Prevalence of Avian Influenza A(H5) and A(H9) Viruses in Live Bird Markets, Bangladesh


We conducted a cross-sectional study in live bird markets (LBMs) in Dhaka and Chittagong, Bangladesh, to estimate the prevalence of avian influenza A(H5) and A(H9) viruses in different types of poultry and environmental areas by using Bayesian hierarchical logistic regression models. We detected these viruses in nearly all LBMs. Prevalence of A(H5) virus was higher in waterfowl than in chickens, whereas prevalence of A(H9) virus was higher in chickens than in waterfowl and, among chicken types, in industrial broilers than in cross-breeds and indigenous breeds. LBMs with ≥1 wholesaler were more frequently contaminated by A(H5) virus than retail-only LBMs. Prevalence of A(H9) virus in poultry and level of environmental contamination were also higher in LBMs with ≥1 wholesaler. We found a high level of circulation of both avian influenza viruses in surveyed LBMs. Prevalence was influenced by type of poultry, environmental site, and trading patterns.

Low pathogenicity avian influenza A(H9N2) virus and highly pathogenic avian influenza A(H5N1) virus are endemic in poultry populations in Bangladesh (1–4). In addition to their adverse effect on poultry production, these viruses have resulted in sporadic influenza cases in humans (2,3). Because there is potential for generating novel reassortant variants between them or with other virus subtypes, their persistent circulation in poultry poses a serious threat to animal and human health globally (5–9).

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DOI: https://doi.org/10.3201/eid2412.180879

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However, such information is generally poorly document-
ed, or even ignored.

To address these issues, we conducted a cross-section-
al survey in the 2 largest cities in Bangladesh, Dhaka and
Chittagong, during February–March 2016. First, we esti-
imated prevalence of influenza A(H5) and A(H9) viruses
in marketed poultry and the LBM environment. We also
accounted for the clustering effect at LBM level by using
Bayesian hierarchical logistic regression models. Second,
we assessed the effect of type of poultry and environmen-
tal site, and the position of LBMs in the poultry value chain
on AIV prevalence.

Materials and Methods

Sample Collection

An LBM was defined as an open space in which ≥2 poultry
stalls sell live poultry at least once a week, and only those
selling >400 poultry/day were considered eligible for this
study. We aimed to sample 40 LBMs, and from each of
these LBMs, 60 birds and 50 environmental sites (sample
size calculations in online Technical Appendix 1, https://
wwwnc.cdc.gov/EID/article/24/12/18-0879-Techapp1.
pdf). We used a stratified cluster sampling design. For
poultry, LBMs, stalls within selected LBMs, and birds
within selected stalls constituted the primary, secondary,
and tertiary sampling units, respectively. For environmen-
tal sites, LBMs constituted primary sampling units and
environmental sites within selected LBMs constituted sec-
ondary sampling units.

We stratified LBMs by city for Dhaka and Chittagong
and, within each city, by poultry sales into large and small
LBMs, hypothesizing that the risk for AIV infection var-
ies between geographic locations and the number of poul-
try traded. Also, simple random sampling with too small a
sample size of LBMs was not likely to capture diversity of
LM types because the distribution of LBMs as a function
of their size tended to be right-skewed; the largest LBMs
were often wholesale markets (24,25). We hypothesized
that samples of different origins have different AIV preva-
ences and thus stratified birds and environmental sites
into 5 types of poultry and 10 types of environmental sites
commonly found to be contaminated with AIV (26) (online
Technical Appendix 2 Tables 1, 2, https://wwwnc.cdc.gov/
EID/article/24/12/18-0879-Techapp2.pdf).

At the first sampling stage, we sampled 40 LBMs. The
number of LBMs selected in Dhaka (n = 26) and Chit-
tagong (n = 14) was proportional to the number of LBMs
eligible in each city (n = 80 for Dhaka and n = 36 for Chit-
tagong). In each city, we further stratified LBMs by size:
50% of the selected LBMs were large, trading the highest
number of poultry (13 largest LBMs in Dhaka and 7 largest
LBMs in Chittagong); 50% were small, randomly selected
from the bottom 50% of eligible LBMs in terms of number
of poultry traded.

At the second sampling stage, we randomly selected
stalls and environmental sites in each LBM independently
for each type of poultry and environmental site. We cre-
ated a list of stalls selling each poultry type for each LBM.
We then selected stalls from these lists by using a random
number generator. Likewise, for each type of environmen-
tal site, we selected sites from a list of sites identified in
each LBM by using a random number generator. We col-
lected 1 swab specimen from each environmental site. We
pooled 5 swab specimens collected from the same LBM
and site type.

At the third sampling stage, for each poultry type, we
randomly selected 5 birds from each of the stalls selected
for that type and collected cloacal and oropharyngeal swab
specimens from each of the selected birds. We pooled 5
swab specimens collected from the same stall and poultry
type separately for cloacal and oropharyngeal swab speci-
mens. We transported samples collected in Chittagong
on the day of sampling to the Chittagong Veterinary and
Animal Sciences University (Chittagong) and samples col-
lected in Dhaka on the day of sampling to the Bangladesh
Livestock Research Institute (Dhaka). Samples were stored
at −80°C until diagnostic laboratory processing.

Sample Screening

We screened pools for AIVs by using a real-time reverse
transcription PCR (RT-PCR) and specific primers and
probes (27, 28). We extracted virus RNA by using the Mag-
MAX RNA Isolation Kit (QIAGEN, Hilden, Germany) and
reverse transcribed and amplified virus RNA by using the
AgPath-ID One-Step RT-PCR (ThermoFisher Scientific,
Waltham, MA, USA). We then screened a pool with a cy-
cle threshold (Ct) <40 for the AIV matrix gene for the H5
and H9 genes. Results were considered positive for the H5
subtype if Ct <38 and positive for the H9 subtype if Ct <40
(27,28). A pool was considered positive for AIV if its Ct
for the AIV matrix gene <38 or if it was positive for any of
the H5 and H9 subtypes. A given group of 5 birds was
considered positive if any of its cloacal and oropharyngeal
pools showed a positive result.

Bayesian Hierarchical Logistic Regression Models

We developed 2-level Bayesian hierarchical logistic regres-
sion models to estimate LBM-level, bird-level, and environ-
mental swab specimen–level prevalence from pooled swab
samples, accounting for lower-level (swab specimens) and
higher-level (LBMs) risk factors. We developed separate
models for poultry and environmental samples to avoid pa-
rameters related to different sampling units interfering with
each other (29). We used LBM type (retail or mixed), city
(Chittagong or Dhaka), and size (small or large) as LBM-
level risk factors. We defined an LBM with only retailers (i.e., trader selling poultry to end-users only) as retail and an LBM with ≥1 wholesaler (i.e., trader selling poultry to other traders) as mixed. Each LBM-level risk factor (LBM type, city, and size) was assessed separately because we could not include them simultaneously in any given model due to the small number of sampled LBMs. In models for poultry samples, we differentiated birds into 1) chicken and waterfowl or 2) broiler, Desi, Sonali (i.e., chicken types), and waterfowl. Waterfowl were not differentiated further because of the small number of pools collected from ducks and geese. In models for environmental samples, we differentiated environmental sites into stall and slaughter areas or classified as environmental area without differentiation. We ran models (online Technical Appendix 1) by using a Markov Chain Monte Carlo simulation in JAGS (30) and R.3.4.2 (31).

**Results**

**Descriptive Results for Pooled Swab Samples**

We collected 477 pairs of cloacal and oropharyngeal pooled samples from 2,384 birds, and 400 environmental pooled samples from 2,000 environmental sites in 40 LBMs in Chittagong and Dhaka. Each pool was composed of 5 swab specimens, except for 1 pair of cloacal and oropharyngeal pools made from 4 swab specimens collected from geese. We collected 12 pairs of cloacal and oropharyngeal pooled samples from all LBMs, except for 11 pairs from 3 LBMs. We sampled chickens in all LBMs (8–12 pairs/LBM), and waterfowl in 25 LBMs (0–4 pairs/LBM). Broilers accounted for most samples (32.1%), followed by Desi (26.6%) and Sonali (25.6%). Ducks accounted for 76% of 75 pool pairs collected from waterfowl and geese accounted for 24%. We collected 10 environmental pools in each LBM (stall areas: 4–8 pools, slaughter areas: 2–6 pools).

Of 47.4% (416/877) pools considered positive for AIV, 6.5% pools were negative for the AIV matrix gene but positive for any of the H9 and H5 subtypes. The H9 subtype (63.2% positive pools) was detected more frequently than the H5 subtype (21.6%), and 12.3% of pools were positive for both subtypes and 27.4% of pools were negative for both subtypes. Although 80.0% of the LBMs had ≥1 A(H5) virus–positive poultry or environmental pool, 97.5% had ≥1 A(H9) virus–positive poultry or environmental pool. We determined the prevalence of pools that were positive for A(H5) and A(H9) viruses according to sample and LBM type (Table 1).

Approximately 33.3% of pools collected from waterfowl were positive for A(H5) virus, whereas only 5.5% of those collected from chickens were positive. In contrast, the prevalence of A(H9) virus–positive pools was higher in chickens (36.3%) than in waterfowl (18.7%). Among waterfowl, ducks (19.3%) and geese (16.7%) had a similar prevalence of A(H9) virus–positive pools, but the prevalence of A(H5) virus–positive pools was higher in ducks (36.8%) than in geese (22.2%). For both H5 and H9 subtypes, the prevalence of positive pools was higher for oropharyngeal samples (8.6% for H5 and 31.9% for H9) than for cloacal samples (3.6% and 9.9%) in all surveyed poultry types (online Technical Appendix 2 Table 1).

Approximately 25% of environmental pools were positive for A(H9) virus, and the prevalence of positive pools was higher in slaughter areas (31.5%), especially knives and boards used for slaughter and processing, than stall areas (20.2%). The prevalence of A(H5) virus–positive environmental pools was lower (10.8%) and did not vary between slaughter and stall areas (online Technical Appendix 2 Table 2).

**Bayesian Model Results**

Convergence was achieved for all models; the Gelman and Rubin statistic was <1.001 and the effective sample size was >10,000 for all parameters. For each AIV subtype, the best models reasonably predicted the number of positive pools (online Technical Appendix 2 Figure 1). In the best H5 models (i.e., lowest deviance information criterion), A(H5) virus prevalence differed according to poultry species (chicken, waterfowl), but not according to the type of environmental site. In contrast, in the best H9 models, A(H9) virus prevalence differed according to type of poultry (broiler, Desi, Sonali, waterfowl) and environmental site (slaughter and stall area). For both subtypes, LBM size and city did not improve model fit when compared with LBM type. For ease of comparison between the 2 AIV subtypes, we report LBM-level, bird-level, and environmental swab specimen–level prevalences of A(H5) and A(H9) viruses on the basis of the best H9 models with LBM type (Tables 1, 2). This reporting did not affect interpretation of results, and we provide estimates obtained with more parsimonious models (online Technical Appendix 2 Tables 3–6).

LBM-level A(H5) virus prevalence was lower in retail LBMs than in mixed LBMs, and the posterior median estimate was ≈100% for mixed LBMs. However, among contaminated LBMs, levels of virus detection in birds and environmental areas did not vary between LBM types, but A(H5) virus prevalence in waterfowl was ≈6 times higher than in chickens (Figure). The prevalence did not vary between chicken breeds or environmental areas.

In contrast to that for A(H5) virus, we found that the posterior median estimate of the LBM-level A(H9) virus prevalence was ≈100% for retail and mixed LBM groups, but the level of virus detection in birds and environmental areas was higher for mixed LBMs than for retail LBMs. A(H9) virus prevalence was highest in broilers and lowest in waterfowl. The prevalence in broilers was 3.8 times as
high as that in waterfowl and 1.6 times as high as that in Desi and Sonali (Figure). The environmental swab specimen–level prevalence was ≥2 times as high for slaughter areas than for stall areas (Figure).

**Discussion**

We detected A(H5) and A(H9) viruses in marketed poultry and environmental sites in nearly all LBMs sampled in Chittagong and Dhaka. The prevalence of A(H5) virus was higher in waterfowl than in chickens, whereas the prevalence of A(H9) virus was higher in chickens than waterfowl and also varied among chicken types, being more prevalent in broilers than in Desi and Sonali breeds. Slaughter areas were more frequently contaminated by A(H9) virus than stall areas. Whereas mixed LBMs were more frequently contaminated by A(H5) virus than were retail LBMs, prevalence of A(H9) virus was higher in mixed LBMs than in retail LBMs for birds and environmental areas.

AIVs were ubiquitous in surveyed LBMs. The LBM-level prevalence of A(H5) virus in Bangladesh was higher than in other AIV-endemic countries, including Egypt (32) and Vietnam (16). For both AIV subtypes, LBM-level prevalence was also higher than in another study conducted in Chittagong (21), which found that 17.5% of LBMs had ≥1 environmental sample pool contaminated by A(H5) virus and 12.5% of LBMs had ≥1 environmental sample pool contaminated by A(H9) virus. This difference might have been caused by different sampling schemes; in our study, we collected a larger number of pools per LBM.

Bird-level prevalence was also higher than that reported in other AIV-endemic countries, including Bangladesh (1,4,16,19). However, care must be taken when comparing these results because studies used different study designs and sample screening protocols over different periods. Bird-level prevalence for contaminated LBMs was much higher than for virologic surveys conducted in backyard and commercial farms in Bangladesh (1,4,33,34). This finding suggests that virus transmission was amplified along the value chain from farms to LBMs. Overcrowding and continuous supply of susceptible birds of different species and breeds might have created conditions promoting the silent transmission of AIVs within these markets (10).

Our results suggest that birds in LBMs with a mixture of wholesalers and retailers were at higher risk for infection than birds in LBMs with primarily retail poultry businesses. Poultry value chains supplying different business types might differ structurally, thereby affecting the risk for

**Table 1. Prevalence of avian influenza A(H5) and A(H9) viruses in pooled poultry and environmental samples, Chittagong and Dhaka, Bangladesh**

<table>
<thead>
<tr>
<th>Sample type</th>
<th>No. pools</th>
<th>A(H5) virus prevalence, % (95% HDI)†</th>
<th>A(H9) virus prevalence, % (95% HDI)†</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Poultry</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Retail LBM</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Broiler</td>
<td>96</td>
<td>5.2 (90–9.4)</td>
<td>37.5 (10.8–22.5)</td>
</tr>
<tr>
<td>Sonali</td>
<td>62</td>
<td>3.2 (14.0–13.2)</td>
<td>32.3 (6.6–14.5)</td>
</tr>
<tr>
<td>Desi</td>
<td>61</td>
<td>3.4 (13–12.2)</td>
<td>29.5 (6.8–13.4)</td>
</tr>
<tr>
<td>Waterfowl</td>
<td>20</td>
<td>8.1 (4.8–6.8)</td>
<td>25.0 (2.8–7.0)</td>
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<tr>
<td>Mixed LBM</td>
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<td></td>
<td></td>
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<tr>
<td>Broiler</td>
<td>57</td>
<td>1.8 (0–4.0)</td>
<td>47.4 (13.1–30.1)</td>
</tr>
<tr>
<td>Sonali</td>
<td>60</td>
<td>10.0 (1.4–5.7)</td>
<td>31.7 (8.0–19.8)</td>
</tr>
<tr>
<td>Desi</td>
<td>66</td>
<td>9.1 (1.3–5.2)</td>
<td>39.4 (8.3–20.4)</td>
</tr>
<tr>
<td>Waterfowl</td>
<td>55</td>
<td>27.3 (6–24.6)</td>
<td>16.4 (3.4–9.7)</td>
</tr>
<tr>
<td><strong>Environmental site</strong></td>
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<td></td>
</tr>
<tr>
<td>Retail LBM</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stall area</td>
<td>101</td>
<td>5.9 (0–10.4)</td>
<td>16.8 (3.2–9.1)</td>
</tr>
<tr>
<td>Slaughter area</td>
<td>99</td>
<td>7.1 (0–10.1)</td>
<td>25.3 (6.2–16.6)</td>
</tr>
<tr>
<td>Mixed LBM</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stall area</td>
<td>102</td>
<td>15.7 (1–11.3)</td>
<td>23.5 (5.2–14.1)</td>
</tr>
<tr>
<td>Slaughter area</td>
<td>98</td>
<td>14.3 (0–11.0)</td>
<td>37.8 (9.9–25.0)</td>
</tr>
</tbody>
</table>

*Desi, which means “local” in Bengali, are indigenous chicken breeds raised in backyard farms. Sonali is a cross-breed of the Rhode Island Red cocks and Fayoumi hens. HDI, high-density interval; LBM, live bird market.

†Bird and environmental swab specimen–level prevalence in contaminated live bird markets. Median values are reported.

**Table 2. Prevalence of avian influenza A(H5) and A(H9) viruses in LBMs, Chittagong and Dhaka, Bangladesh**

<table>
<thead>
<tr>
<th>Sample type</th>
<th>No. LBMs</th>
<th>H5 virus median prevalence, % (95% HDI)</th>
<th>H9 virus median prevalence, % (95% HDI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Poultry</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Retail LBM</td>
<td>20</td>
<td>69.9 (40.2–100.0)</td>
<td>96.4 (85.5–100.0)</td>
</tr>
<tr>
<td>Mixed LBM</td>
<td>20</td>
<td>92.0 (72.3–100.0)</td>
<td>96.0 (84.0–100.0)</td>
</tr>
<tr>
<td>Environmental sites</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Retail LBM</td>
<td>20</td>
<td>76.5 (47.2–100.0)</td>
<td>94.9 (80.5–100.0)</td>
</tr>
<tr>
<td>Mixed LBM</td>
<td>20</td>
<td>93.2 (75.5–100.0)</td>
<td>96.0 (84.0–100.0)</td>
</tr>
</tbody>
</table>

*Prevalence estimates were made by using the best H9 models with LBM type. HDI, high-density interval; LBM, live bird market.
introduced birds being already infected. Wholesalers generally trade a larger number of birds from more diverse geographic origins than do retailers (23) and therefore might have increased likelihood of virus introduction into mixed LBMs. Moreover, because wholesalers might sell birds to retailers in the same LBM (23), virus amplification might be increased through the presence of wholesalers.

The higher prevalence of A(H9) virus in broilers than in Sonali and Desi might result from differences in the structure of their respective value chains (23). Depending on the chicken type, different value chain actors might be involved and their trading practices might differ (23). The amount of time chickens spend with traders, the density at which chickens are kept in flocks of traders, and the frequency of contact with chickens from other flocks might vary with chicken type. The greater number of broilers marketed in surveyed LBMs might mean that broilers are more likely than Desi and Sonali to be sourced from large numbers of flocks, which are then mixed in densely populated trucks during transport to LBMs, promoting AIV transmission. However, these prevalence patterns might also be caused by varying levels of genetic susceptibility to AIV infection (35,36). Further investigations are needed to disentangle the possible influences of trade-related and genetic factors on AIV transmission in these chicken types. The higher level of contamination with A(H9) virus in slaughter areas than in stall areas suggests that, in the absence of appropriate biosecurity measures, slaughtering is likely to expose humans to AIVs by fomite transmission (37).

Co-circulation of A(H5) and A(H9) viruses arouses concerns over evolution of novel reassortant variants (5–8). Detection of both subtypes in some poultry pools suggests that these subtypes co-circulated near each other or in the same host during the study period. Although A(H5) viruses have considerable variability in their ability to infect, cause disease, and be transmitted among waterfowl (38), waterfowl are generally known to be less susceptible to highly pathogenic avian influenza A(H5N1) viruses (39). Therefore, waterfowl could harbor this virus but remain asymptomatic and serve as a potential host.
for genesis of novel AIVs in the presence of other virus subtypes. Also, the high level of A(H9) virus circulation among chickens could provide an ideal environment for virus diversification and selection in the LBM system. The different prevalence patterns in chickens and waterfowl observed suggest that these poultry species should be separated in LBMs and that active surveillance of novel reassortant variants should be implemented.

This study had some limitations. First, our models only accounted for clustering of sampled birds at the LBM level, but not at stall level. It is plausible that clustering of samples at stall level has less influence on AIV infection probability across the study population than clustering at the LBM level because stallholders in a given LBM in Bangladesh are likely to be supplied by the same traders and trade between each another (23). However, potential risk factors at stall level, such as ducks and hygiene level (27), might cause heterogeneous levels of AIV infection across stalls.

Second, our models did not account for the fact that sampling units in each stratum were selected with unequal probabilities. Although we selected different numbers of birds for each poultry type to account for variations in poultry populations, birds were still selected with different probabilities because their populations varied between clusters and strata. This selection might have resulted in larger SEs and thus less precise estimates compared with what could have been obtained with proportional sample sizes. Moreover, the overall prevalence might have been biased toward prevalence in samples selected with higher probabilities.

Third, our models assumed perfect sensitivity and specificity of real-time RT-PCR for pooled samples. The assays used in this study are considered highly sensitive and specific (27,28), and previous studies did not report any differences in virus detection for pooled and individual samples (40–42). Furthermore, virus detection in our study was based on parallel interpretation of cloacal or oropharyngeal sample test results (i.e., positive if ≥1 was positive). However, pools that were negative for the AIV matrix gene but positive for any of the H5 and H9 subtypes indicate that accounting for actual test sensitivity and specificity would enable more robust prevalence estimation. Virus isolation might be attempted for RT-PCR–positive pools to assess the viability of virus material. However, this testing was not attempted in our study. Each pool consisted of swab specimens from different birds or environmental sites. Thus, multiple AIV subtypes and virus species, including Newcastle disease viruses, could be present in the same pool and interfere with growth of each virus in chicken eggs (43). Should such studies be replicated, the collection of individual swab specimens and their pooling at the laboratory is recommended to enable analysis of individual swab specimens that formed a virus-positive pool.

Fourth, we collected samples over a short period to reduce variability that could arise from seasonal variations in AIV prevalence. We focused on winter months, which are often reported to be periods of higher risk for AIV infection (44). Therefore, our estimates only represented AIV prevalence during that period and did not capture seasonal changes.

Contrary to previous cross-sectional studies, our approach enabled us to estimate AIV prevalence not only by poultry species but also by chicken type and account for the type of LBMs in which sampled poultry were marketed. Despite most AIV surveys and surveillance activities being based on multistage sampling, single-level analytic methods are generally used to analyze their results, while ignoring within-market correlation in poultry infection status. Accounting for this effect by incorporating LBM-specific random effects in a hierarchical model, and enabling mutual influence between bird-level, environmental swab specimen–level, and LBM-level parameters, improved the reliability of prevalence estimates (29). When applied to other settings, this approach needs to be adapted on the basis of an understanding of the variety of poultry value chains. Information about LBM locations and about trading practices and numbers and types of poultry sold within these LBMs is rarely readily available and would need to be collected to inform the study design.

In conclusion, LBMs surveyed in Bangladesh were highly contaminated by A(H5) and A(H9) viruses. The level of virus detection was associated with the type of poultry and environmental area and the presence of wholesalers in LBMs. These findings need to be included in the design of risk-based surveillance and control interventions aimed at reducing AIV prevalence, human exposure, and the risk for emergence of novel virus reassortant variants.

Acknowledgments
We thank Eric Brum for his support during study implementation and the participants involved in the study.

This study was supported by the BALZAC research program “Behavioural adaptations in live poultry trading and farming systems and zoonoses control in Bangladesh” (BB/L018993/1) and is 1 of 11 programs supported by the Zoonoses and Emerging Livestock Systems, a joint research initiative between the Biotechnology and Biological Sciences Research Council, the Defence Science and Technology Laboratory, the Department for International Development, the Economic and Social Sciences Research Council, the Medical Research Council, and the Natural Environment Research Council.

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Emerging Infectious Diseases • www.cdc.gov/eid • Vol. 24, No. 12, December 2018

References


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Prevalence of Avian Influenza A(H5) and A(H9) Viruses in Live Bird Markets, Bangladesh

Technical Appendix 1

Sample Size Calculations

The selection of 20 large and 20 small live bird markets (LBMs) was necessary to detect a statistically significant difference between 2 groups of LBMs with assumed LBM-level avian influenza virus prevalences of 50% and 10%, respectively (95% significance, 80% power). Assuming a prevalence of infection of 7% in chickens in a contaminated LBM, detecting the infection in chickens traded in a contaminated LBM with a confidence of 95% required the sampling of 40 chickens.

Assuming a prevalence of infection of 7% in chickens and 14% in waterfowls in a contaminated LBM, sampling 40 chickens and 20 waterfowls from each LBM was needed to detect at least 1 infected chicken and 1 infected duck with a 95% significance. For chicken breeds, we aimed to sample 15 broilers, 15 Desi (local in Bengali, indigenous chicken breeds raised in backyard farms) and 10 Sonali (cross-breed of the Rhode Island Red cocks and Fayoumi hens) in LBMs in which more broilers were sold than Sonali, and 10 broilers, 15 Desi and 15 Sonali in in LBMs in which more Sonali were sold than broilers, hypothesizing that their numbers were proportional to their relative numbers sold in Dhaka and Chittagong LBMs. For waterfowl, we aimed to sample 15 ducks and 5 geese/LBM, or 20 ducks were sampled if geese were not present. Assuming that the probability of a pool of 5 environmental samples being contaminated in a contaminated LBM was 0.3 (and the probabilities of environmental sites being contaminated were independent), it was necessary to collect 10 pools to detect at least 1 contaminated pool with a probability of 0.97.

Bayesian Hierarchical Logistic Regression Model

For a given viral subtype (i.e., H5 or H9), the contamination status of a given LBM $m$, $\omega_m$, was assumed to follow a Bernoulli distribution with parameter $\gamma_{a_n}$, the probability of a
LBM of type $\alpha_m$ being contaminated. In other words, $\gamma_{\alpha_m}$ could be interpreted as the LBM-level prevalence for LBMs of type $\alpha_m$ (Equation 2.1):

$$\omega_m \sim \text{Bernoulli}(\gamma_{\alpha_m}) \quad (\text{Equation 2.1})$$

In each iteration, the contamination status of each LBM was first simulated, such that all pools from the same LBM originated from either a contaminated (i.e., some pools could be positive) or noncontaminated LBM (i.e., all these pools were necessarily negative). Because models were simulated separately for poultry and environmental samples, the interpretation of LBM-level prevalence differed accordingly: the LBM-level prevalence estimated from a model based on poultry (or environmental) samples referred to the proportion of LBMs with at least 1 infected poultry (or contaminated environmental site).

The real-time reverse transcription PCR result of pool $i$ in LBM $m$, $y_{i,m}$, was assumed to follow a Bernoulli distribution with parameter $\theta_{i,m}$ (Equation 2.2), the probability of pool $i$ being contaminated (Equation 2.3):

$$y_{i,m} \sim \text{Bernoulli}(\theta_{i,m}) \quad (\text{Equation 2.2})$$

$$\theta_{i,m} = \omega_m \times \lambda_{i,m} \quad (\text{Equation 2.3})$$

$\lambda_{i,m}$ was the probability of pool $i$ in LBM $m$ being contaminated if this LBM was contaminated. If LBM $m$ was not contaminated ($\omega_m = 0$), all pools from this LBM were negative by real-time reverse transcription PCR ($y_{i,m} = 0$). If LBM $m$ was contaminated ($\omega_m = 1$), $y_{i,m}$ was then simulated by a Bernoulli trial with parameter $\lambda_{i,m}$ (as $\theta_{i,m} = \lambda_{i,m}$). A pool $i$ was positive if at least 1 of its swabs was infected. Therefore, $\lambda_{i,m}$ was expressed as a function of 1) the underlying bird– (or environmental swab)–level prevalence, $\pi_{i,m}$ and 2) the number of birds (or environmental swabs) comprising pool $i$ in LBM $m$, $n_{i,m}$ (Equation 2.4):

$$\lambda_{i,m} = 1 - (1 - \pi_{i,m})^{n_{i,m}} \quad (\text{Equation 2.4})$$
\( \pi_{i,m} \) was assessed through a Bayesian hierarchical logistic regression to account for the hierarchical data structure. \( \pi_{i,m} \) only depended on the type of sample (Equation 2.5):

\[
\logit(\pi_{i,m}) = \delta_m + \sum_j \beta_j \times \psi_{j,i,m} \quad (\text{Equation 2.5})
\]

\( \beta_j \) was a regression coefficient for sample of type \( j \), and \( \psi_{j,i,m} \) an indicator variable, equal to 1 if the pool \( i \) in market \( m \) was of type \( j \), and null otherwise. \( \delta_m \) was the LBM-specific intercept.

At the second level, \( \delta_m \) was assumed to follow the LBM-specific normal distribution (Equation 2.6):

\[
\delta_m \sim \text{Normal}(\mu_m, \sigma_m^2) \quad (\text{Equation 2.6})
\]

The variance \( \sigma_m^2 \) assumed that prevalence varied between LBMs after adjusting for a LBM-level predictor. The mean \( \mu_m \) was modeled as a linear function of a LBM-level intercept, \( \phi \), and 1 of the LBM-level predictors, \( \beta_\alpha \) (Equation 2.7), which was used to differentiate the LBM-level prevalence:

\[
\mu_m = \phi + \beta_\alpha \times \tau_{\alpha,m} \quad (\text{Equation 2.7})
\]

\( \tau_{\alpha,m} \) was an indicator variable, equal to 1 if the market \( m \) was of type \( \alpha \); it was otherwise null. All unknown parameters were specified by weakly informed priors to enable the observed data to be the main contributor to the estimation of the posterior distributions (online Technical Appendix 1 Table).

The models were run by using a Markov chain Monte Carlo simulation in JAGS (1) and R.3.4.2 (2). After a burn-in period of 5,000 iterations, each model was iterated up to the point where convergence was achieved in all parameters on the basis of the Gelman and Rubin statistic (3,4) and the effective sample size (5). Although LBM-level prevalence was estimated directly from each model, the underlying bird- and environmental site-level prevalence was estimated by taking the inverse logit transformation of the corresponding regression coefficients. Median and 95% highest density interval are reported. All possible combinations of lower- and higher-level
regression predictors were tested. Models were compared with each other on the basis of the deviance information criterion (DIC) (6). Half of the variance of the posterior mean deviance was used as an estimate of the effective number of parameters (7). Models with lower DIC were considered to better support the data than those with higher DIC if the DIC difference was >5.

Finally, a posterior predictive check was performed to assess model adequacy. In each iteration, model parameter values were sampled from their joint posterior distribution. The contamination status of each market, and in contaminated markets, the contamination status of each pool were then simulated. The pool-level prevalence was computed and formed the posterior predictive distribution along with those computed from other iterations. This prevalence was compared with the observed pool-level prevalence by using the Bayesian p value, which represents the probability that the former could be equal to or more extreme than the latter (7).

References

   http://www.stats.ox.ac.uk/~nicholls/MScMCMC15/jags_user_manual.pdf


Technical Appendix 1 Table. Weakly informed priors used in models for analyzing prevalence of avian influenza A(H5) and A(H9) viruses in live bird markets, Bangladesh.

<table>
<thead>
<tr>
<th>Equation</th>
<th>Notation</th>
<th>Prior distribution*</th>
</tr>
</thead>
<tbody>
<tr>
<td>2.1</td>
<td>$\gamma$</td>
<td>Beta (1,1)</td>
</tr>
<tr>
<td>2.5</td>
<td>$\beta$</td>
<td>Normal (0, 1,000)</td>
</tr>
<tr>
<td>2.6</td>
<td>$\sigma$</td>
<td>Uniform (0, 100)</td>
</tr>
<tr>
<td>2.7</td>
<td>$B$</td>
<td>Normal (0, 1,000)</td>
</tr>
<tr>
<td>2.7</td>
<td>$\phi$</td>
<td>Normal (0, 1,000)</td>
</tr>
</tbody>
</table>

*Posterior distributions are presented in online Technical Appendix 2 Figures 2, 3 (https://wwwnc.cdc.gov/EID/article/24/12-0879-Techapp2.pdf).

Technical Appendix 1 Figure. Model for analyzing prevalence of avian influenza A(H5) and A(H9) viruses in live bird markets, Bangladesh. The 2-level hierarchical relationship between data and model parameters is presented. Rectangles indicate constants, and circles indicate variables. Solid arrows indicate stochastic dependency, and dashed arrows indicate deterministic dependency. Subscript letters correspond to those in the model description.
Prevalence of Avian Influenza A(H5) and A(H9) Viruses in Live Bird Markets, Bangladesh

Technical Appendix 2.

Technical Appendix 2 Table 1. Prevalence of avian influenza A(H5) and A(H9) viruses in poultry samples from live bird markets, Bangladesh*

<table>
<thead>
<tr>
<th>Sample</th>
<th>No. pools†</th>
<th>Pool-level H5 virus prevalence, %</th>
<th>Pool-level H9 virus prevalence, %</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Cloacal</td>
<td>Oropharyngeal</td>
</tr>
<tr>
<td>Chicken</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Broiler</td>
<td>153</td>
<td>0.7</td>
<td>3.3</td>
</tr>
<tr>
<td>Sonali</td>
<td>122</td>
<td>2.5</td>
<td>6.6</td>
</tr>
<tr>
<td>Desi</td>
<td>127</td>
<td>2.4</td>
<td>5.5</td>
</tr>
<tr>
<td>Subtotal</td>
<td>402</td>
<td>1.7</td>
<td>5.0</td>
</tr>
<tr>
<td>Waterfowl</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Duck</td>
<td>57</td>
<td>14.0</td>
<td>29.8</td>
</tr>
<tr>
<td>Goose</td>
<td>18</td>
<td>11.1</td>
<td>22.2</td>
</tr>
<tr>
<td>Subtotal</td>
<td>75</td>
<td>13.3</td>
<td>28.0</td>
</tr>
<tr>
<td>Total</td>
<td>477</td>
<td>3.6</td>
<td>8.6</td>
</tr>
</tbody>
</table>

*Desi, “local” in Bengali, are indigenous chicken breeds raised in backyard farms. Sonali is a cross-breed of the Rhode Island Red cocks and Fayoumi hens.
†When a given type of poultry was not available, other types of poultry were sampled. All pools contained 5 swab specimens, except for 1 pair of cloacal and oropharyngeal pools from geese, which contained 4 swab specimens.
‡A pool was considered positive if any of its cloacal and oropharyngeal pools tested positive results.

Technical Appendix 2 Table 2. Prevalence of avian influenza A(H5) and A(H9) viruses in environmental samples from live bird markets, Bangladesh

<table>
<thead>
<tr>
<th>Sample</th>
<th>No. pools*</th>
<th>Pool-level H5 virus prevalence, %</th>
<th>Pool-level H9 virus prevalence, %</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stall area†</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water run-off</td>
<td>34</td>
<td>8.8</td>
<td>8.8</td>
</tr>
<tr>
<td>Poultry cage floor</td>
<td>40</td>
<td>15.0</td>
<td>22.5</td>
</tr>
<tr>
<td>Poultry display table</td>
<td>40</td>
<td>10.0</td>
<td>22.5</td>
</tr>
<tr>
<td>Poultry drinking water</td>
<td>40</td>
<td>5.0</td>
<td>27.5</td>
</tr>
<tr>
<td>Poultry waste disposal area/bin</td>
<td>36</td>
<td>19.4</td>
<td>22.2</td>
</tr>
<tr>
<td>Floor in the area where poultry are kept</td>
<td>13</td>
<td>0.0</td>
<td>7.7</td>
</tr>
<tr>
<td>Subtotal</td>
<td>203</td>
<td>10.8</td>
<td>20.2</td>
</tr>
<tr>
<td>Slaughtering area</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Water run-off</td>
<td>40</td>
<td>10.0</td>
<td>35.0</td>
</tr>
<tr>
<td>Floor of slaughtering area</td>
<td>40</td>
<td>12.5</td>
<td>25.0</td>
</tr>
<tr>
<td>Poultry waste disposal area/bin</td>
<td>37</td>
<td>10.8</td>
<td>27.0</td>
</tr>
<tr>
<td>Chopping and slaughtering table</td>
<td>40</td>
<td>10.0</td>
<td>32.5</td>
</tr>
<tr>
<td>Slaughtering and processing knives/board</td>
<td>40</td>
<td>10.0</td>
<td>37.5</td>
</tr>
<tr>
<td>Subtotal</td>
<td>197</td>
<td>10.7</td>
<td>31.5</td>
</tr>
<tr>
<td>Total</td>
<td>400</td>
<td>10.8</td>
<td>25.8</td>
</tr>
</tbody>
</table>

*When a given type of environmental site was not available, other types of environmental site were sampled.
†In each live bird market, 5 of 6 environmental sites were sampled depending on their availability.

Technical Appendix 2 Table 3. LBM-level prevalence of avian influenza A(H5) virus estimated from best H5 models, Bangladesh*

<table>
<thead>
<tr>
<th>Sample type</th>
<th>No. LBMs</th>
<th>Median prevalence, % (95% HDI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Poultry</td>
<td>40</td>
<td>88.7 (69.1–100.0)</td>
</tr>
<tr>
<td>Environmental</td>
<td>40</td>
<td>91.9 (74.7–100.0)</td>
</tr>
</tbody>
</table>

*HDI, high-density interval; LBM, live bird market.
Technical Appendix 2 Table 4. Pool-, bird-, and environmental swab specimen–level prevalence of avian influenza A(H5) virus estimated from best H5 models, Bangladesh*

<table>
<thead>
<tr>
<th>Sample</th>
<th>No. pools</th>
<th>Pool-level prevalence, %</th>
<th>Bird-level median prevalence, % (95% HDI)†</th>
</tr>
</thead>
<tbody>
<tr>
<td>Poultry</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chicken</td>
<td>402</td>
<td>5.5</td>
<td>1.5 (0–4.0)</td>
</tr>
<tr>
<td>Waterfowl</td>
<td>75</td>
<td>33.3</td>
<td>8.9 (0.2–22.1)</td>
</tr>
<tr>
<td>Environmental</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Area</td>
<td>400</td>
<td>12.0</td>
<td>2.6 (0.1–6.9)</td>
</tr>
</tbody>
</table>

*HDI, high-density interval. †Bird- and environmental swab specimen–level prevalence in contaminated live bird markets from the best H5 models.

Technical Appendix 2 Table 5. LBM-level prevalence of avian influenza A(H9) virus estimated from best H9 models, Bangladesh*

<table>
<thead>
<tr>
<th>Sample type</th>
<th>No. LBMs</th>
<th>Median prevalence, % (95% HDI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Poultry</td>
<td>40</td>
<td>98.0 (91.8–100.0)</td>
</tr>
<tr>
<td>Environmental</td>
<td>40</td>
<td>97.6 (90.3–100.0)</td>
</tr>
</tbody>
</table>

*HDI, high-density interval; LBM, live bird market.

Technical Appendix 2 Table 6. Pool-, bird-, and environmental swab specimen–level prevalence of avian influenza A(H9) virus estimated from best H9 models, Bangladesh*

<table>
<thead>
<tr>
<th>Sample</th>
<th>No. pools</th>
<th>Pool-level prevalence, %</th>
<th>Bird-level median prevalence (95% HDI)†</th>
</tr>
</thead>
<tbody>
<tr>
<td>Poultry</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chicken</td>
<td>153</td>
<td>41.2</td>
<td>12.0 (4.8–21.1)</td>
</tr>
<tr>
<td>Sonali</td>
<td>122</td>
<td>32.0</td>
<td>7.3 (2.7–13.5)</td>
</tr>
<tr>
<td>Desi</td>
<td>127</td>
<td>34.6</td>
<td>7.6 (2.8–13.9)</td>
</tr>
<tr>
<td>Waterfowl</td>
<td>75</td>
<td>18.7</td>
<td>3.1 (0.8–6.6)</td>
</tr>
<tr>
<td>Environmental</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Stall area</td>
<td>203</td>
<td>20.8</td>
<td>4.0 (1.0–8.7)</td>
</tr>
<tr>
<td>Slaughtering area</td>
<td>197</td>
<td>30.5</td>
<td>7.7 (2.1–15.8)</td>
</tr>
</tbody>
</table>

*HDI, high-density interval. Desi, "local" in Bengali, are indigenous chicken breeds raised in backyard farms. Sonali is a cross-breed of the Rhode Island Red cocks and Fayoumi hens. †Bird- and environmental swab specimen–level prevalence in contaminated live bird markets from the best H9 models.

A 1A. Retail LBMs  
B 1B. Mixed LBMs  

Technical Appendix 2 Figure 1. Posterior predictive checks of the models presented for analysis of prevalence of avian influenza A(H5) and A(H9) viruses in live bird markets, Bangladesh. A) Retail live bird
markets; B) mixed live bird markets. Dotted lines indicate H5 subtypes, and solid lines indicate H9 subtypes. Diamonds indicate median values, horizontal bars indicate 95% high-density interval of the posterior predictive distribution, and ×s indicate observed pool-level prevalences. p values correspond to the proportion of posterior predictive values that are equal to or more extreme than the observed prevalence. Desi, “local” in Bengali, are indigenous chicken breeds raised in backyard farms. Sonali is a cross-breed of the Rhode Island Red cocks and Fayoumi hens.
Technical Appendix 2 Figure 2. Posterior distribution of parameters used in models for prevalence of avian influenza A(H5) subtype virus in live bird markets, Bangladesh. Desi, “local” in Bengali, are indigenous chicken breeds raised in backyard farms. Sonali is a cross-breed of the Rhode Island Red
cocks and Fayoumi hens. Values along baselines are medians. Solid horizontal bars indicate 95% high density intervals. Each panel (A–L) shows a different situation that is listed at the bottom of each figure panel. A) γ/retail/bird/H5; B) γ/mixedl/bird/H5; C) γ/retail/environment/H5; D) γ/mixed/environment/H5; E) β/Sonalii/H5; F) β/Desii/H5; G) β/waterfowl/H5; H) β/stall/H5; I) B/mixed/bird/H5; J) β/mixed/environment/H5; K) φ/bird/H5; L) φ/environment/H5.
Technical Appendix 2 Figure 3. Posterior distribution of parameters used in models for prevalence of avian influenza A(H9) subtype virus in live bird markets, Bangladesh. Desi, “local” in Bengali, are indigenous chicken breeds raised in backyard farms. Sonali is a cross-breed of the Rhode Island Red.
cocks and Fayoumi hens. Values along baselines are medians. Solid horizontal bars indicate 95% high density intervals. Each panel (A–L) shows a different situation that is listed at the bottom of each figure panel. A) γ/retail/bird/H9; B) γ/mixedl/bird/H9; C) γ/retail/environment/H9; D) γ/mixed/environment/H9; E) β/Desi//H9; F) β/stall/H9; G) β/Sonali/H9; H) β/waterfowl/H9; I) B/mixed/bird/H9; J) β/mixed/environment/H9; K) φ/bird/H9; L) φ/environment/H9.