Unexpected infection outcomes of China-origin H7N9 low pathogenicity avian influenza virus in turkeys

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The China-origin H7N9 low pathogenicity avian influenza virus (LPAIV) emerged as a zoonotic threat in 2013 where it continues to circulate in live poultry markets. Absence of overt clinical signs in poultry is a typical LPAIV infection outcome, and has contributed to its insidious maintenance in China. This study is the first description of H7N9 LPAIV (A/Anhui/1/13) infection in turkeys, with efficient transmission to two additional rounds of introduced contact turkeys which all became infected during cohousing. Surprisingly, mortality was observed in six of eight (75%) second-round contact turkeys which is unusual for LPAIV infection, with unexpected systemic dissemination to many organs beyond the respiratory and enteric tracts, but interestingly no accompanying mutation to highly pathogenic AIV. The intravenous pathogenicity index score for a turkey-derived isolate (0.39) affirmed the LPAIV phenotype. However, the amino acid change L235Q in the haemagglutinin gene occurred in directly-infected turkeys and transmitted to the contacts, including those that died and the two which resolved infection to survive to the end of the study. This polymorphism was indicative of a reversion from mammalian to avian adaptation for the H7N9 virus. This study underlined a new risk to poultry in the event of H7N9 spread beyond China.

A novel H7N9 influenza A virus (IAV) emerged from poultry as a zoonotic infection in China in 20131. A series of annual winter epidemics of respiratory infection in humans have occurred2, with 1589 confirmed cases, including 616 deaths, by 20th September 20173, the majority of which have been associated with poultry contact at live poultry markets (LPMs)4,5. Widespread dissemination of H7N9 within poultry in China is attributed to it being a low pathogenicity avian influenza virus (LPAIV) with no obvious clinical signs in birds, especially in the chicken which is assumed to be the main maintenance host6,7.

Genetic polymorphisms in this H7N9 LPAIV include Q235L (complete gene numbering) in the haemagglutinin (HA) gene of many human isolates which reflects mammalian adaptation. This residue is also identified as position 217 in the H7 HA1 mature peptide which corresponds to position 226 in the H3 numbering convention5. The E627K (PB2 gene) polymorphism has been reported in many human isolates8 with continuing evolution evident at the full-genome level in both human and poultry isolates9–11. The H7N9 internal genes originated from endemic Chinese H9N2 viruses via reassortment1,9. However, despite its widespread circulation in poultry in several Chinese provinces, the emergence of a highly pathogenic (HP)AIV cleavage site (CS) sequence was reported relatively recently during winter 2016–201712,13.

Several published in vivo poultry studies of China-origin H7N9 LPAIV have included chickens14–19, using mainly the prototype human-origin A/Anhui/1/13 H7N9 LPAIV isolate7 for infection, with absence of overt clinical signs reflecting the field situation in China. Inclusion of contact chickens enabled the transmission frequency

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within this species to be assessed, with outcomes ranging from highly efficient or intermediate infection of contacts\textsuperscript{14,15,18} to nil transmission\textsuperscript{16,19}. However, a proportion of H7N9 LPAIV human isolates and LPM isolates from avian and environmental specimens possess the Q235 polymorphism which occurs in avian-origin IAVs\textsuperscript{10,14}. Although H7N9 LPAIV has been isolated occasionally from domestic ducks in Chinese LPMs\textsuperscript{10,14}, experimental investigation has shown that ducks and geese are infected less efficiently \textit{in vivo} with infrequent or poor transmission\textsuperscript{17,19}. Turkeys, however, are an insignificant poultry species in China, which contrasts with the substantial commercial turkey sector in Europe and North America\textsuperscript{20}. Given that any dissemination to new geographic regions might result in infection of non-chicken hosts\textsuperscript{21,22}, this study compared the \textit{in vivo} susceptibility of chickens and turkeys to China-origin H7N9 LPAIV, and evaluated its pathogenicity and transmission within turkeys.

**Materials and Methods**

**Viruses.** A/Anhui/1/13 H7N9 LPAIV was received from the National Institute for Medical Research, UK, and had been previously passed through three rounds (egg passage (EP)3) using 9–11 days-old specific pathogen free (SPF) embryonated chickens’ eggs\textsuperscript{23}. Three further passages at APHA produced the EP6 stock which served as the inoculum in the \textit{in vivo} experiments, and henceforth will be referred to as H7N9 “wild type” (wt). The EP6 H7N9 was full-genome sequenced and confirmed as identical to the original GenBank sequence submission (accession numbers: CY187618–CY187625) except for an amino acid change in the HA, namely N141D (complete gene numbering). Virus isolation (VI) from clinical specimens derived from the \textit{in vivo} experiments in this study was also done in chicken eggs and 12-days-old embryonated turkey eggs\textsuperscript{23}. The EP6 stock and H7N9 LPAIV isolates were titrated in chicken eggs to determine the 50% egg infectious dose (EID\textsubscript{50})\textsuperscript{24}.

**Ethics statement and containment work.** All animal experiments and procedures required approval from the local APHA Animal Welfare and Ethical Review Body to comply with the relevant European and UK legislation\textsuperscript{25} and were carried-out in accord with the Home Office (UK) Project Licence 70/8332. Any infected poultry which began to display severe clinical signs were euthanised and were recorded as a mortality. UK regulations categorise the H7N9 LPAIV as a SAPO 4 and advisory ACDP 3 pathogen because it is a notifiable animal disease agent\textsuperscript{26} and presents a zoonotic risk\textsuperscript{27}. All laboratory and containment work with H7N9 specimens, including infected poultry, was done in licenced BSL 3 facilities\textsuperscript{27,28}.

**Experimental design.** Prior to infection, birds were swabbed and bled for M-gene RRT-PCR and serological testing to exclude previous AIV exposure. Sixteen SPF chickens (White Leghorn, Valo) and 48 turkeys (Commercial White, high health status, Aviagen) were used to attempt a transmission chain by direct-inoculation followed by contact exposure. All birds were 3-weeks-age, with six chickens and 18 turkeys being directly-inoculated (100 µl per bird, via the ocular-nasal route) and referred to as “donor” (D0) birds in that they served to attempt to transmit infection to contact birds. Contacts are referred to as “recipients”, abbreviated as R1 birds for “first contacts” which were introduced for cohousing with D0 birds at 1 day post-infection (dpi) (Fig. 1). Cohousing investigated transmission of infection via the natural contact route and was attempted in four experimental units, summarised as follows:

(a) Chicken (D0) to chicken (R1) transmission, with H7N9 wt used to directly-inoculate one group of D0 chickens housed in a single pen using a single dose of 8log\textsubscript{10} EID\textsubscript{50} per chicken.

(b) Turkey (D0) to turkey (R1) transmission, with three different doses of H7N9 wt used to directly-inoculate three separate groups of D0 turkeys housed in separate pens, namely 8, 6 and 4log\textsubscript{10} EID\textsubscript{50} per turkey (Fig. 1).

The four experimental units were divided for housing into four Perspex pens (120 cm × 120 cm (total floor space 1.44 m\textsuperscript{2}) x 60 cm height) which provided housing density in accord with UK animal welfare recommendations\textsuperscript{29}. Details of infection and cohousing which included attempts to extend the transmission chain beyond the R1 birds are outlined for both species.

\textbf{Figure 1.} Schematic showing the plan for \textit{in vivo} transmission of H7N9 LPAIV. Direct-inoculation of D0 (“donor”) birds followed by introduction of two rounds of R1 and R2 (“recipient”) contact birds into each pen, including the planned removal (by culling) of D0 and R1 birds at 4 and 7 dpi respectively. Cohousing periods (3-days) for D0/R1 and R1/R2 birds are indicated by blue and pink shading respectively, with R2 birds intended to remain (unshaded days) to the conclusion of the study at 15 dpi. See Results for subsequent differences from this plan.
Chicken to chicken transmission, with attempted further transmission to turkeys. An ocular/nasal dose of 8log<sub>10</sub> EID<sub>50</sub> of H7N9 wt was directly administered to six D0 chickens. At 1 dpi an equal number of six R1 chickens were introduced into the pen for cohousing for three days when the six D0 chickens were withdrawn at 4 dpi (3 days post-contact (dpc)). Replacement of the six D0 chickens with six R2 ("second contacts") turkeys at 4 dpi was informed by the transmission findings observed at the same time in the turkey-to-turkey transmission groups (described below): The six R1 chickens were withdrawn at 7 dpi (6 dpc), with the four R2 turkeys remaining to the conclusion of the study at 15 dpi, i.e. 11 dpc.

Turkey to turkey transmission. An ocular/nasal dose of 8log<sub>10</sub> EID<sub>50</sub> of H7N9 wt was directly administered to six D0 turkeys in a single pen, with two additional turkey pens housing six D0 turkeys in each for direct-inoculation at 6log<sub>10</sub> EID<sub>50</sub> and 4log<sub>10</sub> EID<sub>50</sub>. At 1 dpi an equal number of R1 turkeys were introduced to each of the three D0 turkey groups (Fig. 1). For the two pens which contained D0 turkeys directly-inoculated with the two highest H7N9 doses, cohousing continued for three days before the six D0 turkeys were withdrawn at 4 dpi from each pen, culled and replaced with four R2 turkeys to extend the transmission chain, i.e. from R1 to R2 turkeys (Fig. 1). It was planned to withdraw and cull the R1 turkeys from the three pens at 7 dpi (6 dpc), and to allow the R2 turkeys in these two pens to remain to the conclusion of the study at 15 dpi (11 dpc). In the pen containing six D0 turkeys infected with the lowest H7N9 dose (4log<sub>10</sub> EID<sub>50</sub> per turkey), six R1 turkeys were introduced at 1 dpi, with withdrawal and culling at 4 dpi and 7 dpi (6 dpc) respectively with no R2 turkeys introduced due to the inefficient acquisition of infection by the R1 turkeys.

Collection of clinical specimens from chickens and turkeys. Buccal and cloacal swabs were collected daily from all birds and stored in 1 ml of brain heart infusion broth containing antibiotics, namely 1000 IU penicillin G, 10 mg/ml amphotericin B and 1 mg/ml gentamicin (BHI). Organs (see Results) were collected from birds which were either culled according to the experimental plan (Fig. 1) or died, and were cut for 10% (w/v) suspension in BHI<sup>30</sup>. Blood was collected for serum separation from birds at cull or which survived to the conclusion of the experiment at 15 dpi.

RNA extraction and AIV reverse transcription Real-Time PCR (RRT-PCR). RNA was extracted from BHI swab fluids and tissue supernatants by robotic and manual methods respectively<sup>30</sup>. All extracted RNA specimens were tested by the M-gene RRT-PCR, whereby 2 µl RNA was added to a final 25 µl reaction volume containing 12.5 µl (× 2) QuantiFast Probe RT-PCR Master Mix (Qiagen), 0.5 µl (× 50) ROX dye solution and 0.25 µl QuantiFast RT Mix (from same Qiagen kit), the primers and probe<sup>31</sup> at final concentrations of 0.6 µM and 0.2 µM respectively. The proprietary lock nucleic acid probe is available from the Roche universal probe library (UPL) 104<sup>′</sup> with the sequence 5′FAM-CTGGGCAC-BHQ1–3′ (Roche). Thermocycling was performed using an Agilent Mx3000 instrument (Stratagene) as described<sup>32</sup> with the RT inactivation at 95 °C shortened to 5 minutes and the number of PCR cycles reduced to 40. Ct values 36 were considered as AIV positive, while those in the range Ct 36.01–39.99 were sub-threshold, and full-genome sequences obtained by Next Generation Sequencing (NGS) as previously<sup>33</sup>. For molecular pathotyping, RNA was directly extracted from clinical specimens but with the carrier RNA included, followed by GK7.3/GK7.4<sup>34</sup> amplification across the HA gene CS sequence as described<sup>30</sup>. To investigate the L235Q mammalian/avian adaptation polymorphism in the HA gene, two primers specific for H7N9 wt were designed which flanked this region:

L235Q fwd: 5′-ATTCCCGCAGATGACTAAG-3′
L235Q rev: 5′-ACTGCCCTGCTATCTATGTT-3′

This L235Q primer pair substituted for the GK7.3/GK7.4 primers in the above conventional RT-PCR with identical thermocycling to produce a 436 bp amplicon. Products were confirmed and purified for cycle-sequencing with the two corresponding RT-PCR primers<sup>35</sup>. Raw sequence data was obtained in both orientations from the HA CS and L235Q conventional (i.e. gel detection) RT-PCRs, and was assembled by the SeqMan II suite (Lasergene bioinformatics package, version 13; DNASTar, Madison, USA) to provide the consensus sequence. Any instances of possible mixed L/Q235 polymorphisms were initially identified by the SeqMan II software interpretation, followed by visual inspection of any dual chromatogram peaks at the same nucleotide
position. In instances of uncertainty, cycle sequencing of the purified amplicons was repeated to identify residue 235 as either distinct L or Q amino acids, or as a mixed population of both indicated by X.

Analysis of H7N9 genetic polymorphisms. In order to assess the possible significance of any H7N9 genetic polymorphisms which emerged following \textit{in vivo} infection of chickens and turkeys, 7792 sequences of the relevant China-origin H7N9 genes were downloaded from the GISAID database (http://platform.gisaid.org/epi/i3) on 24/1/2018 (deposited since 2013) and included 77 sequences from H7N9 HPAIVs deposited since early 2017. The downloaded sequences were aligned along with the corresponding gene sequences from the isolated progeny viruses by using the MAFFT multiple sequence alignment software (version 7)\textsuperscript{35}.

H7- and AIV-specific serology. Sera derived from clotted blood were incubated at 56 °C. Turkey sera (100 µl) were pre-absorbed by adding 20 µl packed chicken red blood cells and incubated for 30 minutes at ambient temperature with occasional agitation. HI testing for HA-specific antibodies in both poultry species used the homologous H7N9 viral antigen titrated to 4 haemagglutination units\textsuperscript{23}. Sera were also tested by the multi-species influenza A ELISA (IDEXX, France) and the FluA ELISA (ID Vet, France), both of which are blocking ELISAs that detect antibodies against the IAV type-specific NP antigen. The H7-specific antibody ELISA (version 2, ID Vet) was also used.

Intravenous pathogenicity index (IVPI). The IVPI test was performed and scored by standard methods\textsuperscript{23} with egg fluids diluted to a titre of 9log\textsubscript{10} EID\textsubscript{50}/ml\textsuperscript{18} and 100 µl administered intravenously per four-week-old chicken. If a chicken required euthanasia during the 10-day IVPI determination, it was recorded as having been found dead on the next day.

| Species and dose administered to donor (D0) birds | Chickens 8 log\textsubscript{10} EID\textsubscript{50} | Turkeys 8 log\textsubscript{10} EID\textsubscript{50} | Turkeys 6 log\textsubscript{10} EID\textsubscript{50} | Turkeys 4 log\textsubscript{10} EID\textsubscript{50} |
|---|---|---|---|---|
| Directly-infected or contact | D0 | D0 | D0 | D0 |
| Dpi (dpc): all culled (apparently healthy), except E and FD as indicated | 4 | 4 | 7 (6) | 7 (6) |
| Bird ID #: | 1 | 3 | 11 | 12 |
| L235Q polymorphism in HA gene | Q235 | L235 | ND | Q235 |
| Organs | | | | |
| Nasal cavity (turbinates) | − | + | − | − |
| Airways** | − | − | − | − |
| Lung** | − | − | − | − |
| Air Sacs** | − | − | − | − |
| Proventriculus** | − | − | − | − |
| Caecal Tonsil** | + | − | − | − |
| Intestine (enterocytes) | − | − | − | − |
| Pancreas** | − | − | − | − |
| Spleen** | − | − | N/a | + |
| Bursa** | − | − | − | − |
| Thymus* | − | − | − | − |
| Skin* | − | − | − | − |
| Feather follicles* | − | − | − | − |
| Heart** | − | − | − | − |
| Skeletal muscle* | − | − | − | − |
| Brain* | − | − | − | − |
| Kidney** | − | − | − | − |
| Liver** | − | − | − | − |
| Gonads* | − | − | − | − |

Table 1. Detection of IAV NP antigen in organs by immunohistochemistry. Four experimental groups indicated in the header, divided by D0 species and direct dose of H7N9wt (log\textsubscript{10} EID\textsubscript{50}). E = euthanised, FD = found dead. L235Q polymorphisms indicated for individual birds at time of cull or death. * Detected only in endothelial cells; ** Detected in endothelial and parenchymal cells. N/a: not available; ND not done. Additional infected R2 turkeys # 40 and # 142 (see Results) survived to be culled at the end of the study (11 dpc). No detectable IAV-specific NP antigen was detected in these two survivors by IHC.
Results

Infection and transmission of H7N9 LPAIV in chickens and turkeys. Swabs and sera collected from all 60 birds prior to inoculation or exposure confirmed all to be negative for AIV shedding and seronegative by homologous HI and both anti-NP ELISAs. Direct-inoculation of D0 chickens and turkeys with the high $8\log_{10} \text{EID}_{50}$ dose resulted in infection of all six (100%) in both pens (Fig. 2a and b, Supplementary Table S1). Introduction of R1 chickens and turkeys into the respective pens at 1 dpi for co-housing with the D0 birds resulted in 100% infection (100%) in both pens (Fig. 2a and b, Supplementary Table S1).
the same species (Fig. 1) resulted in two of six (33%) R1 chickens becoming infected (Fig. 2a, Supplementary Table S1). By contrast, all R1 turkeys (6/6; 100%) became infected (Fig. 2b). Replacement of D0 turkeys with R2 turkeys at 4 dpi (Fig. 1) resulted in onward transmission of infection to all four (100%) of the latter (Fig. 2b). Successful turkey transmission was observed in a third pen where the lower 6log10 EID50 dose also resulted in 6/6 (100%) D0, 6/6 (100%) R1 and 4/4 (100%) R2 turkeys becoming infected (Fig. 2c, Supplementary Table S1). Mean positive buccal shedding preceded cloacal shedding in D0 and R1 turkeys, but a similar time of mean shedding onset was observed for both tracts in the R2 turkeys (Fig. 2b and c). In a fourth pen, direct inoculation with the lowest (4log10 EID50) resulted in infection in 4/6 (67%) D0 turkeys with only 3/6 (50%) R1 turkeys infected, indicating reduced transmission at this inoculation dose (Supplementary Fig. S1 and Table S1). These experiments showed that >4log10 EID50 was the minimal infectious dose required to establish shedding in all D0 turkeys, with accompanying successful transmission of H7N9 infection to all R1 and then R2 contact turkeys as depicted in the schematic diagram (Fig. 1).

The ready susceptibility to contact infection in turkeys informed the decision to introduce four R2 turkeys at 4 dpi into the pen where chicken D0 to R1 transmission had been attempted and shown to be inefficient. Three of the four introduced R2 turkeys became infected (Fig. 2a and Supplementary Table S1), underlining the susceptibility of this species to contact infection. However, the mean shedding titres were generally lower and delayed in R1 and R2 contact turkey groups where infection was defined on the basis of single instances of positive shedding and/or sub-threshold shedding for two successive days (Fig. 2a, Supplementary Fig. S1 and Table S1).

**Mortality in contact-infected (R1 and R2) chickens and turkeys.** No mortality or overt clinical signs were observed in any of the D0 and R1 infected chickens. However, in the 8log10 EID50 turkey pen, 1/6 (17%) of R1 turkeys died suddenly at 136 hpc (5.67 dpc) without any prior clinical signs (Fig. 3). A further 3/4 (75%) of the R2 turkeys in the same pen died between 88 and 184 hpc (mean time to death (MDT): 144 hpc (6.0 dpc)). In the 6log10 EID50 turkey pen there were no deaths among the six R1 infected turkeys, but 3/4 (75%) of the R2 turkeys died between 100 and 172 hpc (MDT: 140 hpc (5.83 dpc)) (Fig. 3). Five of the six R2 turkeys which died in both pens displayed ruffled feathers and huddled behaviour during the preceding day. Among these five R2 turkeys, one also had closed eyes and was incapable of independent feeding and drinking, while another developed drooping wings and an unsteady gait, hence both were euthanised. The sixth R2 turkey died suddenly, with no obvious clinical signs when inspected 1 hr earlier. The gross pathology of all seven turkeys which died (Fig. 3) was characterised by marked pancreatic haemorrhage (Supplementary Fig. S2). One apparently healthy R1 turkey which was pre-planned for cull at 6 dpc also displayed such gross haemorrhagic lesions of the pancreas (#11, Table 1). There were no clinical signs nor deaths among any of the D0 and R1 infected turkeys in the pen where the initial inoculation dose was 4log10 EID50.

**Organ tropism of H7N9 LPAIV in chickens and turkeys.** H7N9 RNA was detected in the turbinate, intestine and spleen of D0 chickens, culled at 4 dpi (Fig. 4a). The IHC approach, known to be less sensitive than AIV RRT-PCR, revealed uncommon virus-specific staining in the turbinates (Fig. 5) of one D0 chicken and the caecal tonsil of another (Table 1). H7N9 viral RNA was also detected in the turbinates of one R1 infected chicken at 116 hpc (5.67 dpc). H7N9 RNA titres in many organs when culled at 4 dpi included the intestine, turbinates, trachea, lung, air sacs and spleen (Fig. 4b). IHC investigation affirmed systemic viral dissemination which included the pancreas, kidney, heart, liver and brain (Fig. 5 and Table 1). D0 turkeys which were infected with the lower 6log10 EID50 dose included several negative or sub-threshold viral RNA titres in organs (Fig. 4c). IHC also revealed less extensive or absent virus-specific staining in several organs compared to the D0 turkeys that had been infected with 8log10 EID50 thereby reflecting a dose/dissemination effect (Table 1). However, R1 and R2 turkeys in both pens harboured high viral titres in organs collected at time of cull or death (Fig. 4b and c), with IHC confirming systematic dissemination that included notable endothelial tropism (Table 1).

The two R2-infected turkeys which survived to the end of the experiment from the 8log10 and 6log10 EID50 pens were culled at 11 dpc (15 dpi) and had seroconverted by all four antibody tests (Supplementary Table S2). Both displayed a huddled stance and ruffled feathers for 2 days (turkey # 40) and 1 day (turkey #142), commencing at 7 and 8 dpc respectively, followed by resolution. Organs sampled from both recovered R2 turkeys revealed reduced viral RNA titres compared those of R2 turkeys which died (Fig. 4b and c), and this was reflected in the absence of virus-specific IHC staining in both (Table 1 footnote). Examination of the cloacal shedding from these R2 recovered turkeys showed that both experienced a later onset and lower mean titres compared to the six R2 turkeys which had died (Supplementary Fig. S3).

**Seroconversion following H7N9 infection and/or exposure.** D0, R1 and R2 infected birds were bled at cull, i.e. 4 dpi, 6 dpi and 11 dpc respectively. Progressive seroconversion to the homologous H7N9 H1 antigen was observed (Supplementary Table S2). Among all the 31 available post-infection/exposure sera available for homologous H7 H1 testing, 16/31 (52%) were seropositive, while the 32 sera tested by the anti-NP ELISAs included 10/32 (31%) which were seropositive by one of the two (i.e. the ID Vet or IDEXX AIV ELISA; n = 5) or both anti-NP ELISAs (n = 5). Only 4/32 (13%) of sera were H7 seropositive by the H7-specific ELISA.

**Genetic changes observed in H7N9 progeny viruses in infected birds.** Molecular pathotyping of progeny viruses. The observation of unexpected mortality (75%) among the R2 turkeys (Fig. 3) indicated a distinct viral phenotype which contrasted with an absence of overt clinical signs in chickens. This difference prompted molecular pathotyping investigations of 52 swabs and one brain specimen from D0, R1 and R2 turkeys which all revealed an unchanged LPAIV CS sequence compared to the H7N9 wt inoculum, namely PEIPKGRLF...
The 10 chicken swabs obtained from D0 and R1 birds also revealed an unchanged LPAIV CS sequence.

**L235Q polymorphisms in HA gene.** Full-genome sequencing of the four R2 turkey and the one D0 chicken isolate all revealed the L235Q polymorphism to have occurred in the HA of all five (Table 2). Investigation of 51 swabs collected from D0 turkeys inoculated with the $8\log_{10}$ and $6\log_{10}$ EID$_{50}$ doses showed that a mixed L/Q (X) polymorphism emerged followed by Q235 becoming increasingly prevalent by 4 dpi (Fig. 6a and b). All 85 (57 buccal, 28 cloacal) and all 37 swabs (24 buccal, 13 cloacal) sampled at 1–6 dpc and 1–8 dpc from R1 and R2 turkeys respectively possessed Q235, although two of the R1 turkey swabs were an X235 mixture. Investigation of 24 swabs from the infected D0 chickens displayed a very similar emergence of the X235 and Q235 polymorphisms (Fig. 6c). All four buccal swabs from the R1 chickens (sampled 3–6 dpc) possessed the Q235 polymorphism. No other amino acid changes were observed elsewhere in the HA genes of the four isolates obtained from the R2 turkeys and the D0 chicken (Table 2).

**Changes in other non-HA genetic segments.** Additional amino acid changes were identified in other gene segments among three of the four isolates obtained from the R2 turkeys and the D0 chicken, although for the turkey-derived isolates there were some differences depending on whether chicken or turkey eggs were used for
isolation (Table 2). These polymorphisms were identified as changes relative to the H7N9 wt inoculum, with the PA R269K, NA T10I/S and PB1-F2 M1T (possible ablation of peptide) changes having also occurred in other sequenced China-origin H7N9 isolates (Supplementary Table S4). However, the PB2 Y360H and PA N10H polymorphisms appear to be unique to the viral progeny obtained in the current study.

**IVPI.** IVPI tests were done on two H7N9 LPAIV isolates, namely the EP6 inoculum stock (H7N9 wt, chicken egg passaged) and the isolate recovered in turkey eggs from the kidney of a R2 turkey which died, referred to as H7N9 turkey-adapted (ty-adapted) (Table 2). Both the H7N9 wt and ty-adapted isolates registered IVPI scores of 0.00 and 0.39 respectively which were <1.2, therefore confirming the officially-defined LPAIV phenotype for both.

**Discussion**

While H7N9 LPAIV infection of chickens caused no clinical signs as in previous studies\textsuperscript{14-19}, unexpected mortality occurred in 75% of infected R2 turkeys (Fig. 3) following efficient transmission (Fig. 2b and c, Supplementary Table S1). Virulent infection resembled HPAIV pathogenesis and mortality, with H7N9 detection in feather

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**Figure 5.** Examples of IAV-specific IHC staining. Demonstration of virus-specific immunolabelling (brown staining) of the NP in the turbinates, pancreas and kidney of turkeys and chickens where the initial D0 inoculation was 8log\textsubscript{10} EID\textsubscript{50} dose per bird. Organs from D0 and R1 birds were collected at 4 dpi (cull of both species) and 6 dpc (cull for chicken, death for turkey) respectively, with the R2 turkey sampled at death at 4 dpc. The R1 chicken was uninfected and its tissue sections served as a negative control for IHC.
H7N9 dose for initial direct infection of donor (D0) birds | Identifier # for sampled bird (see Table 1) | Sampling of clinical specimens for VI at indicated time of calf or death (dpi or dpc), while * indicates sampling time of live ty | Unchanged H7 LPAIV CS confirmation by NGS of isolates | HA polymorphisms identified in HA by NGS of isolates of isolate | Amino acid changes in other (non-HA) genetic segments identified by NGS of isolates
--- | --- | --- | --- | --- | ---
8log10 EID<sub>50</sub> | D0 ck (* 5) | 4 dpi, cloacal, nasal, tongue, oropharynx, tracheal aspirate, pericardial fluid, heart, liver, spleen, lung, kidney, kidney, lung, liver, spleen | Yes | L235X | PA: N10H
8log10 EID<sub>50</sub> | R2 ty (* 38) | 7 dpi, kidney, FD (IVPI of turkey egg isolate: 0.39) | Yes | L235Q | Chicken egg isolate: NA: T10L; turkey egg isolate: NA: T10S
8log10 EID<sub>50</sub> | R2 ty (* 40) | 9 dpi*, cloacal swab, survived to study end at 11 dpi | Yes | L235Q | Chicken egg isolate: PB2: Y360H; turkey egg isolate: PB2: Y360H
6log10 EID<sub>50</sub> | R2 ty (* 141) | 7 dpi, cloacal swab, EUTH | Yes | L235Q | No changes in other segments
6log10 EID<sub>50</sub> | R2 ty (* 141) | 7 dpi, pancreas, EUTH | Yes | L235Q | Chicken egg isolate: PA: R269K; turkey egg isolate: PA: R269K; PB1-F2: M1T

Table 2. Sequencing findings of H7N9 progeny viruses obtained as virus isolates from a directly-infected chicken (D0) and three infected turkeys (R2). Full-genome sequences were obtained by NGS of the D0 chicken H7N9 isolate (obtained by VI in chicken eggs), while both chicken and turkey eggs were used to isolate H7N9 from the R2 turkeys. Further evidence for an unchanged LPAIV CS during H7N9 in vivo infection and transmission is summarised in Supplementary Table S3. The broader occurrence of the identified polymorphisms in the PB2, PB1-F2, PA and NA genes was investigated by analysing all available China-origin H7N9 gene sequences (Supplementary Table S4). Species abbreviations: ck = chicken; ty = turkey; FD = found dead; EUTH = euthanised; NGS = Next generation sequencing.

follies of D0-infected turkeys (Table 1) implying that viremia may have systemically disseminated the infection, reflecting similarities with viral tropism in feathers during H5N1 HPAIV outbreaks<sup>36,37</sup>. Examination of turkey brain tissues revealed infection (Table 1 and Fig. 4b), thereby also resembling HPAIV tropism<sup>38</sup>. However, investigation of the CS sequences in the turkey progeny viruses revealed the LPAIV CS (monobasic) sequence to be unchanged throughout transmission (Supplementary Table S3), with the LPAIV phenotype (IVPI) confirmed by testing an R2 turkey isolate, namely H7N9 ty-adapted. It was important to establish that no mutation had occurred in the monobasic LPAIV CS during transmission in turkeys, particularly in view of the emergence of H7N9 HPAIV variants during the “fifth wave” in China during winter 2016–2017<sup>12,13</sup>. Inefficient transmission from D0 to R1 chickens reflected other studies which described considerable variability in H7N9 LPAIV transmission within this host<sup>14–16,18,19</sup>.

Little or no mortality is normally observed during most H7 LPAIV turkey outbreaks<sup>39–41</sup>, but up to 40% mortality has been reported in association with secondary bacterial infections and husbandry factors<sup>42,43</sup>. Experimental infection of high health status turkeys with H7 LPAIV’s resulted in up to 40–60% mortality for three of 12 North American isolates<sup>44</sup>, but without the extensive systemic dissemination described in the current study (Figs 4 and 5, Table 1). Because SPF turkeys are unavailable, the H7N9 LPAIV study also used high health status turkeys which were unlikely to be complicated by secondary bacterial infections, with the absence of exudate in the lungs and no Gram staining in tissue sections (data not shown) indicating that secondary infection was unlikely to have contributed to the H7N9 LPAIV virulence. The containment facilities included air supply via HEPA filters which reduced the risk of acquiring exogenous respiratory pathogens. Extensive endothelial tropism was clear, and in a number of organs infection had also spread into the adjacent parenchymal tissues (Table 1), hence this widespread pathogenesis is likely to have contributed to multiple-organ failure in turkeys<sup>45</sup>, as exemplified by gross haemorrhage in the pancreas (Supplementary Fig. S2).

Possible reasons for the unusual virulence in turkeys merit consideration. IAV host adaptation and pathogenesis mechanisms are emphasised in pandemic concerns regarding avian to human adaptation<sup>46–48</sup>, but the context of this avian study differed in that clearly distinct outcomes resulted in two poultry hosts. The L235Q change in the HA was detected soon after infection of D0 birds (both species) and was maintained during efficient transmission to R2 turkeys. This polymorphism is long-known to affect avian to mammalian adaptation of IAVs, whereby Q235L influences a general change in host cell receptor preference from those which possess α-2,3 to α-2,6 sialic acid linkages, known as “avian” and “human” receptors respectively, although H7N9 wt (L235) binds to both α-2,3 and α-2,6 receptors with a slight preference for the former<sup>49</sup>. The L235Q polymorphism appears to be a reversion away from human receptor binding<sup>50</sup>, and was previously reported following chicken infection both in vivo and in ovo<sup>51</sup>. It is speculated that the distribution of α-2,3 receptors in turkeys may enable the greater H7N9 dissemination within turkeys compared to chickens and was supported by the four D0 turkeys sacrificed at 4 dpi which displayed differing organ tropism: The two which received 8log10 EID<sub>50</sub> were already experiencing extensive systemic distribution while the two which received 6log10 EID<sub>50</sub> displayed typical LPAIV turkey tropism largely restricted to the respiratory tract<sup>51</sup>, with the former two having already changed to Q235 while the latter two turkeys possessed L235 (Table 1). A dose effect is the likely cause of these differences, but it is speculated that a greater distribution of avian receptors in multiple turkey tissues may have assisted systemic dissemination of LPAIV infection (with consequences for virulence) following the L235Q switch. Both receptor types broadly occur in the turkey respiratory and enteric tracts<sup>52,53</sup>, but their distribution in other organs remains to be investigated.

Polymorphisms at other residues in the HA are also known to influence its acid-stability which is the optimal pH at which HA-driven fusion of the viral envelope occurs with the endosomal membrane at the early stages of intracellular infection<sup>54</sup>. The fusion pH correlates with phenotypic changes that include the IAV avian to mammalian adaptation switch. For example, HA acid-stability was altered by mutating particular residues in H5N1 HPAIV which changed its transmission and pathogenesis among ducks<sup>55</sup>, which also affected these phenotypic
characteristics in mice and ferrets (reviewed by Russell54). In a purely avian context, the fusion pH of two closely
related H7N3 LPAIVs was studied using a mallard-derived (precursor) isolate and its derivative which caused
turkey outbreaks. The two isolates differed in the HA and NA at one position in each, with the turkey isolate
experiencing viral/endosomal fusion at a higher pH than the mallard isolate which may reflect adaptation to the
avian host species of origin56. However, other than the L235Q switch, the HA protein sequences of the four H7N9
R2 turkey isolates and the chicken D0 isolate remained identical (Table 2), suggesting that HA acid-stability may
be an unlikely factor in explaining differences in transmission, systemic dissemination and pathogenesis in both

**Figure 6.** Changing proportions of polymorphisms at residue 235 in the viral HA gene of directly-inoculated
(D0) birds swabbed at 1–4 dpi. (a and b) presents D0 turkeys inoculated with 8log_{10} and 6log_{10} EID_{50} of H7N9
wt respectively, while (c) presents D0 chickens inoculated with 8log_{10} EID_{50} of H7N9 wt. The total numbers of
swabs of both types investigated at each daily time point are denoted by “n”, with the numbers of buccal and
cloacal swabs distinguished in parenthesis (buccal, cloacal). All cloacal swabs contained Q235 except one in
panel (b) at 4 dpi which contained L235.
hosts. The fusion pH in H7N9 LPAIV decreased due to the L235Q polymorphism 50, but it is unclear whether this altered acid-stability may influence the enhanced dissemination and virulence in turkeys.

Interestingly, acid-stability changes correlated with altered sensitivity of different IAVs (including a reverse-genetic H7N9) to anti-viral host responses which included interferon-induced transmembrane proteins which inhibit viral and cellular membrane fusion 57. More broadly, it is possible that differences between chicken and turkey host responses to H7N9 LPAIV infection may influence the distinct organ tropism and pathogenesis. The possible role of an immune response was underlined by the two R2 turkeys which resolved infection and survived to the end of the study. Both turkeys experienced a delayed onset and generally lower titres of enteric shedding compared to the R2 turkeys which died (Supplementary Fig. S3). It is speculated that the two turkeys produced a successful host response to limit viral replication and tropism, these observations being more akin to a typical LPAIV infection in this species 51, while the turkeys which died did not. In vitro studies have shown that innate immune and apoptotic responses differ in chickens and ducks following H5N1 HPAIV infection which may in turn affect the clear differences in clinical outcomes in these two species 52,53. Infection of chickens, quails and ducks with H7N9 LPAIV resulted in the strongest early IFN-α and TLR-7 responses occurring in the chickens, with the authors noting that this antiviral response correlated with lower viral replication in this species compared to quail 54. However, there is presently limited knowledge concerning further details of turkey host responses to any AIV infection, particularly the role of innate immunity.

Enzootic tropism in many organs which distinguished H7N9 LPAIV infection of turkeys from chickens has already been noted (Table 1), with a recent study showing that innate immunity escape may enable this localisation for some LPAIV strains which may contribute to severe LPAIV infections in galliforme poultry 55. LPAIV tropism in avians is classically restricted to the respiratory and enteric tracts due to the localisation of trypsin as the protease required for HA CS activation during infection 56. However, chicken H9N2 in vitro studies have shown that other proteases may activate the LPAIV CS 53,54, and in some circumstances include the ubiquitous furin protease, normally associated with HPAIV CS activation 55. While there are no reported descriptions of alternative proteases in turkey tissues, it is speculated that non-trypsin proteases may have facilitated H7N9 LPAIV CS activation in multiple visceral organs.

Intravenous inoculation of both H7N9 variants in chickens yielded a higher IVPI score for ty-adapted compared to wt because of one chicken mortality. The four full-genome sequenced R2 progeny viruses from turkeys and the chicken-derived isolate included polymorphisms in the non-HA genetic segments (Table 2), but there was no clear association between these and any known pathogenicity markers, with some of the polymorphisms having been previously observed in China-origin H7N9 (both LPAIVs and HPAIVs) since 2013 (Table S4). Further infection studies with both H7N9 LPAIV wt and ty-adapted variants in turkeys (including intravenous administration, albeit this is not a natural route of infection) may provide clues concerning pathogenesis mechanisms in this host.

In view of the continuing viral evolution and expansion of infection in Chinese poultry, these novel findings will contribute to continuing risk assessments concerning future spread of this H7N9 LPAIV beyond China 12,22. Serological approaches will remain important for active surveillance in chickens 64, while, unusually for LPAIVs, passive surveillance may be also required to monitor for China-origin H7N9 LPAIV incursions into turkeys. In conclusion, this study presented a potentially dangerous notifiable disease scenario where, in the event of China-origin H7N9 LPAIV spreading beyond Asia, undetected maintenance within the chicken sector may be followed by introduction, spread and significant mortality in turkeys.

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**Author Contributions**
M.J.S., A.H.S., S.M.B. and I.H.B. conceived and designed the study. M.J.S., A.H.S., S.T., A.P., S.W., A.M.P.B., D.H. and A.N. did the experiments. M.J.S., A.H.S., S.T., S.M., A.M.P.B. and A.N. interpreted the data. M.J.S., I.H.B. and S.M.B. wrote the paper.

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