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Quantitative Risk Assessment of Developing Salmonellosis through Consumption of Beef in Lusaka Province, Zambia



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Highlights for Review

- Risk of developing salmonellosis through beef generally low in Zambia.
- Consumption patterns have an effect on risk of developing salmonellosis.
- Kitchen cross-contamination increases risk of developing *Salmonellosis*.
- Cooking alone not adequate response to exceptional events of beef contamination.

1 **Quantitative Risk Assessment of Developing Salmonellosis through Consumption of Beef**
2 **in Lusaka Province, Zambia**

3
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Abstract

Based on the Codex Alimentarius framework, this study quantitatively assessed the risk of developing salmonellosis through consumption of beef in Lusaka Province of Zambia. Data used to achieve this objective were obtained from reviews of scientific literature, Government reports, and survey results from a questionnaire that was administered to consumers to address information gaps from secondary data. The Swift Quantitative Microbiological Risk Assessment (sQMRA) model was used to analyse the data. The study was driven by a lack of empirically-based risk estimation despite a number of reported cases of salmonellosis in humans.

A typology of consumers including all age groups was developed based on their beef consumption habits, distinguishing between those with low home consumption, those with medium levels of home consumption, and those with high levels through restaurant consumption. This study shows that the risk of developing salmonellosis in this population, from consuming beef, was generally low. At ID₅₀ of 9.61×10^3 cfu/g and a retail contamination concentration of 12 cfu/g, the risk of developing salmonellosis through the consumption of beef prepared by consumers with low and medium levels of beef consumption was estimated at 0.06% and 0.08%, respectively, while the risk associated with restaurant consumption was estimated at 0.16% per year.

The study concludes that the risk of developing salmonellosis among residents in Lusaka province, as a result of beef consumption, was generally low, mainly due to the methods used for food preparation. Further work is required to broaden the scope of the study and also undertake microbiological evaluation of ready-to-eat beef from both the household and restaurant risk exposure pathways.

Keywords: Beef consumption; Quantitative risk assessment; Salmonellosis; sQMRA; Zambia

55 1.0 Introduction

56 The expanding trade of food and livestock, and increased human travel and migration are a
57 means of spreading infectious diseases, irrespective of national borders (Evans & Leighton,
58 2014). This makes the control of infectious diseases and maintenance of food safety
59 important for all countries. This expansion of trade and human travel may lead to a transfer
60 of diseases to areas where such were not a problem originally. This is because disease
61 spread is usually accompanied with cultural changes including eating habits, mass catering,
62 complex and lengthy food supply procedures, increased international movement, and poor
63 hygiene practices in the native community.

64 One of the most widespread infectious foodborne disease of humans is salmonellosis
65 (Carrasco, Morales-Rueda, & García-Gimeno, 2012; Kagambèga et al., 2013; Teunis et al.,
66 2010). Salmonellosis is a disease of both humans and animals caused by two species of
67 *Salmonella* (*S. enterica* and *S. bongori*) (Kemal, 2014; OIE, 2014). The pathogens can cause
68 enteric fevers, gastroenteritis, and septicemia which are of both socio-economic and public
69 health importance (Ulaya, 2013). The majority of infections are associated with the
70 ingestion of contaminated foods such as beef and beef products, poultry, pork, eggs, milk,
71 cheese, seafood, fruits, juices, and vegetables (Freitas Neto et al., 2010; Jackson et al.,
72 2013); although most infections caused by multidrug-resistant *Salmonella* are acquired
73 through contaminated foods of animal origin (Abouzeed et al., 2000).

74 Although the domestic market for beef is small and under-developed in Zambia, demand for
75 beef products has grown steadily in Lusaka province, the capital region, now home to
76 almost 2.7 million people (CSO, 2015). Shifting consumption patterns are associated with an
77 emerging middle class with increasing purchasing power. There is also an increase in
78 domestic beef production in both commercial and traditional sectors, and a rising import of
79 beef and beef products to cover the increased demand in the country (World Bank, 2011).
80 The increase in production may have negative impacts in terms of food safety, especially in
81 traditional production, as the country does not have enough slaughter facilities (Lubungu,
82 Sitko, & Hichaambwa, 2015). Indeed, Zambia, like other low and middle income countries of
83 Africa, has few formal abattoirs compared to a large number of informal slaughterhouses
84 associated with poor hygienic practices (Haileselassie, Taddele, Adhana, & Kalayou, 2013).
85 There is higher risk of fecal spillage on the meat because of slaughtering on the floor

86 (slaughter slabs). Given this scenario, the chances of producing contaminated carcasses are
87 high, since contamination of carcasses may occur throughout the value chain (from
88 production through to consumption). This might lead to the introduction of *Salmonella* into
89 the food chain if there was an early exposure of domestic animals to the organism that
90 results in long-term persistent infections (Muma, 1998; Isogai *et al.*, 2005; Haileelassie *et*
91 *al.*, 2013; Ndalama & Mdegela 2013).

92 *Salmonella* has been previously detected in human samples in Lusaka; out of the 200 clinical
93 diarrhoea stool samples, 9 (4.5%) were found to be bacteriological culture positive for
94 *Salmonella* (Hang'ombe, 1998). Mwansa *et al.* (2002) reported that of 124 adults and 105
95 children with persistent diarrhea in Zambia, 6 (5%) and 21 (20%) were infected with non-
96 typhoidal *Salmonella* (NTS) species, respectively. In an earlier study at the University
97 Teaching Hospital (UTH), Lusaka, 45 strains of various NTS species were isolated from stool
98 samples, blood, and cerebral spinal fluid (CSF)(Hangombe, 1998). About 93% of the strains
99 were isolated from infants less than two years old. *Salmonella Heidelberg* was the most
100 common species isolated from stool and revealed a multi-drug resistant character. This
101 shows that *Salmonella* is present and that there is a risk of getting salmonellosis once an
102 individual consumes contaminated food, including beef, or gets otherwise exposed (e.g.
103 direct contact with infected animals).

104 Previous studies have also indicated that among the microbiological hazards in the beef
105 value chain, *Salmonella* has a great public health significance (Dhanoa & Fatt, 2009; Kemal,
106 2014; Plym L & Wierup, 2006). Muma *et al.* (1998) isolated *Salmonella* from beef carcasses in
107 a survey involving abattoirs in Lusaka and Copperbelt provinces, whose results
108 demonstrated that there was a high level of contamination on carcasses due to poor
109 hygiene status in abattoirs (Muma, 1998; Ntanga 2013). Further, diarrhoeal cases have been
110 reported in Lusaka, some of which were due to *Salmonella* infections (Mwansa *et al.*, 2002;
111 Hang'ombe *et al.*, 2011).

112 Despite evidence of presence of *Salmonella* species in beef from previous research, very
113 little is known about the risk of salmonellosis through consumption of beef in Lusaka
114 Province and Zambia in general. It is therefore important to assess whether the increase in
115 beef consumption increases public health burdens due to exposure to foodborne hazards.

116 To address this information gap, this study used a Swift Quantitative Microbiological Risk
117 Assesment (sQMRA) model to quantify these risks (Evers & Chardon, 2010).

118 There is a paucity of published literature that demonstrates a quantitative risk of developing
119 salmonellosis through the consumption of beef using sQMRA food safety risk analysis tool.

120 This paper illustrates scenarios where both the household and restaurant risk pathways
121 have been used to assess the risk of developing salmonellosis through the consumption of
122 beef prepared in three different ways.

123

ACCEPTED MANUSCRIPT

124 **2.0 Methodology**

125 **2.1 Study area**

126 This study was conducted in the Lusaka province of Zambia, an area with relatively high beef
127 consumption due to high purchasing power (Sinkala et al., 2014).

128 **2.2 Swift Quantitative Microbiological Risk Assessment (sQMRA) model**

129 The sQMRA-model model was developed by Evers & Chardon, (2010). It is implemented in a
130 Microsoft Excel spreadsheet. Deviating from a full-scale Quantitative Microbiological Risk
131 Assessment (QMRA), where pathogen numbers are followed through the whole food chain,
132 this model starts at retail and ends with the number of human cases of illness. The model is
133 deterministic and includes cross-contamination and preparation (heating) in the kitchen and
134 as well as dose–response relationship. The general setup of the sQMRA tool consists of
135 consecutive questions for values of each of the 11 parameters, always followed by
136 intermediate model output broken down into categories of contamination, cross-
137 contamination and preparation, as show in Figure 2 under the results section. Model input
138 and output are summarized and exposure as well as cases are attributed to the
139 distinguished categories. As a relative risk measure, intermediate and final model outputs
140 are always compared with results from a full-scale QMRA of *Campylobacter* on chicken fillet
141 as shown in Figure 3, 4 and 5 under the results section. The model allows results of the
142 research to be quickly interpreted in terms of public health risk, given that pathogen
143 concentration is determined from the model. It is also more accessible and understandable
144 for scientists that are new to the QMRA research area or are not very mathematically
145 inclined (Evers & Chardon, 2010)

146 **2.3 Study design and Data sources**

147 The study used a cross sectional design which depended on both secondary and primary
148 data sources.

149 **Secondary data:** This was a risk analysis desktop study which mainly depended on review of
150 scientific peer reviewed papers and grey literature (secondary data). The literature review
151 was guided by research questions based on the sQMRA model as shown in Table 1.
152 Literature search was conducted on major electronic databases including Web of Science
153 and Pub Med (NLM) using The Norwegian University of Life sciences (NMBU) library

154 database. Further, grey literature from conference proceedings and reports from
155 government institutions and Non-Governmental Organisations were obtained online using
156 “Google search engine” and “Google scholar”. Search of key terms such as, “Beef
157 consumption, Quantitative risk assessment, Salmonellosis, beef value chain, Zambia”,
158 were used. Guided by questions in table 1, literature which contained relevant data were
159 included in the study and the rest were excluded. This was the main source of data (almost
160 98%).

161 **Primary data:** After an extensive literature review, it was discovered that there were
162 information gaps on serving portions and consumption patterns of beef in Zambia. A survey
163 was therefore undertaken to fill these information gaps. This only formed about 2% of the
164 data.

165 A structured questionnaire was used to address the information gaps on serving portions
166 and consumption patterns. The study had a convenient sample size of hundred (100)
167 respondents. The sampling frame was composed of respondents from two areas with a
168 different socio-economic status (40 low and 60 medium income communities), so as to
169 obtain a representative estimate of average serving portions and consumption patterns.
170 Residential areas were used as a proxy for socioeconomic status using the Central Statistical
171 Office conditions of leaving survey (CSO, 2010; Mweemba & Webb, 2008). Respondents
172 were conveniently identified and interviewed from the butcheries in low and medium/high
173 cost residential areas where they were found buying beef and restaurants where they were
174 found eating beef.

175 **2.4 Ethical Approval**

176 Ethical approval was sought and approved from the School of Veterinary Medicine Board of
177 Graduate Studies Committee and the University Of Zambia, Directorate of Graduate Studies
178 (DRGS).

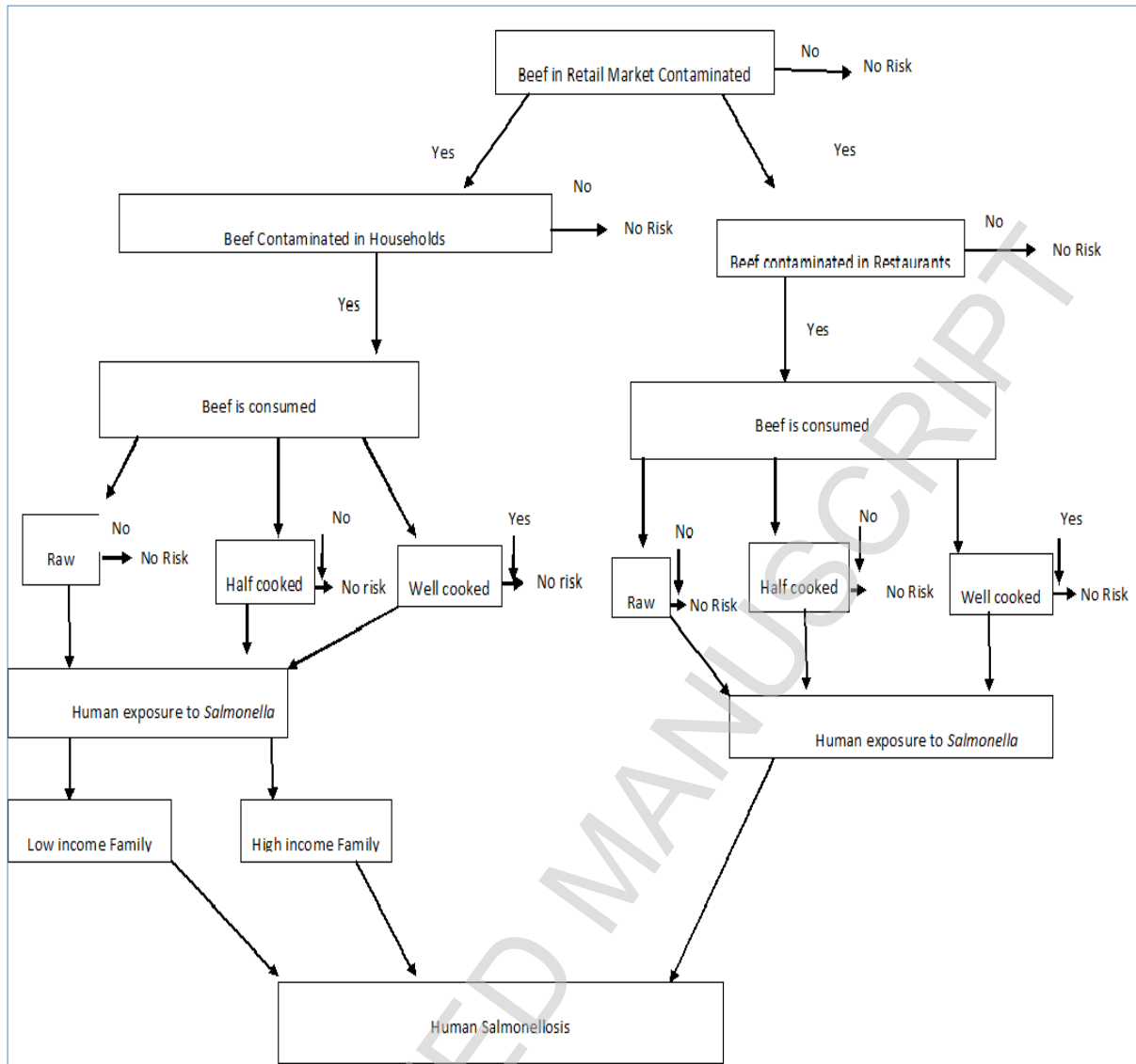
179 **Table 1: Literature review guide based on SQMRA model**

<p>Case definition</p> <ol style="list-style-type: none"> 1. What is the pathogen of interest? 2. What is the food product of interest? 3. What is the population size? 4. What are the population characteristics? 5. What is the consumption period? 	<p>Consumption data</p> <ol style="list-style-type: none"> 1. How many portions are consumed in the population per consumption period? 2. What is the average size of one portion? 3. What percentage of the portions is contaminated at retail? 4. What is the average concentration of colony forming units (cfu) per gram in contaminated portions?
<p>Kitchens cross contamination</p> <ol style="list-style-type: none"> 1. Given contaminated portions, what percentage of the portion will contaminate the environment? E.g. hands and kitchen equipment? 2. Given contaminated portions, what percentage of the cfu's on a portion will contaminate the environment? E.g. hands and kitchen equipment? 3. Given cross contamination, what percentage of cfu's in the environment ends up being ingested? 	<p>Kitchen preparation</p> <ol style="list-style-type: none"> 1. What percentage of the portions is prepared; Done, Half done, Raw 2. What percentage of cfu's on a portion will survive during preparation? -Done, Half done and Raw <p>Infection and illness</p> <ol style="list-style-type: none"> 1. At which dose (number of cfu's) per portion will half of the exposed population get infected? 2. What percentage of infected people will get ill?
<p>Cfu = colony forming units</p>	

180

181 **2.5 Data management and analysis**

182 The data collected from the survey were coded and entered into STATA, SE/ 12 for Windows
 183 (StaaCorp, College Station, TX). Descriptive statistics on average serving portions,
 184 consumption patterns, and kitchen preparation methods of beef were calculated. Data from
 185 the literature review were entered in the Excel version of the sQMRA model developed by
 186 Evers and Chardon, (2010). This model was then run twelve times to come up with results
 187 for the exposure assessment following the household and restaurant risk exposure
 188 pathways as shown in figure 1.



189

190 Figure 1: Conceptual framework of the household and restaurant risk exposure pathways

191

192 3.0 RESULTS

193 3.1 Exposure assessment

194 3.1.1 Case definition

195 The pathogen of interest was *Salmonella* species and the targeted product was beef. The
196 population size of Lusaka province was taken to be 2,669,249 in this model according to
197 Central Statistics Office of Zambia (CSO, 2015). A consumption period of one year was
198 defined to assess the number of people who would get ill in this study (i.e., the number of
199 people who would get ill per year).

200 3.1.2 Consumption data

201 In this study, a portion size was defined as the amount/ size of beef an individual consumes
202 per meal. There was no available beef consumption data in Lusaka province. The study
203 assumed that residents in Lusaka province who were employed consumed beef. According
204 to Labour Force Survey of Zambia, 75% of the 2.67 million Lusaka residents were in formal
205 or informal employment (CSO, 2015). Using the later information, the study therefore
206 logically assumed that 75% of the residents in Lusaka province who were employed
207 consumed beef because of their purchasing power (CSO, 2015; World Bank, 2011). The
208 survey revealed that two portions of beef were served (Lunch and dinner). Hence the
209 number of portions consumed by a population was calculated to be 2,001,937 per
210 consumption period multiplied by 2 servings for lunch and dinner (4,003,874 portions).

211 3.1.3 Serving portions and consumption patterns

212 The results of the survey revealed that the average serving portion of beef per serving at a
213 household level was 60g among low consumers and 83.1g among medium beef consumers,
214 while that for restaurants (high beef consumers) was 192g. Most beef at the household level
215 was prepared and consumed well done (91%); 9% was prepared half done; while no (0%)
216 beef was consumed raw. The consumption patterns from the data showed that 60% of
217 respondents consumed beef once every week, 16% consumed once in every fortnight, 15%
218 consumed beef once a month and 9% consumed beef every day through various forms. At
219 household level, beef was cooked once, but then served in two different periods (2 serving
220 portions-lunch and dinner).

221 **Contamination of raw beef at retail outlets:** Literature review showed a wide range of raw
222 beef contamination at retail outlets from 2.42% to 62% (Ahmad et al., 2013; Kumar, Rao, &
223 Haribabu, 2014; Mrema, Mpuchane, & Gashe, 2006; Sallam, Mohammed, Hassan, &
224 Tamura, 2014; Tafida et al., 2013; Van, Moutafis, Istivan, Tran, & Coloe, 2007; Yang et al.,
225 2010). A similar study in Botswana revealed that retail contamination of beef stood at 20%
226 (Mrema et al., 2006). This study therefore used the data from Botswana because it is a
227 neighbouring country with similar experiences in retail beef handling practices like in many
228 other low and middle income countries in Africa (Haileselassie et al., 2013; Mrema et al.,
229 2006).

230 This study considered only minimum and high concentrations of *Salmonella* and hence the
231 average concentration of colony forming units (cfu) per gram in a contaminated portion of
232 beef was taken to have a minimum value of 3.36cfu/g and a maximum value of 12cfu/g
233 (Ahmad et al., 2013; Ba'aba, 2014; USA -FSIS, 2011).

234 **3.1.4 Kitchen cross-contamination**

235 Due to a lack of literature on *Salmonella* in beef kitchen cross-contamination, *Salmonella* in
236 chicken kitchen cross-contamination was used as a proxy. This is because cross-
237 contamination does not differ regardless of the food product where preparation methods
238 are similar (Evers & Chardon, 2010). The percentage of portions that would contaminate the
239 environment such as the hands and kitchen was therefore set at 45% for restaurants and
240 40% under the household risk exposure pathways (Medeiros, Nascimento, & Robson, 2014).
241 The percentage of cfu on a portion that would contaminate the environment such as hands
242 and kitchen was 30% (Kusumaningrum *et al.*, 2003).

243 The percentage of beef portions that would cross-contaminate the environment such as the
244 hands and household kitchen used in this model was assumed to range from 4 to 32% (12%
245 of dishcloths, 24% of persons' hands, 4% refrigerator door handles, 20% oven door handles,
246 24% counter-tops and 32% draining boards) (Gorman *et al.*, 2002), while the percentage of
247 cfu on a beef portion that would contaminate the environment such as the hands and
248 kitchen in household was assumed to be 16.6% (Gorman *et al.*, 2002). In the household and
249 restaurant risk pathways, it was assumed that 9% and 14% of cfu (value ranges from 0.02 to
250 75%) on a portion would end up being ingested as a result of beef that is prepared half done
251 (Ravishankar *et al.*, 2010).

252 **3.1.5 Kitchen preparation**

253 From the questionnaire survey on beef preparation, the percentage of doneness on the
254 portion of beef at household kitchen level was; 91% well done, 9% half done and 0% raw,
255 while that at restaurant kitchen level was 84% well done, 16% half done (mostly roasted T-
256 bone) and 0% raw. In the reviewed literature the percentage of beef prepared raw was high
257 at 37% (Bogardet al., 2013) which was not realistic to African cultures like that of Zambia.

258 The percentages of microorganisms surviving on a contaminated portion of beef during
259 preparation in both household and restaurant kitchen were 0%, 20% and 100% when beef
260 was prepared well done, half done and raw respectively (Evers & Chardon, 2010). It was
261 assumed to be zero when well done because of overboiling of meat which is normally
262 practiced in Zambia; and 100% when raw due to poor hygiene practices along the beef value
263 chain in developing countries (Haileselassie et al., 2013). Evers & Chardon, (2010) also used
264 0% in well done and 100% when prepared raw, in their sQMRA model.

265 **3.1.6 Infection and illness**

266 In this study, the dose (number of cfu's) per gram of portion that would cause half of the
267 exposed population to get salmonella infection (ID₅₀) was taken to be a minimum of $9.61 \times$
268 10^3 cfu (9,610) and maximum of 5.0×10^4 (Teunis et al., 2010; WHO/FAO, 2002). The study
269 assumed that 100% of the exposed population would get ill when they ingested such doses
270 of *Salmonella* (Blaser & Newman, 1982). The infectious dose of *Salmonella* was assumed to
271 be a minimum of 9.61×10^3 cfu/g and a maximum of 5×10^4 cfu/g (Teunis et al., 2010). The
272 average concentration of cfu's per gram in a contaminated portion of raw beef was a
273 minimum of 3.36 cfu/g and maximum 12 cfu/g (Ahmad et al., 2013; Ba'aba, 2014; Teunis,
274 1997; USA -FSIS, 2011; WHO/FAO, 2002).

275 **3.2 Risk characterisation**

276 A total of 12 simulations which included eight from the household risk pathway (4 for the
277 low beef consumers, 4 for medium beef consumers) and 4 for the restaurant (high beef
278 consumers) risk exposure pathway, were run. Each run produced a summary of the input
279 parameters (Figure 2) and the output model results for the highest risk of developing
280 salmonellosis among the low beef consumers (Figure 3) and medium beef consumers

281 (Figure 4) in a household risk pathway and high beef consumers in a restaurant risk pathway
 282 (Figure 5).

283

INPUT PARAMETERS			
Pathogen:		Salmonella	
Food product:		Meat	
Population size:		2669249	
Pop. Characteristics:		Population of Lusaka	
Consumption period:		One year	
Number	Para-meter	Question	Value
1	N	Portions consumed	4.0E+06
2	M	Portion size in grams	60
3	Sr/+	Prevalence in retail	20%
4	Cr/+	cfu per gram contaminated product	12.0
5	Scc/r	Portions causing cross. cont.	45%
6	Fcc	cfu's from portions to environment	30%
7	Fei	cfu's from environment to ingestion	9.0%
8	Sprd/cc	Portions prepared done	91%
8	Sprh/cc	Portions prepared half-done	9.0%
8	Sprr/cc	Portions prepared raw	0.000%
9	Fprd	cfu's surviving when prep. Done	0%
9	Fprh	cfu's surv. when prep. Half-done	20%
9	Fprr	cfu's surviving when prep. Raw	100%
10	ID50	ID50 (number of cfu's)	9.6E+03
11	Pill/inf	% people infected who get ill	100%
Time stamp:		04/07/2016 16:49	
sQMRA-tool			

284 Figure 2: sQMRA input parameters for the low beef consumer under the household risk
 285 exposure pathway

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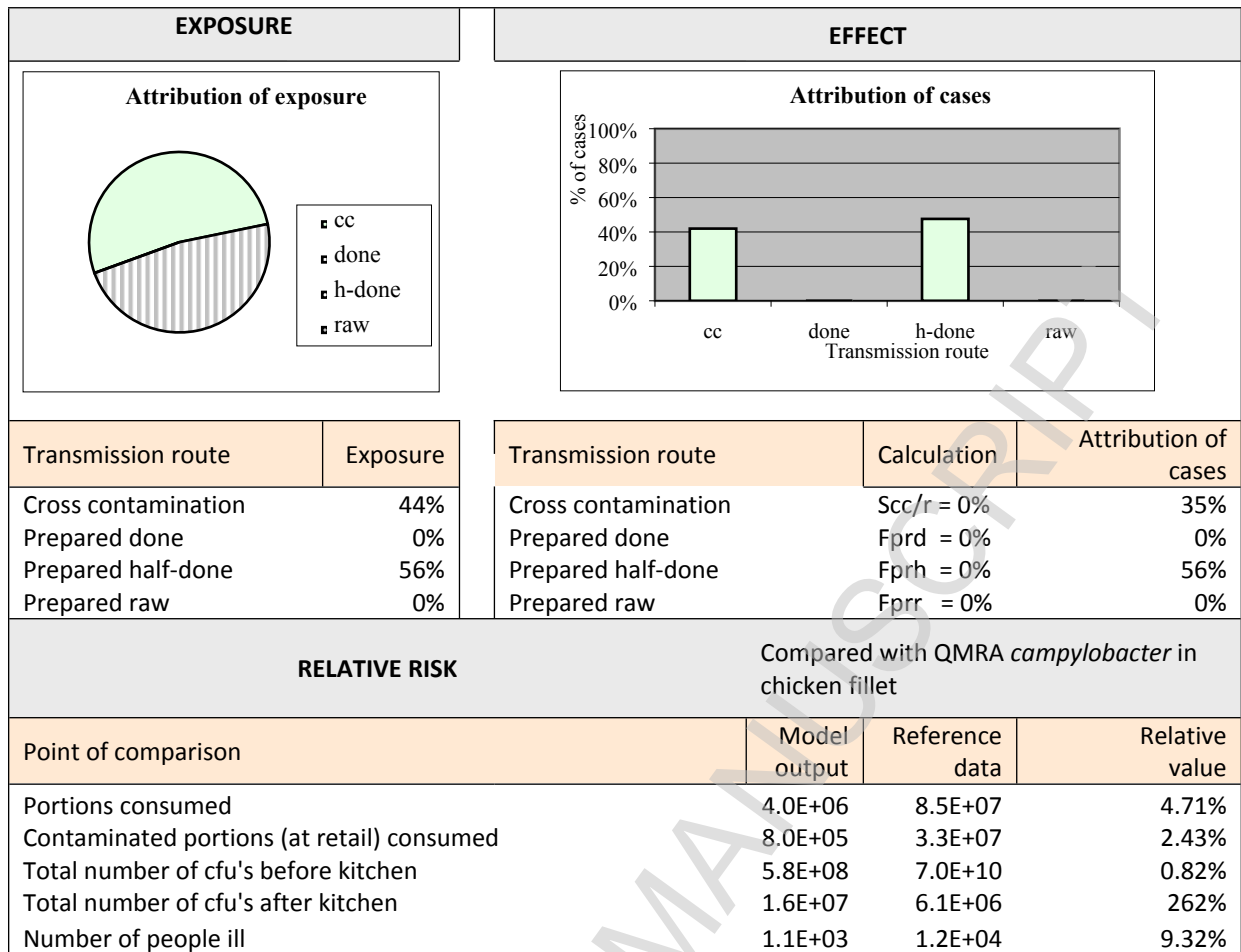
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295 Figure 3: Model output at 12cfu/g and ID50 at 9.61×10^3 cfu (high probability for low beef
 296 consumers under the household risk exposure pathway)

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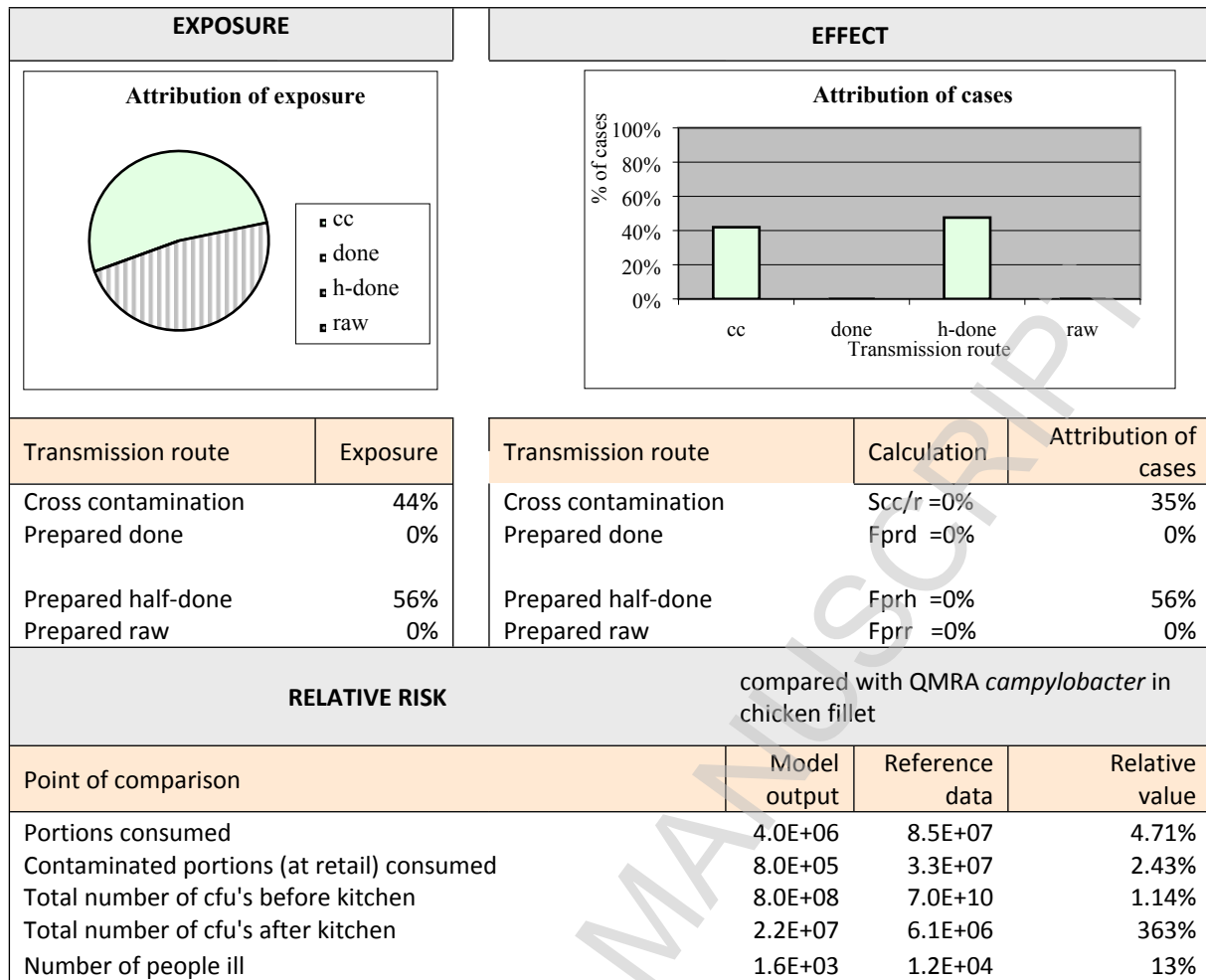
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310 Figure 4: Model output at 12cfu/g and ID50 at 9.61×10^3 cfu (high probability for medium
 311 beef consumers under the household risk exposure pathway)

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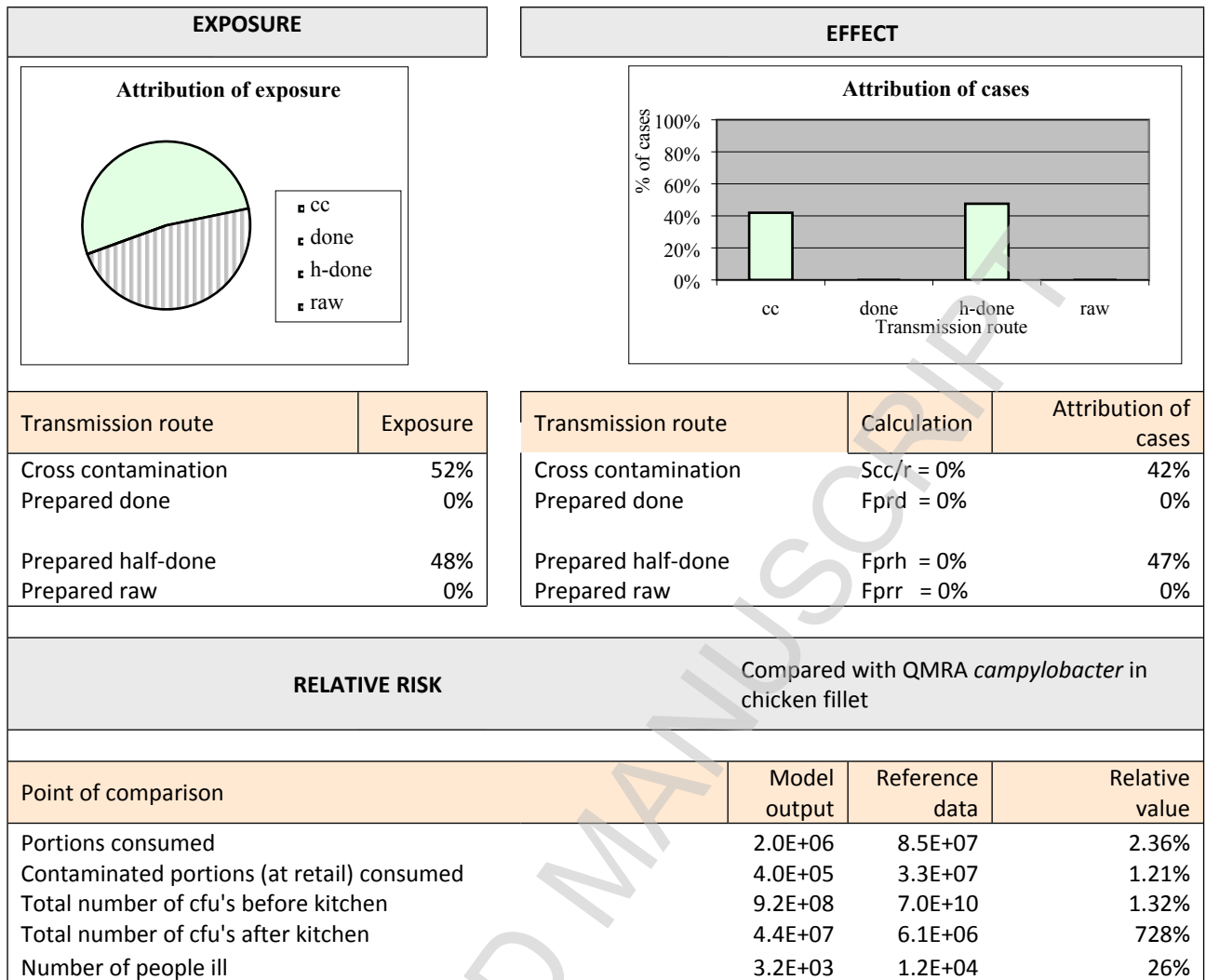
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325 Figure 5: Model output at 12cfu/g and ID50 at 9.61×10^3 cfu (high probability for the high
 326 beef consumers under the restaurant risk exposure pathway)

327

328 Table 2 (risk characterization) summarises the results of all the outputs of the 12
 329 simulations. Of the 4 case scenarios for the low beef consumers (through the household risk
 330 pathway), scenario 3 recorded the highest risk with 1,100 out of a population of 2,001,937
 331 people developing salmonellosis through the consumption of *Salmonella* contaminated
 332 beef, representing a probability of 0.04%.

333 Among the medium beef consumers through the household risk pathway, 1,600 out of a
 334 population of 2,001,937 people risked developing salmonellosis through consumption of
 335 salmonella contaminated beef, representing a probability of 0.05%.

336 Among the heavy consumers of beef (through the restaurant) risk pathway, 3,200 out of a
 337 population of 2,001,937 people risked developing salmonellosis through consumption of
 338 salmonella contaminated beef, representing a probability of 0.16%.

339

340 **Table 2: Summary of the outputs of 12 simulations under household and restaurant risk**
 341 **exposure pathways**

Low beef consumers under household risk pathway				
Scenario	Portion (g)	cfu/g	ID50	Model output (No. People ill per year)
1	60	3.36	9,610	320
2	60	3.36	50,000	62
3	60	12	9,610	1,100
4	60	12	50,000	220
Medium beef consumers under household risk pathway				
1	83.1	3.36	9,610	450
2	83.1	3.36	50,000	86
3	83.1	12	9,610	1,600
4	83.1	12	50,000	310
High beef consumers under restaurant risk pathway				
1	192	3.36	9,610	890
2	192	3.36	50,000	170
3	192	12	9,610	3,200
4	192	12	50,000	610

342

343

344

345

346

347 3.2.1 Uncertainty

348 Like many other risk analysis studies, there were substantial missing data as input
349 parameters in the model. To cover up for these data gaps, a simple survey on the
350 consumption patterns and serving portions of beef in the population was done to get the
351 average serving portions, so as avoid too much reliance on logical assumptions and use of
352 data from other countries. The pathogen numbers were followed through the food chain,
353 which in this case starts at retail and ends with the number of human cases of illness. It
354 would be more robust to follow the pathogen numbers along the entire value chain (farm to
355 folk at a national level), but this would require more resources.

356

357 4.0 Discussion

358 This study was conducted with the aim of assessing the risk of developing salmonellosis
359 through consumption of beef in Lusaka Province of Zambia. The key question was to find
360 out whether beef sold in Lusaka province posed a risk of *Salmonella* infection through
361 consumption of meals prepared at home and those consumed in restaurants. In this study,
362 it was observed that the risk of developing salmonellosis as a result of beef consumption
363 was generally low for both exposures from restaurants and in households. The low risk in
364 the current study was attributed to low serving portions per meal, low consumption
365 patterns and preparation methods of beef both in restaurants and in households.

366 The serving portion of beef has the potential to contribute to risk of *Salmonella* infection in
367 humans. In this study, the average serving portion of beef was 60g and 83.1g per meal for
368 low and middle income households and 192g/meal in restaurants. This contributed to low
369 risks found in this study. The small serving portions could be attributed to the high price of
370 beef on the market and hence most people opted for other livestock products rather than
371 beef. This is in agreement with the previous findings on urban consumption patterns of
372 livestock products in Zambia where consumption patterns of livestock products was
373 influenced by household affluence defined as the low, medium, and high expenditure
374 terciles or income groups (Hichaambwa, 2012). In the same study Hichaambwa showed that
375 within each city, the expenditure share of livestock products increased from the low to the
376 high income group while it marginally decreased in the case of fish (Hichaambwa, 2012).

377 In terms of preparation methods, most of the beef consumed in Lusaka was prepared well
378 done through boiling with only few (16%) in restaurants where T-bone was normally
379 prepared half done. Consumption of T-bone contributed to doubling the risk of developing
380 salmonellosis in the current study through the restaurant pathway. Consumption of raw
381 beef was not a common practice in Zambia hence recording 0% and thus further reducing
382 the risk.

383 Although consumption of well cooked beef does not pose a risk of developing salmonellosis,
384 other ways of getting infection with *Salmonella* is cross-contamination in the kitchen which
385 could occur when handling contaminated beef. Lordache and Tofan (2008) in a study on the
386 cross-contamination of *Salmonella enteritidis* on sterile and non-sterile meat showed that
387 cross-contamination of *Salmonella* could occur in the kitchen environment (Lordache and
388 Tofan, 2008). In the current study, cross-contamination in the kitchen was one of the
389 contributing factors for risk of developing salmonellosis. Results showed that much of the
390 risk was contributed by cross contamination at restaurant level compared to other scenarios
391 when concentration of *Salmonella* in retail beef was 12cfu/g of beef and infectious dose fifty
392 of (ID50) 9.61×10^3 cfu/g. This was in agreement with the observation by Mughini-Gras et al.,
393 (2014) who showed that not using a chopping board for raw meat only (cross-
394 contamination) and consuming raw/undercooked meat were risk factors for infection with
395 *Salmonella* originating from cattle. In the current study, there were low numbers of
396 predicted cases of salmonellosis at high contamination (12cfu/g) and high ID50 (5×10^4
397 cfu/g). This indicated that cooking alone cannot be considered an adequate response to
398 exceptional events of extreme foodborne bacterial pathogen contamination; other factors
399 like cross-contamination could lead to salmonellosis infection even when beef is well
400 cooked (Teunis et al., 2010).

401 In general, the low risk of developing salmonellosis in the current study is in agreement with
402 the observation by Abdunaser *et al.* (2009) who reported the risk of developing
403 salmonellosis in human per 100g serving portion of ground beef to be low (ranging from 0
404 to 2.33×10^{-06}), though it was based on ground beef contrary to the current study which
405 considered beef without specifying whether ground or beef parts.

406 We acknowledge that this model is deterministic and does not allow the variability
407 inherently linked to food-borne diseases to be modelled. However, our model could be a

408 starting platform for further studies on the epidemiology of salmonellosis in Zambia. The
409 model also represents a way of communicating results across regional and cultural/
410 economic borders.

411

412 **Conclusion**

413 The risk of developing salmonellosis from consumption of contaminated beef is generally
414 very low among the beef consumers in Lusaka. This was attributed to low beef consumption
415 and adequate cooking methods.

416

417 **Declaration of interest**

418 Authors declare no conflict of interest.

419

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427 **5.0 References**

- 428 Abdunaser, D., Almabrouk, F., Ashraf, W., Yves, M., Olivier, C., & Moez, S. (2009).
429 Quantitative risk assessment of human salmonellosis linked to the consumption of
430 ground beef. *Iraqi Journal of Veterinary Sciences*, 23(Suppl.2), En263-En273.
- 431 Abouzeed, Y. M., Hariharan, H., Poppe, C., & Kibenge, F. S. B. (2000). Characterization of
432 Salmonella isolates from beef cattle, broiler chickens and human sources on prince
433 Edward Island. *Comparative Immunology, Microbiology and Infectious Diseases*, 23(4),
434 253–266. [http://doi.org/10.1016/S0147-9571\(99\)00079-X](http://doi.org/10.1016/S0147-9571(99)00079-X)
- 435 Ahmad, M. U. D., Sarwar, A., Najeeb, M. I., Nawaz, M., Anjum, A. A., Ali, M. A., & Mansur, N.
436 (2013). Assessment of microbial load of raw meat at abattoirs and retail outlets.
437 *Journal of Animal and Plant Sciences*, 23(3), 745–748.
- 438 Ba'aba, A. I. (2014). *Bacterial load and isolation of salmonella species from cattle carcasses*
439 *at Kano abattoir- Kano State, Nigeria*. Ahmadu Bello University Zaria– Ngeria.
- 440 Blaser, M. J., & Newman, L. S. (1982). A review of human salmonellosis: I. Infective dose.
441 *Reviews of Infectious Diseases*, 4(6), 1096–1106. <http://doi.org/10.2307/4452912>
- 442 Bogard, A. K., Fuller, C. C., Radke, V., Selman, C. A., & Smith, K. E. (2013). Ground Beef
443 Handling and Cooking Practices in Restaurants in Eight States 3. *Journal of Food*
444 *Protection*, 76(12), 2132–2140. <http://doi.org/10.4315/0362-028X.JFP-13-126>
- 445 Carrasco, E., Morales-Rueda, A., & García-Gimeno, R. M. (2012). Cross-contamination and
446 recontamination by Salmonella in foods: A review. *Food Research International*, 45(2),
447 545–556. <http://doi.org/10.1016/j.foodres.2011.11.004>
- 448 CSO. (2010). *Central Statistical Office Living Conditions Monitoring Survey*. Retrieved from
449 <http://www.zamstats.gov.zm/report/Lcms/2006-2010 LCMS Report Final Output.pdf>
- 450 CSO. (2015). *Central Statistical Office Labour Force Survey-Republic of Zambia*.
- 451 Dhanoa, A., & Fatt, Q. K. (2009). Non-typhoidal Salmonella bacteraemia: epidemiology,
452 clinical characteristics and its' association with severe immunosuppression. *Annals of*
453 *Clinical Microbiology and Antimicrobials*, 8, 15. <http://doi.org/10.1186/1476-0711-8-15>
- 454 Evans, B. R., & Leighton, F. A. (2014). A history of One Health. *Scientific and Technical*
455 *Review of the Office of International Epizootics*, 33(2), 413–420.
- 456 Evers, E. G., & Chardon, J. E. (2010). A swift Quantitative Microbiological Risk Assessment
457 (sQMRA) tool. *Food Control*, 21(3), 319–330.

- 458 <http://doi.org/10.1016/j.foodcont.2009.06.013>
- 459 Freitas Neto, O. de, Penha Filho, R., Barrow, P., & Berchieri Junior, A. (2010). Sources of
460 human non-typhoid salmonellosis: a review. *Revista Brasileira de Ciência Avícola*.
461 <http://doi.org/10.1590/S1516-635X2010000100001>
- 462 Gorman, R., Bloomfield, S., & Adley, C. C. (2002). A study of cross-contamination of food-
463 borne pathogens in the domestic kitchen in the Republic of Ireland. *International*
464 *Journal of Food Microbiology*, 76(1–2), 143–150. [http://doi.org/10.1016/S0168-](http://doi.org/10.1016/S0168-1605(02)00028-4)
465 [1605\(02\)00028-4](http://doi.org/10.1016/S0168-1605(02)00028-4)
- 466 Haileselassie, M., Taddele, H., Adhana, K., & Kalayou, S. (2013). Food safety knowledge and
467 practices of abattoir and butchery shops and the microbial profile of meat in Mekelle
468 City, Ethiopia. *Asian Pacific Journal of Tropical Biomedicine*, 3(5), 407–412.
469 [http://doi.org/10.1016/S2221-1691\(13\)60085-4](http://doi.org/10.1016/S2221-1691(13)60085-4)
- 470 Hang'ombe, B. M., Ulaya, W., Mwansa, J. C. L., Mubita, C., Isogai, N., Mulenga, E., ... Isogai,
471 E. (2011). Feasibility of using dot blot hybridization to detect *Salmonella* invA, spiC and
472 sipC directly from clinical specimens. *African Journal of Microbiology Research*, 5(6),
473 582–585. <http://doi.org/10.5897/AJMR10.478>
- 474 Hangombe, B. M. (1998). Incidence and characterisation of *Salmonella* Enteritidis in Poultry
475 products and human diarrhoea cases in Lusaka District, Zambia.
- 476 Hichaambwa, M. (2012). *Urban Consumption Patterns of Livestock Products in Zambia and*
477 *Implications for Policy by Munguzwe Hichaambwa Urban Consumption Patterns of*
478 *Livestock Products in Zambia and Implications for Policy*.
- 479 Iordache, L., & Tofan, C. (2008). The cross-contamination of *Salmonella* enteritidis on sterile
480 and non-sterile meat. *Journal of Agroalimentary Processes and Technologies*, 14, 337–
481 340.
- 482 Isogai, E., Silungwe, M., Sinkala, P., Chisenga, C., Mubita, C., Syakalima, M., ... Yasuda, J.
483 (2005). Rapid detection of *Salmonella* on commercial carcasses by using isothermal and
484 chimeric primer-initiated amplification of nucleic acids (ICAN)-enzyme-linked
485 immunosorbent assay (ELISA) in Zambia. *International Journal of Applied Research in*
486 *Veterinary Medicine*, 3(4), 367–371.
- 487 Jackson, B. R., Griffin, P. M., Cole, D., Walsh, K. A., & Chai, S. J. (2013). Outbreak-associated
488 salmonella enterica serotypes and food commodities, United States, 1998–2008.
489 *Emerging Infectious Diseases*, 19(8), 1239–1244.

- 490 <http://doi.org/10.3201/eid1908.121511>
- 491 Kagambèga, A., Lienemann, T., Aulu, L., Traoré, A. S., Barro, N., Siitonen, A., & Haukka, K.
492 (2013). Prevalence and characterization of *Salmonella enterica* from the feces of cattle,
493 poultry, swine and hedgehogs in Burkina Faso and their comparison to human
494 *Salmonella* isolates. *BMC Microbiology*, *13*, 253. [http://doi.org/10.1186/1471-2180-13-](http://doi.org/10.1186/1471-2180-13-253)
495 253
- 496 Kemal, J. (2014). A Review on the Public Health Importance of Bovine Salmonellosis. *Journal*
497 *of Veterinary Science & Technology*, *5*(2). <http://doi.org/10.4172/2157-7579.1000175>
- 498 Kumar, P., Rao, J., & Haribabu, Y. (2014). Microbiological Quality of Meat Collected from
499 Municipal Slaughter Houses and Retail Meat Shops from Hyderabad Karnataka Region,
500 India. *APCBEE Procedia*, *8*(Caas 2013), 364–369.
501 <http://doi.org/10.1016/j.apcbee.2014.09.001>
- 502 Kusumaningrum, H. D., Riboldi, G., Hazeleger, W. C., & Beumer, R. R. (2003). Survival of
503 foodborne pathogens on stainless steel surfaces and cross-contamination to foods.
504 *International Journal of Food Microbiology*, *85*(3), 227–236.
505 [http://doi.org/10.1016/S0168-1605\(02\)00540-8](http://doi.org/10.1016/S0168-1605(02)00540-8)
- 506 Lubungu, M., Sitko, N. J., & Hichaambwa, M. (2015). *Analysis of Beef Value Chain in Zambia :
507 Challenges and Opportunities of Linking Smallholders to Markets by Indaba Agricultural
508 Policy Research Institute (IAPRI) Analysis of Beef Value Chain in Zambia : Challenges
509 and Opportunities of Linking Smallholde*. Retrieved from
510 <http://fsg.afre.msu.edu/zambia/wp103.pdf>
- 511 Medeiros, D. G. G. D. A., Nascimento, M. C. L. R. de R. Do, & Robson, E. F. M. (2014).
512 *Salmonella* spp . detection in chicken meat and cross- contamination in an industrial
513 kitchen. *African Journal of Microbiology Research*, *8*(11), 1130–1139.
514 <http://doi.org/10.5897/AJMR2013.6487>
- 515 Mrema, N., Mpuchane, S., & Gashe, B. A. (2006). Prevalence of *Salmonella* in raw minced
516 meat , raw fresh sausages and raw burger patties from retail outlets in Gaborone ,
517 Botswana. *Journal of Food Control*, *17*, 207–212.
518 <http://doi.org/10.1016/j.foodcont.2004.09.019>
- 519 Mughini-Gras, L., Enserink, R., Friesema, I., Heck, M., Van Duynhoven, Y., & Van Pelt, W.
520 (2014). Risk factors for human salmonellosis originating from pigs, cattle, broiler
521 chickens and egg laying hens: A combined case-control and source attribution analysis.

- 522 *PLoS ONE*, 9(2). <http://doi.org/10.1371/journal.pone.0087933>
- 523 Muma, J. B. (1998). *Application of Hazard Analysis Critical Control Point(HACCP) concept to*
524 *study cattle slaughterhouse hygiene and carcass contamination in Zambia*. University of
525 Zambia.
- 526 Mwansa, J., Mutela, K., Zulu, I., Amadi, B., & Kelly, P. (2002). Antimicrobial sensitivity in
527 enterobacteria from AIDS patients, Zambia. *Emerging Infectious Diseases*, 8(1), 92–93.
528 <http://doi.org/10.3201/eid0801.010018>
- 529 Mweemba, M. A. J., & Webb, E. (2008). Residential area as proxy for socio-economic status,
530 paediatric mortality and birth weight in Lusaka, Zambia. *Journal of Tropical Pediatrics*,
531 54(6), 406–409. <http://doi.org/10.1093/tropej/fmn041>
- 532 Ndalama E , Mdegela RH, N. (2013). Assessment of hygienic practices and faecal
533 contamination of beef at Vingunguti slaughterhouse in Dar es salaam, Tanzania.
534 *Tanzania Veterinary Journal*, 28(2), 23–29.
- 535 NTANGA, PIUS, D. (2013). *Assessment Of Microbial Contamination In Beef From Abattoir to*
536 *Retail Meat Outlets In Morogoro Municipality, Tanzania*. Sokoine University of
537 Agriculture.
- 538 Plym L, F., & Wierup, M. (2006). Salmonella contamination: a significant challenge to the
539 global marketing of animal food products. *Revue Scientifique et Technique*
540 *(International Office of Epizootics)*, 25(2), 541–554.
- 541 Ravishankar, S., Zhu, L., & Jaroni, D. (2010). Assessing the cross contamination and transfer
542 rates of Salmonella enterica from chicken to lettuce under different food-handling
543 scenarios. *Journal of Food Microbiology*, 27(6), 791–794.
544 <http://doi.org/10.1016/j.fm.2010.04.011>
- 545 Sallam, K. I., Mohammed, M. A., Hassan, M. A., & Tamura, T. (2014). Prevalence, molecular
546 identification and antimicrobial resistance profile of salmonella serovars isolated from
547 retail beef products in mansoura, egypt. *Journal of Food Control*, 38(1), 209–214.
548 <http://doi.org/10.1016/j.foodcont.2013.10.027>
- 549 Sinkala, Y., Simuunza, M., Pfeiffer, D. U., Munang, H. M., Mulumba, M., Kasanga, C. J., ...
550 Mweene, A. S. (2014). Challenges and Economic Implications in the Control of Foot and
551 Mouth Disease in Sub-Saharan Africa : Lessons from the Zambian Experience.
552 *Veterinary Medicine International*, 2014(Article ID 373921), 12.
553 <http://doi.org/http://dx.doi.org/10.1155/2014/373921>

- 554 Tafida, S. Y., Kabir, J., Kwaga, J. K. P., Bello, M., Umoh, V. J., Yakubu, S. E., ... Hendriksen, R.
555 (2013). Occurrence of Salmonella in retail beef and related meat products in Zaria,
556 Nigeria. *Food Control*, 32(1), 119–124. <http://doi.org/10.1016/j.foodcont.2012.11.005>
- 557 Teunis, P. F. M. (1997). *Infectiousgastro-enteritis-Opportunities for dose response modelling*.
558 Teunis, P. F. M., Kasuga, F., Fazil, A., Ogden, I. D., Rotariu, O., & Strachan, N. J. C. (2010).
559 Dose-response modeling of Salmonella using outbreak data. *International Journal of*
560 *Food Microbiology*, 144(2), 243–249. <http://doi.org/10.1016/j.ijfoodmicro.2010.09.026>
- 561 Ulaya, W. D. (2013). *Determination Of Virulence Factors In Salmonella Isolates Of Human ,*
562 *Poultry And Dod Origin In Lusaka District , Zambia*.
- 563 USA -FSIS. (2011). *National Prevalence Estimate of Pathogens in Domestic Beef*
564 *Manufacturing Trimmings (Trim) December 2005- January 2007*.
- 565 Van, T. T. H., Moutafis, G., Istivan, T., Tran, L. T., & Coloe, P. J. (2007). Detection of
566 Salmonella spp. in retail raw food samples from vietnam and characterization of their
567 antibiotic resistance. *Applied and Environmental Microbiology*, 73(21), 6885–6890.
568 <http://doi.org/10.1128/AEM.00972-07>
- 569 WHO/FAO. (2002). *Risk assessments of Salmonella in eggs and broiler chickens*.
570 *Interpretative summary* (Vol. Microbiolo).
- 571 World Bank. (2011). *What Would it Take for Zambia’s Beef and Dairy Industries to achieve*
572 *their Potential ? Finance and Private Sector Development Unit, Africa Region*. Retrieved
573 from
574 [https://openknowledge.worldbank.org/bitstream/handle/10986/2771/623770ESW0Gr](https://openknowledge.worldbank.org/bitstream/handle/10986/2771/623770ESW0Gray000public00BOX361530B.pdf?sequence=1)
575 [ay000public00BOX361530B.pdf?sequence=1](https://openknowledge.worldbank.org/bitstream/handle/10986/2771/623770ESW0Gray000public00BOX361530B.pdf?sequence=1)
- 576 Yang, B., Qu, D., Zhang, X., Shen, J., Cui, S., Shi, Y., ... Meng, J. (2010). Prevalence and
577 characterization of Salmonella serovars in retail meats of marketplace in Shaanxi,
578 China. *International Journal of Food Microbiology*, 141(1–2), 63–72.
579 <http://doi.org/10.1016/j.ijfoodmicro.2010.04.015>
- 580