associated with stagnant water and flooding (e.g. leptospirosis, anthrax, cryptosporidiosis).

Figure 2.10 Areas where rainfall per day increases by 10% or more between 2000 and 2050 (Ericksen et al., 2011)

![Figure 2.10](image1)

Figure 2.11 Flood frequency (Ericksen et al., 2011)

![Figure 2.11](image2)

The next map aggregates different thresholds that can stress production, including flips in: growing period, reliable crop-growing days, annual average temperature, annual average maximum temperature, maximum temperature exceeds 30 centigrade, changes in variability in rainfall, and increase in rainfall.
In terms of exposure to multiple climate threats, southern Africa has the largest area exposed (across Namibia, Angola, Zambia, Botswana, Mozambique and South Africa) with multiple threats, followed by northeastern Brazil, Mexico, Guyana, Nicaragua, and small areas in Tanzania, Ethiopia, the DRC, Uganda, India, and Pakistan, as well as the Middle East.

While studies are starting to emerge on the likely effect of climate change on human disease, and changes in spatial dynamics of some animal diseases (e.g. blue tongue) are believed to be influenced by climate change, there is still little strong evidence on impacts of climate change on zoonotic disease in dynamic systems with multiple drivers. For example, as countries get warmer and wetter biological mechanisms would suggest that many diseases increase. However, if countries simultaneously get richer or invest more wisely in health care the net impact may be disease decrease (Perry et al., 2011).

The effects of climate change on livestock and non-vector-borne disease have, with some exceptions, received little attention. The climate-livestock-poverty nexus was reviewed by Thornton et al. (2008) and this section is largely based on their findings. Climate change may affect livestock disease through several pathways:

- **Pathogens**: higher temperatures and greater humidity generally increase the rate of development of parasites and pathogens that spend part of their life cycle outside the host. Changes to wind can affect spread of pathogens. Flooding that follows extreme climate events provides suitable conditions for many water-borne pathogens. Drought and desiccation are inimical to most pathogens.

- **Vectors**: vector-borne diseases are especially sensitive to climate change. Changes in rainfall and temperature regimes may affect both the distribution and the abundance of disease vectors, as can changes in the frequency of extreme events (outbreaks of Rift Valley fever have been linked to ENSO, for example).

- **Hosts**: some will be exposed to new pathogens and vectors as their range increases and impacts can be severe. Climate stress (heat, inadequate food and water) can also lower immunity.

Of the 13 priority zoonoses, food-borne zoonoses, leptospirosis and trypanosomosis are likely to show high climate sensitivity. However, cysticercosis, tuberculosis, rabies, brucellosis and echinococcosis are unlikely to show high climate sensitivity. It is less clear how climate change will affect the epidemiology of other priority zoonoses, although some evidence suggests there may be important negative impacts.

- **Food-borne zoonoses**: A recent extensive literature review concluded that campylobacteriosis and salmonellosis were most likely to increase with air temperature; campylobacteriosis and non-cholera vibrio infections with water temperature; cryptosporidiosis followed by campylobacteriosis with increased frequency with precipitation; and cryptosporidiosis followed by non-cholera vibrio in association with precipitation events. Listeria sp. was not associated...
with temperature thresholds, extreme precipitation events, or temperature limits (ECDC, 2012).

- **Leptospirosis**: Leptospirosis is considered one of the more climate sensitive diseases. Flooding and heavy rainfall have been associated with numerous outbreaks of leptospirosis around the world. With global climate change, extreme weather events such as cyclones and floods are expected to occur with increasing frequency and greater intensity and may potentially result in an upsurge in the disease incidence as well as the magnitude of leptospirosis outbreaks (Lau et al., 2010).

- **Trypanosomosis**: While climate will modify (generally decrease, but not everywhere) habitat suitability for the tsetse fly, the demographic impacts on trypanosomosis risk through bush clearance are likely to outweigh those brought about by climate change (Thornton et al., 2006).

- **Q fever** is transmitted in aerosols and climate change could affect survivability. Toxoplasmosis has rodent hosts and rodent populations are sensitive to climate change. Climate change and other environmental changes have the potential to expand the geographic range of the vectors and leishmaniasis transmission in the future. Anthrax is often associated with a combination of heavy rain and warm temperatures following a drought that encourages spores to germinate. These extreme events will be more common with climate change.
Chapter 3: Literature review of zoonoses of importance

Summary
We undertook a systematic literature review of the 13 zoonoses identified as important. Eight of these are 'endemic classical zoonoses' that is, zoonoses that are typically present across a wide range of communities at most times (although perhaps showing annual and inter-annual variability). In the case of bacterial food-borne zoonoses, we identified five diseases, which ranked highest on a number of recent assessments of impact (salmonellosis, listeriosis, toxoplasmosis, campylobacteriosis and disease caused by diarrhoeagenic Escherichia coli). We also considered three epidemic or outbreak-associated zoonoses (rabies, leishmaniasis and anthrax) and hepatitis E, an emerging disease, which may have an important zoonotic transmission.

For each disease we present information on prevalence, epidemiology, geographical hotspots and some key research questions and we estimated an 'endemic disease burden' score for all the countries for which information existed.

3.1 Methodology for systematic literature review
We generated search terms that incorporated certain key words, identified and screened abstracts, reviewed full papers and synthesised required information. Diseases were initially considered on the basis of their appearance in the top 13 list of zoonoses generated as explained in the previous chapter. These were: Taenia solium cysticercosis, leptospirosis, anthrax, brucellosis, echinococcosis, hepatitis E, leishmaniasis, Q fever, rabies, toxoplasmosis, trypanosomosis, tuberculosis, and food borne infections (caused by Salmonella spp., Listeria monocytogenes, diarrhoeagenic Escherichia coli, and Campylobacter spp.).

PubMed (www.ncbi.nlm.nih.gov/pubmed/) and CABDIRECT (www.cabdirect.org/) were used in doing the searches but also Google (www.google.co.ke/) including the Google scholar. Unpublished materials including student theses (mainly from the University of Nairobi) were accessed by visits. Related articles appearing during the active searches in PubMed were also utilized in sourcing for extra details, as well as from relevant references cited in the main papers reviewed. When available, the cited original papers were retrieved, reviewed, and relevant information retrieved. This also applied to major review papers providing a summary of the needed information and references for the original papers available. We used the same data as summarized in the review paper if the original paper was not available.

Search terms were formulated, by disease, and by country or region; different combinations, were either relaxed or broadened to capture more articles or were restricted to refine or limit the number of resulting articles. Different Boolean operators were used (including AND, OR, parenthesis) for specific PubMed and CABDIRECT searches. Phrases guided by key words were used for the Google searches. We also applied wildcard symbols, mainly * to broaden the results in some of the searches.

The first step involved screening the abstracts by title, abstracts not relevant for the project objectives were left out. The searches were originally limited to the last 10 years, but we also considered old studies if the search results were initially few. We considered studies conducted in Africa, South Asia and South East Asia. Some studies from the Middle East were also included. Those abstracts that were considered relevant (based on the title) were extracted into a word document and subsequently reviewed by a second person. Full papers linked to the relevant abstracts were extracted and reviewed. Prevalence information, if available, was extracted from abstract in cases where the full paper could not be accessed. Sources providing no information on the number of samples / subjects analysed were not considered- as the basis for the calculation of the prevalence estimates could not
be established. Also excluded were papers with missing geographical locations for the specific studies.

An excel® database was developed to capture information extracted during the review process (including the different search terms used). Variables extracted included: country where the study was done / or where the results apply, geo-spatial location (the specific location or coordinates if given), number of herds studied, number of samples analysed, the specific diagnostic test(s) done, subjects (livestock species, food, humans), individual prevalence, herd prevalence, year data were collected and a description of the study population. Where multiple surveys were reported in one study, each survey was listed separately (e.g. if prevalence was estimated in cattle and sheep these were considered as two different surveys each with an associated sample size, species and prevalence). We distinguish between “community studies” which are conducted in the community and can be considered representative of it, and “high risk studies” which were conducted in high risk populations (sick people in hospitals, malnourished children, cattle which failed ante-mortem inspection, samples taken during an outbreak etc.). Maps were generated using data from community studies only.

We defined geographical hotspots as those which had a high prevalence confirmed in multiple surveys. The number of studies needed to consider estimates reliable varied from pathogen to pathogen depending on the number of studies available and is given in the results section of the different diseases.

In order to estimate the ‘top twenty countries’ for endemic disease burden we collated the geographical hotspots. We standardized scores for each disease so the country that had highest prevalence had a score of ten, and so on. We then summed scores for each disease by country.

To develop the maps of endemic zoonoses we collected on-locational or descriptive data on zoonoses using systematic literature review and details were documented in a MS Excel spreadsheet. The study area covered entire Africa and Asia (South central Asia and South East Asia). The constitution of the spatially referenced database was performed by introducing locational or spatial data in the form of coordinates into the spreadsheet. These coordinates were approximated from the ‘Geospatial location’ section of the database and were sourced from existing GIS databases and occasionally from websites such as Google maps and www.longitude-latitude-maps.com.

The spreadsheet was then imported into the GIS software package ArcGIS v.10 (ESRI, Redlands). This software package allows for the seamless linkage of MS Excel spreadsheets to the GIS by using the coordinates columns, and these are imported as event data. The zoonoses locations were then mapped and the column ‘prevalence’ from the descriptive data used to map the magnitude of the prevalence as a percentage. For visualization purposes, mapping was done with a base map of agricultural farming systems on the background.

3.2 Results for systematic literature review- endemic zoonoses

We conducted a systematic review of brucellosis, tuberculosis, leptospirosis, trypanosomosis, cysticercosis, and Q fever and some bacterial food-borne diseases.

- **Brucellosis – the deceptive disease** – causes fever and occasionally chronic disease in people; mainly abortion and infertility in cattle, shoats and pigs
- **Tuberculosis – white plague** – a major cause chronic illness in people, causes wasting and illness mainly in cattle
- **Leptospirosis – swamp fever** – causes fever and occasionally jaundice in people and fever and infertility in cattle and pigs; wildlife important reservoirs
- **Q fever – the most contagious disease** – causes fever and occasionally death in people, carried by cattle, shoats, pets and wildlife, causes abortion in shoats
• Cysticercosis – *pork worm* – most common cause of adult-onset epilepsy in poor, pig-keeping communities, leads to carcass condemnation in pigs
• Trypanosomosis – *sleeping sickness* – cause of acute and chronic illness in people, historically caused severe epidemics, the most important disease of cattle in sub Saharan Africa; wildlife important reservoirs
• Bacterial food-borne disease – *the forgotten zoonoses* – major cause of gastrointestinal disease in people; some but not all cause illness in animals. Several have wildlife interface
• Echinococcosis – *cystic disease* - a major cause of illness in people and loss in sheep and goats from condemnation of carcasses

We obtained information from 1098 surveys covering around six million animals, ten million people and six thousand food or environment samples. Endemic zoonoses impose an important burden in all regions, although the distribution varies according to disease. Trypanosomosis is found only in sub-Saharan Africa, cysticercosis is rare (though not absent) from cultures where pigs are not kept, and brucellosis is associated with high populations of ruminants. Zoonotic food-borne diseases, the most important zoonoses, are at much higher prevalence in poultry and pigs than ruminants. Table 3.1 summarises the prevalence for important zoonoses by region and for all developing countries. It gives the overall prevalence (humans, livestock, wildlife, other animals) and the prevalence for humans and livestock separately.

Table 3.1 Prevalence (%) of important zoonoses by region

<table>
<thead>
<tr>
<th></th>
<th>North Africa, Near East</th>
<th>East Africa</th>
<th>Southern Africa</th>
<th>West Africa</th>
<th>South Asia</th>
<th>SE Asia</th>
<th>All developing</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brucellosis*</td>
<td>13%</td>
<td>8%</td>
<td>14%</td>
<td>16%</td>
<td>16%</td>
<td>2%</td>
<td>12%</td>
</tr>
<tr>
<td>Tuberculosis^</td>
<td>9</td>
<td>8</td>
<td>5</td>
<td>7</td>
<td>17</td>
<td>0.2</td>
<td>7</td>
</tr>
<tr>
<td>Leptospirosis*</td>
<td>30</td>
<td>24</td>
<td>17</td>
<td>28</td>
<td>27</td>
<td>24</td>
<td>24</td>
</tr>
<tr>
<td>Q fever*</td>
<td>19</td>
<td>11</td>
<td>4</td>
<td>13</td>
<td>19</td>
<td>1</td>
<td>19</td>
</tr>
<tr>
<td>Cysticercosis^</td>
<td>Few pigs</td>
<td>12</td>
<td>23</td>
<td>16</td>
<td>14</td>
<td>12</td>
<td>14</td>
</tr>
<tr>
<td>Trypanosomosis^</td>
<td>Not present</td>
<td>9</td>
<td>12</td>
<td>10</td>
<td>N/A</td>
<td>N/A</td>
<td>10</td>
</tr>
<tr>
<td>Food-borne disease</td>
<td>25</td>
<td>27</td>
<td>21</td>
<td>30</td>
<td>18</td>
<td>25</td>
<td>25</td>
</tr>
<tr>
<td>Overall</td>
<td>15</td>
<td>10</td>
<td>16</td>
<td>15</td>
<td>25</td>
<td>22</td>
<td>16</td>
</tr>
<tr>
<td>Human</td>
<td>15</td>
<td>15</td>
<td>11</td>
<td>10</td>
<td>19</td>
<td>11</td>
<td>16</td>
</tr>
<tr>
<td>Livestock</td>
<td>15</td>
<td>10</td>
<td>16</td>
<td>16</td>
<td>17</td>
<td>18</td>
<td>15</td>
</tr>
</tbody>
</table>

*based mainly on seroprevalence, indicates current or recent infections (last 1-2 years)
^ based on parasitological tests, indicates current infections

**Brucellosis – the deceptive disease**

**Pathogen**

The most important species of *Brucella* are zoonotic: *B. abortus*, responsible for bovine brucellosis; *B. melitensis*, the main etiologic agent of ovine and caprine brucellosis and an increasing cause of cattle brucellosis; and *B. suis*, causing pig brucellosis.

**Studies**

259 studies were assessed covering 476,067 animals and 31,842 people and 537 food samples. 248 studies were from communities and 11 from high-risk groups (mainly people in hospitals).
Tests
Commonly used tests for brucellosis detect antibodies produced in response to infection. A combination of tests may be used to improve accuracy or ability to detect. The antibodies tested for generally persist for several months (IgG) or several years (IgM). Positive tests (to both antibodies) indicate the animal is currently sick, is chronically infected or has been infected in the last year or so. Hence positive tests are roughly equivalent to annual cases.

Prevalence
In community surveys, the prevalence was 13% in shoats, 13% in bovines, 7% in camels, and 5% in other species (chickens, pigs, dogs). Among livestock-keepers/abattoir workers prevalence was 11%, and among suspect hospital patients, 7%. A large study in India found that 2% of patients in the general hospital population tested positive for brucellosis.

Epidemiology
The main risks for people are occupational (contact with livestock) and consumption of dairy products. In some areas, brucellosis may be maintained in reservoir wild animal hosts (African buffaloes and North American bison) in other cases diseases spills-over to wildlife and if eliminated in cattle brucellosis will die out in wildlife. Brucellosis is more problematic in intensive systems than extensive and pasture-based systems.

Hot spots
Brucellosis is mainly a problem where ruminants are important (Africa and South Asia). Shoat-keeping communities are most at risk from *B. melitensis* considered the most pathogenic form. Countries with multiple surveys (>=4) and high prevalence (>15%) include in descending order: Togo, Mali, Ivory Coast, Zambia, Niger, India, Sudan, Cameroon and Burundi (human and animal combined)

Impact
Sero-positive animals have higher rates of abortion, stillbirth, infertility, calf mortality and lameness. This is associated with lower milk yields (around 25% milk loss in aborted cows). Usually, infected females will abort only once, although they may remain infected their entire life. The losses are estimated at 6-10% of the annual value produced per animal (Mangen et al., 2002).

Agricultural losses have been estimated at $427 million per year for sub-Saharan Africa and $600 million for Latin America (Mangen et al. 2002; Seleem et al., 2009)

Human brucellosis usually presents as an acute febrile illness, often mistaken for malaria or typhoid. Chronic complications are not uncommon.

Key research questions
- Improving diagnosis in people, given widespread under-diagnosis and confusion with malaria
- Public-private partnerships for control – promising studies suggest that by combining human health investment and livestock sector investment, brucellosis can be controlled in a cost-effective way
- Role of wildlife in maintaining infection – wildlife have an important role in some circumstances. The extent of this is not known, nor are effective strategies for managing disease in wildlife populations
- Reducing risky behaviours around husbandry and consumption- much of the risk from brucellosis can be reduced by simple precautions applied to handling cattle and food.
- Developing a vaccine for *B. suis*
- Effective vaccination which can be distinguished from infection to aid in control
Figure 3.1 Brucellosis prevalence in community surveys
**Tuberculosis – the white death**

**Pathogen**
Worldwide and historically, most human tuberculosis (TB) is caused by *Mycobacterium tuberculosis*. *M. bovis* is responsible for cattle tuberculosis. It affects a wide range of animals and is responsible for zoonotic TB in humans. In west Africa, *M. africanum* causes up to half of human tuberculosis – it has characteristics intermediate between *M. tuberculosis* and *M. bovis* the agent responsible for bovine tuberculosis. Atypical mycobacteria are found in the soil and environment and can infect both people and animals.

**Studies**
110 surveys were assessed covering 336,152 livestock and 5,829 humans. 89 studies were community-based and 12 in high-risk populations.

**Tests**
The standard method for detection of bovine tuberculosis is the tuberculin test, which involves the intradermal injection of bovine tuberculin protein derivatives (PPD) and the subsequent detection of swelling at the site of injection. This may be performed using bovine tuberculin alone or as a comparative test using avian and bovine tuberculin. More recently, a gamma interferon test has been developed. Meat inspection is also used to detect tubercular lesions in cattle, but in developing countries is not very accurate. A large study in Ethiopia found routine inspection detected 3.5% carcasses with lesions whereas detailed meat inspection procedures identified 10.2% carcasses, a more than three fold difference (Biffa et al).

Positive tests are roughly equivalent to prevalence (or animals currently sick with TB).
Prevalence

Overall, 7.4% of livestock were positive. Overall prevalence was as follows: Bovines: 8%, camels 11%, shoats 2%, pigs 15%, wildlife 5%.

There were an estimated 12 million cases of human TB (prevalence) in 2010 (WHO, 2011). Twenty-two high burden countries account for approximately 80% of all new TB cases. An extensive literature exists on the prevalence of human TB but there is little information on what proportion is zoonotic, and our review concentrated on this. Table 3.2 summarises more recent studies from developing countries: on average 10.5% of human TB cases were associated with M. bovis. Our study suggests a higher overall prevalence than previous best estimates (3.1%)5 but a strongly bimodal distribution: zoonotic TB is either very important or minor in a given context.

Table 3.2 Studies since 1999 on proportion of zoonotic TB

<table>
<thead>
<tr>
<th>Country</th>
<th>Study</th>
<th>MTBC</th>
<th>M. bovis</th>
<th>% M. bovis</th>
<th>Reference abbreviation</th>
<th>Year</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cameroon</td>
<td>15 district hospitals in Ouest</td>
<td>455</td>
<td>1</td>
<td>0.20%</td>
<td>Niobe-Eyangoh</td>
<td>2003</td>
</tr>
<tr>
<td>Djibouti</td>
<td>Unknown</td>
<td>85</td>
<td>1</td>
<td>1.20%</td>
<td>Koeck</td>
<td>2002</td>
</tr>
<tr>
<td>Egypt</td>
<td>Fever hospitals in cities</td>
<td>67</td>
<td>1</td>
<td>1.50%</td>
<td>Cooksey</td>
<td>2002</td>
</tr>
<tr>
<td>Ghana</td>
<td>Korle-Bu teaching hospital</td>
<td>64</td>
<td>2</td>
<td>3.10%</td>
<td>Addo</td>
<td>2007</td>
</tr>
<tr>
<td>Guinea-b</td>
<td>Unknown</td>
<td>229</td>
<td>4</td>
<td>1.70%</td>
<td>Käilénius</td>
<td>1999</td>
</tr>
<tr>
<td>Madagascar</td>
<td>Antananarivo, Ansirabe, Fianarantsoa, Mahajanga</td>
<td>400</td>
<td>5</td>
<td>1.30%</td>
<td>Rosolofo-Razanamparany</td>
<td>1999</td>
</tr>
<tr>
<td>Nigeria</td>
<td>2 hospitals Ibadan</td>
<td>60</td>
<td>3</td>
<td>5.00%</td>
<td>Cadmus</td>
<td>2006</td>
</tr>
<tr>
<td>Nigeria</td>
<td>Lagos</td>
<td>91</td>
<td>4</td>
<td>4.40%</td>
<td>Idigbe</td>
<td>1986</td>
</tr>
<tr>
<td>Nigeria</td>
<td>3 hospitals Jos</td>
<td>50</td>
<td>10</td>
<td>20.00%</td>
<td>Mawak</td>
<td>2006</td>
</tr>
<tr>
<td>Tanzania</td>
<td>Arusha</td>
<td>34</td>
<td>7</td>
<td>20.60%</td>
<td>Cleaveland</td>
<td>2007</td>
</tr>
<tr>
<td>Tanzania</td>
<td>Pastoralist North &amp; South</td>
<td>38</td>
<td>7</td>
<td>18.40%</td>
<td>Kazwala</td>
<td>2001</td>
</tr>
<tr>
<td>Tanzania</td>
<td>Arusha</td>
<td>34</td>
<td>7</td>
<td>20.60%</td>
<td>Mfinanga</td>
<td>2004</td>
</tr>
<tr>
<td>Uganda</td>
<td>Kampala</td>
<td>344</td>
<td>1</td>
<td>0.30%</td>
<td>Asiiimwe</td>
<td>2008</td>
</tr>
<tr>
<td>Uganda</td>
<td>Kampala</td>
<td>234</td>
<td>1</td>
<td>0.40%</td>
<td>Niemann</td>
<td>2002</td>
</tr>
<tr>
<td>Uganda</td>
<td>Karamoja</td>
<td>10</td>
<td>3</td>
<td>30.00%</td>
<td>Oloya</td>
<td>2007</td>
</tr>
<tr>
<td>Uganda</td>
<td>Mbarara</td>
<td>69</td>
<td>0</td>
<td>0.00%</td>
<td>Byarugaba</td>
<td>2009</td>
</tr>
<tr>
<td>Bangladesh</td>
<td>Clinical</td>
<td>350</td>
<td>0</td>
<td>0.00%</td>
<td>Nakajima</td>
<td>2010</td>
</tr>
<tr>
<td>India</td>
<td>TB meningitis</td>
<td>34</td>
<td>24</td>
<td>64.90%</td>
<td>Shan</td>
<td>2006</td>
</tr>
<tr>
<td>India</td>
<td>EPTB hospital adjusted for prev EPTB in population</td>
<td>155</td>
<td>22</td>
<td>64.90%</td>
<td>Jain</td>
<td>2011</td>
</tr>
<tr>
<td>India</td>
<td>EPTB hospital adjusted for prev EPTB in population</td>
<td>115</td>
<td>53</td>
<td>64.90%</td>
<td>Prasad</td>
<td>2005</td>
</tr>
<tr>
<td>Pakistan</td>
<td>Hospital, Lahore</td>
<td>42</td>
<td>5</td>
<td>11.90%</td>
<td>Nawaz</td>
<td>2012</td>
</tr>
</tbody>
</table>

MTBC= Mycobacterium tuberculosis complex  EPTB=Extra-pulmonary tuberculosis

Prev. = prevalence

Cattle can also be affected by M. tuberculosis and can in turn shed this in secretions and excretions. Cattle positive for M. tuberculosis to be a problem in South Asia: in one of the studies we reviewed, taking place in India, 7.1% of pharyngeal swabs from cattle were positive for M. tuberculosis. This

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5 Historically, M bovis was responsible for 5-30% of TB cases in the US, UK and Netherlands (Olmstead and Rhode, 2011; Cousins, 2001). Currently, zoonotic TB can be high in specific circumstances in California, during 1980-1997, 34% of culture-confirmed TB cases in were caused by M. bovis. However, many experts consider the role of zoonotic TB to be minor or negligible. The most authoritative review estimated that worldwide 3.1% of human TB cases are caused by M. bovis (Cosivi et al., 1998).
suggests that the burden of ‘zoonotic’ TB may be under-estimated as ‘human TB’ may be acquired from cattle.

*M. bovis* infects a range of African wildlife (high levels of infection have been found in the Kruger Park, Zambia and Serengeti). This is a potential source of infection to livestock and people, as well as a threat to wildlife. Where *M. bovis* is established in wildlife hosts (e.g. badgers in the UK or possums in New Zealand) eradication is very difficult.

**Epidemiology**
Commonly found risk factors are close contact of animals (intensive and peri-urban systems), increasing herd size and presence of wildlife reservoirs. Important risk factors for zoonotic TB people are close contact with animals and consumption of raw milk. Prevalence appears to be higher on intensive farms.

**Impact**
Muller summarizes a range of early reviews from Europe and North America before control was widespread. Infected cattle lost 10% of milk production and 4% of meat production and infected cows had one fewer calf. Unfortunately, good economic data is missing from developing countries but similar losses could be anticipated. TB lesions are also an important reason for carcass condemnation but it seems likely that routine meat inspection misses most cases (Biffa et al., 2010).

Zoonotic TB has a similar course in people as non-zoonotic. Overall, one third of the world’s population is currently infected with TB. Of those infected with TB that do not receive treatment, about 5-10% will develop TB disease some time in their lives. Zoonotic TB is more likely to present as extra-pulmonary, and prevalence of extra-pulmonary TB is a crude proxy for zoonotic TB.

Agricultural losses worldwide have been estimated at $3 billion (Garnier et al., 2003).

**Geographical hotspots**
Zoonotic TB is mainly a problem where cattle are important (Africa and South Asia). Dairying communities are most at risk. Countries with multiple surveys (>=2) and high prevalence (>5%) include in descending order: Bangladesh, Burkina Faso, Pakistan, Ghana, Kenya, Mali, Cameroon, Chad, India, and Ethiopia.

Human TB (zoonotic 3-10%) is mainly localized in high burden countries: India, China, Indonesia, Nigeria, Bangladesh, and Pakistan.

**Key research questions:**
- *M. tuberculosis* (human TB) appears to be common in livestock in some areas (especially India). What is the significance for transmission?
- A zoonotic reservoir has been suspected for *M. africanum* – so far little evidence but not fully investigated
- TB is one of the most important and common human diseases. There is much uncertainty on the proportion of this attributable to *M. bovis*, and our review suggested that the proportion is higher than in previous estimates and most of a problem in South Asia
- Impact of *M. bovis* on cattle in Africa and South Asia. Much of the information on impact is derived from earlier studies in Europe or North America and may not be applicable to developing countries
- Wildlife-livestock interface in hotspots (Tanzania, Ethiopia, Zambia and South Africa)
- Understanding relation between intensification and disease: Cattle TB appears to increase with intensification and urban farming
- Controlling cattle *M. bovis* and *M. tuberculosis* in cultures that do not permit culling of cattle
Figure 3.2 Tuberculosis prevalence in community studies

Legend
Prevalence %
- Below 5
- 6 - 15
- 16 - 30
- Above 30

Legend
Camels
- Below 1
- About 1
- Above 1

Small ruminants
- Below 1
- About 1
- Above 1

TB-pigs
- Below 6
- More than 6
Pathogen
Leptospirosis is an infectious disease caused by pathogenic organisms belonging to the genus *Leptospira*. There are many serovars (>250) but typically only around 10-20 are found in a given region. Serovars can be grouped into 25 serogroups.

Tests
Microscopic agglutination test is the gold standard and was used by most of our studies. Paired serum samples are used to identify current or recent infection. Antibodies may persist for several years. Most of the surveys in our review were community based. For these, a positive result indicates current infection, chronic infection or infection in the last few years.

Prevalence
109 surveys were assessed covering 52,534 animals and 83,596 people. In community surveys the prevalence was 34% in swine, 29% in bovines, 14% in small ruminants, 16% in wildlife and 24% in people.

Among patients presenting with fever of unknown origins around 20% (7-57%) had leptospirosis. Among patients with suspected leptospirosis 60-90% had positive diagnoses.

Epidemiology
Infected animals often become carriers. Wildlife are affected and can be important reservoir hosts. Risk factors for humans include presence of rodents, farm animals and floods. Risk factors for animals include smaller farms and extensive (pasture-grazing systems).

Geographical hotspots
Leptospirosis is mainly a problem in tropical countries where stagnant water can be found and where cattle, pigs or rodents are frequent. SE Asia has been regarded as a hot spot but fewer studies have been carried out in Africa.

Countries with multiple surveys (>=4) and high prevalence (>20%): Ethiopia, Vietnam, Nigeria, Egypt, and Malaysia.

Impact
In people, leptospirosis most often presents as a febrile illness. Around 5-10% of cases may develop jaundice or other complications and among these case fatality may reach 20%. SE Asia is considered a hot spot and in some areas is the second most common cause of fever after malaria.

In livestock, leptospirosis is associated with abortion, still-birth, infertility and milk reduction in cattle and swine.

There is little good data on losses associated with leptospirosis in developing countries. In Australia, total loss was estimated at 2.2% at herd level (Holroyd, 1980).

In Vietnam, infection with some serovars correlated with one less live pig per litter, equivalent to 8% loss of production (Boqvist et al., 2002).

Key research questions
- Prevalence and incidence in Africa: leptospirosis has been considered most problematic in SE Asia, this study suggests it may be more important than suspected in Africa
- Leptospirosis as a misdiagnosis in people: like brucellosis and Q fever, leptospirosis is often under-diagnosed and better tests as well as awareness raising among the medical community and public is needed
- Context specific vaccination – vaccination is effective but needs to be adapted for the serovars present
- Understanding whether livestock or wildlife are main reservoir: in some studies livestock appear to be the most important reservoir, in others rodents. This has implications for control
- Risk reduction: human behaviour is important in decreasing risk
- Impact of climate change on extreme wet weather events and hence leptospirosis: leptospirosis is strongly associated with flooding and stagnant water

Figure 3.3 Leptospirosis prevalence in community studies

Trypanosomosis – sleeping sickness and ‘the malaria of cattle’

Pathogens
Tsetse-transmitted trypanosomosis is an infectious disease unique to Africa and caused by various species of blood parasites. The disease affects both people (Rhodesian and Gambian sleeping sickness) and animals (nagana).

Test
The most common test in animals is direct microscopic examination of blood for parasites. Hence positive tests correspond to current infections.

Prevalence
103 studies were assessed covering 109,443 animals and 99,808 people. These were mostly parasitological studies so they represent current infections. In community studies, there was a prevalence of 10% among domestic animals and 5% among wild animals. Among humans (either suspect hospital patients or in focal areas for trypanosomosis) prevalence was 6%.
The WHO reports human trypanosomosis as highly spatially distributed: in the last 10 years, over 70% of reported cases occurred in the Democratic Republic of Congo (DRC). Angola, Central African Republic, Chad, Sudan and Uganda make up most of the remaining burden.

Epidemiology

Trypanosoma brucei gambiense (T.b.g.) is found in west and central Africa; it currently accounts for over 95% of reported cases of sleeping sickness and causes a chronic infection (Gambian sleeping sickness). Most transmission is anthroponotic and can be controlled effectively through interventions targeted at human reservoirs; however, animal reservoirs have a role in the epidemiology. Pigs are an animal reservoir and recently have been associated with the persistence and epidemics of sleeping sickness in Uganda, Equatorial Guinea and Cameroon. Other domestic animals and wildlife are also implicated.

T. brucei rhodesiense (T.b.r.) is found in eastern and southern Africa. Nowadays, this form (Rhodesian sleeping sickness) represents less than 5% of reported cases and causes an acute infection.

Agricultural expansion, deforestation and the removal of wildlife reduce the natural habitats and wildlife hosts of tsetse. Moreover, applications of insecticides to cotton and other crops may also reduce tsetse numbers and it is generally agreed that agricultural expansion/intensification is likely to reduce trypanosomosis challenge, at least in the short term (Bourne and Wint, 1994). However, in the short term there may be an upsurge in disease as tsetse, lacking alternative wildlife hosts, feed more on cattle.

Impacts

Acute sleeping sickness is a serious disease in people.

Trypanosomosis has serious health impacts in livestock. However, the non-zoonotic T. congolense and T. vivax are less pathogenic than zoonotic T. rhodesiense. Swallow (1999) summarises a number of studies and estimates reduced productivity of around 10-20% across a range of parameters (mortality, calving rate, milk, draft power).

Geographical hotspots

Countries with multiple surveys (>=4) and high prevalence (>8%): Sudan, Mozambique, Tanzania, Ethiopia, Cameroon, Nigeria, and Burkina Faso

Key research areas:

- Arrest the northerly advance through Uganda of the zoonotic parasite T. brucei rhodesiense, which threatens to converge with T. brucei gambiense
- Farmer-and community-based management of disease: while technical highly effective, sustainability remains elusive
- Trypanocide resistance: an emerging problem across Africa which may also threaten the efficacy of human drugs
- Pen-side tests to allow better and more timely treatments avoiding cattle losses and slowing development of resistance
- Impact of climate, agricultural intensification and demography on disease dynamics
- Factors leading to massive human outbreaks as occurred historically at the start of the 20th century and to a lesser extent in the 1960s
Figure 3.4 Trypanosomosis (trypanosomiasis) prevalence in community surveys

Trypanosomiasis in Bovine

Legend
Prevalence %
- Below 2.5
- 2.6 - 5
- Above 5

Agro-pastoral and pastoral systems
Mixed crop livestock extensive
Mixed crop livestock intensifying
Other urban, forest and landless systems

Trypanosomiasis in Other Species

Legend
Prevalence %
Donkey
- Below 5
- Above 5

Small ruminants
- Below 5

Humans
- Below 5
- Above 5

Pigs
- Below 1
- 1 - 3
**Cysticercosis – pig tapeworm**

**Pathogen**
Cysticercosis is a systemic parasitic infestation caused by the pork tapeworm (*Taenia solium*).

**Tests**
The tests commonly used for cysticercosis in pigs include meat inspection, lingual inspection and antigen ELISA tests. These indicate current infections. In humans, stool samples are also used to identify current tapeworm infections and imaging to identify brain cysts.

**Prevalence**
125 studies were assessed covering 349,923 pigs and 10,385,132 people. In community studies, the average prevalence in pigs was 17%. Among humans the prevalence in community studies was 11% (this combines people infected with *Taenia solium* as well as the much rarer cases of human cysticercosis). Among hospital patients and epileptics the prevalence was 12%.

**Epidemiology**
Humans are at risk not from consumption of pork with cysts but from consumption of tapeworm eggs shed by themselves or another human carrier. The disease persists in poor, pig-keeping communities where pigs have access to human faeces. Intensification would be expected to reduce prevalence of the disease.

**Geographical**
Hot spots: Rwanda, Congo, Chad, Togo, Nigeria, and Ghana (Geerts et al., 2004)

**Impacts**
In some countries, pigs with visible infections (by lingual palpation or mucous membrane inspection) fetch a lower price. This was estimated as a 30% reduction in price in the Cameroon (Praet et al., 2003). A study in Tanzania estimated the price of healthy pigs at $45 and infected at $21: a reduction of 46% (CIRAD, 2012). Heavily infected pigs may be condemned during meat inspection; however, in many countries either smallholder pigs are not always inspected at slaughter or inspection is inadequate.

Cysticercosis is believed to be the most common cause of adult onset epilepsy in poor, pig-keeping communities.

In Cameroon, the cost of treatment of one cysticercosis patient (wage loss not included) was estimated at Euro 260 (Praet et al., 2003)

**Key research questions**
- Eradication of cysticercosis from an ecosystem: with the new vaccine as well as effective therapeutics, cysticercosis is eradicable but there have been no serious investments in Africa or Asia
- Pen-side tests for diagnosis of cysticercosis in pigs: a lateral flow test has been recently developed but requires serum (whole blood would be more convenient)
- More comprehensive and effective meat inspection: as for TB, it appears that because of financial incentives and dysfunctional systems, current meat inspection is not effective in poor countries
- Hotspots among marginalised pig-keeping groups
Figure 3.5 Cysticercosis prevalence in community surveys
**Q fever – the most contagious disease**

**Pathogen**
Q fever is an infectious disease of animals and humans caused by a species of bacteria (*Coxiella burnetii*).

**Tests**
Most tests detect antibodies. Antibodies may persist for several years. Most of the surveys in our review were community based. For these, a positive result indicates current infection, chronic infection or infection in the last few years.

**Prevalence**
We accessed 81 surveys covering 27,252 animals and 11,023 people. In community studies, prevalence was as follows: bovines 28%, other animals (cats, dogs, horses and poultry) 26%, shoats 15%. Among febrile patients in hospitals, 0-40% (average 8%) had antibodies to Q fever.

**Epidemiology**
*Coxiella burnetii* is most frequently found in ruminants (cattle, sheep, and goats) but can also be detected in wildlife and companion animals. According to the literature (although not in our review) sheep appear to be infected most frequently, followed by goats and less frequently, cattle. Human cases are often associated with proximity to small ruminants (particularly at parturition or during abortions) and dry, windy weather. At least in Europe, here is no conclusive evidence in support of a link between an increased density of animals and/or farms and spillover from infected farms to humans (EFSA, 2010).

**Geographical hotspots**
Because Q fever has been investigated in few countries it is difficult to identify hotspots. Countries with high sheep populations would be expected to be at higher risk. Countries with multiple surveys (>=4) and high prevalence (>15%) include in descending order: Nigeria, Zimbabwe, India, and Egypt.

**Impacts**
Animals that carry this organism and shed it into the environment usually do not show any signs of disease. Infected ewes and does may abort or give birth to weak offspring. There is little data available on the economic impacts.

In people around 50% of infections may be asymptomatic; other patients have influenza-like symptoms, a minority have atypical pneumonia or hepatitis. In around 5% of patients, chronic infection establishes.

**Key research questions**
- Prevalence studies in more countries
- Economic impacts of Q fever in livestock
- Factors leading to outbreaks of Q fever in human populations
- High risk groups: pastoralists appear to have very high levels of Q fever – more studies are needed on prevalence and prevention in this group
- Q fever as an emerging disease
- Vaccination to manage Q fever in high risk populations
Figure 3.6 Q fever prevalence in community surveys

Legend
Prevalence %
- Below 5
- 5 - 15
- 15 - 25
- Above 25

Agricultural and pastoral systems
Mixed agro-pastoral extensive
Mixed crop-livestock extensive
Other-urban, forests, and landless systems
**Bacterial food-borne disease – the forgotten zoonoses**

**Pathogen**
In this category we include the bacterial zoonotic diseases, which are transmitted mainly through food. We reviewed *Salmonella*, toxigenic *Escherichia coli*, *Listeria*, *Campylobacter* and *Toxoplasma* which are among the most important causes of food-borne disease as well as hepatitis E, an emerging zoonosis. Other zoonoses of somewhat lesser importance not reviewed are: *Staphylococcus aureus*, *Bacillus cereus*, and *Clostridium spp*.

In this section, we do not include the previously considered classical endemic zoonoses that are often food-borne (brucellosis, Q fever, zoonotic tuberculosis) but have other important transmission pathways. We did not consider non-zoonotic diseases associated with animal source foods (typhoid, rotavirus disease, scarlet fever, giardiasis, shigellosis etc.). We call these ‘forgotten’ zoonoses because health experts, decision makers and the public are often unaware of the important role zoonoses play in food-borne infections.

**Tests**
A variety of tests were used. In most cases, a positive result indicates current, recent or chronic infection.

**Prevalence**
We accessed 258 surveys covering 27,425 animals and 263,995 people and 4,208 food or environmental samples. In community studies, prevalence was as follows: bovines 16%, shoats 20%, pigs 30%, poultry 36%, other animals 17%, food 31%. Among people in the community prevalence was 21% and among high-risk groups prevalence was 20%.

**Geographical hotspots**
Countries with multiple surveys (>=4) and high prevalence (>15%) include in descending order: Tanzania, South Africa, Gambia, Vietnam, Nigeria, Senegal, India, and Egypt.

**Impacts**
Some of these diseases can have impacts in animals (salmonellosis, listeriosis, toxoplasmosis). However, in many cases strains pathogenic to people are not pathogenic to animals which means farmers have less incentives for control.

In people, food-borne disease is an important cause of illness and economic loss. There is no good information on the proportion of gastrointestinal disease burden associated with food-borne zoonoses in developing countries. In developed countries, the proportion varies from 30-50% (Grace et al., 2008). Food-borne pathogens also cause other health problems, less common but more serious (e.g. kidney failure, septicaemia, abortion, encephalitis etc.); around 2-3% of people with acute food-borne zoonoses may also go on to develop serious complications (Lindsay, 1997). The health burden of these is considered to at least equal the burden due to gastrointestinal illness.

**Key research questions**
- Impact of food-borne bacterial diseases in livestock
- Managing food safety in the informal sector where most of the poor buy and sell but food safety regulation is not working
- Gender and food safety – much of food purchase, processing, and handling is done by women but they are often not engaged in food-safety programs
- Relation between food safety and food security
- Attribution – how much food-borne illness is due to zoonotic disease or agricultural products
- High risk groups – the young, old, pregnant and immunosuppressed are especially vulnerable to food-borne disease and special targeting is needed to reach them
Figure 3.8 Food borne disease prevalence in community surveys

Figure 3.9 Toxoplasmosis prevalence in community surveys
Echinococcosis – cystic disease

Pathogen
Cystic echinococcosis (CE) in humans is caused by the larval stage of *E. granulosus*, *E. ortleppi*, *E. intermedius* or *E. canadensis*. All these parasites have canines (usually domestic dogs), as definitive hosts and a variety of ungulates, particularly farm animals, as intermediate hosts. Man is generally an aberrant intermediate host in which the hydatid cyst develops, usually in the liver or lungs as a space-occupying lesion, which can result in considerable morbidity.

Prevalence
We did not review echinococcosis in depth; however, a comprehensive assessment has recently been carried out by Budke et al. (2006).

Epidemiology
Cystic echinococcosis (CE) is a condition of livestock and humans that arises from eating infective eggs of the cestode *Echinococcus granulosus*. Dogs are the primary definitive hosts for this parasite, with livestock acting as intermediate hosts and humans as aberrant intermediate hosts.

Geographical hotspots
More than 90% of human cases occur in the 8 endemic regions in North Africa-near East and China. In descending order: China (Tibetan plateau), Turkey, India, Iraq, Iran, and Afghanistan.

Impact
A preliminary estimate of the annual global burden of CE has suggested approximately 1 million DALYs are lost due to this disease (Budke et al., 2006). This is likely to be a substantial underestimate (Craig et al., 2007). In addition the losses to the global livestock industry is around $2 billion lost annually and cost of illness is around the same.

We conducted a systematic review for toxoplasmosis but have not included the information here because of some epidemiological complexities. For toxoplasmosis, high prevalence may be associated with less risk, because the most vulnerable group (pregnant women) are exposed before they are pregnant. For hepatitis E there is some uncertainty over the extent of zoonotic transmission.

Key research questions
- Control of echinococcosis in remote and insecure regions
- Relation between toxoplasmosis prevalence and risk: as toxoplasmosis is most serious if encountered by a naïve pregnant women, it may be that cultures where exposure to toxoplasmosis is very high (e.g. France) have less disease burden than cultures where exposure is low
- Prevalence of toxoplasmosis
- Changing behaviours that increase exposure to toxoplasmosis and echinococcosis
- Zoonotic component of hepatitis E
3.3 Top twenty countries for endemic zoonoses burden

Twenty-eight countries appeared in the ‘geographical hotspots’ listing. To be considered a geographical hotspot for a disease, a country had multiple surveys (human and animal combined, but only community surveys) with a high average prevalence (table 3.2). (The cut-off prevalence varied with disease reflecting that for different diseases, different prevalences are considered high\(^6\)). The country with the highest prevalence is ranked as 1. The ranking for each disease is shown in Table 3.2 as well as the number of surveys and cut-off prevalence the ranking was based on.

<table>
<thead>
<tr>
<th>Table 3.2 Geographical hotspots for zoonotic disease (country)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brucellosis</td>
</tr>
<tr>
<td>------------</td>
</tr>
<tr>
<td>1 Togo</td>
</tr>
<tr>
<td>2 Mali</td>
</tr>
<tr>
<td>3 Ivory Coast</td>
</tr>
<tr>
<td>4 Zambia</td>
</tr>
<tr>
<td>5 Niger</td>
</tr>
<tr>
<td>6 India</td>
</tr>
<tr>
<td>7 Sudan</td>
</tr>
<tr>
<td>8 Cameroon</td>
</tr>
<tr>
<td>9 Burundi</td>
</tr>
</tbody>
</table>

To calculate the countries with the highest burden of endemic zoonoses we gave each country a weighting according to its ranking for average prevalence for each disease (the country with the highest prevalence for the disease got a weighting of ten). Weights were added across the diseases (so a higher score represents a higher average prevalence summed across all the endemic zoonoses considered). The countries with highest burden of endemic zoonoses are shown in Table 3.3. (This ranking is probably biased towards countries with better university and research infrastructure as they conduct and publish more studies: for example, there are many more studies from Nigeria than from the Central African Republic).

<table>
<thead>
<tr>
<th>Table 3.3 Countries with most zoonotic disease hotspots</th>
</tr>
</thead>
<tbody>
<tr>
<td>Country</td>
</tr>
<tr>
<td>---------------</td>
</tr>
<tr>
<td>Nigeria</td>
</tr>
<tr>
<td>Ethiopia</td>
</tr>
<tr>
<td>Tanzania</td>
</tr>
<tr>
<td>Togo</td>
</tr>
<tr>
<td>India</td>
</tr>
<tr>
<td>Mali</td>
</tr>
<tr>
<td>Vietnam</td>
</tr>
<tr>
<td>Sudan</td>
</tr>
<tr>
<td>Bangladesh</td>
</tr>
<tr>
<td>Burkina Faso</td>
</tr>
</tbody>
</table>

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\(^6\) To give an extreme example, for a rare disease like rabies one in 100,000 animals might be considered a high prevalence, while for a common disease like brucellosis one in 5 animals might be considered high.
3.4 Outbreak zoonoses

We retrieved papers on three of the outbreak zoonoses that appeared in the ‘top 13’ zoonoses listing. These were: rabies, anthrax and leishmaniasis.

**Rabies**

Twenty-one papers were accessed on rabies. These were not useful for assessing prevalence or cases but were consistent with the geographical patterns of rabies. Most rabies cases are concentrated in high-risk countries in Africa and Asia. (Bangladesh, India, Myanmar, Pakistan, China, Egypt, Sudan, Ethiopia, Tanzania, Ghana).

**Anthrax**

Thirty-five papers were accessed on anthrax. They were not useful for assessing prevalence or cases but were generally consistent with the geographical patterns of anthrax. There are approximately 10-100 thousand human incidences annually throughout the world with significant numbers of cases in Chad, Ethiopia, Zambia, Zimbabwe and India.

**Leishmaniasis**

Paper retrieval was not useful in assessing prevalence. Most leishmaniasis is zoonotic, but anthropogenic transmission is more important in outbreaks, 90% of human visceral leishmaniasis cases occurring in South Asia, Sudan, Ethiopia, and Brazil and 90% of cutaneous leishmaniasis cases occurring in Afghanistan, Algeria, Iran, Saudi Arabia, Syria, Brazil, Colombia, Peru, and Bolivia.

Other important outbreak pathogens that were not in the ‘top 13’ list but did appear in the ‘top 25’ were:

- Rift Valley fever virus
- Hanta virus
- Ebola virus
- Chickungunya virus.

Avian influenza was also in this list, a zoonosis that is endemic in some regions (Indonesia, South China, Egypt and possibly elsewhere), but in most countries occurs as outbreaks, which are controlled (rich countries) or burn out (poor countries) (Bett et al., in press).

These five pathogens (including avian influenza) are all caused by viruses and are characterised by high case fatality but low burden of disease. Together they cause around 15,000 deaths a year which is trivial in comparison to the top 13 zoonoses (causing 2.5 million deaths a year). Currently humans are mainly spill-over hosts and there is no sustained anthropogenic transmission (human-to-human). However, if these pathogens were to mutate to allow easy human-to-human transmission while maintaining their high case fatality the impacts would be enormous. Hence, these diseases are of interest not so much because of their burden of disease but because of potential to become diseases with higher burden. Smallpox, bubonic plague, HIV-AIDS, malaria and measles are examples of former zoonoses that jumped species with civilisation-altering impacts (Wolfe et al., 2007).

Perry and Grace (2009) argue that many negative impacts of zoonoses and emerging diseases are from inappropriate responses by authorities, farmers and general public rather than disease itself. This was especially evident in the avian influenza pandemic, when outbreaks led to large changes in purchasing behaviour, which probably had little impact on mitigating risk. Similarly, the reluctance to support commodity-based trade is prejudicial to developing countries without any commensurate benefit in reducing human health risk.
The map was extracted from HealthMap (www.healthmap.org). It shows all disease reports between Jan 1st 2010 and May 2nd 2012 of the endemic zoonoses considered in this report. The sources used were: ProMed, FAO, OIE, Eurosurveillance, Google.

**Figure 3.10 Outbreaks of five important zoonotic diseases 2010-2012 as aggregated and reported by HealthMap (www.HealthMap.org)**

(Extracted from HealthMap, www.healthmap.org)

In all 200 reports are cited, on the HealthMap site, distributed as follows: leptospirosis (124), trypanosomosis (24), brucellosis (20), Q fever (12) and bovine tuberculosis (10). Between Jan 1st 2010 and Dec 31st 2010 there were only three reports for brucellosis and all were in people. As explained in Chapter 2 this implies under-reporting of actual new cases by several orders of magnitude.
3.5 Global burden of disease
The original Global Burden of Disease Study (GBD) was commissioned by the World Bank in 1991 to provide a comprehensive assessment of the burden of 107 diseases and injuries and ten selected risk factors for the world. Burden of disease is calculated using the disability-adjusted life year (DALY). This time-based measure combines years of life lost due to premature mortality and years of life lost due to time lived in states of less than full health. The GBD represents the most authoritative source of information on human illness.

There are some challenges in using the GBD to assess zoonoses.
- Firstly, zoonoses (especially in poor countries) are widely unreported, and under-reporting is relatively greater for zoonoses than for non-zoonotic diseases of comparable prevalence (Schelling et al., 2007). As the GBD report is based on national information for levels of mortality and cause of illness, this under-reporting is reflected in the GBD.
- Secondly, several zoonoses with considerable burdens are not included in the GBD assessment. For example, rabies, echinococcosis, cysticercosis, leptospirosis, and brucellosis.
- Thirdly, the GBD is organised around diseases and not pathogens or transmission pathways. For example, diarrhoeal diseases, among the highest causes of morbidity and mortality in poor countries, comprise one category. Although the majority of important diarrhoeal pathogens are zoonotic (Schlundt et al., 2004) it is not currently possible to identify the zoonotic component of diarrhoeal disease from GBD figures.

In order to use the GBD to estimate disease we made some assumptions:
- Tuberculosis: we took the conservative estimate of Cosivi (1998) who estimated worldwide the proportion of TB caused by M. bovis at 3.1%. This literature review suggests the proportion is higher. A higher proportion is also consistent with historical data.
- Diarrhoeal diseases: we assumed 33% of diarrhoea disease is due to zoonotic pathogens. In developed countries, several reviews (Schlundt et al., 2004, Flint et al., 2005) argue the majority of gastrointestinal disease burden is due to zoonotic pathogens (>50%). However, given the lack of evidence for developing countries we took a conservative estimate of 33%.
- Trypanosomosis, Chagas disease, leishmaniasis: Japanese encephalitis: are all considered as zoonotic.
- Schistosomiasis: cases in regions where the zoonotic species *Schistosoma japonicum* predominates are considered zoonotic.
- Dengue is not included. Although dengue is a zoonosis and the sylvatic cycle (monkey-mosquito) has important implications for disease eradication, most transmission is human-to-human.
- Tetanus is not included. Tetanus is a sapro-zoonoses and the load of toxins in the environment is largely the result of contamination with ruminant faeces. However, most human burden is from contact with the environment and not animals.
- Respiratory disease is a major cause of human sickness and death and a certain proportion is due to zoonotic diseases such as Q fever. We did not include these as no reliable estimates could be found.

For zoonoses recorded in the GBD, 68% of the burden is made up of just 13 countries (Figure 3.1). There is a very high correlation (99%) between protein energy malnutrition and burden of zoonoses indicating the strong relation between poverty, dependence on livestock, and zoonotic disease.

---

7 Protein-energy malnutrition is a nutritional deficiency resulting from either inadequate energy (caloric) or protein intake and manifesting in either marasmus or kwashiorkor. Marasmus is characterised by wasting of body tissues, particularly muscles and subcutaneous fat, and is usually a result of severe restrictions in energy intake. Kwashiorkor affects mainly children, is characterised by oedema (particularly ascites), and is usually the result of severe restrictions in protein intake. However, both Types can be present simultaneously (marasmic kwashiorkor) and mask malnutrition due to the presence of oedema.
Figure 3.11 Health burden of zoonoses in million disability adjusted life years (DALY)

From Global Burden of Disease, World Health Organisation, 2008
3.6 Comparing systematic literature review to in country literature search

Introduction and summary

In this review we extracted papers mainly in English (with a minority in French) from medical and agricultural databases available online. Systematic literature reviews which only include some languages and which depend on major databases risk missing important information. Hence, we conducted a study in Vietnam to review literature for three of the key zoonoses (Data provided by HSPH).

154 papers were identified of which 117 were in Vietnamese and only 27 in English.

Methods

We used a large range of Vietnamese scientific journals, library documents, as well as meetings with key researchers on zoonoses, and open sources.

- Vietnamese journals on preventive medicine, practical medicine, public health, veterinary sciences and techniques, agriculture and rural development
- Institution libraries: Vietnam Medical Information Centre (MOH), National Institute of Hygiene and Epidemiology, National Institute of Malaria Research, National Institute of Animal Husbandry.
- University libraries: Hanoi Medical University, Hanoi School of Public Health, Hanoi University of Agriculture
- Key researchers/research groups from institutes and universities
- Conferences proceedings
- Web sites: Ministry of Health, Ministry of Agriculture and Rural Development

The process of creating the search strategy consisted of two steps: (i) identification of key concepts characterizing the research questions and (ii) generation of a list of search terms that reflected the key concept. The main concept identified was zoonotic diseases in Vietnam. For this concept a number of subject terms and keyword terms were identified, which was then combined for the search:

The overall search term components considered to define “zoonotic diseases” AND “Vietnam” for the search, were: (i) population surveyed (human or animal), (ii) prevalence and (iii) laboratory techniques. Diseases were searched by their common names, as well as the names of the causative agents. The keywords were used both English and Vietnamese, for examples:

- English: “Cysticercosis” AND “Vietnam” or “Taeniasis” AND “Vietnam”
- Vietnamese: “Sán dây lợn” or “Bệnh lợn gào”

English papers: we searched online databases of Science Direct, Pubmed and Web of Science with keywords of disease name or names of the pathogens in the fields of title/keywords/abstract.

All the electronic copies and hard copies papers were scanned and reviewed from their title and abstract to see if the papers are relevant to research on zoonotic diseases in Vietnam. After 2 screening rounds, many zoonoses were identified from research done. However, due to the time constraint and to respond to the TOR, we decided to select 3 zoonotic diseases, including cysticercosis (pig tapeworm), leptospirosis, and Salmonellosis for in-depth review. For each of the paper related to the selected diseases, we collected key information on i) location of the study, ii) on human or animal or both, iii) robustness of research design, iv) analysis method, v) prevalence. It happened also when a paper reporting different values for different sample analysis (e.g. milk, serum or both), these were treated separately to have different prevalences of the targeted samples.
Results
We found 50 papers, project reports and student's thesis study related to cysticercosis, including 47 in Vietnamese and 3 in English; 64 papers, project reports and student's thesis study related to salmonellosis, including 40 in Vietnamese and 24 in English; 40 papers, project reports and student's thesis study related to leptospirosis, including 30 in Vietnamese and 10 in English.

Conducting an in-country review covering Vietnamese journals, libraries and conference proceedings dramatically increased the number of papers and samples. It also revealed papers on less commonly studied aspects (e.g. cysticercosis in animals other than pigs and leptospirosis in wildlife) which were missed by the systematic, web-based, mainly English language review.

For four of the six comparisons, the prevalence estimated by systematic and in-country review were similar, but for two there was a marked discrepancy (27% prevalence of cysticercosis in people versus 4% and 57% prevalence of leptospirosis in livestock and companion animals (domestic) versus 14% in the systematic and in-country reviews respectively). The much smaller number of studies in the systematic review makes it likely that these are less accurate.

However, while 60% of the papers in Vietnamese were judged to have a ‘moderate’ or ‘weak’ methodology, only 37% of the papers in English were so judged.

Table 3.4 Comparing data on cysticercosis from a systematic review and an in-country review

<table>
<thead>
<tr>
<th></th>
<th>Systematic review</th>
<th>In country review</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>People</td>
<td>Pigs</td>
</tr>
<tr>
<td>Studies (number)</td>
<td>6</td>
<td>1</td>
</tr>
<tr>
<td>Community studies (number)</td>
<td>5</td>
<td>1</td>
</tr>
<tr>
<td>Prevalence (%)</td>
<td>27.4</td>
<td>9.9</td>
</tr>
<tr>
<td>Samples for prevalence (no.)</td>
<td>1,434</td>
<td>323</td>
</tr>
</tbody>
</table>

Table 3.5 Comparing data on leptospirosis from a systematic review and an in-country review

<table>
<thead>
<tr>
<th></th>
<th>Systematic review</th>
<th>In country review</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>People</td>
<td>Domestic</td>
</tr>
<tr>
<td>Studies (number)</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>Community studies (number)</td>
<td>1</td>
<td>3</td>
</tr>
<tr>
<td>Prevalence (%)</td>
<td>12.8</td>
<td>57.0</td>
</tr>
<tr>
<td>Samples for prevalence (no.)</td>
<td>961</td>
<td>456</td>
</tr>
</tbody>
</table>

Table 3.6 Comparing data on salmonellosis from a systematic review and an in-country review

<table>
<thead>
<tr>
<th></th>
<th>Systematic review</th>
<th>In country review</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>People</td>
<td>Animals</td>
</tr>
<tr>
<td>Studies (number)</td>
<td>1</td>
<td>11</td>
</tr>
<tr>
<td>Community studies (number)</td>
<td>0</td>
<td>11</td>
</tr>
<tr>
<td>Prevalence (%)</td>
<td>n/a</td>
<td>13.1</td>
</tr>
<tr>
<td>Samples for prevalence (no.)</td>
<td>n/a</td>
<td>6831</td>
</tr>
</tbody>
</table>
Figure 3.12 Leptospirosis and cysticercosis in Vietnam identified from in-country review.
3.7 Interpretation of the review of endemic and outbreak zoonoses

As well as the likely existence of large amounts of missed literature because literature review was based on English/French publications indexed on online databases, other weaknesses and potential biases include:

- Many zoonoses have never been looked for in many places, and published literature reflects research infrastructure as well as disease prevalence.
- Zoonoses which are more difficult to detect or test for (e.g. campylobacteriosis, listeriosis) are under-represented because they are rarely investigated.
- We generally used data from the last 10 years. If data was very scarce we extended some searches backwards. However, given rapid changes in farming systems as well as changes in diagnostic techniques earlier studies may not reflect the current situation.
- Studies indicate infection and not necessarily clinical disease.
- Some surveys don’t distinguish to species level making it impossible to distinguish between zoonotic and non-zoonotic pathogens.
- Surveys focus on presence rather than transmission: so *Mycobacterium tuberculosis* will usually be classified as non-zoonotic although it is possible that the human victim acquired the infection from livestock.
- Most surveys only report on one pathogen.
- Varying sensitivity and specificity of surveys because of different tests used making direct comparison difficult.
- Often little information on sampling and some community surveys may have considerable selection bias although authors claim sampling was representative.
- In general, studies with fewer samples and country estimates based on fewer studies appear to over-estimate prevalence. This may be because researchers focus on areas they think may have a problem even if this is not always reported in the survey.
- Some areas are over-represented (near universities) and many others under-represented.
- Some countries are under-represented because of less capacity in assessing zoonotic diseases.
- In the current analysis, small samples have as much weight as large samples. We plan to construct weighted prevalences by sample size and extrapolate to populations at risk.
- We class slaughterhouse surveys as ‘community’ that is representative of the livestock population. In some places, sick animals are less likely to be slaughtered but in others they are more likely.

Suggestions for overcoming these problems and improving our understanding of zoonoses of importance to the poor:

- Further analyse the data collected to allow better extrapolation to agro-ecosystems and investigate risk factors.
- Conduct large scale, probabilistic, stratified surveys to accurately determine prevalence of key pathogens.
- Conduct surveys in regions where pathogens are likely to be present but data is lacking (risk-targeted).
- Collect economic and behavioural data to understand the impact and risk factors for key pathogens.
- Develop better, cheaper diagnostics that can detect multiple pathogens in multiple species.
- Support bio-repositories for pathogens with meta-data allowing investigation of epidemiology and risk factors.
Chapter 4: Updated map of emerging zoonotic disease events

Summary
The study searched for papers on zoonotic disease emergence events since 2004. Out of 43 new or newly identified events, most are viral and zoonotic from wild animal hosts. Although, the mappable zoonotic new events (n = 30) are globally spread across every continent, there may be clusters in northeast US, South America, Europe and South East Asia. These trends likely reflect surveillance differences with perhaps a trend to higher representation in developing countries reflecting increased attention over this period. Combined with existing data on zoonotic EID events from 1940-2004 (n = 202), the clearest potential hotspots are USA and Western Europe, which may reflect historical surveillance differences. Countries with most events are USA, UK, Australia, and France. (Chapter prepared by IOZ)

4.1 Introduction
Novel pathogens continue to emerge worldwide, the majority of which are zoonotic. ILRI engaged Zoological Society of London and Ecohealth Alliance to produce updated high resolution maps of zoonotic human emerging infectious disease (EID) events, by 1) extraction from the database used in Jones et al. (2008), and 2) further collection of EID events to update the database until 2012.

4.2 Methodology for updating map of emerging zoonotic disease events
We review the zoonotic human EID events in Jones et al. (2008), and methods for data collection and mapping of new events.

1) Jones et al. (2008) EID database. The data from Jones et al. (2008) covers 335 human EID events occurring between 1940 and 2004, 216 of which were zoonotic. In the Jones et al. (2008) analysis, an EID event was defined as the first emergence of a pathogen in a human population as a result of either increasing incidence, virulence, geographic range, drug resistance, or any other cited factor in case reports and literature. Single case reports were excluded, as were those with uncertain data quality. Separate pathogen strains were not considered as separate events, with the exception of drug resistant strains. For the maps presented in Jones et al. (2008), the location of each EID event was geo-referenced to point localities where possible, based on locations given in data sources (e.g. a village, or a hospital). These data were then converted to give each pathogen a location in a one decimal degree global grid. However, 95 of these events could only be traced to larger areas such as subnational units (e.g. states) or whole countries. To standardise for further analysis in the paper, one grid cell in these areas was selected via a random draw to represent the location of a particular EID. One of the random draws was selected for the maps presented in the paper.

For the purposes of this report, we mapped all those events that had a point locality, and where only subnational unit location information was available, we assigned an EID event to the weighted centroid of that area calculated using the ‘Generate Centroid’ function in HawthsTools 3.27 for ArcMap 9.3. Where only country information was present, these data were excluded from the maps. A total of 172 events were mapped (Fig. 1a).

2) Further data collection of EID events. We made slightly different assumptions when collecting new data than in the original Jones et al. (2008) protocols, based on updated criteria and suggested changes to the definition of “emerging” since 2008. We only considered emerging pathogens to be those completely novel to humans, or having novel virulence in humans or novel drug resistance. Geographic range, incidence and any miscellaneous factors were excluded. Furthermore, single case reports are not excluded, in contrast to the previous map. Separate strains or subspecies are once
again only considered in the context of virulence and drug resistance. We have also extended our criteria to cover issues not addressed by Jones et al. (2008). We excluded any events from non-natural infections (for example, accidental inoculation of laboratory workers), and we accept novel pathogens not yet given certified scientific names. Additionally, we explicitly define an event as zoonotic if there is evidence the pathogen has an animal host or vector, or if it is cited in the literature as a likely zoonosis. Much assumption of zoonotic status is based upon transmission/natural hosts of closely related pathogens. Zoonoses were sub-classified by host type into one of four groups: 1) wildlife hosts only, 2) non-wildlife (i.e. domestic) hosts only, 3) both host types, 4) unknown. Finally, we accepted any diagnostic method (e.g. serology, microbial culture, DNA) as evidence of human infection (or animal infection for evidence as a zoonosis).

Initial leads on new EID events were obtained by various search methods including; 1) searches in peer-reviewed journal articles using Web of Science v5.0 (search terms = [novel OR new] AND human AND pathogen AND [emerging OR outbreak]); 2) searches in the ProMED reporting archives (search terms = [novel OR new] AND human); and 3) expert suggestion (EcoHealth Alliance and colleagues). Initial searches were conducted in articles from 2004 onwards, although no date restriction was made in accepting EID events or their supporting references. From initial leads, further information was found via follow-up of references. Once a potential EID event was identified, we extensively traced literature on the pathogen both backwards and forwards to find the first chronological case (not the first chronological report). Data were not recorded if the pathogen could not be assigned a single, feasible spatiotemporal origin (for example, many novel human viruses have recently been discovered that were then found in populations worldwide and/or historical samples, giving no clear emergence time or location). Following Jones et al. (2008), we recorded multiple fields of interest including details of pathogen taxonomy, transmission mode, known hosts, pathogenicity, data quality and any miscellaneous notes, in addition to geographic location. This information was generally unavailable for pathogens described very recently (in the past 10 years or so) because they have not yet been studied in detail, though no exclusions were made based on missing supplementary data.

4.3 Results: new zoonotic emerging disease events
We identified 43 new EID events, 34 of which were zoonotic (Fig. 4.1b). Mapping of the new EID events followed that above for the previously gathered events. Point localities were all mapped were exact coordinates were given in the source and the exact coordinates were found using Google Earth v6.2. If only textual geospatial information was given, the locality of weighted centroids was assigned to EID events with only sub-national unit and EIDs with only country localities were excluded. In total, 30 new EID events were retained.

The majority of these new EID event pathogens are viruses, with several bacteria and only two protozoa (Table 1). Three events were due to drug-resistant strains, and only one due to a newly virulent strain. Although data was collected with a perspective to continuing from where Jones et al. (2008) left off, 15 EID events occurred pre-2004, 7 of which are from the 1980s or 1990s. This likely includes previously unidentifiable outbreaks that have only been classified using modern techniques. 17 of the 30 EID event pathogens in Table 1 have identified reservoir hosts or at least known animal infections (11 wild, 2 domestic, 4 both host types). However, most are recently discovered, therefore natural hosts for the remaining 13 will no doubt take time to locate and confirm.

4.4 Maps of zoonotic emerging disease events
To remap the combined previous and the new EID events, we created a one decimal degree grid using HawthsTools 3.27, and used ArcMap 9.3’s ‘Spatial Join’ tool to assign each EID event point to a single grid cell. New zoonotic EID events were mapped and labelled (Fig. 4.2) (n=30). Although the
events are globally spread across every continent, there may be potential clusters of new or newly identified EID events in the Northeast US, South America, Continental Europe, and Southeast Asia. Only one grid cell contained more than one event, which was in Cameroon, containing Human T-lymphotrophic viruses 3 and 4, which were discovered simultaneously (Table 1). Previous EID events were mapped in combination with the new events (n=202) (Fig. 4.3). Again, events are present across most of the inhabited world. In contrast to the new events, the clearest potential hotspots are now the USA, and Western Europe, which likely reflects historical differences in surveillance and reporting. As in Jones et al., the maximum number of events in any grid cell was 6, occurring in Central London, UK. Separate maps were also produced for those events with wild hosts, and non-wild hosts (Fig. 4.4 4.5). Similar patterns were present for each, with the most noticeable difference being that very few EID events from non-wild hosts occurred in Africa or South America. Maps were also produced to illustrate breakdown of events in each grid cell by zoonotic host categorisation, drug resistance, and type of data (see Figure 3).

Figure 4.1. Tree diagrams illustrating data structure for EID events from 1) Jones et al. 2008, and 2) this update, with respect to zoonotic status, spatial data quality of entries, and whether data was accepted or rejected for the new maps contained in this report.
Figure 4.2 New zoonotic disease events identified in 2012 and not previously mapped

2012 Update Zoonotic EID Events  ★  1  ★  2
Figure 4.3 Previous zoonotic emerging disease events were mapped in combination with the new events.

**Zoonotic EID Events**

- 1
- 2-3
- 4-5
- 6

![Map of zoonotic EID events worldwide with legends indicating the number of events.]
Figure 4.4 Zoonotic emerging disease events with wildlife hosts

Zoonotic EID events (Wild hosts)
Figure 4.5 Zoonotic emerging disease events with non-wildlife hosts

Zoonotic EID events (Non-wild hosts)  • 1  • 2  • 3-4
Figure 4.6 Maps of all zoonotic emerging infectious disease events (n = 202) stratified by potential variables of interest. Size of circles denotes number of events in each one degree grid cell, and colour denotes breakdown of events in terms of type of zoonotic host
Figure 4.7 Maps of all zoonotic emerging infectious disease events (n = 202) stratified by potential variables of interest. Size of circles denotes number of events in each one degree grid cell, and colour denotes breakdown of events in terms of drug-resistant events.

b)

Zoonotic EID Events

- ● 1
- ○ 2-3
- ○ 4-5
- ○ 6

- ▲ Drug-Res
- ▼ Non Drug-Res
Figure 4.8 Maps of all zoonotic emerging infectious disease events (n = 202) stratified by potential variables of interest. Size of circles denotes number of events in each one degree grid cell, and colour denotes breakdown of events in terms of spatial data resolution (see also Figure 1).
4.5 Conclusion on zoonotic emerging infectious disease vents
We have geo-located and mapped a total of 202 zoonotic emerging infectious disease events, 30 of which were newly collected during this study. High-resolution maps are provided for all events, as well as events stratified by type of zoonotic host, and other potential classifiers of interest. Potential trends in hotspot distribution could reflect surveillance differences. It is also possible that the higher representation of developing countries within the new events may reflect increasing research focus on developing countries or improving diagnostic technology.

(Note
Is the world becoming sicker or are we just better able to detect disease? The last decades have seen dramatic improvements in biological disease detection with dozens of new potential pathogens anticipated by 2020. At the same time innovations in information management are increasing awareness of disease outbreaks. Perry et al. (2011) explore this in a recent review and conclude that there is overall evidence for increased emergence of disease in recent decades, and not just improvements in diagnosis and surveillance. The current increase in disease emergence is not historically unprecedented: major epidemiological transitions also occurred during the Neolithic when livestock were domesticated on a wide-scale, during the age of exploration when Old World pathogens were introduced to the New World, and to a lesser extent with increased global travel in the nineteenth century).
Chapter 5: Poor livestock keepers in livestock systems

Summary
We also updated the regional maps based on the national poverty data showing:
- Agro-ecosystems
- Numbers of cattle, sheep, goats and pigs
- Human population
- Poor livestock keepers

The maps and tables are shown in the this section as well as a summary of regional characteristics relevant to the review.

Figure 5.1 Poor livestock keepers (million) by region
East Africa
Countries: Sudan, Ethiopia, Eritrea, Djibouti, Somalia, Kenya, Uganda, Rwanda, Burundi, Tanzania

Fig 5.2: Eastern Africa region: Farming systems (Herrero et al 2009, Notenbaert et al 2009)

Table 5.1: Species population by farming system in Eastern Africa (Herrero et al 2009)

<table>
<thead>
<tr>
<th>Farming system</th>
<th>Agro-pastoral and pastoral</th>
<th>Mixed extensive</th>
<th>Mixed intensifying</th>
<th>Others</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bovines</td>
<td>32,239,100</td>
<td>59,221,300</td>
<td>13,481,300</td>
<td>4,479,000</td>
<td>109,420,700</td>
</tr>
<tr>
<td>Goats</td>
<td>35,603,000</td>
<td>31,020,500</td>
<td>6,699,100</td>
<td>3,082,780</td>
<td>76,405,380</td>
</tr>
<tr>
<td>Sheep</td>
<td>34,404,700</td>
<td>29,999,200</td>
<td>4,893,750</td>
<td>2,254,030</td>
<td>71,551,680</td>
</tr>
<tr>
<td>Pigs</td>
<td>85,169</td>
<td>433,390</td>
<td>543,184</td>
<td>183,700</td>
<td>1,245,443</td>
</tr>
<tr>
<td>People</td>
<td>30,608,700</td>
<td>76,115,200</td>
<td>37,649,000</td>
<td>14,669,900</td>
<td>159,042,800</td>
</tr>
<tr>
<td>Poor livestock keepers</td>
<td>12,125,100</td>
<td>31,719,400</td>
<td>11,281,600</td>
<td>2,740,620</td>
<td>57,866,720</td>
</tr>
</tbody>
</table>

East Africa is characterised by:
- Poor livestock keepers 36% of the population; cattle are relatively important.
- Mixed extensive systems predominate but agro-pastoral /pastoral are more important than most other regions. Pastoralists have high vulnerability to zoonoses.
- Zoonoses with a wildlife interface are important.
- Rapidly dairy development in highlands: bringing risks of brucellosis, tuberculosis and milk-borne diseases.
- Rapid growth in pig production in Uganda brings risks of emerging disease such as Ebola.
- Intra-regional trade important for the horn of Africa (shoats), and of interest to Ethiopia.
- High zoonoses burden in Ethiopia and Tanzania.
- Insecurity in Somalia and possibly South Sudan with implications for zoonoses.
Southern Africa Region

Countries: Zambia, Malawi, Mozambique, Zimbabwe, Botswana, Swaziland, Lesotho, South Africa

Fig 5.3: Southern Africa region: Farming systems (Herrero et al 2009)

Table 5.2: Species population by farming system in SA (Herrero et al 2009)

<table>
<thead>
<tr>
<th>Farming system</th>
<th>Agro-pastoral/pastoral</th>
<th>Mixed extensive</th>
<th>Mixed intensifying</th>
<th>Others</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bovines</td>
<td>9,611,320</td>
<td>8,532,650</td>
<td>2,307,420</td>
<td>1,746,690</td>
<td>22,198,080</td>
</tr>
<tr>
<td>Goats</td>
<td>7,300,510</td>
<td>6,164,890</td>
<td>643,834</td>
<td>1,301,330</td>
<td>15,410,564</td>
</tr>
<tr>
<td>Sheep</td>
<td>9,611,320</td>
<td>8,532,650</td>
<td>2,307,420</td>
<td>1,746,690</td>
<td>22,198,080</td>
</tr>
<tr>
<td>Pigs</td>
<td>696,286</td>
<td>1,300,700</td>
<td>840,703</td>
<td>215,223</td>
<td>3,052,912</td>
</tr>
<tr>
<td>People</td>
<td>17,757,100</td>
<td>32,350,900</td>
<td>5,730,630</td>
<td>16,532,100</td>
<td>72,370,730</td>
</tr>
<tr>
<td>Poor livestock keepers</td>
<td>6,286,600</td>
<td>10,182,200</td>
<td>783,379</td>
<td>1,650,570</td>
<td>18,902,749</td>
</tr>
</tbody>
</table>

Southern Africa is characterised by:
- Poor livestock keepers 26% of population. Lowest in SSA; Cattle relatively important.
- Agro-pastoral/pastoral more important. Pastoralists have high vulnerability to zoonoses.
- Zoonoses with a wildlife interface are important.
- Significant commercial ranching and farming, with better animal health and zoonoses control. Some countries with export potential.
- Better animal health services and disease reporting systems than most other SSA regions.
- Wide regional variation in farming systems, zoonoses and response capacity.
West Africa region

Countries: Mauritania, Mali, Niger, Chad, Senegal, The Gambia, Guinea Bissao, Guinea, Sierra Leone, Liberia, Ivory Coast, Burkina Faso, Ghana, Togo, Benin, Nigeria, Cameroon

Fig 5.4: West Africa region: Farming systems (Herrero et al 2009)

Table 5.3: Species population by farming system in WA (Herrero et al 2009)

<table>
<thead>
<tr>
<th>Farming system</th>
<th>Agro-pastoral/pastoral</th>
<th>Mixed extensive</th>
<th>Mixed intensifying</th>
<th>Others</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bovines</td>
<td>13,458,000</td>
<td>28,220,300</td>
<td>2,403,490</td>
<td>1,034,530</td>
<td>45,116,320</td>
</tr>
<tr>
<td>Goats</td>
<td>19,936,400</td>
<td>37,553,900</td>
<td>16,816,900</td>
<td>2,603,100</td>
<td>76,910,300</td>
</tr>
<tr>
<td>Sheep</td>
<td>13,841,500</td>
<td>29,747,800</td>
<td>8,505,920</td>
<td>2,024,590</td>
<td>54,119,810</td>
</tr>
<tr>
<td>Pigs</td>
<td>810,597</td>
<td>2,598,830</td>
<td>2,518,430</td>
<td>1,367,550</td>
<td>7,295,407</td>
</tr>
<tr>
<td>People</td>
<td>18,540,400</td>
<td>80,711,100</td>
<td>72,559,700</td>
<td>13,249,700</td>
<td>185,060,900</td>
</tr>
<tr>
<td>Poor livestock keepers</td>
<td>11,343,700</td>
<td>38,161,100</td>
<td>19,231,600</td>
<td>2,011,420</td>
<td>70,747,820</td>
</tr>
</tbody>
</table>

West Africa is characterised by:

- Poor livestock keepers 38% of the population; highest in SSA. Goats (followed by sheep) relatively most important species.
- Mixed extensive more important, but mixed intensifying more important than other regions of SSA. Pastoralism/agro-pastoralism mainly in the Sahel.
- Cattle important for traction, but tsetse a major barrier. Trypanotolerant cattle important in the some areas. Little dairying and reliance on imported dairy products. Traditional dairying in the Sahel has high risk of milk-borne zoonoses because of cultural practices.
- Also high imports of poultry. Slow growth in livestock production.
- High zoonoses burden in Nigeria and sub-humid coastal countries.
- Insecurity in several countries.
North Africa region
Countries: Western Sahara, Morocco, Algeria, Tunisia, Libya, Egypt

Fig 5.5: North Africa Region: Farming systems (Herrero et al 2009)

Table 5.4: Species population by farming system in NA (Herrero et al 2009)

<table>
<thead>
<tr>
<th>Farming system</th>
<th>Agro-pastoral pastoral</th>
<th>Mixed extensive</th>
<th>Mixed intensifying</th>
<th>Others</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bovines</td>
<td>2,008,120</td>
<td>1,595,520</td>
<td>2,367,870</td>
<td>462,762</td>
<td>6,434,272</td>
</tr>
<tr>
<td>Goats</td>
<td>6,325,060</td>
<td>1,667,820</td>
<td>1,881,460</td>
<td>578,935</td>
<td>10,453,275</td>
</tr>
<tr>
<td>Sheep</td>
<td>23,411,000</td>
<td>10,130,600</td>
<td>5,593,700</td>
<td>1,606,520</td>
<td>40,741,820</td>
</tr>
<tr>
<td>Pigs</td>
<td>2,321</td>
<td>477</td>
<td>26</td>
<td>0</td>
<td>2,824</td>
</tr>
<tr>
<td>People</td>
<td>23,570,000</td>
<td>14,934,900</td>
<td>51,341,000</td>
<td>5,623,850</td>
<td>95,469,750</td>
</tr>
<tr>
<td>Poor livestock keepers</td>
<td>4,946,850</td>
<td>916,512</td>
<td>2,088,880</td>
<td>203,899</td>
<td>8,156,141</td>
</tr>
</tbody>
</table>

Northern Africa is characterised by:
- Poor livestock-keepers 8% of the population. Lowest in Africa.
- Relatively high urbanisation and higher development indices.
- Sheep by far the most important species, followed by goats and then cattle. Only region where sheep pre-dominate. Pigs rare.
- Pastoralism/agro-pastoralism the most important system. Unlike elsewhere in Africa, mixed extensive is the least important.
- Intensive production important in many countries – especially poultry.
- Extremely high poultry numbers and density along the Nile allow avian influenza to persist and a risk factor for other poultry diseases.
Central and SE Africa

Countries: Central African Republic, Equatorial Guinea, Gabon, Congo, Congo DR, Angola, Namibia

(this region is included in West Africa for the review of endemic zoonoses)

Fig 5.6: Central and SE Africa region- Farming systems (Herrero et al 2009)

Table 5.5: Species population by farming systems in Central and SE Africa (Herrero et al 2009)

<table>
<thead>
<tr>
<th>Farming system</th>
<th>Agro-pastoral pastoral</th>
<th>Mixed extensive</th>
<th>Mixed intensifying</th>
<th>Others</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bovines</td>
<td>4,353,250</td>
<td>2,291,670</td>
<td>344,621</td>
<td>1,899,710</td>
<td>8,889,251</td>
</tr>
<tr>
<td>Goats</td>
<td>2,911,600</td>
<td>1,730,230</td>
<td>775,076</td>
<td>2,920,870</td>
<td>8,337,776</td>
</tr>
<tr>
<td>Sheep</td>
<td>2,826,090</td>
<td>678,948</td>
<td>212,043</td>
<td>625,979</td>
<td>4,343,060</td>
</tr>
<tr>
<td>Pigs</td>
<td>985,813</td>
<td>835,169</td>
<td>358,327</td>
<td>2,728,110</td>
<td>4,907,419</td>
</tr>
<tr>
<td>People</td>
<td>8,888,550</td>
<td>7,730,480</td>
<td>10,530,100</td>
<td>31,979,900</td>
<td>59,129,030</td>
</tr>
<tr>
<td>Poor livestock</td>
<td>3,657,390</td>
<td>2,660,500</td>
<td>2,891,810</td>
<td>9,223,170</td>
<td>18,432,870</td>
</tr>
</tbody>
</table>

Central and South Eastern Africa is characterised by:

- Poor livestock-keepers 31% of the population. Second highest after W Africa. Generally similar to W Africa in species, systems and zoonoses.
- Goats the most numerous, followed by cattle then sheep.
- Pastoralism/agro-pastoralism the most important system. Mixed extensive, and mixed intensifying both important.
- Pigs important, especially in Congo DR.
- Central African rain forests a hotspot for bio-diversity, human incursion, game meat utilisation and disease emergence.
- Insecurity problems persisting in Congo DR.
South Central Asia Region

Countries: Iran, Afghanistan, Pakistan, Nepal, India, Bhutan, Bangladesh (our review of endemic zoonoses included Sri Lanka)

Fig 5.7: South Central Asia region: Farming systems (Herrero et al 2009)

Table 5.6: Species population by farming system in SC Asia (Herrero et al 2009)

<table>
<thead>
<tr>
<th>Farming system</th>
<th>Agro-pastoral pastoral</th>
<th>Mixed extensive</th>
<th>Mixed intensifying</th>
<th>Others</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bovines</td>
<td>10,545,400</td>
<td>125,910,000</td>
<td>204,813,000</td>
<td>16,553,900</td>
<td>357,822,300</td>
</tr>
<tr>
<td>Goats</td>
<td>29,862,700</td>
<td>66,316,600</td>
<td>95,724,300</td>
<td>9,702,000</td>
<td>201,605,600</td>
</tr>
<tr>
<td>Sheep</td>
<td>37,978,700</td>
<td>40,641,100</td>
<td>39,511,400</td>
<td>5,621,940</td>
<td>123,753,140</td>
</tr>
<tr>
<td>Pigs</td>
<td>185,107</td>
<td>4,873,080</td>
<td>6,977,520</td>
<td>1,212,390</td>
<td>13,248,097</td>
</tr>
<tr>
<td>People</td>
<td>51,652,300</td>
<td>281,271,000</td>
<td>630,135,000</td>
<td>44,731,600</td>
<td>1,007,789,900</td>
</tr>
<tr>
<td>Poor livestock keepers</td>
<td>11,086,000</td>
<td>54,073,900</td>
<td>57,803,200</td>
<td>5,890,470</td>
<td>128,853,570</td>
</tr>
</tbody>
</table>

South Central Asia is characterised by:
- Poor livestock-keepers 13% of the population. A relatively lower proportion or the population are poor livestock keepers than in West and East Africa but in absolute numbers about the same.
- Mixed intensifying systems most important followed by mixed extensive.
- Cattle are the most important species followed by goats and then sheep.
- Important dairy (mainly buffalo) and draft sectors but production low. Milk-borne zoonoses very important but transmission more from contact than consumption.
- Pigs are localised mainly in the North East of India but here they are very important.
- Relatively stronger government services than much of SSA.
- Insecurity problems persisting in several countries.
South East Asia region
Countries: Myanmar, Thailand, Laos, Vietnam, Cambodia, Malaysia, Indonesia, Brunei, Philippines, Papua New Guinea

Fig 5.8: South East Asia Region: Farming systems (Herrero et al 2009)

Table 5.7: Species population by farming system (Herrero et al 2009)

<table>
<thead>
<tr>
<th>Farming system</th>
<th>Agro-pastoral pastoral</th>
<th>Mixed extensive</th>
<th>Mixed intensifying</th>
<th>Others</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Goats</td>
<td>209,463</td>
<td>2,658,470</td>
<td>9,412,130</td>
<td>2,107,110</td>
<td>14,387,173</td>
</tr>
<tr>
<td>Sheep</td>
<td>675</td>
<td>616,862</td>
<td>6,016,720</td>
<td>226,156</td>
<td>6,860,413</td>
</tr>
<tr>
<td>Pigs</td>
<td>584,026</td>
<td>4,160,260</td>
<td>10,307,800</td>
<td>3,275,040</td>
<td>18,327,126</td>
</tr>
<tr>
<td>People</td>
<td>1,678,350</td>
<td>11,542,900</td>
<td>28,397,400</td>
<td>9,013,500</td>
<td>50,632,150</td>
</tr>
<tr>
<td>Poor livestock keepers</td>
<td>654,932</td>
<td>7,726,660</td>
<td>11,278,000</td>
<td>4,160,070</td>
<td>23,819,662</td>
</tr>
</tbody>
</table>

South East Asia is characterised by:
- Poor livestock-keepers high proportion of overall population but concentrated in Indonesia, Vietnam and the Philippines. China has high numbers but poverty decreasing rapidly (not included in this map).
- Mixed intensifying systems most important followed by mixed extensive systems.
- High urbanisation, high demand for animal source foods, stabilising populations.
- Pockets of deprivation and high vulnerability to zoonoses: hill tribes in Thailand and Vietnam, Papua New Guinea and Timor Leste.
- Pigs are the most important followed by goats then sheep. Only region where pigs predominate.
- Cattle are few in number but growing rapidly in some countries (especially Muslim).
- High zoonoses burden in Viet Nam, Myanmar, Philippines and Indonesia.
- Very high density of monogastrics, poor biosecurity, wildlife interfaces, backyard close to intensive systems all favour disease emergence and persistence.
Chapter 6: Conclusions

This final short chapter pulls together some of the more important points and conclusions around zoonoses, poverty, poor livestock keepers, emerging markets and livestock system changes.

Objectives of controlling zoonoses
The review distinguishes between different categories of zoonoses and suggests that endemic zoonoses are of most concern where the objective is lowering the burden of human disease and increasing the productivity and profitability of livestock for poor people. Among the most important and most neglected of endemic zoonoses are food-borne zoonoses. Outbreak zoonoses are of concern when there is an objective of reducing vulnerability of neglected populations. Emerging zoonoses are of concern when the object is foresight, and understanding disease emergence in order to try and avert pandemics of major impact. Because a small number of zoonoses are responsible for the majority of human and animal burden, targeting these zoonoses is likely to be an effective use of scarce resources.

Lack of evidence
The report draws attention to the lack of evidence on zoonoses presence, prevalence, drivers and impact. There are obviously major problems around disease reporting systems in developing countries, despite considerable support at the time of the avian influenza pandemic. A question is whether to invest more in existing systems or to explore alternative ways to generate evidence about disease? We suggest the latter. Our report revealed showed literature is one of the best ways of understanding what diseases are present and their impact. Moreover, valuable information exists in the grey literature, which is not currently easily available. However, more and better information is needed which can only be obtained through field surveys. Recent advances in technology (bio-repositories, genomics, e-technologies, etc) offer opportunities for radically improving our understanding of zoonoses epidemiology and control.

Hotspots of zoonoses, poverty and emerging markets
An underlying hypothesis of the report was that hot spots for zoonoses and poverty exist, and that targeting these hotspots has good prospects for alleviating health burdens while improving livelihoods. The study confirmed that a relatively small number of countries have a disproportionate share of poor livestock keepers and zoonoses burden (notably India, Ethiopia, and Nigeria). However, the association between zoonoses as a barrier to emerging markets for smallholders was less obvious (because countries with rapidly evolving markets tend to have fewer poor livestock keepers and better control of human disease). The relation between poverty and livestock keeping with emerging zoonotic events was not obvious, possibly because of the unpredictable nature of disease emergence, relatively poor detection and a possible relation between emergence and intensive livestock-keeping (associated with rich countries). We conclude, controlling zoonoses could substantially reduce the human disease burden and support the livelihoods of poor farmers, but the benefits in terms of increasing access to emerging markets require further research.

Opportunities
We identified gaps and opportunities for research to reduce the burden of disease for the zoonoses and regions in the report. These include: better understanding of the implications for intensification and emerging markets on zoonoses; models for zoonoses control in emerging markets; ecosystem models for management of zoonoses with a wildlife interface; improvement of surveillance for existing and new diseases; understanding the impacts multiple burdens of zoonoses in order to better allocate resources; technologies and innovation for detection, diagnosis, prevention, treatment and response.
References


Bourn, D and Wint, W. 1994. Livestock, land use and agricultural intensification in sub-Saharan Africa. Pastoral development network discussion paper, ODI.


Cardoen, S., Van Huffel, X. et al., 2009. Foodborne Pathogens and Disease, 6(9): 1083-1096.


EFSA, 2010, Scientific Opinion on Q fever, European Food Safety Authority Journal, 8 (5): 1595

ENHanCE, undated. Quantitative and Qualitative Approaches to the Prioritisation of Diseases, Position paper 1, National Center for Zoonoses Research, University of Liverpool UK

no. 5, CGIAR Research Program on Climate Change, Agriculture and Food Security (CCAFS). Copenhagen, Denmark. Available online at: www.ccafs.cgiar.org


Grace, D., McDermott J., 2011. Livestock epidemics and disasters. In Kelman et al., ed Handbook of Hazards and Disaster Risk Reduction, Routledge,


doi:10.1371/journal.pone.0013965


IAEA, 2002, Characteristics and Parameters of Family Poultry Production in Africa.
International Atomic Energy Authority, Vienna


Mangen, M.J., Otte, J., Pfeiffer, D., Chilonda, P., 2002, Bovine brucellosis in Sub-Saharan Africa: Estimation of sero-prevalence and impact on meat and milk offtake potential, Food and Agriculture Organisation of the United nations, Rome


Perry, B., Grace, D., 2009. The impacts of livestock diseases and their control on growth and development processes that are pro-poor. Philosophical Transactions of the Royal Society B, 364, 2643-2655


## ANNEX ONE – PRIORITY SATION OF ZOONOSES

<table>
<thead>
<tr>
<th>Disease</th>
<th>Expert opinion</th>
<th>Deaths - annual</th>
<th>Affected</th>
<th>Impact livestock</th>
<th>Farm intervention</th>
<th>other</th>
<th>Score</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cysticercosis</td>
<td>OIE, Perry</td>
<td>50000</td>
<td>50000000</td>
<td>High</td>
<td></td>
<td>1</td>
<td>5</td>
</tr>
<tr>
<td>Gastrointestinal (zoonotic)</td>
<td>GBD, Rosetta</td>
<td>1466666.6-67</td>
<td>233333333</td>
<td>High</td>
<td></td>
<td>1</td>
<td>5</td>
</tr>
<tr>
<td>Leptospirosis</td>
<td>OIE, Perry</td>
<td>123000</td>
<td>1700000</td>
<td>High</td>
<td></td>
<td>1</td>
<td>5</td>
</tr>
<tr>
<td>Hepatitis E</td>
<td></td>
<td>300000</td>
<td>14000000</td>
<td>Low</td>
<td></td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td>Tuberculosis (zoonotic)</td>
<td>GBD, OIE, Perry</td>
<td>100000</td>
<td>554500</td>
<td>High</td>
<td></td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td>Rabies</td>
<td>OIE, Perry</td>
<td>55000</td>
<td>70000</td>
<td>Medium</td>
<td></td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td>Leishmaniasis</td>
<td>OIE, GBD, Rosetta</td>
<td>47000</td>
<td>2000000</td>
<td>Low</td>
<td></td>
<td>1</td>
<td>4</td>
</tr>
<tr>
<td>Brucellosis</td>
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<td>15000</td>
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<td>Chagas</td>
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<td>Hanta virus</td>
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<td>Avian influenza virus</td>
<td>OIE</td>
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<td>145</td>
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<td>BSE</td>
<td>OIE</td>
<td>182</td>
<td>188</td>
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<td>3</td>
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<td>Buffalo pox</td>
<td>Perry</td>
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<td>common?</td>
<td>High</td>
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<td>Chlamyphila psittaci</td>
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<td>2250</td>
<td>22000</td>
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<td>Japanese encephalitis</td>
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<td>Rift valley fever</td>
<td>OIE, Perry</td>
<td>45</td>
<td>150</td>
<td>Medium</td>
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<td>Mange</td>
<td>Perry</td>
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<td>common?</td>
<td>High</td>
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<td>Lassa fever</td>
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<td>Lyme</td>
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<td>Pneumonia zoonotic</td>
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<td>GBD, Perry</td>
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<td>Low</td>
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<td>OIE, Perry</td>
<td>2000</td>
<td>100000</td>
<td>Low</td>
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<td>Brucella melitensis</td>
<td>OIE, Perry</td>
<td>..</td>
<td>..</td>
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<td>Brucella suis</td>
<td>OIE, Perry</td>
<td>..</td>
<td>..</td>
<td>High</td>
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<td>Enzootic abortion of ewes (ovine chlamydiosis)</td>
<td>OIE</td>
<td>negligible</td>
<td>rare</td>
<td>High</td>
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<td>OIE</td>
<td>negligible</td>
<td>low</td>
<td>High</td>
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<td>Nipah</td>
<td>Perry</td>
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<td>Medium</td>
<td></td>
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<td>2</td>
</tr>
<tr>
<td>Orf</td>
<td>Perry</td>
<td>negligible</td>
<td>common?</td>
<td>Medium</td>
<td></td>
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<td>2</td>
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<td>OIE</td>
<td>Unknown</td>
<td>Medium</td>
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<tr>
<td>Disease</td>
<td>Source</td>
<td>Incidence</td>
<td>Prevalence</td>
<td>Risk</td>
<td>Control Measure</td>
<td>Description</td>
<td></td>
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<td>----------------------------------------------</td>
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<td>Toxocara vitulorum</td>
<td>Perry</td>
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<td>unknown</td>
<td>High</td>
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<td>High</td>
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<td>16</td>
<td>High</td>
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<td>OIE</td>
<td>165</td>
<td>5000</td>
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<td>Kyasanur forrest</td>
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<td>20</td>
<td>500</td>
<td>Low</td>
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<td>Marburg virus</td>
<td></td>
<td>9</td>
<td>11</td>
<td>Low</td>
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<td>Pasteurella multocida</td>
<td>OIE</td>
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<td>High</td>
<td></td>
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<tr>
<td>West Nile fever</td>
<td>OIE</td>
<td>100</td>
<td>100000</td>
<td>Low</td>
<td>1</td>
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<td>Equine encephalomyelitis (Eastern)</td>
<td>OIE</td>
<td>2</td>
<td>6</td>
<td>Low</td>
<td>1</td>
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<tr>
<td>Equine encephalomyelitis (Western)</td>
<td>OIE</td>
<td>low</td>
<td>Low</td>
<td></td>
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<td>Newcastle disease</td>
<td>OIE</td>
<td>Negligible</td>
<td>Low</td>
<td>Medium</td>
<td>1</td>
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<tr>
<td>Burkholderia mallei</td>
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<td>negligible</td>
<td>Low</td>
<td>Medium</td>
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<tr>
<td>Clostridium botulinum</td>
<td>Perry</td>
<td>100</td>
<td>1000</td>
<td>Low</td>
<td>0</td>
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<tr>
<td>New world screwworm (Cochliomyia hominivorax)</td>
<td>OIE</td>
<td>negligible</td>
<td>Low</td>
<td>Medium</td>
<td>0</td>
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<tr>
<td>Old world screwworm (Chrysomya bezziana)</td>
<td>OIE</td>
<td>negligible</td>
<td>Low</td>
<td>Medium</td>
<td>0</td>
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<tr>
<td>Tick borne encephalities</td>
<td></td>
<td>300</td>
<td>15000</td>
<td>Medium</td>
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<td>Tularaemia</td>
<td>OIE</td>
<td>1000</td>
<td>50000</td>
<td>Low</td>
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Agunloye 2002. Leptospiral Agglutinating antibodies in sheep and goats in South West Nigeria. Israel Journal of Veterinary Medicine, 57, 2,
Akinbami et al. 2010. Seroprevalence of Toxoplasma gondii antibodies amongst pregnant women at the Lagos State University Teaching Hospital, Nigeria. Niger postgrad Med 17(2):164-7
Bachou et al 2006. Bacteraemia among severely malnourished children infected and uninfected with the human immunodeficiency virus-1 in Kampala, Uganda. BMC Infect Dis. 8:160
Bedard et al 1993. A prevalence study on bovine tuberculosis and brucellosis in Malawi. Preventive Veterinary Medicine 16(3), 193-205


Bhatti 2008
Bhatti 2009
Bhatti 2010


Boa et al. 2006. Epidemiological survey of swine cysticercosis using ante-mortem and post-mortem examination tests in the southern highlands of Tanzania. Veterinary Parasitology 139, 249–255


Boonsilp et al. 2011. Molecular detection and speciation of pathogenic Leptospira spp. in blood from patients with culture-negative leptospirosis. BMC Infect Dis 11(1):338


Bouzidi et al. 2012. Salmonella contamination of laying-hen flocks in two regions of Algeria. Food Research International 45 (2) 897–904
Cadmus et al. 2010. Mycobacterium bovis, but also M. africanum present in raw milk of pastoral cattle in North-central Nigeria. Trop Anim Health Prod, 42:1047–1048
Caron M, Kazanji M 2008. Hepatitis E virus is highly prevalent among pregnant women in Gabon, central Africa, with different patterns between rural and urban areas. Virol J;5:158
Cleaveland et al. 2007. Mycobacterium bovis in rural Tanzania: risk factors for infection in human and cattle populations. Tuberculosis (Edinb);87(1):30-43.


Cooper et al 2005. Identification of genotype 3 hepatitis E virus (HEV) in serum and fecal samples from pigs in Thailand and Mexico, where genotype 1 and 2 HEV strains are prevalent in the respective human populations. JOURNAL OF CLINICAL MICROBIOLOGY. 1684–1688


Downie, K., PHD thesis


Downie-Ngini,, K 2011. PHD thesis


Feresu 1982. Serological survey of leptospiral antibodies in cattle in Zimbabwe


Grace, 2006. Epidemiology and Control of Cattle Trypanosomosis in Villages under Risk of Trypanocide Resistance in West Africa. Institute for Parasitology and Tropical Veterinary Medicine. Freie Universitat Berlin, Germany. PHD thesis
Halliday et al 2011
http://cid.oxfordjournals.org/content/30/1/214.full.pdf+html
Idigbe
Ismail et al 2006
Ito et al, 2002
Jain


Kaud et al 2010. Epidemiology of Brucellosis among farm animals. Nature and Science, 8(5)

Kawaguchi et al 2008

Kazwala


Kobbe et al., 2009
Koeck 2002. [Epidemiology of resistance to antituberculosis drugs in Mycobacterium tuberculosis complex strains isolated from adenopathies in Djibouti. Prospective study carried out in 1999]. Med Trop (Mars); 62(1):70-2


Madiannikov et al. 2010. 20. Coxiella burnetii, the causal agent of Q Fever in cattle, sheep and goats in Bogor and Bali. Journal of Veterinary, 180-187


Mahatmi et al. DETECTION OF COXELLA BURNETII, THE CAUSAL AGENT OF Q FEVER IN CATTLE, SHEEP AND GOATS IN BOGOR AND BALI. Journal of Veterinary, 180-187


Makita et al. 2011. Herd prevalence of bovine brucellosis and analysis of risk factors in cattle in urban and peri-urban areas of the Kampa la economic zone, Uganda. B MC Veterinary Research, 7, 60


Mathur et al. 2001. Sero-epidemiology of hepatitis E virus (HEV) in urban and rural children of North India
McDermott and Arimi 2002. Brucellosis in sub-Saharan Africa: Epidemiology, control and impact. Veterinary Microbiology 90, 1-4, 111-134
Meenakshisundaram et al. 2010.SERO-PREVALENCE OF LEPTOSPIROSIS IN SMALL RUMINANTS IN VIRUDHUNAGAR DISTRICT OF TAMIL NADU.Tamilnadu J. Veterinary & Animal Sciences 6 (3) 136-137.
Mfinanga
101


Mohamed et al 2009. Phenotypic and Molecular Typing of Tuberculous and Nontuberculous Mycobacterium Species from Slaughtered Pigs in Egypt. Journal of Veterinary Diagnostic Investigation vol. 21 no. 1 48-52


Mtové et al 2010. Invasive Salmonellosis among Children Admitted to a Rural Tanzanian Hospital and a Comparison with Previous Studies. PLoS Oneb 16;5(2


Muller et al. 2008. Molecular characterization of Mycobacterium bovis isolated from cattle slaughtered at the Bamako abattoir in Mali. BMC Veterinary Research, 2008


Nawaz


Nguyen et al 2002. Result of survey on taeniasis and cysticercosis humans and pigs in Bac Ninh and Bac Kan provinces.


Niemann et al 2002. Mycobacterium africanum subtype II is associated with two distinct genotypes and is a major cause of human tuberculosis in Kampala, Uganda. Journal of Clinical Microbiology Vol. 40 No. 9 pp. 3398-3405


Nilleri et al 1989. The focus of human African trypanosomosis in Moissala (Chad): prospective study of 16 villages using the direct card agglutination test (Testryp CATT) and the ion exchange mini-column (mAECT)]. Med Trop 49(3), 253


Njitchouang et al 2010. Analysis of the domestic animal reservoir at a micro-geographical scale, the Fontem sleeping sickness focus (South-West Cameroon) Journal of Cell and Animal Biology, 4(5), 73-80


Nonga and Muhairwa 2010. Prevalence and antibiotic susceptibility of thermophilic Campylobacter isolates from free range domestic duck (Cairina moschata) in Morogoro municipality, Tanzania. Trop Anim Health Prod;42(2):165-72
Nsengiyumwa et al. 2003. Cysticercosis as a Major Risk Factor for Epilepsy in Burundi, East Africa. Epilepsia, 44(7):950–955,
Nsengwa via Downie
Nzano et al. 2010. Multiple contaminations of chickens with Campylobacter, Escherichia coli and Salmonella in Yaounde (Cameroon). Journal of infection in developing countries http://scholar.qsensei.com/content/1ny333
Ogunsanmi et al. 2000. EFFECTS OF MANAGEMENT, SEASON, VEGETATION ZONE AND BREED ON THE PREVALENCE OF BOVINE TRYPANOSOMOSIS IN SOUTHWESTERN NIGERIA
Ohaeri, 2010. Prevalence of Trypanosomosis in Ruminants in Parts of Abia State, Nigeria
Olaya et al. 1999. 120. Prevalence of antibodies against spotted fever, murine typhus, and Q fever Rickettsiae in humans living in Zambia. American Journal of Tropical Medicine and Hygiene 1999 Vol. 61 No. 1 pp. 70-72
Oloya et al. 2006. Responses to tuberculin among Zebu cattle in the transhumance regions of Karamoja and Nakasongola district of Uganda. Trop Anim Health Prod 38:275–283
Olulofemi et al. 2008. PREVALENCE OF TSETSE FLY AND BOVINE TRYPANOSOMOSIS IN THE BIOLOGICAL CONTROL OF TSETSE FLY PROJECT (BICOT) WITHIN LAFIALOCAL GOVERNMENT AREA OF NASARAWA STATE, NIGERIA
Pillet et al. The challenge of controlling human African trypanosomosis in a remote and unstable area of the Democratic Republic of Congo
PREVALENCE OF HUMAN Taenia solium CYSTICERCOSIS IN MAYO-DANAY DIVISION (CAMEROON)
Raghunath et al 1993. Isolation of Campylobacter from human and other sources in Bombay
Saleh et al. 2010
Samad et al. 1997. Sero-epidemiological studies on Toxoplasma gondii infection in man and animals in Bangladesh. The Southeast Asian journal of tropical medicine and public health, 28, Issue: 2, Pages: 339-343
Samdi et al. 2010. Periodic Variation in Trypanosoma Infection Rates in Trade Small Ruminants at Slaughter in Kaduna Central Abattoir
Seck et al. 2010. The prevalence of African animal trypanosomoses and tsetse presence in Western Senegal
Selvaraj et al. 2011
Shan
Sharma et al 1979. Sero-epidemiologic investigations on brucellosis in the states of Uttar Pradesh (U.P.) and Delhi (India).
Sharoufi et al 2009. Molecular characterization of Mycobacterium bovis strains isolated from cattle slaughtered at two abattoirs in Algeria. BMC Veterinary Research 2009, 5:4
Simo et al 2006. High prevalence of Trypanosoma brucei gambiense group 1 in pigs from the Fontem sleeping sickness focus in Cameroon. Vet Parasitol. 139(1-3):57-66
Somers et al 2006. Taenia solium taeniasis and cysticercosis in three communities in North Vietnam. Tropical Medicine & International Health, 11, 1, 65-72
Sovya, 2005
Stoszek et al 2006. 11. High prevalence of hepatitis E antibodies in pregnant Egyptian women. Transactions of the Royal Society of Tropical Medicine and Hygiene 100 No. 2 pp. 95-101
Subharr et al
Subhar et al, 2001


Tawewat Deemagarn


Thai et al., 2006

Theis et al., 1994


Van Den et al 2006. An update of the bovine trypanosomosis situation at the edge of Hluhiwe-Imfolozi Park, Kwazulu-Natal Province, South Africa


Vanderick and Mbornyigabo 1972 (quoted by Zoli et al 2003)


Vavativithya et al 1990. 34. Importance of salmonellae and Campylobacter jejuni in the etiology of diarrheal disease among children less than 5 years of age in a community in Bangkok, Thailand.

Vekemans et al 1999. Potential source of human exposure to Mycobacterium bovis in
Victoriano et al 2009. Leptospirosis in the Asia Pacific region. BMC Infectious Diseases 9: 147
Waiswa et al 2003. Domestic animals as reservoirs for sleeping sickness in three endemic foci in South-eastern Uganda. Annals of Tropical Medicine and Parasitology, 97(2), 149-155(7)
Walín et al 2008
Wanyangu et al 1987
Woldemariam et al 2006. Prevalence and distribution of Salmonella in apparently healthy slaughtered sheep and goats in Debre Zeit, Ethiopia
Xu et al 2010. Seroprevalence of Cysticercosis in Children and Young Adults Living in a Helminth Endemic Community in Leyte, the Philippines. Journal of Tropical Medicine Volume 2010 (2010), Article ID 603174, 6 pages
Yadav and Sethi 1979. Sero-epidemiological studies on coxiellosis in animals and man in the state of Uttar Pradesh and Delhi (India). Int J Zoonoses. 6(2):67-77
Yodnopaklow, P., Mahuntussangapong, A., 2000, Trop Med Int Health t 250-255


Nguyễn Văn Dương, Phan Trường Chinh, and Lê Hữu Nghị (1997). "Results of investigating some epidemiology characteristics of Leptospirosis of pig raised in Nghe An, Quang Nam, Quang Ngai and Daklak." Journal of Industrial and Rural Food 8: 352-353. (in Vietnamese)


ANNEX 4 REFERENCES FOR IN-COUNTRY LITERATURE REVIEW (VIET NAM)


Ngan, N.T.T., Nguyen Ngoc "A study on the occurrence of Leptospirosis in the goat population in Northern Vietnam."


Truong Quang, Truong Ha Thai (2007). Changes of the enteric bacterial fauna and role of Salmonella in the syndrome of diarrhea in piglets from 2 to 4 months old. Journal of veterinary science 2007, XIV(6), 52-57. (in Vietnamese)


Nguyen Canh Dung (2011). The role of E.coli, Salmonella in causing diarrhea in pig in Lam Dong province. Veterinary technology science magazine, vol. XVIII, No 1, p.56-64. (in Vietnamese)


Vo Bich Thuy, Tran Thi Hanh (2002). The result of determining of Salmonella sp biochemical characteristic isolated from food derived from animal in Hanoi. Veterinary technology science magazine. No. 4, p:19-24. (in Vietnamese)


Nguyen Van Suu (2005). The result of determining the biochemical characteristics and the factors cause disease of Salmonella bacterium isolated from calves, buffalo calves suffering diarrhea in North. p: 33-40. (in Vietnamese)


Tran Quang Dien. The suitation of Salmonella gallinarum-pullorumtreen infection on chicken in the North p: 39-41. (in Vietnamese)


Tong Vu Thang. The study of relationship between Salmonella pollution on mixed food, litter and the rate of Salmonella infection on eggs in six raising chicken farms in Ho Chi Minh. Veterinary technology science magazine, 2008, vol. XV, No 1, p:62-68. (in Vietnamese)

Luu Quynh Huong, Tran Thi Hanh. The result of determining the type of Salmonella on retail chicken in Hanoi. Veterinary technology science magazine, 2006, vol. XIII, No 1, p:50-53. (in Vietnamese)

Dinh Nam Lam, Phan Ngoc Anh. First step of supervisi ng Salmonella infection on duck in Can Tho, p: 6-12. (in Vietnamese)

Do Trung Cu. The result of isolation and determining the factors causing paratyphoid diseases of Salmonella bacterium in pig in North, p: 10-18. (in Vietnamese)

Tran Thi Hanh, Dang Thi Thanh Son. The rate of Salmonella isolates S.typhimemrium, S.enteritidis in chicken in raising chicken farm in North. p: 27-34. (in Vietnamese)


Luu Quynh Huong, Tran Thi Hanh. The rate of prevalence of Salmonella bacterium in raw chicken meat in Hanoi. p: 50-54. (in Vietnamese)

Do Trung Cu, Tran Thi Hanh. Determining the factors causing diseases of Salmonella typhimurium isolated from pig diarrhea in North, p: 33-37. (in Vietnamese)


Truong Quang. Diseases caused by Salmonella infect on the technological index of Luong phuong domestic hybrid chicken. p: 20-25. (in Vietnamese)

To Lien Thu. The sultation of antibiotics resistance of Salmonella bacterium is E.coli isolated from pork and chicken meat in North, p: 29-35. (in Vietnamese)

Truong Quang, Tieu Quang An. Determining the rate of infection and isolates of Salmonella gallinarum pullorum in Luong phuong purebred chicken and domestic hybrid chicken. p: 15-19. (in Vietnamese)

Hoang Thi Phi Phuong, Tran Thi Hanh. Study the influence of E.coli and Salmonella in food on the weaning piglet. p:41-46. (in Vietnamese)

C. Le Bas, Tran Thi Hanh, Nguyen Tien Thanh. Analysis the epidemiology of Salmonella enteric on pork in the process of slaughter in Vietnam by serotyping and _i_n di tr__ng xung Veterinary technology science magazine, 2007, vol. XIV, no 6, p:33-45. (in Vietnamese)

Phung Quoc Chuong. The result of susceptibility to antibiotics of Salmonella bacterium isolated from pets in Daklak. p:47-53. (in Vietnamese)

Dao Thi Vi Hoa, Dao Xuan Vinh. Stability of Salmonella typhi ty2 using to produce the vaccine of Vi polysaccharide. Preventive medicine magazine, vol. XXIII, no 6(105), p:30-35. (in Vietnamese)


Nguyen Xuan Duong, Phan Luc, Pham Sy Lang, Nguyen Van Duc, Truong Van Dung (2007). Situation of intestinal cestode infection of ducks in some regions of Red River delta. Vet Sci Techn, 14, 6, 72-75 (in Vietnamese)


Phan Anh Tuan, Tran Thi Kim Dung (2010). Sero-infection rate and risk factors for infection with cysticercus cellulosae. J Malaria Parasit Dis Control, 1, 59-64 (in Vietnamese)


Luong Van Huan (1998). Parasitic worms in pigs in some southern provinces and control measures. Sci Econ Magazine, 1, 5-7 (in Vietnamese)


Doan Thi Hanh Nguyen (2000). Epidemiologic characteristics, clinical and result of treatment in patients infect cysticercosis, Scientific conference on technology and environment, 262-266 (in Vietnamese)

Nguyen Van Doanh, Le Quang Cuong (2005). A study of some epidemiologic characteristics of epilepsy in Thai Bao, a community where cysticercosis is endemic. Journal of practical medicine, 12, 45-47 (in Vietnamese)


Nguyen Van Chap, Hoang Ky, Nghiem Quoc Hung, Kieu Duc Hung (1999). Base on 20 patients were diagnosed of cerebral cysticercosis by CT scanner (Nhận 20 b

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ANNEX 5 Summary details of the 30 new zoonotic EID events collected that were mapped (greater spatial resolution than country-level) with major references.

<table>
<thead>
<tr>
<th>Pathogen</th>
<th>Type</th>
<th>Location</th>
<th>Year</th>
<th>Refs</th>
</tr>
</thead>
<tbody>
<tr>
<td>unnamed <em>Brucella</em> spp.</td>
<td>bacteria</td>
<td>Lima, Peru</td>
<td>1985</td>
<td>[1]</td>
</tr>
<tr>
<td><em>Ngari virus</em></td>
<td>virus</td>
<td>Kassala, Sudan</td>
<td>1988</td>
<td>[2,3]</td>
</tr>
<tr>
<td><em>Baboon cytomegalovirus</em></td>
<td>virus</td>
<td>Pittsburgh, Pennsylvania, USA</td>
<td>1992</td>
<td>[4,5]</td>
</tr>
<tr>
<td><em>Castelo dos Sonhos virus</em></td>
<td>virus</td>
<td>Castelo dos Sonhos, Brazil</td>
<td>1995</td>
<td>[6]</td>
</tr>
<tr>
<td><em>Araraquara virus</em></td>
<td>virus</td>
<td>Araraquara, Brazil</td>
<td>1996</td>
<td>[6,7]</td>
</tr>
<tr>
<td><em>Babesia venatorum</em></td>
<td>protozoa</td>
<td>Romagna, Italy</td>
<td>1998</td>
<td>[8,9]</td>
</tr>
<tr>
<td><em>Iquitos virus</em></td>
<td>virus</td>
<td>Iquitos, Peru</td>
<td>1999</td>
<td>[10]</td>
</tr>
<tr>
<td><em>Anajatuba virus</em></td>
<td>virus</td>
<td>Anajatuba, Brazil</td>
<td>2000</td>
<td>[12,13]</td>
</tr>
<tr>
<td><em>Clostridium difficile 027/BI/NAP</em> (gatifloxacin &amp; moxiflacin resistant)</td>
<td>bacteria</td>
<td>Pennsylvania state, USA</td>
<td>2001</td>
<td>[14]</td>
</tr>
<tr>
<td><em>Rickettsia parkeri</em></td>
<td>bacteria</td>
<td>Tidewater, Virginia, USA</td>
<td>2002</td>
<td>[15,16]</td>
</tr>
<tr>
<td><em>Nam Dinh virus</em></td>
<td>virus</td>
<td>Nam Dinh province, Vietnam</td>
<td>2003</td>
<td>[17,18]</td>
</tr>
<tr>
<td><em>Chapare virus</em></td>
<td>virus</td>
<td>near Cochamba, Bolivia</td>
<td>2003</td>
<td>[19]</td>
</tr>
<tr>
<td><em>Juquitiba virus</em></td>
<td>virus</td>
<td>Juquitiba, Brazil</td>
<td>2003</td>
<td>[20,21]</td>
</tr>
<tr>
<td><em>Campylobacter jejuni SA clone</em> (tetracycline resistant)</td>
<td>bacteria</td>
<td>Vermont state, USA</td>
<td>2003</td>
<td>[22]</td>
</tr>
<tr>
<td><em>Bartonella melophagi</em></td>
<td>bacteria</td>
<td>Ohio state, USA</td>
<td>2004</td>
<td>[23]</td>
</tr>
<tr>
<td><em>Brucella inopinata</em></td>
<td>bacteria</td>
<td>Portland, Oregon, USA</td>
<td>2005</td>
<td>[24]</td>
</tr>
<tr>
<td><em>Bartonella alsatica</em></td>
<td>bacteria</td>
<td>Alsace region, France</td>
<td>2005</td>
<td>[25]</td>
</tr>
<tr>
<td><em>Human T-cell lymphotropic virus 3</em></td>
<td>virus</td>
<td>Southern Cameroon</td>
<td>2005</td>
<td>[26]</td>
</tr>
<tr>
<td><em>Human T-cell lymphotropic virus 4</em></td>
<td>virus</td>
<td>Southern Cameroon</td>
<td>2005</td>
<td>[26]</td>
</tr>
<tr>
<td><em>Melaka virus</em></td>
<td>virus</td>
<td>Melaka, Malaysia</td>
<td>2006</td>
<td>[27]</td>
</tr>
<tr>
<td><em>Kampar virus</em></td>
<td>virus</td>
<td>Kampar, Malaysia</td>
<td>2006</td>
<td>[28]</td>
</tr>
<tr>
<td><em>Severe fever with thrombocytopenia syndrome bunyavirus</em></td>
<td>virus</td>
<td>Dingyuan county, China</td>
<td>2006</td>
<td>[29,30]</td>
</tr>
<tr>
<td><em>Neoehrlichia mikurensis</em></td>
<td>bacteria</td>
<td>Middle Franconia, Germany</td>
<td>2007</td>
<td>[31,32]</td>
</tr>
<tr>
<td><em>Plasmodium falciparum</em> (artemisinin resistant)</td>
<td>protozoa</td>
<td>Pailin, Cambodia</td>
<td>2007</td>
<td>[33,34]</td>
</tr>
<tr>
<td><em>Dandenong virus</em></td>
<td>virus</td>
<td>Dandenong, Australia</td>
<td>2007</td>
<td>[35,36]</td>
</tr>
<tr>
<td><em>Crimean-Congo Hemorrhagic Fever virus AP92 (newly virulent)</em></td>
<td>virus</td>
<td>Istanbul, Turkey</td>
<td>2007</td>
<td>[37]</td>
</tr>
<tr>
<td><em>Bundibugyo ebolavirus</em></td>
<td>virus</td>
<td>Bundibugyo, Uganda</td>
<td>2007</td>
<td>[38,39]</td>
</tr>
<tr>
<td><em>Lujo virus</em></td>
<td>virus</td>
<td>Lusaka, Zambia</td>
<td>2008</td>
<td>[40,41]</td>
</tr>
<tr>
<td><em>Titi monkey adenovirus</em></td>
<td>virus</td>
<td>Davis, California, USA</td>
<td>2009</td>
<td>[42]</td>
</tr>
</tbody>
</table>
Sources


Figure 2. Global maps of zoonotic EID events at a one decimal-degree level, where size of circles denotes number of events in grid cell. Maps depict a) labelled new events collected in this update (n = 30), b) all combined events from Jones et al. 2008 and this update (n = 202), and stratified maps of only those events with c) wildlife hosts (n = 121), and d) non-wildlife hosts (n = 91). Events with both host types are included in both c) and d), and events with unknown hosts are included in neither. Majoor CJ, Magis-Escurra C, van Ingen J, Boeree MJ, van Soolingen D. Epidemiology of Mycobacterium bovis disease in humans, the Netherlands, 1993–1997. Emerg Infect Dis [serial on the Internet]. 2011 Mar [date cited]. http://dx.doi.org/10.3201/eid1703.101111