



Chaters, G.L. et al. (2019) Analysing livestock network data for infectious disease control: an argument for routine data collection in emerging economies. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 374(1776), 20180264.

There may be differences between this version and the published version. You are advised to consult the publisher's version if you wish to cite from it.

<http://eprints.gla.ac.uk/181834/>

Deposited on: 15 March 2019

Enlighten – Research publications by members of the University of Glasgow_
<http://eprints.gla.ac.uk>

1 **Analysing livestock network data for infectious diseases control: an argument for routine data**
2 **collection in emerging economies**

3

4 G.L. Chaters^{1,a}, P.C.D. Johnson^{1,a}, S. Cleaveland¹, J. Crispell², W.A. de Glanville¹, T. Doherty³, L.
5 Matthews¹, S. Mohr¹, O.M. Nyasebwa⁶, G. Rossi³, L.C.M. Salvador^{3,4,5}, E. Swai⁶, R.R.Kao^{3b}

6 ¹ Boyd Orr Centre for Population and Ecosystem Health, Institute of Biodiversity, Animal Health and
7 Comparative Medicine, University of Glasgow, Glasgow G12 8QQ, UK

8 ² School of Veterinary Medicine, University College Dublin, Ireland

9 ³ Royal (Dick) School of Veterinary Studies and Roslin Institute, University of Edinburgh, Easter Bush
10 Campus, Midlothian, Scotland, UK

11 ⁴ Department of Infectious Diseases, University of Georgia, Athens, Georgia, USA

12 ⁵ Institute of Bioinformatics, University of Georgia, Athens, USA

13 ⁶ Department of Veterinary Services, Ministry of Livestock and Fisheries, Tanzania;

14

15 **Abstract**

16 Livestock movements are an important mechanism of infectious disease transmission. Where these are
17 well recorded, network analysis tools have been used to successfully identify system properties,
18 highlight vulnerabilities to transmission and inform targeted surveillance and control. Here we highlight
19 the main uses of network properties in understanding livestock disease epidemiology, discuss statistical
20 approaches to infer network characteristics from biased or fragmented datasets. We use a “hurdle
21 model” approach that predicts (i) the probability of movement and (ii) the number of livestock moved
22 to generate synthetic ‘complete’ networks of movements between administrative wards, exploiting
23 routinely collected government movement permit data from northern Tanzania. We demonstrate that
24 this model captures a significant amount of the observed variation. Combining the cattle movement
25 network with a spatial between-ward contact layer we create a multiplex, over which we simulated the
26 spread of ‘fast’ ($R_0=3$) and ‘slow’ ($R_0=1.5$) pathogens, and assess the effects of random versus targeted
27 disease control interventions (vaccination and movement ban). The targeted interventions substantially
28 outperform those randomly implemented for both fast and slow pathogens. Our findings provide

^a These authors contributed equally to this work

^b Corresponding author: rowland.kao@ed.ac.uk

29 motivation to encourage routine collection and centralisation of movement data to construct
30 representative networks.

31

32 **1. Introduction**

33 The “static network” concept of a population as a set of “individuals” (nodes) with immutable contacts
34 (links) between them is now well-established in infectious disease modelling. The network
35 representation occurs naturally because the “individual” is typically well-defined (e.g. a person, animal,
36 city, herd, or farm) and the number of potentially infectious contacts per individual is usually few [1-
37 5]. While there are a few studies for human diseases that include comprehensive, explicit network data
38 [6], more frequently these are either generated indirectly (for example, using mobile phone data or
39 gravity models to predict commuter flow [7-10]), or are explicit but at small geographical scales [11,
40 12]. In contrast, in Great Britain (GB) cattle movement data have been recorded for individuals on a
41 daily basis for almost two decades [13]. This data richness has presented both challenges and
42 opportunities for the application of network analyses in infectious disease epidemiology [4, 5]. Similar
43 livestock data now exist in many other countries [14-20]. However, they remain rare in emergent
44 economies where disease burden is often high and zoonotic risk is more pronounced due to the high
45 proportion of people who live and work in close contact with livestock [21]. About one billion of the
46 world’s poorest people (earning < US\$2 per day) depend at least partially on livestock for their
47 livelihoods [22], making the trade of livestock and the freedom to move livestock to access natural
48 resources vital in many impoverished communities [23-25]. In many regions, such as Sub-Saharan
49 Africa, there are frequent but poorly recorded cross-border movements [26-28] and, when coupled with
50 poor within-country knowledge of livestock movements, this creates risks for international pathogen
51 transmission.

52

53 Though network analyses would be greatly aided by systems for comprehensive routine recording of
54 between-farm and market movement, as occurs in GB and elsewhere, in countries with developing
55 infrastructure collecting these data can be onerous and costly and requires well-evidenced justification.
56 Here, we provide an overview of the role of network analysis in epidemiology, paying particular

57 attention to the challenges of exploiting extensive but fragmented data. These insights are used to
58 analyse livestock movements in northern Tanzania, where there is a high burden of livestock disease
59 including zoonoses [29-35], no formal livestock traceability system implemented at a national level,
60 and limited resources for disease control. We demonstrate the utility of our network by identifying
61 nodes to target disease control and surveillance interventions, considering both fast and slowly
62 transmitting pathogens, and interrogate their efficacy through simulation, demonstrating substantial
63 potential benefits in reducing disease spread.

64

65 **2. Fundamental network concepts applied to livestock diseases**

66

67 **2.1 Centrality measures and transmission patterns**

68 Network centrality measures originated in social science [36], and are used to quantify the importance
69 of nodes and links in a network, with obvious applications to identifying disease risks [19, 37-41].
70 Common measures include degree centrality (the number of links associated with a node^a), betweenness
71 centrality (the number of times a node or link is traversed by the shortest paths between all other node
72 pairs), and eigenvector centrality (loosely, a measure of how connected a node is to well-connected
73 neighbours).^b Network centrality measures have been used to analyse livestock movement data from
74 many countries, with each using different types of data source [4, 17, 40-43]. One example showing the
75 relevance of all three of these centrality measures comes from the analysis of the costly [44] 2001 foot-
76 and-mouth disease (FMD) epidemic in GB. First, a small number of “cull ewes” were sold and
77 transported long distance across GB; these were responsible for seeding virus into many otherwise low
78 risk areas [45]. These seeding movements are a characteristic of “small world” network behaviour [1]
79 with the long-range movements acting as links with high betweenness centrality [45, 46]. Second,
80 Longtown auction market (the largest in GB) played a dominant role in spreading disease [47],
81 demonstrating the importance of high degree centrality. Third, since the epidemic, prohibition of direct

^a For directed networks like livestock movements, where transmission is overwhelmingly in the direction of the movement only, the geometric mean of in- and out-degree can be used.

^b See the supplementary information for a disease-relevant interpretation.

82 market-to-market livestock movements means that some farms now act as “middlemen” between
83 markets, representing a risk that could be effectively targeted to restrict disease spread [4, 48]. This
84 role, linking highly connected nodes, is a well-recognised feature of high eigenvector centrality.

85

86 **2.2. Network Dynamics**

87

88 In a static network, the infection pressure from a single individual is reduced over time as each daughter
89 infection ‘uses up’ the link it was infected over [49, 50]. Further, the components of the network (groups
90 of nodes which can reach each other) are well defined. In dynamic networks, links can shift between
91 individuals over time (rewiring), nodes can appear or disappear and the components of the network can
92 change in size and composition. Rewiring a link away from an infected individual has the potential to
93 expose another susceptible individual, thus increasing the probability of disease persistence [51, 52].

94 Link dynamics also greatly complicate measures of network structure. For example, for an *SIS* infection
95 process on a static network, where susceptible individuals (*S*) can become infected (*I*) and eventually
96 recover to susceptible again, the eigenvector centrality scores of the nodes of the network contact matrix
97 represents the expected proportion of time those nodes are infected over the long term^a. This is the case
98 so long as the probability of recovery before re-infection is high (e.g. if the density of infected nodes is
99 always low, or the recovery time is substantially shorter than the time between infected generations).

100 However, livestock movements vary daily, seasonally and from year-to-year. Contact patterns between
101 farms and therefore eigenvector centrality measures can change dramatically depending on the season
102 and stochastic progression of the epidemic. This influences epidemic spread [4, 13, 18, 51], an effect
103 also seen in human diseases [14, 53]. Individual variability in disease progression and severity will
104 also influence disease generation times and therefore what movements are likely to cause infection
105 spread. Thus, predictions of node importance and targeting can depend strongly both on the dynamic

^a For an irreducible positive definite matrix (e.g. a contact matrix where all nodes belong to a single strong component), the Perron-Frobenius theorem applies and the matrix is guaranteed to have a unique largest eigenvalue (and positive eigenvector). For directed networks, strong connectivity amongst all nodes is required (all nodes can reach each other reciprocally, i.e. are members of the same strong component). Where this is not the case, eigenvector centrality is not well-defined, and other network measures need to be considered (for example by using singular values).

106 properties of the network and the properties of the underlying disease, making the identification of
107 general principles for the targeting of control more challenging (e.g. [54]; also Supplementary
108 Information).

109

110 Livestock movements are also an example where the actual contact occurs episodically. Episodic
111 behaviour is a subject of considerable study in the network literature, especially where there are patterns
112 of concentrated bursts (“burstiness”) separated by long waiting periods [55-57]. While an infection
113 may itself cause episodic activity, it is most frequently studied as a property of the underlying network.
114 Episodic activity has been shown to slow an epidemic on simulated [58] and real networks [59] but can
115 also increase epidemic speed, for example, due to observed correlations between the topology of the
116 network and the frequency of episodic contacts [60]. Epidemic spread also depends on within-node
117 infection dynamics; in a simulated avian influenza outbreak, patterns of recorded vehicle movement
118 between farms could either slow or accelerate pathogen spread, depending on the disease parameters
119 and detection threshold at the farm level [61].

120

121 Infection events themselves can also change the network structure. If the perceived jeopardy is
122 sufficiently high, rumours of pathogen spread may change contact patterns [62, 63]. For livestock,
123 farmers may be inclined to sell infected animals due to their condition, or may be restricted from selling
124 animals until the farm is officially declared disease-free [64]. In human disease, modelling analyses that
125 included changes in the contact process over the course of the recent West African Ebola epidemic were
126 used to inform changes in policy [65], highlighting the relevance for detailed datasets on contact
127 patterns and their changes over time, both routinely and in response to an outbreak [66].

128

129 **2.3 The role of pathogen sequence data for relating transmission networks to livestock networks**

130

131 Although livestock movements tell us about potentially infectious contacts, the relationship between
132 these contacts and the transmission network of actual infectious contacts is only partially understood.
133 Duration of contact, heterogeneity in immune response, and environmental conditions are some of the

134 factors that could affect which livestock movements transmit infection. The growing availability of high
135 coverage pathogen sequence data provides an unprecedented opportunity to quantify this relationship
136 [67, 68]. A number of tools have been developed to estimate transmission from genetic data [69-78]
137 and new tools continue to be developed [69, 73, 79]. However there remain many challenges [80-84].
138 A key limitation is that pathogen evolution needs to occur on a similar or faster timescale to the disease
139 generation time in order to infer direction of transmission [80]. Considering larger epidemiological units
140 (e.g. farms rather than animals) can alleviate this problem, since the generation time will be
141 concomitantly longer [73, 74, 77]. Epidemiological information is still required to estimate transmission
142 from genetic data and contact network data is important when trying to identify the most likely
143 transmission events [85, 86], but there are few tools to formally integrate these [87]. Phylodynamic
144 approaches that leverage all available data could provide new insights into pathogen transmission and
145 result in more targeted and improved control interventions, but they must overcome the challenge of
146 appropriate weighting of the often biased and/or fragmented data. Nevertheless, even limited genetic
147 data integrated into transmission models can improve epidemiological insights [88] and in situations
148 where other data are fragmented or sparse, sequence data can greatly strengthen the understanding of
149 transmission and inform control.

150

151 **Section 3. Exploiting network properties**

152

153 **3.1. Evaluating system resilience**

154

155 Invasion of a livestock network by an infectious pathogen has the capacity to impair or destroy the
156 function of individual nodes, either by the direct impact on livestock, or by the restrictions resulting
157 from control efforts. The impact on network structure can be considerable, *in extremis* resulting in the
158 destruction of the network as a functioning entity. For infectious diseases, interventions such as
159 movement restrictions, culling or prolonged herd testing are all designed to reduce transmission, but
160 will also have varying degrees of impact on livestock movements and potentially impair the nodes role
161 in the network. Such changes have economic impact [89, 90] and, if sufficiently harmful, can result in

162 node removal and/or substantial long-term harm to the network. Resilience of a network typically
163 focuses on its ability to recover, retain the same structure, and adapt to maintain system functionality
164 when exposed to disturbances [91-93]. One approach to eliminate disease, such as during the 2001
165 FMD epidemic, is to disrupt the network by preventing trade for a period (link removal). These
166 movement restrictions, however, can result in excessive livestock welfare issues, welfare culls, and
167 significant long term industry damage [94]. Less disruptively, lasting adjustments (link rewiring) can
168 minimize the impact of highly influential nodes, whilst maintaining overall trade function. An example
169 of this is the implementation of high biosecurity and compartmentalisation in some poultry companies
170 to isolate themselves from disease incursion despite close physical proximity to infected farms,
171 allowing operations to continue in the face of national restrictions [95].

172

173 Minimising the number of affected nodes, or protecting particular ones, may be important for resilience.
174 In dynamic networks, slowing the rate at which contacts occur can slow the rate of pathogen spread and
175 maintain communication between nodes [4], improving the networks resilience. Conversely, reducing
176 contact rates can also increase pathogen spread [61]. Additional complications arise when considering
177 multiple layers of a network and multiple diseases that spread on it. Ultimately, targeting control
178 measures that consider the spread of multiple pathogens on a network could be more efficient and
179 robust. Additionally, prior to designing and imposing changes on a network, particularly in economies
180 where livelihoods are heavily dependent on a functioning livestock movement network, the network's
181 resilience to proposed changes should be assessed.

182

183 **3.2. Exploiting network data to improve surveillance**

184

185 The concepts of network resilience can be used to improve surveillance. Albert *et al.* showed the extent
186 to which different types of complex network can be resilient to breakdown (which makes disease
187 difficult to control) or vulnerable to breakdown (which makes disease easier to control) [96]. Nodes (or
188 links) can be removed from a network randomly or using targeted measures such as removing nodes
189 that are highly ranked by one or more centrality measure. In terms of surveillance, random and targeted

190 node removal can be compared to non-targeted and targeted surveillance [4]. Network analysis can thus
191 provide an analytical framework to predict which farms to test in targeted surveillance strategies and
192 estimate net gains in performance. While generic network analysis can be valuable [5], it can be made
193 more robust by an understanding of the characteristics of the real system [97] and the dynamics of the
194 considered pathogen [48]. Network analysis has been used to inform targeted surveillance strategies in
195 many livestock systems [43, 97-100], leading to considerable gains in surveillance efficiency [101,
196 102]. Analyses of GB livestock networks have identified highly connected premises with a high risk of
197 both becoming infected with and spreading disease [38], and have used simulations to show how
198 targeted surveillance could reduce the size of potential epidemics [4]. For Swedish cattle and pigs, a
199 bespoke metric was identified to consider the timing and sequence of possible incoming and outgoing
200 infection chains [14]. This metric was subsequently expanded to consider the size of the in- and out-
201 components and then used to analyse the German pig trade movements network to identify high-risk
202 farms [15]. Such data are not typically available in low resource settings; having such network
203 knowledge could enable the use of cost-efficient, network measure-targeted surveillance for disease
204 control, but needs justification for the additional cost and effort required.

205

206 **3.3. Multiplexes, multi-layer networks and multi-host pathogen systems**

207

208 Complex systems are inherently multi-dimensional, with components linked via a complex set of often
209 directed and weighted interactions, giving rise to diverse and unpredictable behaviours [103]. For
210 infectious diseases, these can arise when spread occurs by more than one mechanism (e.g. animal trade,
211 airborne, fomites, sharing a resource or insect vectors), resulting in a multiplex, or where transmission
212 occurs across more than one species, an example of a multi-layer network. Both can compromise
213 disease control [104], especially when there are biases in available data or ability to exert control [105].
214 The multiplex representation was first developed in the social sciences to represent different types of
215 inter-personal relationships [106]. It has since been used in a variety of contexts, including ecological
216 systems [107], air transport [108], behavioural biology [109], and epidemiology [110]. In one livestock
217 example, a study of a dairy system in northern Italy explicitly accounted for two independent

218 transmission routes: cattle and veterinarian movements. This study found that at the local scale
219 veterinarian movements explained the spread of *Mycobacterium avium* subspecies *paratuberculosis*
220 better than cattle movements and geographic distance failed to capture the impact of veterinarian visits
221 [111, 112]. This highlights a need to identify the potentially multiple transmission routes beyond
222 discrete livestock movements when collecting data to construct a livestock network that is
223 representative of a transmission network.

224

225 Many pathogens are multi-host and therefore the network multi-layer. This complication often has
226 severe implications for humans, livestock and wildlife [113]. Unfortunately most analytical frameworks
227 of resilience are unsuitable for multi-dimensional systems [114], and network resilience can be
228 influenced by interdependence with other networks [115]. Recent work using percolation theory to
229 study the vulnerability of a system of interdependent networks [116] shows the overlap between
230 network layers can improve network resilience and this makes diseases harder to eradicate [117]. By
231 disentangling system dynamics from system structure, network characteristics can be identified that
232 influence resilience [115]. A well-known exemplar is the transmission of *Mycobacterium bovis*, the
233 cause of bovine tuberculosis (bTB), between cows and European badgers (*Meles meles*), where the role
234 of different layers can be quantified by exploiting their spatial patterns (Figure S1) [64]. At finer
235 granularities, radio-collar data were used to quantify inter- and intra-species contacts for cattle and
236 badgers [118]; adding a layer of indirect contacts based on badger latrines locations to this network
237 showed better correspondence to badger-to-badger transmission patterns [119].

238

239 **4. Movement networks where there is limited resource for explicit traceability**

240 There are many examples where livestock movement data have facilitated the planning of disease
241 control and surveillance [42, 120, 121][17, 19, 122]. Conversely, an absence of movement information
242 can obstruct disease control [45, 123]. In settings where comprehensive tracing systems are absent, a
243 variety of methods have been used to quantify livestock movement patterns and construct movement
244 networks. These include the use of GPS collar data to describe mobility patterns of pastoral herds and

245 overlaps with wildlife areas [27, 43, 124, 125], household and market surveys [126], transport vehicle
246 records [127] and international movement permits [28, 128].

247

248 Movement permits are used in many countries to certify livestock health and/or to regulate movement
249 taxes, and have been used to quantify livestock flow and construct movement networks [128, 129]. The
250 often ephemeral and patchy nature of these records, due to poor archiving or non-compliance [130], can
251 result in substantial non-random “missingness” that is difficult to quantify. In these cases, movement
252 permits have been used in conjunction with household and/or market survey data to estimate the risk of
253 disease introduction and target surveillance and vaccination campaigns, also illustrating the importance
254 of a regional disease control approach [28, 122, 131, 132]. Such analyses have identified traders as key
255 targets for disease control [130], demonstrated the effects of cattle movement on regional disease
256 transmission [133], identified increased risks of bTB with increased herd introductions [41] and, with
257 serology data, identified the role of between-village cattle movements in transmitting Rift Valley fever
258 virus [134].

259

260 Biased network samples can make reconstruction of network characteristics difficult. This was
261 addressed in GB by extrapolating from a small biased network sample via statistical associations
262 between common factors in the network study and a national population survey [135].

263

264 Another approach to network construction is to impose an underlying model on observed population
265 densities. Specifically, if census data (populations and locations) are available or can be estimated,
266 gravity [136] and radiation [137] models provide two ways of creating network models of population
267 mobility. While there is ongoing research regarding their relative merits [138], they share the property
268 of describing movement in terms of relative population size and a measure of distance. Gravity models,
269 for example, describe the probability of a movement occurring in inverse proportion to spatial distance
270 from each hub.

271

272 **Section 5. Evaluating network-based control strategies for livestock movements in Tanzania.**

273 **Introduction to the study**

274 Tanzania provides an exemplar of a rapidly developing emerging economy. In northern Tanzania there
275 is a heavy reliance upon livestock for food, traction power, income, savings and social status.
276 Movements can be over long distances, often on foot, and occasionally over international boundaries
277 with multiple levels of market activity [26, 85, 139, 140]. The pathogen burden is often high, and this
278 impacts productivity, creates herd/flock instability and, in the case of zoonoses, directly affects human
279 health [30, 32, 33, 141-146]. In addition to protecting human health, reducing the burden of endemic
280 livestock pathogens to improve livestock health and productivity is recognised as a route away from
281 poverty and necessary to meet global food demands [23, 147-153]. Livestock sales are also a major
282 source of income in rural communities [154-156]. In addition to trade between markets, livestock can
283 be sold privately, borrowed or gifted between households and are regularly moved to access natural
284 resources [41, 157, 158]. A reduction in endemic livestock disease is therefore paramount to improving
285 livelihoods in such emerging economies.

286

287 Historically there has been no formal, centralised system for identifying and tracing the movement of
288 individual animals in Tanzania, however a paper movement permit certifying livestock health is
289 officially required whenever animals are traded, recording movements *from* markets, though not
290 movements *to* markets. These data are not digitised and the receipt books are stored at administrative
291 Zonal Veterinary Centres in Tanzania. The aims of this study were to: quantify cattle and small ruminant
292 movements in a large (97000 km²) area of northern Tanzania (Arusha, Manyara and Kilimanjaro
293 regions) using archived, routinely collected government movement permit data; infer livestock
294 movement networks; and build this information into livestock disease simulations to inform
295 surveillance and control.

296

297 **Methods**

298 Summary methods are presented here; for full details see the supplementary material.

299

300 Data source and transcription

301 Access was granted to archived government movement permit receipt books at the Northern Zonal
302 Veterinary Office, Arusha. Movement permit receipt books were selected for analysis from 2009, 2011,
303 2013 and 2015. Origin, destination, number of each species (cattle, sheep or goat) moved, and date were
304 manually entered into spreadsheets from 50% of the available permits (30,946 permits), of which
305 19,438 (63%) permits yielded complete data. Only cattle movements are analysed here.

306

307 Statistical Modelling

308 Cattle movements were aggregated temporally by month and spatially at the ward level, because origins
309 and destinations often could not be located at a finer scale. A ward is an administrative unit of mean
310 area 243 km² and mean human population of 12,000 across the 398 wards in the study regions [159].
311 We aimed to infer the inter-ward cattle movement network within the study area; movements to outside
312 the study area and within wards were excluded (local movements from markets are less likely to
313 generate a movement permit due to non-compliance). The resulting data set recorded the movement of
314 86,195 cattle from 98 origin wards to 239 destination wards over the 4 sampled years.

315

316 Due to the large number of non-randomly missing permits, it was not possible to use the movement
317 data directly. Instead, the network was inferred by statistical modelling of the observed movements.
318 First, to distinguish true from artefactual absence of movements (months where an origin ward sent out
319 no cattle) a zero-inflated negative binomial (ZINB) generalised linear model (GLM) was fitted to each
320 origin ward, so that in subsequent modelling steps movements would be imputed in place of false
321 zeroes. Next, inter-ward livestock movement was modelled using a hurdle model. The movement
322 between each pair of wards in a given month is represented by a two-step processes: the binary event
323 of any cattle being moved, modelled by a binomial generalised linear mixed-effects model (GLMM);
324 and the number of animals moved, modelled by a zero-truncated negative binomial (ZTNB) GLMM.
325 Each part of the hurdle model allowed movement to depend multiplicatively on the distance between
326 origin and destination wards and their “masses” (human and cattle population sizes), in addition to other
327 characteristics (Table S1). The combined models can therefore be viewed as a gravity model of the

328 livestock movement network. Unexplained spatial and temporal variation was modelled by fitting
329 random effects for origin and destination ward and for the 48 months.

330

331 Simulated networks

332 The fitted model was used to simulate monthly movements amongst the 398 wards for one year, with
333 the number of movements inflated twofold to account for using a 50% subsample of the data.

334

335 Network measures

336 The simulated data were used to create an observed year-aggregated, static, directed, weighted cattle
337 movement network. A spatial contact layer, connecting all adjacent wards, was added to the market
338 movements network as a simplified means of accounting for contacts and movements between wards
339 that are not represented by the movement permit data. Social network analysis was applied to the
340 resulting multiplex network to identify nodes with high in-degree, out-degree, betweenness and
341 eigenvector centrality where disease control interventions could be targeted.

342 Simulating disease outbreaks and control on the network

343 The spread of a 'fast' ($R_0 = 3$) and 'slow' ($R_0 = 1.5$) pathogen was simulated on the multiplex to assess
344 the effects of disease control interventions on the spread of pathogens with varying infectiousness [166].
345 This was achieved by running a stochastic *SIR* compartmental model within each ward. The total
346 number of cattle in the susceptible (*S*), infectious (*I*) and recovered (*R*) compartments was updated
347 daily, while cattle were moved monthly between wards. The two sources of simulated cattle movement
348 were long distance movements via the market network and short distance movements between adjacent
349 wards to account for unobserved local movements (for a full description see Supplementary
350 Information; an animation of a simulated fast epidemic is available as a supplementary file). Two types
351 of intervention were trialled: proactive vaccination of 70% of the cattle in a ward before the start of the
352 epidemic, and a reactive ban on cattle movements one month after the start of the epidemic. Vaccine
353 interventions were applied to all wards, or targeted at 20 (5%) of wards that were selected randomly,
354 based on their total cattle population size or based on their network centrality measures. The network
355 centrality measures used for targeting interventions were betweenness centrality, eigenvector centrality,

356 and geometric mean degree. The market movement ban was either implemented in all 111 wards that
357 generated outward cattle movements in the simulations and were therefore assumed to have a market,
358 or were targeted in a subset of 20 of these wards, the same number as in the targeted vaccination
359 interventions, and based on the same selection criteria.

360

361 **Results**

362 The two parts of the hurdle model explained a substantial proportion (binomial: 40%; ZTNB: 24%) of
363 the spatial and temporal variation in cattle movement, with movement being more probable over shorter
364 distances and into wards containing a secondary market, and the number of animals moved being most
365 strongly associated with the agro-ecological system of the origin wards and the presence of a primary
366 or secondary market in the origin or destination ward (Table S1; Figure S2). All variables were retained
367 in the hurdle model that was used to simulate the monthly cattle market movements.

368

369 Network and node measures

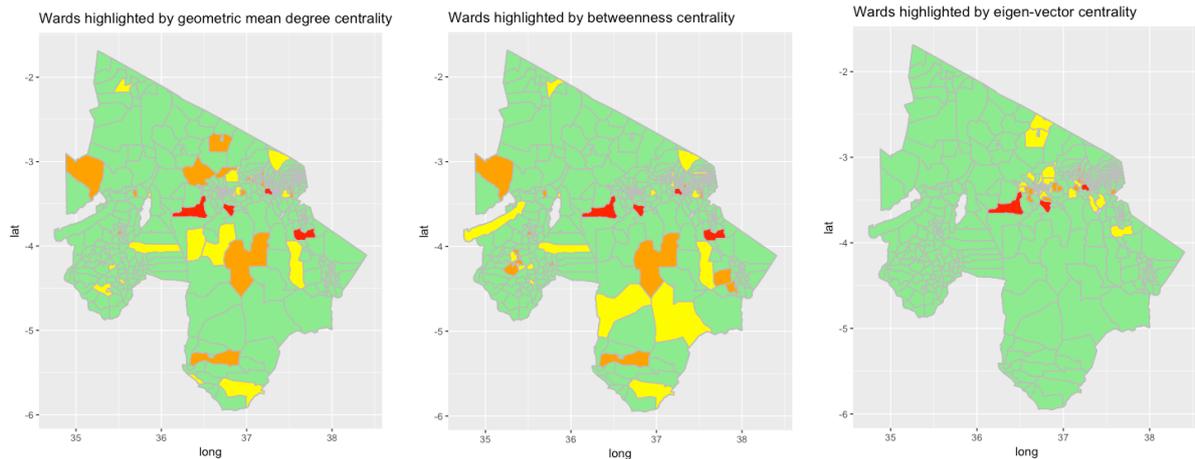
370 The multiplex network is fully strongly connected (all wards can be reached by all other wards) and
371 displays ‘small world’ properties. The spatial network layer connects all adjacent wards and the permit-
372 related movements reduce the network diameter (longest path length between two wards) from 18 on
373 the spatial network to 12 (see supplementary material Table S2 for cattle market, spatial and multiplex
374 networks summary statistics).

375

376 The distributions of the three node centrality measures that were investigated (betweenness,
377 eigenvector, and geometric mean degree) were strongly right-skewed. This indicates that the multiplex
378 may be sensitive to targeted disease control interventions at the highly influential nodes. Figure 1 shows
379 the geographical distribution of the top-ranked wards for each centrality measure, showing the potential
380 for substantial differences in the effectiveness of targeting controls based on centrality measures due to
381 their geographical distribution.

382

383



384

385 **Figure 1.** Spatial distribution of wards with highest centrality measures in the northern Tanzania
 386 livestock movement network, colour shows position in each centrality measure rank, out of 398: red,
 387 top 1%; orange, 1-5%; yellow, 5-10%.

388

389 **Simulated movements and pathogen transmission**

390 Mean reductions in population cumulative incidence (PCI) after 1 year for the fast and slow pathogens
 391 for each intervention scenario are shown in Figure 2. Reductions are relative to PCI reached after 1 year
 392 with no intervention (fast: 24%; slow: 1.7%). The higher the reduction in PCI, the more effective the
 393 intervention. The list of trialled interventions and associated PCI are given in Tables S3 and S4. All
 394 simulated interventions had greater reduction in PCI for the fast pathogen example compared to the
 395 slow, although the ranking of intervention efficacy was similar for both fast and slow pathogens. The
 396 movement ban implemented in all 111 market wards (high economic and logistical costs) performed
 397 only slightly better than when targeted in only 20 wards using network measures, and network-based
 398 targeting was more effective than selecting wards using population size or randomly, although there
 399 was no substantial difference in performance between the network measures. Vaccination applied to all
 400 wards achieved a 100% reduction in PCI for both fast and slow pathogens, while the best-performing
 401 targeted intervention, degree centrality, achieved reductions in PCI of 58% (fast) and 31% (slow). The
 402 “common sense” intervention of targeting using the total number of cattle performed almost as well as
 403 degree centrality, and similarly to the second-best network measure, betweenness, but was much less
 404 efficient, requiring $3.5 \times$ more vaccine doses than degree centrality. Targeting vaccination using

405 eigenvalue centrality performed relatively poorly, particularly against the slow disease, where its
406 performance was comparable to selecting wards randomly.

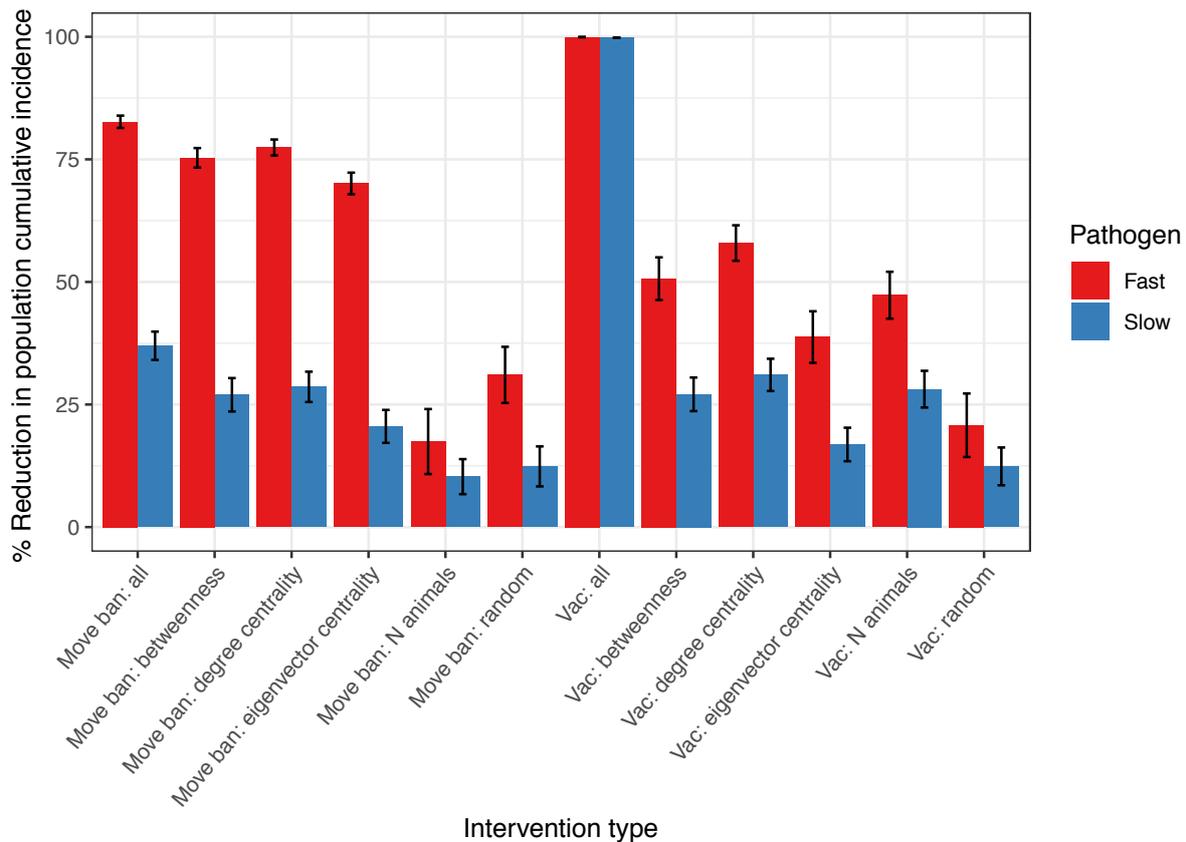
407

408

409

410

411



413

414 **Figure 2.** Mean (\pm SE) percentage reduction in population cumulative incidence (PCI) after 1 year for
 415 simulated ‘fast’ and ‘slow’ transmitting pathogens on the northern Tanzania cattle multiplex network
 416 for two types of intervention (market movement ban or vaccination at 70% coverage) applied using six
 417 strategies: applied to all wards; targeted to 5% ($n = 20$) of wards using each of three network centrality
 418 measures (betweenness, degree, and eigenvector centrality); targeted to the 5% of wards with the
 419 highest cattle population size; and applied to 5% of wards selected randomly. The greater the reduction
 420 in PCI, the more effective the intervention is at reducing total number of cases. Mean PCI under each
 421 scenario is calculated as the geometric mean of 237 simulated epidemics (full data: Tables S3 and S4).

422

423 Discussion

424 It is well established that the network analysis of livestock movements can be used to better understand
 425 and control diseases of commercial and zoonotic importance in higher income countries where livestock
 426 industries tend to be highly structured and movement data are centrally collected and digitised. It is less

427 clear that such approaches are valuable in lower income countries, where movement data are typically
428 unavailable and the cost-benefit ratio less compelling. By exploiting movement permit data collected
429 for health certification and tariff purposes, we have shown that even highly fragmented information
430 about movement patterns can be used to infer network structure. By simulation, we show that the
431 resultant inferred network has the potential to advance strategic understanding. These simulations
432 corroborate that simple network measures can be used to identify good targets for surveillance and
433 disease control that would be appropriate for a range of diseases and reduce the impact of infectious
434 disease at considerably reduced cost and effort. These results could be used to form simple and practical
435 guidelines that could be exploited immediately if, for example, a movement ban was initiated and
436 government needed guidance on where their limited re-enforcement resources should be targeted,
437 although they should not be used for more specific predictions without further data and analysis. They
438 also provide a foundation for deeper research effort, highlighting where the collection of additional
439 empirical data would be useful. For example, the substantial changes in network metrics that result
440 when spatial spread between wards is incorporated highlight the need to augment movement data with
441 more extensive information about local patterns of contact. The homogeneous mixing assumption used
442 at the within-ward level has previously been shown to be useful for developing strategic understanding,
443 even in highly spatially driven scenarios [160], but more detailed recommendations would require
444 modelling of within-ward heterogeneity supported by higher resolution data. This assumption may be
445 less realistic for small urban wards where cattle are tethered, though in larger pastoral and agro-pastoral
446 wards, shared natural resource points might make homogeneous mixing more appropriate (G.L.C.,
447 unpublished data and [158]). Similarly, while the assumption that cattle-to-market movements occur
448 from adjacent wards is consistent with two authors' expert knowledge of livestock management practice
449 (O.M.N. and E.S.), verification with further data collection is an important next step. Finally, simulated
450 movements are dynamically generated based on the random variation generated within the stochastic
451 simulation models. We have not investigated in our dataset evidence of dynamic patterns such as
452 changing network patterns over time because the patchy missingness in our data limits the complexity
453 of the movement model. If more complete data became available for analysis it would be beneficial to
454 assess the evidence for link rewiring throughout the year as this could indicate where control measures

455 should be targeted at specific times. Further potential model deficits include the similar impact of
456 targeting control measures when comparing across centrality measures. This may in part because of the
457 relative crudeness of the disease model; in a more sophisticated model, where the timescales and
458 frequencies of links were considered in greater detail, more substantial differences might be apparent.
459 Similarly, a more explicit model of spatial spread might also prove discriminatory. Finally, the addition
460 of pathogen sequence data where these are available, would provide valuable confirmation of the role
461 of network structure.

462

463 **Conclusions**

464 Despite this demonstration of the value of our inferred network approach, we note that data generation
465 was the result of substantial, time consuming effort, and the resultant inferred network, while useful,
466 has limitations as noted above. Mobile broadband technology is becoming increasingly accessible and
467 coupled with the availability of inexpensive scanning devices, the adoption of routine, robust digitised
468 data recording should be achievable. In this paper, we have shown the benefits of having this data to be
469 potentially substantial. This will be particular pertinent in emerging economies such as Tanzania, where
470 changes in industry structure are likely to have unanticipated disease impacts and will require regular
471 monitoring.

472

473 **Acknowledgements**

474 We are grateful to Rigobert Tarimo and Sambeke Kiruswa for movement permit data entry, The
475 Ministry of Livestock and Fisheries, Tanzania for access to the movement permit data, Stefan Widgren
476 for assistance with the SimInf package, and three anonymous reviewers whose comments greatly
477 improved this manuscript. The movement permit study was supported by the UK BBSRC Zoonoses
478 and Emerging Livestock Systems (ZELS) Initiative BB/L018926/1.

479

480 Study protocols were approved by the ethical review committees of the Kilimanjaro Christian Medical
481 Centre (KCMC/832) and National Institute of Medical Research (NIMR/2028) in Tanzania and in the

482 UK by the ethics review committee of the College of Medical, Veterinary and Life Sciences, University
483 of Glasgow.

484

485

486 **References.**

487

- 488 1. Watts, D.J. and S.H. Strogatz, *Collective dynamics of 'small-world' networks*. Nature,
489 1998. **393**(6684): p. 440-2.
- 490 2. Keeling, M.J., *The effects of local spatial structure on epidemiological invasions*.
491 Proceedings of the Royal Society of London Series B-Biological Sciences, 1999.
492 **266**(1421): p. 859-867.
- 493 3. Liljeros, F., et al., *The web of human sexual contacts*. Nature, 2001. **411**(6840): p.
494 907-8.
- 495 4. Kao, R.R., et al., *Demographic structure and pathogen dynamics on the network of*
496 *livestock movements in Great Britain*. Proc Biol Sci, 2006. **273**(1597): p. 1999-2007.
- 497 5. Robinson, S.E., M.G. Everett, and R.M. Christley, *Recent network evolution*
498 *increases the potential for large epidemics in the British cattle population*. J R Soc
499 Interface, 2007. **4**(15): p. 669-74.
- 500 6. Hufnagel, L., D. Brockmann, and T. Geisel, *Forecast and control of epidemics in a*
501 *globalized world*. Proc Natl Acad Sci U S A, 2004. **101**(42): p. 15124-9.
- 502 7. Brockmann, D., L. Hufnagel, and T. Geisel, *The scaling laws of human travel*.
503 Nature, 2006. **439**(7075): p. 462-5.
- 504 8. Balcan, D., et al., *Multiscale mobility networks and the spatial spreading of infectious*
505 *diseases*. Proc Natl Acad Sci U S A, 2009. **106**(51): p. 21484-9.
- 506 9. Viboud, C., et al., *Synchrony, waves, and spatial hierarchies in the spread of*
507 *influenza*. Science, 2006. **312**(5772): p. 447-451.
- 508 10. Wesolowski, A., et al., *Connecting Mobility to Infectious Diseases: The Promise and*
509 *Limits of Mobile Phone Data*. J Infect Dis, 2016. **214**(suppl_4): p. S414-S420.
- 510 11. Gardy, J.L., et al., *Whole-genome sequencing and social-network analysis of a*
511 *tuberculosis outbreak*. N Engl J Med, 2011. **364**(8): p. 730-9.
- 512 12. Meyers, L.A., M.E. Newman, and B. Pourbohloul, *Predicting epidemics on directed*
513 *contact networks*. J Theor Biol, 2006. **240**(3): p. 400-18.
- 514 13. Green, D.M. and R.R. Kao, *Data quality of the Cattle Tracing System in Great*
515 *Britain*. Vet Rec, 2007. **161**(13): p. 439-43.
- 516 14. Noremark, M., et al., *Network analysis of cattle and pig movements in Sweden:*
517 *Measures relevant for disease control and risk based surveillance*. Preventive
518 Veterinary Medicine, 2011. **99**(2-4): p. 78-90.
- 519 15. Korschake, M., et al., *On the Robustness of In- and Out-Components in a Temporal*
520 *Network*. Plos One, 2013. **8**(2).
- 521 16. Dutta, B.L., P. Ezanno, and E. Vergu, *Characteristics of the spatio-temporal network*
522 *of cattle movements in France over a 5-year period*. Preventive Veterinary Medicine,
523 2014. **117**(1): p. 79-94.
- 524 17. Natale, F., et al., *Network analysis of Italian cattle trade patterns and evaluation of*
525 *risks for potential disease spread*. Preventive Veterinary Medicine, 2009. **92**(4): p.
526 341-350.

- 527 18. Bajardi, P., et al., *Dynamical patterns of cattle trade movements*. PLoS One, 2011.
528 6(5): p. e19869.
- 529 19. VanderWaal, K.L., et al., *Network analysis of cattle movements in Uruguay:
530 Quantifying heterogeneity for risk-based disease surveillance and control*. Preventive
531 Veterinary Medicine, 2016. **123**: p. 12-22.
- 532 20. Leon, E.A., et al., *A description of cattle movements in two departments of Buenos
533 Aires province, Argentina*. Preventive Veterinary Medicine, 2006. **76**(1-2): p. 109-
534 120.
- 535 21. Klous, G., et al., *Human-livestock contacts and their relationship to transmission of
536 zoonotic pathogens, a systematic review of literature*. One Health, 2016. **2**: p. 65-76.
- 537 22. FAO, *The State of Food and Agriculture; Livestock in the balance*. 2009: Rome, Italy.
538 p. 32-53.
- 539 23. Perry, B. and D. Grace, *The impacts of livestock diseases and their control on growth
540 and development processes that are pro-poor*. Philosophical Transactions of the
541 Royal Society B-Biological Sciences, 2009. **364**(1530): p. 2643-2655.
- 542 24. Grace, D., et al., *Mapping of poverty and likely zoonoses hotspots Zoonoses 2012:*
543 Department for International Development.
- 544 25. ILRI. *Why livestock matter 2018* 11/07/2018]; Available from:
545 <https://www.ilri.org/whylivestockmatter>.
- 546 26. Aklilu, Y., *Livestock Marketing in Kenya and Ethiopia: A Review of Policies and
547 Practice*. 2008, Tufts University.
- 548 27. Musemwa, L., et al., . *The Impact of Climate Change on Livestock The Impact of
549 Climate Change on Livestock Production amongst the Resource-Poor Farmers of
550 Third World Countries: A Review*. . Asian J. Agric. Rural Dev., 2012. **2**,; p. 621-631.
- 551 28. Apolloni, A., et al., *Towards the description of livestock mobility in Sahelian Africa:
552 Some results from a survey in Mauritania*. Plos One, 2018. **13**(1).
- 553 29. Hummel, P.H., *Incidence in Tanzania of Cf Antibody to Coxiella-Burneti in Sera from
554 Man, Cattle, Sheep, Goats and Game*. Veterinary Record, 1976. **98**(25): p. 501-505.
- 555 30. Schoonman, L. and E.S. Swai, *Herd- and animal-level risk factors for bovine
556 leptospirosis in Tanga region of Tanzania*. Tropical Animal Health and Production,
557 2010. **42**(7): p. 1565-1572.
- 558 31. Crump, J.A., et al., *Etiology of Severe Non-malaria Febrile Illness in Northern
559 Tanzania: A Prospective Cohort Study*. Plos Neglected Tropical Diseases, 2013. **7**(7).
- 560 32. Assenga, J.A., et al., *Epidemiology of Brucella infection in the human, livestock and
561 wildlife interface in the Katavi-Rukwa ecosystem, Tanzania*. BMC Veterinary
562 Research, 2015. **11**.
- 563 33. Sumaye, R.D., et al., *Inter-epidemic Acquisition of Rift Valley Fever Virus in Humans
564 in Tanzania*. Plos Neglected Tropical Diseases, 2015. **9**(2).
- 565 34. Wensman, J.J., et al., *A study of Rift Valley fever virus in Morogoro and Arusha
566 regions of Tanzania - serology and farmers' perceptions*. Infect Ecol Epidemiol,
567 2015. **5**: p. 30025.
- 568 35. Cash-Goldwasser, S., et al., *Risk Factors for Human Brucellosis in Northern
569 Tanzania*. American Journal of Tropical Medicine and Hygiene, 2018. **98**(2): p. 598-
570 606.
- 571 36. Wasserman, S. and K. Faust, *Social network analysis : methods and applications*.
572 Structural analysis in the social sciences. 1994, Cambridge ; New York: Cambridge
573 University Press. xxxi, 825 p.
- 574 37. Bell, D., J.S. Atkinson, and J.W. Carlson, *Centrality measures for disease
575 transmission networks*. Social Networks, 1999. **21**(1): p. 1-21.

- 576 38. Christley, R.M., et al., *Infection in social networks: using network analysis to identify*
577 *high-risk individuals*. Am J Epidemiol, 2005. **162**(10): p. 1024-31.
- 578 39. Natale, F., et al., *Evaluation of risk and vulnerability using a Disease Flow Centrality*
579 *measure in dynamic cattle trade networks*. Preventive Veterinary Medicine, 2011.
580 **98**(2-3): p. 111-118.
- 581 40. Palisson, A., A. Courcoul, and B. Durand, *Role of Cattle Movements in Bovine*
582 *Tuberculosis Spread in France between 2005 and 2014*. PLoS One, 2016. **11**(3): p.
583 e0152578.
- 584 41. Sintayehu, D.W., et al., *Disease transmission in animal transfer networks*. Prev Vet
585 Med, 2017. **137**(Pt A): p. 36-42.
- 586 42. Buttner, K., et al., *Efficient interruption of infection chains by targeted removal of*
587 *central holdings in an animal trade network*. PLoS One, 2013. **8**(9): p. e74292.
- 588 43. VanderWaal, K., et al., *Optimal surveillance strategies for bovine tuberculosis in a*
589 *low-prevalence country*. Scientific Reports, 2017. **7**.
- 590 44. Haydon, D.T., R.R. Kao, and R.P. Kitching, *The UK foot-and-mouth disease outbreak*
591 *- the aftermath*. Nat Rev Microbiol, 2004. **2**(8): p. 675-81.
- 592 45. Gibbens, J.C., et al., *Descriptive epidemiology of the 2001 foot-and-mouth disease*
593 *epidemic in Great Britain: the first five months*. Veterinary Record, 2001. **149**(24): p.
594 729+.
- 595 46. Shirley, M.D. and S.P. Rushton, *Where diseases and networks collide: lessons to be*
596 *learnt from a study of the 2001 foot-and-mouth disease epidemic*. Epidemiol Infect,
597 2005. **133**(6): p. 1023-32.
- 598 47. Kao, R.R., *The role of mathematical modelling in the control of the 2001 FMD*
599 *epidemic in the UK*. Trends Microbiol, 2002. **10**(6): p. 279-86.
- 600 48. Kao, R.R., et al., *Disease dynamics over very different time-scales: foot-and-mouth*
601 *disease and scrapie on the network of livestock movements in the UK*. J R Soc
602 Interface, 2007. **4**(16): p. 907-16.
- 603 49. Keeling, M.J. and B.T. Grenfell, *Individual-based perspectives on $R(0)$* . J Theor Biol,
604 2000. **203**(1): p. 51-61.
- 605 50. Green, D.M., I.Z. Kiss, and R.R. Kao, *Parameterization of individual-based models:*
606 *comparisons with deterministic mean-field models*. J Theor Biol, 2006. **239**(3): p.
607 289-97.
- 608 51. Enright, J. and R.R. Kao, *Epidemics on dynamic networks*. Epidemics, 2018. **24**: p.
609 88-97.
- 610 52. Kao, R.R., *Networks and Models with Heterogeneous Population Structure in*
611 *Epidemiology*, in *Network Science: Complexity in Nature and Technology*, E. Estrada,
612 et al., Editors. 2010, Springer.
- 613 53. Takaguchi, T., N. Masuda, and P. Holme, *Bursty communication patterns facilitate*
614 *spreading in a threshold-based epidemic dynamics*. PLoS One, 2013. **8**(7): p. e68629.
- 615 54. Holme, P. and N. Masuda, *The Basic Reproduction Number as a Predictor for*
616 *Epidemic Outbreaks in Temporal Networks*. PLoS ONE, 2015.
- 617 55. Barabasi, A.L., *The origin of bursts and heavy tails in human dynamics*. Nature, 2005.
618 **435**(7039): p. 207-11.
- 619 56. Vazquez, A., et al., *Modeling bursts and heavy tails in human dynamics*. Physical
620 Review E, 2006. **73**(3).
- 621 57. Oliveira, J.G. and A.L. Barabasi, *Human dynamics: Darwin and Einstein*
622 *correspondence patterns*. Nature, 2005. **437**(7063): p. 1251.
- 623 58. Min, B., K.I. Goh, and A. Vazquez, *Spreading dynamics following bursty human*
624 *activity patterns*. Phys Rev E Stat Nonlin Soft Matter Phys, 2011. **83**(3 Pt 2): p.
625 036102.

- 626 59. Iribarren, J.L. and E. Moro, *Branching dynamics of viral information spreading*. Phys
627 Rev E Stat Nonlin Soft Matter Phys, 2011. **84**(4 Pt 2): p. 046116.
- 628 60. Karsai, M., et al., *Small but slow world: how network topology and burstiness slow*
629 *down spreading*. Phys Rev E Stat Nonlin Soft Matter Phys, 2011. **83**(2 Pt 2): p.
630 025102.
- 631 61. Nickbakhsh, S., et al., *Implications of within-farm transmission for network*
632 *dynamics: consequences for the spread of avian influenza*. Epidemics, 2013. **5**(2): p.
633 67-76.
- 634 62. Epstein, J.M., et al., *Coupled contagion dynamics of fear and disease: mathematical*
635 *and computational explorations*. PLoS One, 2008. **3**(12): p. e3955.
- 636 63. Funk, S., M. Salathe, and V.A. Jansen, *Modelling the influence of human behaviour*
637 *on the spread of infectious diseases: a review*. J R Soc Interface, 2010. **7**(50): p. 1247-
638 56.
- 639 64. Green, D.M., et al., *Estimates for local and movement-based transmission of bovine*
640 *tuberculosis in British cattle*. Proc Biol Sci, 2008. **275**(1638): p. 1001-5.
- 641 65. Drake, J.M., et al., *Ebola cases and health system demand in Liberia*. PLoS Biol,
642 2015. **13**(1): p. e1002056.
- 643 66. Chowell, G. and H. Nishiura, *Characterizing the transmission dynamics and control*
644 *of ebola virus disease*. PLoS Biol, 2015. **13**(1): p. e1002057.
- 645 67. Cottam, E.M., et al., *Transmission pathways of foot-and-mouth disease virus in the*
646 *United Kingdom in 2007*. PLoS Pathog, 2008. **4**(4): p. e1000050.
- 647 68. Kao, R.R., et al., *Supersize me: how whole-genome sequencing and big data are*
648 *transforming epidemiology*. Trends in microbiology, 2014. **22**(5): p. 282-291.
- 649 69. De Maio, N., C.H. Wu, and D.J. Wilson, *SCOTTI: Efficient Reconstruction of*
650 *Transmission within Outbreaks with the Structured Coalescent*. PLoS Comput Biol,
651 2016. **12**(9): p. e1005130.
- 652 70. Hall, M., M. Woolhouse, and A. Rambaut, *Epidemic Reconstruction in a*
653 *Phylogenetics Framework: Transmission Trees as Partitions of the Node Set*. PLoS
654 Comput Biol, 2015. **11**(12): p. e1004613.
- 655 71. Jombart, T., et al., *Reconstructing disease outbreaks from genetic data: a graph*
656 *approach*. Heredity (Edinb), 2011. **106**(2): p. 383-90.
- 657 72. Jombart, T., et al., *Spatiotemporal dynamics in the early stages of the 2009 A/H1N1*
658 *influenza pandemic*. PLoS Curr, 2009. **1**: p. RRN1026.
- 659 73. Lau, M.S., et al., *A Systematic Bayesian Integration of Epidemiological and Genetic*
660 *Data*. PLoS Comput Biol, 2015. **11**(11): p. e1004633.
- 661 74. Morelli, M.J., et al., *A Bayesian inference framework to reconstruct transmission*
662 *trees using epidemiological and genetic data*. PLoS Comput Biol, 2012. **8**(11): p.
663 e1002768.
- 664 75. Numminen, E., et al., *Two-phase importance sampling for inference about*
665 *transmission trees*. Proc Biol Sci, 2014. **281**(1794): p. 20141324.
- 666 76. Worby, C.J., et al., *Reconstructing transmission trees for communicable diseases*
667 *using densely sampled genetic data*. Ann Appl Stat, 2016. **10**(1): p. 395-417.
- 668 77. Ypma, R.J., et al., *Unravelling transmission trees of infectious diseases by combining*
669 *genetic and epidemiological data*. Proc Biol Sci, 2012. **279**(1728): p. 444-50.
- 670 78. Ypma, R.J., W.M. van Ballegooijen, and J. Wallinga, *Relating phylogenetic trees to*
671 *transmission trees of infectious disease outbreaks*. Genetics, 2013. **195**(3): p. 1055-
672 62.
- 673 79. Pybus, O.G., A.J. Tatem, and P. Lemey, *Virus evolution and transmission in an ever*
674 *more connected world*. Proc Biol Sci, 2015. **282**(1821): p. 20142878.

- 675 80. Biek, R., et al., *Measurably evolving pathogens in the genomic era*. Trends Ecol Evol, 676 2015. **30**(6): p. 306-313.
- 677 81. Frost, S.D.W., et al., *Eight challenges in phylodynamic inference*. Epidemics, 2015. 678 **10**: p. 88-92.
- 679 82. Meehan, C.J., et al., *The relationship between transmission time and clustering 680 methods in Mycobacterium tuberculosis epidemiology*. . bioRxiv, 2018.
- 681 83. Romero-Severson, E., et al., *Timing and order of transmission events is not directly 682 reflected in a pathogen phylogeny*. Molecular biology and evolution, 2014. **31**(9): p. 683 2472-2482.
- 684 84. Worby, C.J., M. Lipsitch, and W.P. Hanage, *Within-host bacterial diversity hinders 685 accurate reconstruction of transmission networks from genomic distance data*. . PLoS 686 computational biology,, 2014. **10**(3).
- 687 85. Di Nardo, A., N.J. Knowles, and D.J. Paton, *Combining livestock trade patterns with 688 phylogenetics to help understand the spread of foot and mouth disease in sub- 689 Saharan Africa, the Middle East and Southeast Asia*. . Revue Scientifique et 690 Technique-OIE,, 2011. **30**(1).
- 691 86. VanderWaal, K.L., et al., *Linking social and pathogen transmission networks using 692 microbial genetics in giraffe (Giraffa camelopardalis)*. . Journal of Animal Ecology, 693 2014. **83**(2): p. 406-414.
- 694 87. Rasmussen, D.A., E.M. Volz, and K. Koelle, *Phylodynamic inference for structured 695 epidemiological models*. PLoS Comput Biol, 2014. **10**(4): p. e1003570.
- 696 88. Viana, M., et al., *Integrating serological and genetic data to quantify cross-species 697 transmission: brucellosis as a case study*. Parasitology, 2016. **143**(7): p. 821-834.
- 698 89. Knight-Jones, T.J. and J. Rushton, *The economic impacts of foot and mouth disease - 699 what are they, how big are they and where do they occur?* Prev Vet Med, 2013. 700 **112**(3-4): p. 161-73.
- 701 90. Smith, R.L., et al., *Minimization of bovine tuberculosis control costs in US dairy 702 herds*. Prev Vet Med, 2013. **112**(3-4): p. 266-75.
- 703 91. Holling, C.S., *Resilience and stability of ecological systems*. Annual Review of 704 Ecology and Systematics, 1973. **4**: p. 1-23.
- 705 92. Holling, C.S., *Engineering resilience versus ecological resilience*, in *Engineering 706 within ecological constraints*. 1996, National Academy: Washington D.C., USA. p. 707 31-44.
- 708 93. Carpenter, S., et al., *From Metaphor to Measurement: Resilience of What to What?* 709 Ecosystems, 2001. **4**(8).
- 710 94. Anderson, I., *Foot and Mouth Disease 2001: Lessons to be Learned Inquiry*. 2002, 711 London: The Stationary Office.
- 712 95. Nickbakhsh, S., et al., *A metapopulation model for highly pathogenic avian influenza: 713 implications for compartmentalization as a control measure*. Epidemiol Infect, 2014. 714 **142**(9): p. 1813-25.
- 715 96. Albert, R., H. Jeong, and A.L. Barabasi, *Error and attack tolerance of complex 716 networks*. Nature, 2000. **406**(6794): p. 378-82.
- 717 97. Rossi, G., et al., *Epidemiological modelling for the assessment of bovine tuberculosis 718 surveillance in the dairy farm network in Emilia-Romagna (Italy)*. Epidemics, 2015. 719 **11**: p. 62-70.
- 720 98. C, D., et al., *Introduction to network analysis and its implications for animal disease 721 modelling*. . Revue Scientifique et Technique (International Office of Epizootics) 722 2011. **30**: p. 425 – 436.
- 723 99. ME., C., *Infectious disease transmission and contact networks in wildlife and 724 livestock*. . Philos. Trans. R. Soc. B Biol. Sci. , 2015. **370**: p. 20140107–20140107.

- 725 100. Dube, C., et al., *A Review of Network Analysis Terminology and its Application to*
726 *Foot-and-Mouth Disease Modelling and Policy Development*. *Transboundary and*
727 *Emerging Diseases*, 2009. **56**(3): p. 73-85.
- 728 101. Bessell, P.R., et al., *Developing a framework for risk-based surveillance of*
729 *tuberculosis in cattle: a case study of its application in Scotland*. *Epidemiology and*
730 *Infection*, 2012: p. 1-10.
- 731 102. Salvador, L.C.M., et al., *Risk-based strategies for surveillance of tuberculosis*
732 *infection in cattle for low-risk areas in England and Scotland*. *Epidemiol Infect*, 2018.
733 **146**(1): p. 107-118.
- 734 103. San Miguel, M., et al., *Challenges in complex systems science*. *European Physical*
735 *Journal-Special Topics*, 2012. **214**(1): p. 245-271.
- 736 104. Webster, J.P., A. Borlase, and J.W. Rudge, *Who acquires infection from whom and*
737 *how? Disentangling multi-host and multi-mode transmission dynamics in the*
738 *'elimination' era*. *Philosophical Transactions of the Royal Society B-Biological*
739 *Sciences*, 2017. **372**(1719).
- 740 105. Godfray, H.C.J., et al., *A restatement of the natural science evidence base relevant to*
741 *the control of bovine tuberculosis in Great Britain†*. *Proceedings of the Royal Society*
742 *of London B: Biological Sciences*, 2013. **280**(1768): p. 20131634.
- 743 106. Kivelä, M., et al., *Multilayer networks*. *Journal of Complex Networks*, 2014. **2**(3): p.
744 203-271.
- 745 107. Pilosof, S., et al., *The multilayer nature of ecological networks*. *Nature Ecology &*
746 *Evolution*, 2017. **1**(4).
- 747 108. Cardillo, A., et al., *Modeling the multi-layer nature of the European Air Transport*
748 *Network: Resilience and passengers re-scheduling under random failures*. *European*
749 *Physical Journal-Special Topics*, 2013. **215**(1): p. 23-33.
- 750 109. Barrett, L., S.P. Henzi, and D. Lusseau, *Taking sociality seriously: the structure of*
751 *multi-dimensional social networks as a source of information for individuals*.
752 *Philosophical Transactions of the Royal Society B-Biological Sciences*, 2012.
753 **367**(1599): p. 2108-2118.
- 754 110. Brooks-Pollock, E., et al., *Eight challenges in modelling infectious livestock diseases*.
755 *Epidemics*, 2015. **10**: p. 1-5.
- 756 111. Rossi, G., et al., *The Potential Role of Direct and Indirect Contacts on Infection*
757 *Spread in Dairy Farm Networks*. *Plos Computational Biology*, 2017. **13**(1).
- 758 112. Rossi, G., et al., *Modelling farm-to-farm disease transmission through personnel*
759 *movements: from visits to contacts, and back*. *Scientific Reports*, 2017. **7**.
- 760 113. Haydon, D.T., et al., *Identifying reservoirs of infection: A conceptual and practical*
761 *challenge*. *Emerging Infectious Diseases*, 2002. **8**(12): p. 1468-1473.
- 762 114. Sole, R.V. and J.M. Montoya, *Complexity and fragility in ecological networks*.
763 *Proceedings of the Royal Society B-Biological Sciences*, 2001. **268**(1480): p. 2039-
764 2045.
- 765 115. Gao, J., B. Barzel, and A.L. Barabasi, *Universal resilience patterns in complex*
766 *networks (vol 530, pg 307, 2016)*. *Nature*, 2016. **536**(7615): p. 238-238.
- 767 116. Gao, J.X., et al., *Percolation of a general network of networks*. *Physical Review E*,
768 2013. **88**(6).
- 769 117. Cellai, D., et al., *Percolation in multiplex networks with overlap*. *Physical Review E*,
770 2013. **88**(5).
- 771 118. Bohm, M., M.R. Hutchings, and P.C.L. White, *Contact Networks in a Wildlife-*
772 *Livestock Host Community: Identifying High-Risk Individuals in the Transmission of*
773 *Bovine TB among Badgers and Cattle*. *Plos One*, 2009. **4**(4).

- 774 119. Silk, M.J., et al., *Quantifying direct and indirect contacts for the potential*
775 *transmission of infection between species using a multilayer contact network.*
776 Behaviour, 2018.
- 777 120. Bigras-Poulin, M., et al., *Network analysis of Danish cattle industry trade patterns as*
778 *an evaluation of risk potential for disease spread.* Preventive Veterinary Medicine,
779 2006. **76**(1-2): p. 11-39.
- 780 121. Kiss, I.Z., D.M. Green, and R.R. Kao, *The network of sheep movements within Great*
781 *Britain: Network properties and their implications for infectious disease spread.* J R
782 Soc Interface, 2006. **3**(10): p. 669-77.
- 783 122. Motta, P., et al., *Implications of the cattle trade network in Cameroon for regional*
784 *disease prevention and control.* Sci Rep, 2017. **7**: p. 43932.
- 785 123. Government, M.f.P.I.N.Z. *National Animal Identification and Tracing.* 2018;
786 Available from: [https://www.mpi.govt.nz/growing-and-harvesting/livestock-and-](https://www.mpi.govt.nz/growing-and-harvesting/livestock-and-animal-care/national-animal-identification-and-tracing/)
787 [animal-care/national-animal-identification-and-tracing/](https://www.mpi.govt.nz/growing-and-harvesting/livestock-and-animal-care/national-animal-identification-and-tracing/).
- 788 124. Handcock, R.N., et al., *Monitoring Animal Behaviour and Environmental Interactions*
789 *Using Wireless Sensor Networks, GPS Collars and Satellite Remote Sensing.* Sensors,
790 2009. **9**(5): p. 3586-3603.
- 791 125. Raizman, E.A., et al., *Feasibility study on the spatial and temporal movement of*
792 *Samburu's cattle and wildlife in Kenya using GPS radio-tracking, remote sensing and*
793 *GIS.* Preventive Veterinary Medicine, 2013. **111**(1-2): p. 76-80.
- 794 126. Poolkhet, C., et al., *Social network analysis used to assess the relationship between*
795 *the spread of avian influenza and movement patterns of backyard chickens in*
796 *Ratchaburi, Thailand.* Research in Veterinary Science, 2013. **95**(1): p. 82-86.
- 797 127. Kim, Y., et al., *Livestock trade network: potential for disease transmission and*
798 *implications for risk-based surveillance on the island of Mayotte.* Scientific Reports,
799 2018. **8**.
- 800 128. Lindstrom, T., et al., *A Bayesian Approach for Modeling Cattle Movements in the*
801 *United States: Scaling up a Partially Observed Network.* Plos One, 2013. **8**(1).
- 802 129. Dube, C., et al., *Comparing Network Analysis Measures to Determine Potential*
803 *Epidemic Size of Highly Contagious Exotic Diseases in Fragmented Monthly*
804 *Networks of Dairy Cattle Movements in Ontario, Canada.* Transboundary and
805 Emerging Diseases, 2008. **55**(9-10): p. 382-392.
- 806 130. Poolkhet, C., et al., *Social network analysis of cattle movement in Kampong Cham,*
807 *Kampong Speu and Takeo, Cambodia.* Acta Tropica, 2016. **159**: p. 44-49.
- 808 131. Wongsathapornchai, K., et al., *Assessment of the likelihood of the introduction of foot-*
809 *and-mouth disease through importation of live animals into the Malaysia-Thailand-*
810 *Myanmar peninsula.* American Journal of Veterinary Research, 2008. **69**(2): p. 252-
811 260.
- 812 132. Selby, R., et al., *Cattle movements and trypanosomes: restocking efforts and the*
813 *spread of Trypanosoma brucei rhodesiense sleeping sickness in post-conflict Uganda.*
814 Parasites & Vectors, 2013. **6**.
- 815 133. Dean, A.S., et al., *Potential Risk of Regional Disease Spread in West Africa through*
816 *Cross-Border Cattle Trade.* Plos One, 2013. **8**(10).
- 817 134. Nicolas, G., et al., *Description and analysis of the cattle trade network in the*
818 *Madagascar highlands: Potential role in the diffusion of Rift Valley fever virus.* Acta
819 Tropica, 2013. **126**(1): p. 19-27.
- 820 135. Nickbakhsh, S., et al., *Generating social network data using partially described*
821 *networks: an example informing avian influenza control in the British poultry*
822 *industry.* BMC Vet Res, 2011. **7**: p. 66.

- 823 136. Xia, Y., O.N. Bjornstad, and B.T. Grenfell, *Measles metapopulation dynamics: a*
824 *gravity model for epidemiological coupling and dynamics*. *Am Nat*, 2004. **164**(2): p.
825 267-81.
- 826 137. Simini, F., et al., *A universal model for mobility and migration patterns*. *Nature*, 2012.
827 **484**(7392): p. 96-100.
- 828 138. Masucci, A.P., et al., *Gravity versus radiation models: on the importance of scale and*
829 *heterogeneity in commuting flows*. *Phys Rev E Stat Nonlin Soft Matter Phys*, 2013.
830 **88**(2): p. 022812.
- 831 139. Bouslikhane, M., *CROSS BORDER MOVEMENTS OF ANIMALS AND ANIMAL*
832 *PRODUCTS AND THEIR RELEVANCE TO THE EPIDEMIOLOGY OF ANIMAL*
833 *DISEASES IN AFRICA*. . 2015, O.I.E.
- 834 140. Muyunda, C., *Hidden value on the hoof: Cross-border livestock trade in East Africa*. .
835 2009, Common Market for Eastern and Southern Africa Comprehensive African
836 Agriculture Development Programme.
- 837 141. Biggs, H.M., et al., *Leptospirosis and Human Immunodeficiency Virus Co-Infection*
838 *Among Febrile Inpatients in Northern Tanzania*. *Vector-Borne and Zoonotic*
839 *Diseases*, 2013. **13**(8): p. 572-580.
- 840 142. Halliday, J., et al., *Bringing together emerging and endemic zoonoses surveillance:*
841 *shared challenges and a common solution*. *Philosophical Transactions of the Royal*
842 *Society B-Biological Sciences*, 2012. **367**(1604): p. 2872-2880.
- 843 143. Heinrich, N., et al., *High seroprevalence of Rift Valley FEVER AND EVIDENCE*
844 *FOR ENDEMIC circulation in Mbeya region, Tanzania, in a cross-sectional study*.
845 *PLoS Negl Trop Dis*, 2012. **6**(3): p. e1557.
- 846 144. Karimuribo, E.D., et al., *Prevalence of brucellosis in crossbred and indigenous cattle*
847 *in Tanzania*. *Livest. Res. Rural Dev*. 19, 2007
- 848 145. Machangu, R.S., G. Mgode, and D. Mpanduji, *Leptospirosis in animals and humans*
849 *in selected areas of Tanzania*. *Belgian Journal of Zoology*, 1997. **127**: p. 97-104.
- 850 146. Vanderburg, S., et al., *Epidemiology of Coxiella burnetii Infection in Africa: A*
851 *OneHealth Systematic Review*. *Plos Neglected Tropical Diseases*, 2014. **8**(4).
- 852 147. Allen, L.H., *Interventions for Micronutrient Deficiency Control in Developing*
853 *Countries: Past, Present and Future*. . *J. Nutr.* , 2003. **133**: p. 3875S–3878S.
- 854 148. Coker, R., et al., *Towards a conceptual framework to support one-health research for*
855 *policy on emerging zoonoses*. *Lancet Infect Dis*, 2011. **11**(4): p. 326-31.
- 856 149. Kelly, A.M. and R.R. Marshak, *Veterinary medicine, global health*. *J Am Vet Med*
857 *Assoc*, 2007. **231**(12): p. 1806-8.
- 858 150. Muma, J.B., Mwacalimba, K.K., Munang'andu, H.M., Matope, G., Jenkins, A.,
859 Siamudaala, V., Mweene, A.S., Marcotty, T.,, *The contribution of veterinary medicine*
860 *to public health and poverty reduction in developing countries*. . *Vet. Ital.* , 2014. **50**:
861 p. 117–29.
- 862 151. Pradere, J.P., *Improving animal health and livestock productivity to reduce poverty*.
863 *Rev Sci Tech*, 2014. **33**(3): p. 735-44, 723-34.
- 864 152. Randolph, T.F., et al., *Invited Review: Role of livestock in human nutrition and health*
865 *for poverty reduction in developing countries*. *Journal of Animal Science*, 2007.
866 **85**(11): p. 2788-2800.
- 867 153. Steinfeld, H., et al., *Livestock's long shadow, Environmental issues and options*.
868 2006.
- 869 154. Covarrubias, K., et al., *Livestock and livelihoods in rural Tanzania and A descriptive*
870 *analysis of the 2009 National Panel Survey*. 2012.
- 871 155. Pica-Ciamarra, U., et al., *Linking smallholders to livestock markets: Combining*
872 *market and household survey data in Tanzania*. 2011.

- 873 156. Williams T.O., S.B.a.O.I., *Improving livestock marketing and intra-regional trade in*
874 *West Africa: determining appropriate economic incentives and policy framework*. .
875 2006, ILRI (International Livest. Res. Institute),: Nairobi, Kenya.
- 876 157. Coppolillo, P.B., *The landscape ecology of pastoral herding: Spatial analysis of land*
877 *use and livestock production in East Africa*. Human Ecology, 2000. **28**(4): p. 527-560.
- 878 158. VanderWaal, K., et al., *Seasonality and pathogen transmission in pastoral cattle*
879 *contact networks*. Royal Society Open Science, 2017. **4**(12).
- 880 159. Statistics, T.N.B.o. *Tanzania in Figures 2012*. 2012 09/14/2018]; Available from:
881 <http://www.nbs.go.tz>.
- 882 160. Keeling, M.J., et al., *Modelling vaccination strategies against foot-and-mouth*
883 *disease*. Nature, 2003. **421**(6919): p. 136-42.
- 884 161. Büttner, K., Krieter, J., Traulsen, A., Traulsen, I., 2013. Static network analysis of a pork supply
885 chain in Northern Germany—Characterisation of the potential spread of infectious diseases via
886 animal movements. Prev. Vet. Med. 110, 418–428.
887 <https://doi.org/10.1016/j.prevetmed.2013.01.008>
888