A modelling study to estimate the health burden of foodborne disease:

cases, general practice consultations and hospitalisations in the UK, 2009

Sarah J O’Brien1,2

Tricia L Larose3,4

Goutam K Adak5

Meirion R Evans6

Clarence C Tam3,7

On behalf of the Foodborne Disease Attribution Study Group\*

Authors’ affiliations

1 = University of Liverpool Institute of Infection and Global Health

2 = NIHR Health Protection Research Unit in Gastrointestinal Infections

The Farr Institute@HeRC, University of Liverpool, 2nd Floor, Block F, Waterhouse Buildings, 1-5 Brownlow Street, Liverpool, L69 3GL

3 = Department of Infectious Disease Epidemiology, London School of Hygiene and Tropical Medicine, Keppel Street, London, WC1E 7HT

4 = Department of Public Health and General Practice, Norwegian University of Science and Technology, Håkon Jarls gate 11, 7491 Trondheim, Norway

5 = Department of Gastrointestinal, Emerging & Zoonotic Infections, Public Health England Centre for Infectious Disease Surveillance and Control, 61 Colindale Avenue, London, NW9 5EQ

6 = Institute of Primary Care and Public Health, Cardiff University, Temple of Peace and Health, Cathays Park, Cardiff, CF10 3NW

7 = Saw Swee Hock School of Public Health, National University of Singapore, 16 Medical Drive, Singapore 117597, Singapore

\* Additional members are Paul Cook, John M Cowden, Kathryn A Jackson, Brian Smyth

Correspondence to:

Prof Sarah J O’Brien

University of Liverpool

The Farr Institute@HeRC

University of Liverpool

2nd Floor, Block F Waterhouse Buildings

1-5 Brownlow Street

Liverpool

L69 3GL

Tel: +44 151 795 8301

E-mail: [s.j.obrien@liverpool.ac.uk](mailto:s.j.obrien@liverpool.ac.uk)

Key words = Foodborne disease, *Salmonella*, *Campylobacter*, norovirus, *Clostridium perfringens*

Word Count (Main Text) =4,158

**Abstract**

**Objective**: To generate estimates of the burden of UK-acquired foodborne disease accounting for uncertainty.

**Design:** A modelling study combining data from national public health surveillance systems for laboratory-confirmed infectious intestinal disease (IID) and outbreaks of foodborne disease, and two prospective, population-based studies of IID in the community. The underlying datasets covered the time period 1993 to 2008. We used Monte Carlo simulation and a Bayesian approach, using a systematic review to generate Bayesian priors. We calculated point estimates with 95% credible intervals (Crl).

**Setting:** United Kingdom (UK), 2009

**Outcome measures:** Pathogen-specific estimates of the number of cases, general practice consultations and hospitalisations for foodborne disease in the UK in 2009.

**Results:** Bayesian approaches gave slightly more conservative estimates of overall health burden (approximately 511,000 cases versus 566,000 cases). *Campylobacter* is the most common foodborne pathogen, causing 280,400 (95% CrI: 182,503 – 435,693) food-related cases and 38,860 (95% CrI: 27,160 – 55,610) general practice (GP) consultations annually. Despite this, there are only around 562 (95% CrI: 189 – 1,330) food-related hospital admissions due *Campylobacte*r, reflecting relatively low disease severity. *Salmonella* causes the largest number of hospitalisations, an estimated 2,490 admissions (95% CrI: 607 – 9,631), closely followed by *E. coli* O157 with 2,233 admissions (95% CrI: 170 – 32,159). Other common causes of foodborne disease include *Clostridium perfringens*, with an estimated 79,570 cases annually (95% CrI: 30,700 – 211,298), and norovirus with 74,100 cases (95% CrI: 61,150 – 89,660). Other viruses and protozoa ranked much lower as causes of foodborne disease.

**Conclusions:** The three models yielded similar estimates of the burden of foodborne illness in the UK and show that continued reductions in *Campylobacter*, *Salmonella*, *Escherichia coli* O157, *C. perfringens* and norovirus are needed to mitigate the impact of food-borne disease.

Word count = 283

**Article summary:**

**Article focus**

* Food safety is a global public health priority.
* Developing better methods for robust assessment of the burden of foodborne disease has been the focus of international efforts for more than a decade.
* We estimated the burden of foodborne disease using UK datasets and taking account of uncertainty.

**Key messages**

* *Campylobacter, Salmonella, E. coli* O157*, Clostridium perfringens* and norovirus are all important causes of foodborne illness in the UK, with *Salmonella* and *E. coli* O157 accounting for over two-thirds of all hospitalisations.
* The findings should be useful to policymakers in prioritising interventions.

**Strengths and limitations of this study**

* This is the first burden of foodborne illness modelling study to incorporate both empirical data and prior information from a systematic review together with Bayesian methodology for estimating the proportion of IID that is transmitted through contaminated food.
* Our estimates are based on high quality datasets, including directly observed, pathogen-specific incidence data.
* Our methods take full account of parameter uncertainties.
* There are several data gaps which need to be filled including pathogen-specific mortality estimates, and information on morbidity in vulnerable populations such as immunocompromised people, older people, and pregnant women.

## INTRODUCTION

Food safety is a global priority.[1] To have maximum impact, the design and funding of food safety interventions need to take account of the overall burden of foodborne disease and the contribution made by each pathogen. Developing better methods for estimating the true burden of foodborne disease has been the focus of international efforts for over a decade.[1-8] This is problematic for various reasons: people usually present with non-specific symptoms of infectious intestinal disease (IID), only a fraction of cases are confirmed by laboratory testing, and not all are reported to national public health surveillance. IID then needs to be attributed to transmission route (foodborne, waterborne, animal-to-person, person-to-person or environment-to-person), which can be difficult if robust epidemiological information is lacking. In a recent population-based, prospective study in the UK (known as the IID2 Study) we found that IID affected around one in four people each year (approximately 17 million cases in 2009).[9] We used novel methods to estimate, for each pathogen, the proportion of IID attributable to food and the health burden of UK-acquired foodborne disease.

**METHODS**

**Data sources**

The Infectious Intestinal Disease Studies

Two population-based studies of IID have taken place in the UK (Box). The first (IID1 Study) was conducted in England in 1993-6,[9] and the second (IID2 Study) took place across the whole of the UK in 2008-9.[10-11] Both comprised i) a prospective cohort study of people living in the community and ii) a prospective study of patients presenting to general practice with symptoms of IID. Samples were obtained for laboratory testing from symptomatic cases in the cohort and from patients presenting to general practice and tested using comprehensive microbiology algorithms.[11,12] The case definitions were identical in both studies, and incidence rates of IID in the community and general practice (GP) consultation rates for IID were calculated. Data on health care use, captured by questionnaires, gave estimates of hospitalisation rates.

**Box: Sample sizes in the IID1 and IID2 Studies [9,10]**

**IID1, England, August, 1993 – January 1996**

Prospective Cohort Study: N = 9,776

General Practice Presentation Study: N = 4,026

**IID2, United Kingdom, April 2008 – August 2009**

Prospective Cohort Study: N = 7,033

General Practice Presentation Study: N = 991

Outbreak Surveillance Data

The four UK national surveillance centres provided data on general outbreaks of IID occurring between 1 January 2001 and 31 December 2008 (N=2,965) (Table 1). There were substantial changes to outbreak reporting in 2009. Prior to 2009 Public Health England (PHE) collected data on all gastrointestinal infection outbreaks no matter what the transmission route i.e. foodborne, waterborne, person-to-person, environment-to-person and animal-to-person. In 2009 PHE limited the collection of outbreak data on “non-foodborne outbreaks” to “gastrointestinal outbreaks including illnesses associated with recreational water exposure, environmental exposure at outdoor events e.g. contact with mud, contact with animals or their faeces and outbreaks of Vero cytotoxin-producing *Escherichia coli* (VTEC) mediated through person-to-person transmission. Thus the data collected up to December 2008 represented a sub-set of outbreaks rather than all outbreaks. This affected the proportion of illnesses assessed as foodborne because the denominators of outbreaks and cases in outbreaks changed substantially as a result of changes in reporting definitions. This was particularly problematic for pathogens like norovirus and *Cryptosporidium*.

**Table 1: Summary of outbreak data for food attribution by pathogen, UK 2001-2008**

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | FOODBORNE OUTBREAKS | | |  | CASES IN FOODBORNE OUTBREAKS | | |  |
| Organism | Foodborne | All outbreaks | % |  | Cases | All cases | % | Source |
| **Bacteria** |  |  |  |  |  |  |  |  |
| *C. perfringens* | 45 | 60 | 75.0% |  | 1691 | 1964 | 86.1% | Outbreak surveillance |
| *Campylobacter* | 31 | 44 | 70.5% |  | 373 | 761 | 49.0% | Outbreak surveillance |
| *E. coli o157* | 25 | 86 | 29.1% |  | 564 | 1041 | 54.2% | Outbreak surveillance |
| *Listeria* | 2 | 2 | 100.0% |  | 6 | 6 | 100.0% | Outbreak surveillance |
| *Salmonella* | 266 | 308 | 86.4% |  | 7128 | 7892 | 90.3% | Outbreak surveillance |
| *Shigella* | 4 | 11 | 36.4% |  | 65 | 310 | 21.0% | Outbreak surveillance |
| **Protozoa** |  |  |  |  |  |  |  |  |
| *Cryptosporidium* | 4 | 65 | 6.2% |  | 415 | 1375 | 30.2% | Outbreak surveillance |
| *Giardia* | 1 | 7 | 14.3% |  | 106 | 159 | 66.7% | Outbreak surveillance |
| **Viruses** |  |  |  |  |  |  |  |  |
| Adenovirus | -- | -- | -- |  | -- | -- | -- | No outbreaks reported |
| Astrovirus | 0 | 18 | 0.0% |  | 0 | 283 | 0.0% | Outbreak surveillance |
| Norovirus | 61 | 2228 | 2.7% |  | 1500 | 58,855 | 2.5% | Outbreak surveillance |
| Sapovirus | -- | -- | -- |  | -- | -- | -- | No outbreaks reported |
| Rotavirus | 1 | 136 | 0.7% |  | 30 | 2338 | 1.3% | Outbreak surveillance |

For each outbreak, information was available on the following: outbreak setting, number of cases affected, number of cases hospitalised, main mode(s) of transmission, pathogen identified and, for outbreaks involving contaminated foods, the implicated food vehicle (where this was ascertained). For this study, point source or disseminated outbreaks involving contaminated food, and outbreaks involving contaminated food with subsequent person-to-person transmission, were considered to be foodborne. In total, there were 446 outbreaks involving foodborne transmission that were available for analysis.

Systematic Literature Review

### We conducted a systematic literature review according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines [13]. We searched four databases (MEDLINE, EMBASE, Web of Science and FoodBase – the UK Food Standards Agency’s research projects database). The full methodology for our systematic review, and a summary of results have been reported previously [14]. We also compared the list of articles that we identified through the systematic review with a list of case-control studies included in a separate, independently published review of case-control study methods for enteric infection [15]. We identified 32 articles published between 01 January 2001 and 31 December 2011 with relevant information that allowed us to determine the percentage of cases of IID attributable to foodborne transmission (see also Technical appendix). The Bayesian priors were based on uniform distributions, which essentially assume that any value within a specified range is equally likely. The lower and upper bounds of the distribution were determined by the lowest and highest estimates from studies found in the literature review. So for example, the reported range for foodborne *Campylobacter* was between 42% and 80%, and these percentages formed the lower and upper bounds used for the uniform prior (see Technical Appendix).

### Modelling approach

We developed a model to estimate the number of cases, general practice consultations and hospital admissions of UK-acquired foodborne disease due to 13 major enteric pathogens: *Clostridium perfringens, Campylobacter, Escherichia coli* O157, *Listeria, Salmonella* (non-typhoidal)*, Shigella, Cryptosporidium, Giardia,* adenovirus, astrovirus, norovirus, rotavirus and sapovirus. The basic model was:

where *F­p*, *Gp* and *Hp* represent, respectively, the estimated number of UK-acquired foodborne disease cases, GP consultations or hospital admissions for pathogen *p* in 2009. *cp* is the UK incidence of infectious intestinal disease (IID) due to pathogen *p*, and *gp* is the GP consultation rate for IID due to pathogen *p*. The constant, *N*, is the mid-2009 population of the UK. The two parameters, *πp* and *γp*, represent, respectively, the proportion of IID cases due to pathogen *p* that are transmitted through food, and the proportion of cases due to pathogen *p* that are hospitalised. We assumed that foodborne cases were equally likely to consult a GP or be hospitalised as non-foodborne cases.

We used various data sources to inform model parameters. The data available for each pathogen are summarised in the Technical Appendix online (Tables A1 to A3). We used two modelling approaches: a Monte Carlo simulation approach and a Bayesian approach. In the Monte Carlo approach, the parameters *πp* and *γp* were defined by Beta distributions fitted to empirical bootstrap samples of UK outbreak data; in the Bayesian approach, these parameters were modelled as binomial quantities and given priors informed by published studies and hospitalisation data from previous studies in the UK. Model details are given in the Technical Appendix.

### Pathogen-specific rates of IID *(cp , gp)*

We obtained data from the IID2 Study on population incidence and GP consultation rates for IID, and their associated uncertainty, for the above pathogens.[10] For *Shigella*, no cases were found in IID2 so we applied the reporting ratio from IID1 (the ratio of community cases to laboratory-confirmed cases reported to national surveillance) to the number of cases reported in 2009 and divided this by the mid-2009 UK population to obtain the overall shigellosis rate.[11] Similarly, we estimated GP consultation rates by applying the reporting ratio from IID1 (the ratio of GP consultations to laboratory-confirmed cases reported to national surveillance) to the number of laboratory reports in 2009. We accounted for uncertainty in incidence estimates by sampling 100,000 times from the distribution of reporting ratios estimated in IID1. For *Listeria*, no incidence data were available from IID1 or IID2 so we used the number of laboratory reports for listeriosis in 2009 as a conservative population incidence estimate.

### Proportion of cases transmitted through food *(πp)*

Estimating the proportion of cases transmitted through food

We used data on outbreaks reported to national surveillance systems between January 2001 and December 2008 to estimate the proportion of cases transmitted through food. For each pathogen, we computed empirical estimates for *πp* by obtaining 4,999 bootstrap samples of the proportion of cases in outbreaks that resulted from foodborne transmission. We then fitted a Beta function to the resulting distribution using maximum likelihood. For *Cryptosporidium* and *Giardia*, this approach gave an unrealistically high estimate for the proportion of cases transmitted through food because, of the few outbreaks that were reported, those involving foodborne transmission were larger. For these two pathogens, we used instead the proportion of outbreaks that were foodborne as an estimate of *πP*, as was done in a previous study.[5] For adenovirus and sapovirus, for which no outbreaks were reported, we used parameters derived from analysis of rotavirus and norovirus outbreaks respectively. For pathogens for which all outbreaks or no outbreaks were foodborne, we specified limits to the fitted Beta distributions as described in the Technical Appendix. The *a* and *b* parameters from the fitted Beta distributions were then used in the Monte Carlo simulations (see Model 1 below).

**Prior distributions for the proportion of cases transmitted through food (πp)**

We obtained prior distributions for the *πp* parameters from the systematic literature review. We divided retrieved articles into two categories: food attribution studies (Group A) and others (Group B). In Group A studies the proportion of cases transmitted through food was estimated for several pathogens, through expert elicitation or retrospective data reviews. Group B studies were primarily pathogen-specific case-control studies, or studies using microbiological typing for source attribution. For Group A and Group B studies, we defined uniform distributions for *πp*, based on the minimum and maximum estimates of the proportion of cases transmitted through food in these studies, for pathogens with at least two published studies. Where the observed proportion from outbreak data fell outside the limits of this uniform distribution, we arbitrarily allowed the lower or upper limit of the distribution to extend by 0.1 beyond the observed value.

**Pathogen-specific hospitalisation (γp)**

Data on hospitalisations were available only for outbreaks reported in England and Wales. For each reported outbreak, excluding those in hospitals or residential institutions, we computed the proportion of cases hospitalised by causative organism. We based hospitalisation estimates on all outbreaks with available data, as we found no major differences in hospitalisation between foodborne and other outbreaks. To account for uncertainty in these parameters, we fitted Beta distributions to bootstrapped data as detailed above for *πp*, but additionally weighting by outbreak size (see Technical Appendix). For adenovirus and sapovirus, we used parameters derived from analysis of rotavirus and norovirus outbreaks respectively. Bootstrap estimates with fitted Beta distributions by pathogen are shown in the Technical Appendix.

**Prior distributions for pathogen-specific hospitalisation (γp)**

We used pathogen-specific, Beta-distributed priors for *γp*. The Beta parameters were informed by an analysis of hospitalisation data from the IID1 and IID2 Studies (see Figure A1, Technical Appendix).

Estimating food-related IID cases, GP consultations and hospitalisations (Fp, Gp, Hp)

We obtained estimates of the number of foodborne cases, GP consultations and hospitalisations using three different approaches. In Model 1, we used Monte Carlo simulation to draw values at random from each parameter distribution. In Model 2, we used a Bayesian approach that included parameters for the prior distributions of *γp* from the IID1 and IID2 studies, and for *πp* from Group A studies as described above. These priors were used, together with the outbreak data, to obtain posterior distributions for these parameters, which were then used in the model. This model could not be applied to sapovirus, because none of the identified studies had information about this pathogen. Model 3 had the same structure as Model 2, except that Bayesian priors for *πp* from Group B studies were used instead. This model was applied to *Campylobacter*, *E. coli* O157, *Listeria* and *Salmonella*, for which sufficient data from published studies were available.A full description of model parameters is given in the Technical Appendix.

For each model, we carried out 100,000 simulations, discarding the first 10% and retaining the model outputs for every 10th simulation. We checked model convergence graphically by plotting parameter values over time to verify adequate mixing, plotting autocorrelograms and comparing density plots for outcome variables by tertile of the simulation chain. We summarised model outputs using the median and central 95% of the posterior distributions to obtain point estimates and 95% credible intervals for the number of food-related cases, GP consultations and hospitalisations by pathogen. We conducted the analyses using Stata 12.1, WinBUGS and Microsoft Excel software. We used the winbugsfromstata module in Stata to carry out the simulations.[16]

**Ethical considerations**

An Ethics Committee favourable opinion was not required. These were secondary analyses of previously collected, publicly available data. All datasets used were completely anonymous and there was no risk of disclosure of personal data.

## RESULTS

### Proportion of cases attributable to foodborne transmission

Table 1 summarises the outbreak data used for estimating the proportion of cases due to foodborne transmission from outbreak data. The identified studies used to inform Bayesian uniform priors are summarised in the Technical Appendix online (Tables A2 and A3). Figure 1 shows the empirical bootstrap distributions for the estimated proportion of cases due to foodborne transmission based on outbreak data. For most pathogens, the Beta distribution provided a reasonable fit to the bootstrapped distribution, with the exception of *Giardia*, for which data were sparse, and rotavirus, for which the estimated proportion foodborne was very small. *Salmonella* and *C. perfringens* had the largest estimated proportion of cases attributable to foodborne transmission, each approximately 90%. Around 50% of *Campylobacter* and *E. coli* O157 cases were estimated to result from foodborne transmission, although there was considerable uncertainty in these estimates as evidenced by the long tails in these distributions. Foodborne transmission accounted for less than 5% of norovirus cases, while approximately 65% of *Giardia* cases, 30% of *Cryptosporidium* cases, and 20% of *Shigella* cases were food-related.

### Proportion of cases hospitalised

Table 2 summarises the data sources used to inform hospitalisation parameters. Figure 2 shows the estimated hospitalisation proportions in reported outbreaks by pathogen, based on the medians of Beta distributions fitted to outbreak data. Hospitalisation was particularly high for *E. coli* O157 (23%). By contrast, less than 2% of cases due to *C. perfringens*, *Campylobacter*, *Giardia*, norovirus and rotavirus were hospitalised.

**Table 2: Summary of hospitalisation data by pathogen, UK 1993-2008**

|  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  | HOSPITALISATION IN OUTBREAKS | | | | |  | HOSPITALISATION IN IID1 AND IID2 STUDIES | | | |
| Organism | Hospitalised | Affected | % | Outbreaks with data | Source |  | Hospitalised | Affected | %1 | Source |
| **Bacteria** |  |  |  |  |  |  |  |  |  |  |
| *C. perfringens* | 2 | 1,120 | 0.2% | 21 | Outbreak surveillance |  | 2 | 78 | 2.6% | IID1 & IID2 |
| *Campylobacter* | 2 | 424 | 0.5% | 29 | Outbreak surveillance |  | 5 | 441 | 1.1% | IID1 & IID2 |
| *E. coli O157* | 197 | 877 | 22.5% | 70 | Outbreak surveillance |  | 0 | 2 | 33.3% | IID1 & IID2 |
| *Listeria* | -- | -- | -- | -- | All outbreaks occurred in hospitals |  | -- | -- | -- | No cases identified |
| *Salmonella* | 419 | 5,527 | 7.6% | 217 | Outbreak surveillance |  | 4 | 114 | 3.5% | IID1 & IID2 |
| *Shigella* | 4 | 153 | 2.6% | 8 | Outbreak surveillance |  | 0 | 11 | 8.3% | IID1 |
| **Protozoa** |  |  |  |  |  |  |  |  |  |  |
| *Cryptosporidium* | 31 | 836 | 3.7% | 46 | Outbreak surveillance |  | 0 | 50 | 2.0% | IID1 & IID2 |
| *Giardia* | 1 | 137 | 0.7% | 5 | Outbreak surveillance |  | 1 | 34 | 2.9% | IID1 & IID2 |
| **Viruses** |  |  |  |  |  |  |  |  |  |  |
| Adenovirus | -- | -- | -- | -- | No outbreaks reported |  | 0 | 79 | 1.3% | IID1 & IID2 |
| Astrovirus | 2 | 88 | 2.3% | 7 | Outbreak surveillance |  | 1 | 67 | 1.5% | IID1 & IID2 |
| Norovirus | 80 | 12,333 | 0.6% | 342 | Outbreak surveillance |  | 2 | 201 | 1.0% | IID1 & IID2 |
| Sapovirus | -- | -- | -- | -- | No outbreaks reported |  | 0 | 77 | 1.3% | IID2 |
| Rotavirus | 20 | 1,211 | 1.7% | 59 | Outbreak surveillance |  | 1 | 64 | 1.6% | IID2 |

### 1Where no hospitalisations were observed, the hospitalised percentage was calculated assuming the next case observed would have been hospitalised (see Technical Appendix)

### Cases, GP consultations and hospital admissions attributable to foodborne transmission (Model 1)

Table 3 presents estimates of food-related cases, GP consultations and hospital admissions in 2009 from 100,000 Monte Carlo simulations. *Campylobacter* was the most common foodborne pathogen, accounting for 286,000 food-related cases (95% CrI: 131,105 – 532,400) and 39,750 GP consultations (95% CrI: 18,890 – 69,540), but ranked third as a cause of food-related hospital admissions (1,376 admissions) behind *Salmonella* (2,536 admissions) and *E. coli* O157 (2,141 admissions). Foodborne norovirus accounted for 3,240 GP consultations (95% CrI: 1,985 – 5,162), but fewer than 500 hospital admissions. Similarly, other pathogens such as *C. perfringens* and a number of the viruses, while contributing large numbers of cases and GP consultations, were responsible for a modest number of food-related hospital admissions. It should be noted, however, that there was a large degree of uncertainty around these estimates, as demonstrated by the wide 95% credible intervals.

**Table 3: Estimates of food-related cases, GP consultations and hospitalisations by pathogen, UK 2009 (Model 1)**

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Organism** | **Cases** | **(95% CrI)** |  | **GP consultations** | **(95% CrI)** |  | **Hospital admissions** | **(95% CrI)** |
| Bacteria |  |  |  |  |  |  |  |  |
| *C. perfringens* | 79,165 | (29,310 - 208,688) |  | 12,610 | (5,707 - 27,890) |  | 165 | (20 - 843) |
| *Campylobacter* | 286,000 | (131,105 - 532,400) |  | 39,750 | (18,890 - 69,540) |  | 1,376 | (289 - 4,607) |
| *E. coli* O157 | 9,536 | (644 - 146,495) |  | 324 | (36 - 2,973) |  | 2,141 | (143 - 33,237) |
| *Listeria* | 169 | (100 – 215) |  | 169 | (100 – 215) |  | -- | -- |
| *Salmonella* | 33,640 | (8,286 - 135,798) |  | 10,030 | (4,019 - 24,299) |  | 2,536 | (608 - 10,400) |
| *Shigella* | 1,274 | (90 - 11,990) |  | 684 | (84 - 2,145) |  | 32 | (2 - 378) |
| Protozoa |  |  |  |  |  |  |  |  |
| *Cryptosporidium* | 2,035 | (354 - 10,129) |  | 588 | (140 - 2,010) |  | 72 | (12 - 395) |
| *Giardia* | 11,250 | (2,239 - 52,878) |  | 1,322 | (286 - 4,960) |  | 88 | (17 - 415) |
| Viruses |  |  |  |  |  |  |  |  |
| Adenovirus | 11,920 | (3,706 - 28,909) |  | 987 | (293 - 2,536) |  | 191 | (51 - 559) |
| Astrovirus | 2,362 | (594 - 7,180) |  | 180 | (41 - 576) |  | 70 | (15 - 262) |
| Norovirus | 73,420 | (50,320 - 104,000) |  | 3,240 | (1,985 - 5,162) |  | 470 | (270 - 779) |
| Rotavirus | 14,850 | (4,698 - 35,330) |  | 1,603 | (494 - 3,856) |  | 237 | (64 - 688) |
| Sapovirus | 40,770 | (26,661 - 60,230) |  | 2,457 | (1,496 - 3,947) |  | 261 | (145 - 445) |
|  |  |  |  |  |  |  |  |  |
| TOTAL | 566,391 |  |  | 73,944 |  |  | 7,639 |  |

### Cases, GP consultations and hospital admissions attributable to foodborne transmission (Models 2 and 3)

Estimates of food-related cases, GP consultations and hospital admissions based on the Bayesian approach used in Model 2 are presented in Table 4. *Campylobacter* was the most common foodborne pathogen, causing 280,400 (95% CrI: 182,503 – 435,693) food-related cases and 38,860 (95% CrI: 27,160 – 55,610) general practice (GP) consultations annually. Despite this, there were only 562 (95% CrI: 189 – 1,330) *Campylobacter*-related hospital admissions. *Salmonella* caused the largest number of hospitalisations, an estimated 2,490 admissions (95% CrI: 607 – 9,631), closely followed by *E. coli* O157 with 2,233 admissions (95% CrI: 170 – 32,159). Other common causes of foodborne disease included *C. perfringens*, with an estimated 79,570 cases annually (95% CrI: 30,700 – 211,298), and norovirus with 74,100 cases (95% CrI: 61,150 –89,660). For Model 2, there were insufficient data from the studies we identified to enable estimation of foodborne sapovirus. For *Campylobacter*, *E. coli* O157, *Listeria* and *Salmonella*, further estimates from Model 3 are presented in Table 5. The estimates from the three different models are compared in Figures 3a, 3b and 3c.

**Table 4: Estimates of food-related cases, GP consultations and hospitalisations by pathogen, UK 2009 (Model 2)**

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Organism** | **Cases** | **(95% CrI)** |  | **GP consultations** | **(95% CrI)** |  | **Hospital admissions** | **(95% CrI)** |
| Bacteria |  |  |  |  |  |  |  |  |
| *C. perfringens* | 79,570 | (30,700 - 211,298) |  | 12,680 | (6,072 - 27,040) |  | 186 | (38 - 732) |
| *Campylobacter* | 280,400 | (182,503 - 435,693) |  | 38,860 | (27,160 - 55,610) |  | 562 | (189 - 1,330) |
| *E. coli* O157 | 9,886 | (748 - 142,198) |  | 342 | (37 - 3,030) |  | 2,233 | (170 - 32,159) |
| *Listeria* | 183 | (161 – 217) |  | 183 | (161 – 217) |  | -- | -- |
| *Salmonella* | 33,130 | (8,178 - 128,195) |  | 10,060 | (4,137 - 24,710) |  | 2,490 | (607 - 9,631) |
| *Shigella* | 1,204 | (181 - 8,142) |  | 602 | (341 - 1,060) |  | 33 | (4 - 270) |
| Protozoa |  |  |  |  |  |  |  |  |
| *Cryptosporidium* | 2,773 | (562 - 12,200) |  | 800 | (233 - 2,386) |  | 94 | (18 - 436) |
| *Giardia* | 7,877 | (1,467 - 36,059) |  | 883 | (197 - 3,288) |  | 47 | (4 - 332) |
| Viruses |  |  |  |  |  |  |  |  |
| Adenovirus | 8,253 | (4,734 – 13,780) |  | 677 | (345 – 1,278) |  | 62 | (30 – 118) |
| Astrovirus | 3,470 | (1,368 - 9,991) |  | 262 | (93 - 812) |  | 11 | (3 - 42) |
| Norovirus | 74,100 | (61,150 - 89,660) |  | 3,276 | (2,240 - 4,729) |  | 332 | (248 - 440) |
| Rotavirus | 10,295 | (6,049 - 16,730) |  | 1,102 | (629 - 1,870) |  | 95 | (48 - 177) |
| Sapovirus1 | -- | -- |  | -- | -- |  | -- | -- |
|  |  |  |  |  |  |  |  |  |
| TOTAL | 511,141 |  |  | 69,727 |  |  | 6,145 |  |

1For sapovirus, no data were identified in the literature review on the proportion of cases attributable to food, so this model could not be applied

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Organism** | **Cases** | **(95% CrI)** |  | **GP consultations** | **(95% CrI)** |  | **Hospital admissions** | **(95% CrI)** |
| *Campylobacter* | 279,900 | (183,100 - 433,098) |  | 38,820 | (27,010 - 55,580) |  | 561 | (189 - 1,343) |
| *E. coli* O157 | 9,536 | (644 - 146,495) |  | 324 | (36 - 2,973) |  | 2,141 | (143 - 33,237) |
| *Listeria* | 166 | (92 - 214) |  | 166 | (92 - 214) |  | --1 | -- |
| *Salmonella* | 33,130 | (8,178 - 128,195) |  | 10,060 | (4,137 - 24,710) |  | 2,490 | (607 - 9,631) |
|  |  |  |  |  |  |  |  |  |
| TOTAL | 322,732 |  |  | 49,370 |  |  | 5,192 |  |

**Table 5: Estimates of food-related cases, GP consultations and hospitalisations by pathogen, UK 2009 (Model 3)**

1For Listeria, the number of hospital admissions could not be calculated, as all reported outbreaks occurred in hospitals

**Comparing the models**

In general, the results from all three approaches were similar for food-related cases and GP consultations. For most organisms, the Bayesian estimates from Model 2 benefited from greater precision. There were differences in the number of food-related hospital admissions estimated by the Monte Carlo and Bayesian approaches for some organisms, notably *Campylobacter*, rotavirus, adenovirus and astrovirus. The differences reflect discordance between outbreak data and data from the IID studies in terms of the hospitalisation rate for these organisms. Where differences were observed, the Bayesian approach gave more conservative estimates of the number of food-related hospital admissions.

For the four pathogens with sufficient data from the literature review to generate estimates from Model 3 (*Campylobacter*, *E. coli* O157, *Listeria* and *Salmonella*), estimates were similar to those from Model 2; however, *Listeria* estimates carried greater uncertainty, because of wide disagreement between the two identified studies regarding the proportion of listeriosis attributable to foodborne transmission. It was impossible to calculate listeriosis hospitalisations because all reported *Listeria* outbreaks occurred in hospitals.

## DISCUSSION

To our knowledge this is the first study to incorporate both empirical data and prior information from a systematic review using Bayesian methodology for estimating the proportion of IID that is transmitted through contaminated food.*Campylobacter* is the most common foodborne pathogen in the UK, causing between 182,503 and 435,693 food-related cases and between 27,160 and 55,610 GP consultations annually (based on Model 2 results). Despite this, the number of *Campylobacter*-related hospital admissions is comparatively small, reflecting a generally lower level of acute disease severity compared with other pathogens. By contrast, *Salmonella* and *E. coli* O157 cause the largest number of hospitalisations, an estimated 2,490 and 2,233 admissions respectively (Model 2), although uncertainty around these estimates is high. Other common causes of foodborne illness include *C. perfringens*, responsible for nearly 80,000 cases annually and norovirus, responsible for nearly 75,000 cases. Other viral agents rank lower as causes of foodborne illness.

Our analysis updates previous estimates for England and Wales in 2000 and expands upon them by accounting for uncertainty.[5] Because of substantial differences in the analyses, the two sets of estimates are not directly comparable. Other studies investigating the burden of foodborne illness caused by a wide range of pathogens have been carried out in Australia, the United States (US) and the Netherlands.[1,2,6,17] In the US and Australian studies norovirus was one of the commonest causes of foodborne disease. In the US study, it was also the second most common cause of food-related hospital admissions. Approximately one quarter of norovirus IID cases in those two studies were attributed to foodborne transmission, whereas our estimate for the UK is less than 5%. A likely reason for this discrepancy is the definitions of outbreaks that are incorporated in the various modelling studies. Some datasets contain only outbreaks transmitted through food whilst others, like ours (until 2009), contained all outbreaks of IID no matter what the route of transmission. This means that the proportion of norovirus cases transmitted through food is likely to be overestimated in datasets that contain only outbreaks transmitted through food.

A major strength of our analysis is the availability of directly observed, pathogen-specific incidence data from the recent IID2 Study in the UK,[10] which precludes the need to adjust for under-ascertainment and requires fewer assumptions about health care usage. The use of methods to account fully for parameter uncertainties is an additional strength, and is useful for highlighting areas where data are sparse. This is particularly true for hospitalisation estimates, for which there is a dearth of reliable data. We investigated other sources of hospitalisation data, such as electronic records of in-patient admissions. However, these data lack specific diagnostic codes for certain key pathogens, including *E. coli* O157, and a large fraction of admissions are classified under non-specific diagnostic codes. We therefore used outbreak data to estimate hospitalisation. A potential limitation is that severe cases requiring hospitalisation might be more reliably recorded in outbreak reports, whereas milder cases might be missed. There might genuinely be higher hospitalisation rates in outbreaks than sporadic cases because of higher dose exposures or different populations might be affected in outbreaks. Alternatively, outbreaks with more hospitalised cases might be more likely to be investigated and reported. This would tend to overestimate hospitalisation rates. Such a bias is possible in the *E. coli* O157 data, where estimates for hospitalisations were considerably higher than for GP consultations. Alternatively, the severity of this disease could mean that cases are admitted directly to hospital without first consulting a GP. Our Bayesian models additionally incorporated prior information on hospitalisation rates from IID1 and IID2. For most pathogens, the two types of models gave similar results. However, the number of hospitalisations in both sets of data was small, reflected in the large degree of uncertainty in the estimates. For rotavirus and astrovirus, the Bayesian model gave somewhat lower estimates of hospital admissions, which might indicate that hospitalisations for these two pathogens are over-reported in outbreak data or that they were under-ascertained in the IID studies. Additionally, outbreaks might occur in specific age groups or individuals with underlying conditions, or be due to high dose exposure. Outbreak reports, however, contain limited information on the populations affected.

Using outbreak data to attribute cases of IID to foodborne transmission relies on certain assumptions, principally that outbreak cases reflect the epidemiology in the wider community. Another potential limitation is that there might be a bias towards investigation or reporting of foodborne outbreaks compared with outbreaks transmitted through other routes, like person-to-person transmission. This, however, does not seem to be the case: there has been a gradual decrease in the proportion of reported outbreaks involving foodborne transmission, which reflects both a reduction in incidence of certain foodborne pathogens, particularly *Salmonella*, and greater investigation of outbreaks in other settings, particularly viral outbreaks in hospitals and residential institutions.[18,19]

Our study focused on foodborne illness burden in the general UK population. Some pathogens, however, are a particular problem among certain high-risk groups, such as *Listeria* among immunocompromised patients and pregnant women, and rotavirus among children under five years. Our analysis was not designed to estimate burden in these subgroups, because our data sources contain limited information on these groups, and the size of some of these high-risk populations is uncertain. However, further studies to estimate burden in these groups is warranted.

We were unable to include other relevant pathogens such as toxoplasmosis, hepatitis A, hepatitis E and non-O157 STEC in our analysis, due to a lack of relevant data in the UK. In a Dutch study *Toxoplasma gondii* caused the highest foodborne disease burden as measured by disability-adjusted life years, reflecting the importance of congenital toxoplasmosis.[16]

Our modelling approach meant we could use data from various sources to incorporate the best available information from the UK and elsewhere. Comparing models with and without prior information indicates where there is disagreement between data sources and enables uncertainty in all the relevant parameters to be accounted for. Uncertainty in these models reflects not simply statistical uncertainty in individual parameters, but disagreement between data sources and availability of information from previous studies. Information from previous studies on the proportion of IID transmitted through food was captured using Bayesian uniform priors. This is probably conservative, as it presupposes that every value within the specified limits is equally likely. For most pathogens, however, the number of available studies was small and using more informative priors was difficult to justify. The exception was *Campylobacter*, for which 14 studies contained relevant data. Even so, using data from risk factor studies presents problems in interpretation. Study design, methods and risk factors investigated varied widely. Consequently, variability between studies in the importance of food-related risk factors is high. The choice of Bayesian priors in estimation is necessarily a subjective process, as it depends on analysts’ confidence in the available information. Establishment of a process to develop greater international consensus on the choice of priors for individual pathogens could help to refine future estimates. Better baseline estimates would also inform predictions of the likely increase in foodborne disease due to climate change.[20]

We did not estimate deaths attributable to foodborne illness, due to the lack of reliable data sources on pathogen-specific mortality rates. Death certificates rarely provide information on specific gastrointestinal pathogens, while deaths in outbreaks are rare and may not be recorded if they occur sometime after the event. More generally, mortality estimates would be difficult to interpret. Deaths attributed to foodborne disease are not necessarily the same as preventable deaths. More focused epidemiological studies on mortality following IID would be helpful.

Our estimates measure foodborne disease burden only in the acute phase of illness. For some pathogens, the long-term consequences of illness can add considerably to their burden, for example *E. coli* O157-associated haemolytic uraemic syndrome (HUS) and *Campylobacter*-associated Guillain-Barré syndrome (GBS).[21,22] Moreover, our estimates are based only on the number of cases of illness, and take no account of the consequences of illness in different sectors of the population. Further studies using additional measures of disease burden and taking into account long-term health consequences are therefore required.

Modeling is not necessarily a substitute for acquiring good quality primary data but it is very useful for pointing to important data gaps and major areas of uncertainty where primary data collection might be focused.

Controlling foodborne disease is an important policy issue. Given the burden of illness caused, there needs to be a continued focus on reducing illness due to *Campylobacter*, *Salmonella*, *C. perfringens* and norovirus.

**References**

1. Kuchenmüller T, Abela-Ridder B, Corrigan T, Tritscher A. World Health Organization initiative to estimate the global burden of foodborne diseases. Rev Sci Tech. 2013;32(2):459-67.
2. Scallan E, Hoekstra RM, Angulo FJ *et al*. Foodborne illness acquired in the United States--major pathogens. Emerg Infect Dis. 2011;17(1):7-15.
3. Scallan E, Griffin PM, Angulo FJ et al. Foodborne illness acquired in the United States--unspecified agents. Emerg Infect Dis. 2011;17(1):16-22.
4. Mead PS, Slutsker L, Dietz V *et al*. Food-related illness and death in the United States. Emerg Infect Dis. 1999;5(5):607-25.
5. Adak GK, Long SM, O'Brien SJ. Trends in indigenous foodborne disease and deaths, England and Wales: 1992 to 2000. Gut. 2002;51(6):832-41.
6. Hall G, Kirk MD, Becker N *et al*. Estimating foodborne gastroenteritis, Australia. Emerg Infect Dis. 2005;11(8):1257-64.
7. Vaillant V, de Valk H, Baron E *et al*. Foodborne infections in France. Foodborne Pathog Dis;2(3):221-32.
8. Flint JA, Van Duynhoven YT, Angulo FJ *et al*. Estimating the burden of acute gastroenteritis, foodborne disease, and pathogens commonly transmitted by food: an international review. Clin Infect Dis. 2005;41:698–704.
9. Tam CC, Rodrigues LC, Viviani L *et al*; IID2 Study Executive Committee. Longitudinal study of infectious intestinal disease in the UK (IID2 study): incidence in the community and presenting to general practice. Gut. 2012;61(1):69-77.
10. Wheeler JG, Sethi D, Cowden JM *et al*. Study of infectious intestinal disease in England: rates in the community, presenting to general practice, and reported to national surveillance. The Infectious Intestinal Disease Study Executive. BMJ. 1999;318(7190):1046-50.
11. O'Brien SJ, Rait G, Hunter PR *et al*. Methods for determining disease burden and calibrating national surveillance data in the United Kingdom: the second study of infectious intestinal disease in the community (IID2 study). BMC Med Res Methodol. 2010;10:39. doi: 10.1186/1471-2288-10-39.
12. Tompkins DS, Hudson MJ, Smith HR *et al*. A study of infectious intestinal disease in England: microbiological findings in cases and controls. Commun Dis Public Health. 1999;2(2):108-13.
13. Liberati A, Altman DG, Tetzlaff J *et al*. The PRISMA statement for reporting systematic reviews and meta-analyses of studies that evaluate health care interventions: explanation and elaboration. Journal of clinical epidemiology. 2009;62(10):e1-34.
14. Tam CC, Larose T, O'Brien SJ. Costed extension to the second study of infectious intestinal disease in the community: Identifying the proportion of foodborne disease in the UK and attributing foodborne disease by food commodity. UK: Food Standards Agency; 2014. Report No.: Project B18021 (FS231043). Available at < http://www.food.gov.uk/sites/default/files/IID2%20extension%20report%20-%20FINAL%2025%20March%202014\_0.pdf> Dated accessed 31 July 2015.
15. Fullerton KE, Scallan E, Kirk MD *et al*. Case-control studies of sporadic enteric infections: a review and discussion of studies conducted internationally from 1990 to 2009. Foodborne Pathog Dis. 2012;9(4):281-92. doi: 10.1089/fpd.2011.1065.
16. Thompson JR, Palmer TM, Moreno S. Bayesian analysis in Stata using WinBUGS. The Stata Journal 2006;6(4):530-549.
17. Havelaar AH, Haagsma JA, Mangen MJ *et al*. Disease burden of foodborne pathogens in the Netherlands, 2009. Int J Food Microbiol. 2012;156(3):231-8. doi: 10.1016/j.ijfoodmicro.2012.03.029.
18. Tam CC, O'Brien SJ, Tompkins DS *et al*. Changes in causes of acute gastroenteritis in the United Kingdom over 15 years: microbiologic findings from 2 prospective, population-based studies of infectious intestinal disease. Clin Infect Dis. 2012;54(9):1275-86. doi: 10.1093/cid/cis028.
19. Gormley FJ, Little CL, Rawal N *et al*. A 17-year review of foodborne outbreaks: describing the continuing decline in England and Wales (1992-2008). Epidemiol Infect. 2011;139(5):688-99. doi: 10.1017/S0950268810001858.
20. Stephen DM, Barnett AG. Effect of temperature and precipitation on salmonellosis cases in South-East Queensland, Australia: an observational study. BMJ Open 2016; 6:e010204. doi: 10.1136/bmjopen-2015-010204.
21. Tariq L, Haagsma J, Havelaar A. Cost of illness and disease burden in The Netherlands due to infections with Shiga toxin-producing *Escherichia coli* O157. J Food Prot. 2011;74(4):545-52. doi: 10.4315/0362-028X.JFP-10-252.
22. Havelaar AH, de Wit MA, van Koningsveld R *et al*. Health burden in the Netherlands due to infection with thermophilic *Campylobacter* spp. Epidemiol Infect. 2000;125(3):505-22.

**Authors' contributions**

The study was conceived by CCT and SJO’B; TLL conducted the systematic literature review; GKA and MRE provided expert interpretation of the outbreak surveillance data used in the analysis; CCT conducted the analysis; SJO’B and CCT wrote the initial draft manuscript; SJO’B, TLL, GKA, MRE and CCT interpreted the results and made substantial contributions to revising the manuscript.

**Acknowledgements**

The authors would like to thank John Cowden and Brian Smyth for expert interpretation of outbreak data and the information scientists at the four UK national surveillance centres for providing the outbreak data on which some of the analyses are based.

**Funding statement**

This work was funded by the Food Standards Agency grant number FS231043, awarded to SJO’B as Project Lead Contractor. SJO’B is also supported by the National Institute for Health Research Health Protection Research Unit in Gastrointestinal Infections at the University of Liverpool [Grant number NIHR HPRU 2012-10038].

**Disclaimer**

The research was part-funded by the National Institute for Health Research Health Protection Research Unit (NIHR HPRU) in Gastrointestinal Infections at the University of Liverpool in partnership with Public Health England (PHE), University of East Anglia, University of Oxford and the Institute of Food Research. The views expressed are those of the authors and not necessarily those of the NHS, the NIHR, the Department of Health or Public Health England.

**Data Sharing**

Data from the IID1 study are available from the UK Data Service (http://ukdataservice.ac.uk/; DOI: 10.5255/UKDA-SN-4092-1). Data from the IID2 study are available from the UK Data Service (http://dx.doi.org/10.5255/UKDA-SN-7820-1). Anonymous outbreak surveillance data are available on request from Public Health England (eFOSS, https://bioinfosecure.phe.org.uk/efoss), Health Protection Scotland and the Public Health Agency of Northern Ireland. Data from the literature review are in the public domain and available as cited in the main manuscript and/or Technical Appendix.

**Competing interests**

None