



HAL
open science

Experimental infection of highly pathogenic avian influenza virus H5N1 in black-headed gulls (*Chroicocephalus ridibundus*)

Antonio Ramis, Geert van Amerongen, Marco van de Bildt, Loneke Leijten,
Raphael Vanderstichel, Albert Osterhaus, Thijs Kuiken

► To cite this version:

Antonio Ramis, Geert van Amerongen, Marco van de Bildt, Loneke Leijten, Raphael Vanderstichel, et al.. Experimental infection of highly pathogenic avian influenza virus H5N1 in black-headed gulls (*Chroicocephalus ridibundus*). *Veterinary Research*, 2014, 45 (1), pp.84. 10.1186/s13567-014-0084-9 . hal-01290581

HAL Id: hal-01290581

<https://hal.science/hal-01290581>

Submitted on 18 Mar 2016

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

RESEARCH

Open Access

Experimental infection of highly pathogenic avian influenza virus H5N1 in black-headed gulls (*Chroicocephalus ridibundus*)

Antonio Ramis^{1*}, Geert van Amerongen², Marco van de Bildt², Loneke Leijten², Raphael Vanderstichel³, Albert Osterhaus² and Thijs Kuiken²

Abstract

Historically, highly pathogenic avian influenza viruses (HPAIV) rarely resulted in infection or clinical disease in wild birds. However, since 2002, disease and mortality from natural HPAIV H5N1 infection have been observed in wild birds including gulls. We performed an experimental HPAIV H5N1 infection of black-headed gulls (*Chroicocephalus ridibundus*) to determine their susceptibility to infection and disease from this virus, pattern of viral shedding, clinical signs, pathological changes and viral tissue distribution. We inoculated sixteen black-headed gulls with 1×10^4 median tissue culture infectious dose HPAIV H5N1 (A/turkey/Turkey/1/2005) intratracheally and intraesophageally. Birds were monitored daily until 12 days post inoculation (dpi). Oropharyngeal and cloacal swabs were collected daily to detect viral shedding. Necropsies from birds were performed at 2, 4, 5, 6, 7, and 12 dpi. Sampling from selected tissues was done for histopathology, immunohistochemical detection of viral antigen, PCR, and viral isolation. Our study shows that all inoculated birds were productively infected, developed systemic disease, and had a high morbidity and mortality rate. Virus was detected mainly in the respiratory tract on the first days after inoculation, and then concentrated more in pancreas and central nervous system from 4 dpi onwards. Birds shed infectious virus until 7 dpi from the pharynx and 6 dpi from the cloaca. We conclude that black-headed gulls are highly susceptible to disease with a high mortality rate and are thus more likely to act as sentinel species for the presence of the virus than as long-distance carriers of the virus to new geographical areas.

Introduction

Historically, highly pathogenic avian influenza viruses (HPAIV) were rarely found in wild birds, and, if they were, they typically did not cause clinical disease [1]. However, in late 2002, HPAIV H5N1 outbreaks were described in Hong Kong parks causing disease and death in wild local and migratory birds; concurrently with these outbreaks, HPAIV H5N1 were isolated from dead chickens and human beings [2]. It was shown through antigenic analysis that these new isolates of HPAIV H5N1 presented a reactivity pattern different from that of HPAIV H5N1 isolated in 1997 and 2001 and caused systemic infection in experimentally infected domestic ducks, with high virus titres and lesions in multiple organs, particularly in the brain [3].

Since 2002, disease and mortality from natural HPAIV H5N1 infection have been described for several gull species: black-headed gull (*Chroicocephalus ridibundus*) [2], great black-headed gull (*Larus ichthyaetus*) and brown-headed gull (*Larus brunnicephalus*) [4]. Also, experimental infections have been performed in laughing gulls (*Larus atricilla*) [5] and herring gulls (*Larus argentatus*) [6] showing that these species were highly susceptible (100%) to different HPAIV H5N1 isolates (A/Whooper Swan/Mongolia/244/05 and A/Duck Meat/Anyang/01), presenting a severe and short (4 to 10 dpi) clinical course, consisting mainly of neurologic signs with a high mortality rate (up to 100% in some cases). The main target organs for viral replication were central nervous system (CNS), pancreas, and adrenal gland, and the duration of the viral shedding after inoculation was in a range of 1 to 10 dpi depending on the isolate, the gull species, and inoculation route. In these experiments,

* Correspondence: antonio.ramis@uab.cat

¹CreSA and Departament de Sanitat i Anatomia Animals, Universitat Autònoma de Barcelona, Barcelona, Spain

Full list of author information is available at the end of the article

virus dynamics and in-depth pathogenesis were not investigated. Lastly, in an experimental infection in common gulls (*Larus canus*) with a HPAIV H5N1 isolate (A/Gull/Chany/P/06) isolated from healthy birds of this species, the virus was shown to replicate in respiratory and digestive tracts and to be excreted from pharynx and cloaca for 2 weeks after the infection, with no clinical signs in infected birds [7].

The black-headed gull is the most ubiquitous gull in Europe and Asia, is a peridomestic species that often occurs in urban and agricultural environments [8] and is known to be susceptible to HPAIV H5N1 infection [2]. Also, black-headed gulls are listed by the European Food Safety Authority [9] as having higher probability to be exposed to Asian lineage HPAIV H5N1 during migration outside the European Union, because of their gregarious behaviour and their tendency to use both wild areas and built-up areas. However, it is known neither how long black-headed gulls can shed HPAIV H5N1 nor how susceptible they are to disease from HPAIV H5N1 infection. These are important considerations to decide whether black-headed gulls are more likely to act as sentinels for the presence of the virus in the environment, or to act as long-distance vectors of the virus to other geographical areas [10]. Therefore, we performed an experimental infection of black-headed gulls using HPAIV A/turkey/Turkey/1/2005 (H5N1) to determine the susceptibility of this host species to infection and disease from this virus; the pattern of pharyngeal and cloacal viral shedding; the clinical signs and pathological changes; and lastly the virus distribution in different tissues. We chose this virus isolate to allow comparison of the results between black-headed gulls and 6 different wild species of Anseriformes [10], which jointly with Charadriiformes—to which the black-headed gull belongs—have traditionally been considered the natural reservoir for avian influenza viruses; and also because this isolate had spread quickly from Asia to Europe in 2005–2006 [11].

Material and methods

Virus preparation

A virus stock of HPAIV A/turkey/Turkey 1/2005 (H5N1) was prepared by propagation in Madin-Darby canine kidney (MDCK) cells and titrated according to standard methods [12]. The fluid had a titre of $1 \times 10^{8.1}$ median tissue culture infectious dose (TCID₅₀)/mL and was diluted with phosphate-buffered saline (PBS) to obtain a final titre of 10^4 TCID₅₀/mL. All the experiments with HPAIV H5N1 were performed under Biosafety level 3 + conditions.

Experimental design

We used 22 black-headed gulls (2–3 weeks old) obtained from a breeding colony site at Blauwestad, The Netherlands. The gulls were housed in negative isolator

units for 15 days to acclimatize them to the new environment. Two days before the experimental inoculation, they were tested for antibodies against influenza viruses by use of a commercially available nucleoprotein-based ELISA test (European Veterinary Laboratory, Woerden, The Netherlands) to assure they were not previously infected. Sixteen birds were inoculated with 1×10^4 TCID₅₀ HPAIV H5N1, 1.5 mL intratracheally and 1.5 mL intraoesophageally, which was the same virus, inoculum dose, and inoculation route as in a previous study [10]. We used this dose to increase the chance of inducing subclinical infection and to simulate field circumstances. In addition, 6 birds that served as negative controls were sham-inoculated in the same manner with PBS-diluted sterile allantoic fluid. Daily, a qualified veterinarian scored clinical signs of disease in all birds according to a standardized list. Also daily, the body weight of each bird was measured, and cloacal and pharyngeal swabs were collected in 1 mL transport medium. Birds were euthanatized by exsanguination under isoflurane anesthesia. After euthanasia, blood samples were taken and necropsies and tissue sampling were performed. The original plan was to euthanatize four HPAIV H5N1-inoculated birds on each of days 2, 4, 6 and 12 post inoculation, two sham-inoculated birds at 3 days post inoculation (dpi) and four sham-inoculated birds at 12 dpi. Because some H5N1-inoculated birds died spontaneously before the planned day of euthanasia, we were unable to completely follow the original plan, and the timing of necropsies and sampling for H5N1-inoculated birds was as follows: four birds at 2 dpi; five birds at 4 dpi; two birds at 5 dpi; two birds at 6 dpi; one bird at 7 dpi; and two birds at day 12 dpi (Table 1).

Pathologic and immunohistologic examination

Necropsies and tissue sampling were performed according to a standard protocol. After fixation in 10% neutral-buffered formalin and embedding in paraffin, 3- μ m-thick tissue sections were made and stained by two methods: hematoxylin and eosin staining for histologic evaluation and an immunohistologic method that used a monoclonal antibody against nucleoprotein for influenza A virus as a primary antibody for detection of influenza viral antigen [12]. The positive control was a lung of a HPAIV (H5N1) infected ferret; negative controls were substitution of the primary antibody by an irrelevant monoclonal antibody of the same isotype, and testing of tissues from sham-inoculated black-headed gulls. The following tissues were examined: brain (cerebrum, cerebellum and brain stem), trachea, bronchus, lung, air sac, oesophagus, proventriculus, duodenum, pancreas, jejunum, ileum, cecum, colon, liver, gallbladder, cloaca, cloacal bursa, spleen, thymus, kidney (cranial and caudal lobes), gonad, adrenal gland, feathered skin, myocardium and skeletal muscle.

Table 1 Follow up of infected gulls

Day post inoculation	Number of birds per day											
	1 dpi	2 dpi	3 dpi	4 dpi	5 dpi	6 dpi	7 dpi	8 dpi	9 dpi	10 dpi	11 dpi	12 dpi
Total alive	16	16	12	12	7	5	3	2	2	2	2	2
With neurologic signs	0	0	1	2	1	3	2	2	2	2	2	2
Died spontaneously	0	0	0	1	2	0	1	0	0	0	0	0
Euthanatized according to programme	0	4	0	4	0	2	0	0	0	0	0	2
Total necropsied	0	4	0	5	2	2	1	0	0	0	0	2

Number of birds showing neurological signs, spontaneous deaths, and programmed euthanasias in H5N1-inoculated black-headed gulls during the course of the experiment.

Virus quantitation by real time RT-PCR

Total nucleic acids were isolated on a MagnaPure LC system using the LC Total Nucleic Acid Isolation Kit (Roche Diagnostics, Almere, The Netherlands) according to the manufacturer's protocol. Influenza A virus was detected using a real-time PCR assay [13]. Amplification and detection were performed on an ABI7500 (Applied Biosystems, Foster City, CA, USA) with the Taqman Fast Virus 1-Step Master Mix.

Virus titration

Tissue samples were weighed and homogenized in 1 mL of transport medium with a FastPrep-24 homogenizer (MP Biomedicals, Eindhoven, The Netherlands). Virus titres were determined by serial 10-fold dilution of the homogenized tissue samples and swabs on MDCK cells as described previously [14].

Statistics

The area under the curve for viral loads from pharyngeal and cloacal swabs within gulls, as measured by RT-PCR (Ct value) and virus isolation (10^x TCID₅₀), were compared using the Wilcoxon matched-pairs signed-ranks

test [15]. A multilevel mixed-effects linear regression, with gull weight (g) as the dependent variable, was fit using maximum likelihood estimation, and included a random effect for gull and first order auto-regressive (AR1) structured residual errors between repeated weight measurements. A fractional polynomial [16], using two terms, was used to estimate weight over time. Normality and homoscedasticity of residuals, at both levels of the model, were evaluated [17], and statistical significance was set at $P < 0.05$.

Results

Virus detection in pharyngeal and cloacal swabs

Virus was detected in the pharyngeal swabs, cloacal swabs, or both, of all 16 H5N1-inoculated birds, indicating that they were all productively infected (Table 2). Based on the area under the curve, there were larger quantities of virus detected in pharyngeal swabs than in cloacal swabs using RT-PCR and virus isolation ($P < 0.001$) (Figure 1A, B). By RT-PCR, nearly all pharyngeal swabs tested positive from 1 dpi to the end of the experiment at 12 dpi. The mean Ct-value was highest at 1 dpi (Ct value 22) and declined slowly on subsequent

Table 2 Virus detection in pharyngeal and cloacal swabs of H5N1-inoculated gulls during the course of the experiment

Day	No. tested	Pharyngeal swab				Cloacal swab			
		RT-PCR		Virus isolation		RT-PCR		Virus isolation	
		No. positive	Ct range	No. positive*	Titre range	No. positive	Ct range	No. positive*	Titre range
1	16	16	19-27	16	1.5-5.5	6	27-32	1	< 0.8-0.8
2	16	16	21-29	15	1.2-3.5	16	22-30	4	1.5-2.8
3	12	11	21-29	10	0.8-4.2	6	28-31	1	< 0.8-0.8
4	12	12	22-30	9	1.8-3.8	7	26-33	2	0.8-2.2
5	7	7	23-32	6	1.8-3.8	6	24-31	3	1.2-2.2
6	5	5	20-34	4	0.8-3.5	5	20-30	2	2.2-2.5
7	3	3	23-28	3	1.5-3.2	3	27-32	0	n.a.
9	2	2	31	0	n.a.	2	28-31	0	n.a.
11	2	2	30-33	0	n.a.	2	31-34	0	n.a.
12	2	2	30-31	0	n.a.	2	31-32	0	n.a.

Virus isolation was attempted on all PCR-positive samples.
n.a., not applicable.

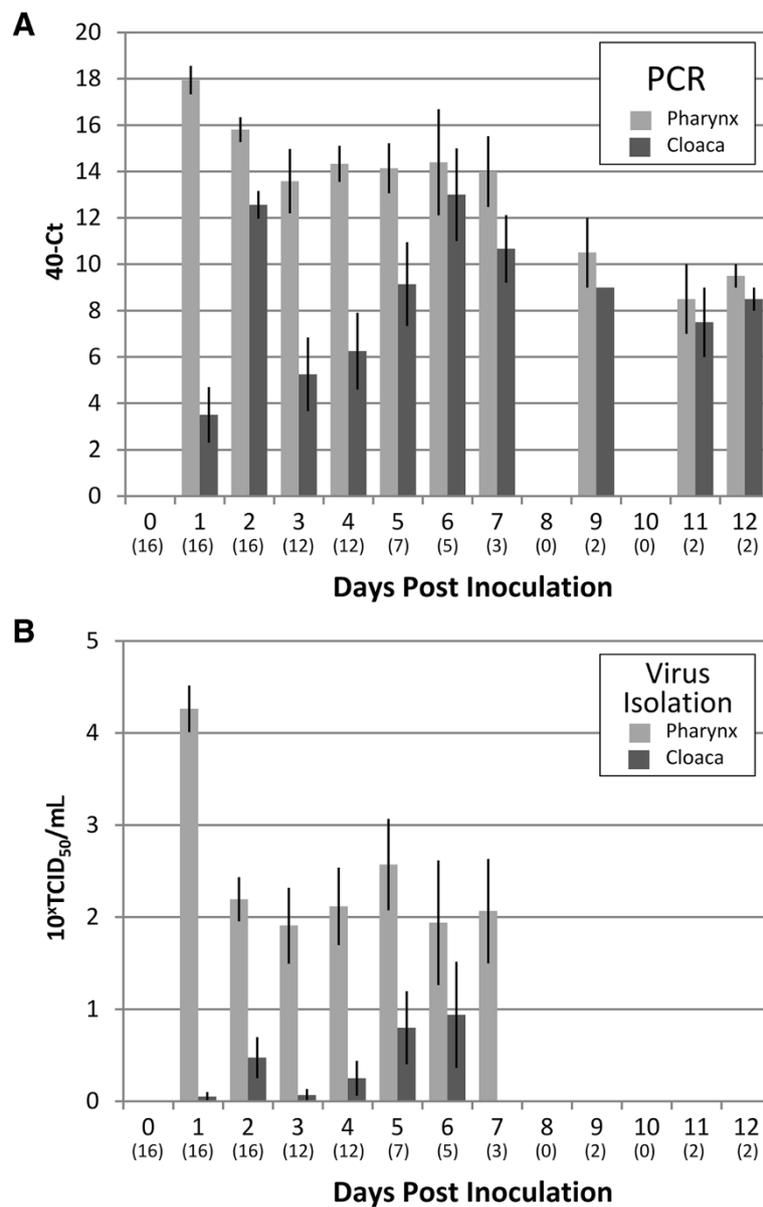


Figure 1 Viral detection in pharyngeal and cloacal swabs using RT-PCR and virus isolation. Bar graph with standard error estimates of results of viral detection in pharyngeal and cloacal swabs taken from gulls inoculated on day 0 with HPAIV H5N1 virus and followed for 12 days. **(A)** Real-time RT-PCR, expressed as 40 (minimum cycling threshold [Ct] value considered negative) – actual Ct value. **(B)** Virus isolation, expressed as 10^x × median tissue culture infectious dose per mL (TCID₅₀/mL), with a titre < 10^{0.8} TCID₅₀/mL considered negative. Numbers below each day (in parentheses) represent total number of birds tested on that day. Each bar represents the average value for all birds tested on that day.

days. Virus was isolated from RT-PCR-positive pharyngeal swabs from most birds from 1 to 7 dpi, but could no longer be isolated on subsequent days. The mean virus titre in pharyngeal swabs was highest (10^{4.26} TCID₅₀) at 1 dpi and between about 10² and 10³ TCID₅₀ from 2 to 7 dpi. By RT-PCR, most cloacal swabs tested positive from 1 dpi to the end of the experiment at 12 dpi, but the Ct values were generally higher (i.e. less virus) than in pharyngeal swabs. The mean Ct-value slowly decreased until 6 dpi (Ct value 27) and slowly increased on subsequent days. Virus

was isolated only from a small proportion of RT-PCR-positive cloacal swabs from 1 to 6 dpi, with mean virus titres between detection limit and just above 10² TCID₅₀, and could not be isolated on subsequent days. No virus was detected by RT-PCR in any pharyngeal or cloacal swabs of sham-inoculated birds.

Clinical signs

During the first 3 dpi, no clinical signs were observed in H5N1-inoculated birds, except one bird (#49) that showed

less activity than normal and slight nonspecific neurological signs at 3 dpi: regularly fluffing of feathers and walking slightly to one side. At 4, 5 and 6 dpi, all H5N1-inoculated birds were less active than sham-inoculated birds, showing increased recumbency and regularly fluffing of feathers. Four of them (#46, 47, 49, 51) showed obvious neurological signs, consisting mainly of torticollis, circling, loss of balance and head tremors. Respiratory or digestive signs, such as diarrhoea, were not observed in any H5N1-inoculated birds. From 7 dpi onwards, the two remaining H5N1-inoculated birds (#47, 51) partially recovered, alternating between normality and neurological signs (Table 1). At 4 dpi, the H5N1-inoculated bird that showed clinical signs at 3 dpi (#49) died spontaneously. Two birds (#46, 50) at 5 dpi and one bird (#48) at 7 dpi, all three H5N1-inoculated, also died spontaneously. No clinical signs or spontaneous mortality were observed in any of the sham-inoculated birds. Based on the statistical model: the body weight of H5N1-inoculated birds was 26 g lower than sham-inoculated birds ($P = 0.029$) after inoculations; within the H5N1-inoculated group, the body weight of birds that died was 26 g lower after inoculations than that of birds that survived ($P = 0.047$). Regardless of their inoculation status, gulls were losing weight during the first week post inoculation, and this trend was reversed after 10 dpi ($P < 0.001$, Figure 2).

Gross lesions

No relevant gross lesions were observed in any H5N1-inoculated or sham-inoculated birds. In general, all birds

showed good body condition and had abundant subcutaneous and perivisceral fat deposits.

Histopathology and immunohistochemistry

The evaluation of histological lesions and the presence of virus antigen in the different cells and tissues of evaluated organs of H5N1-inoculated birds are summarized in Table 3. On day 2, air sacs in all birds (#36, 37, 38, 39) had marked diffuse infiltration of heterophils and multiple variably-sized well-demarcated aggregates of heterophils and macrophages. The adrenal gland in one bird (#37) had multiple foci of necrotic adrenocortical and chromaffin cells. Virus antigen expression was observed in epithelial cells (#36, 37, 38, 39) and unspecified inflammatory cells (#36, 37, 38, 39) of the air sacs, and in adrenocortical cells (#37) of the adrenal gland, localized to lesions in these tissues. Virus antigen expression also was observed in individual tracheal and bronchial ciliated epithelial cells (#38) (Figure 3A, B), air capillary cells (#36, 37), and thymic epithelial cells in absence of histological lesions.

On day 4, all birds (#40, 41, 42, 43, 49) had airsacculitis and air sac granulomas similar to those on day 2. The trachea in one bird (#42) had many heterophils in the lumen. The pancreas in two birds (#40, 49) had multiple foci of necrotic acinar cells with scarce associated inflammatory cells (Figure 3C). The cerebrum in all birds (#40, 41, 42, 43, 49) had multiple randomly distributed foci of necrotic neurons, characterized by chromatolysis and satellitosis (Figure 3E). These foci of necrosis were associated with predominantly perivascular infiltration

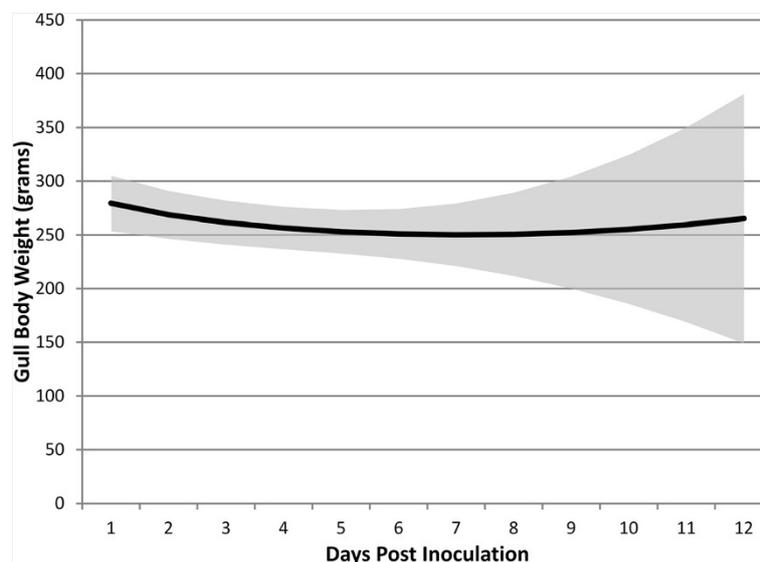


Figure 2 Estimated gull body weights (g) throughout the study period. It was determined by the fractional polynomial terms in the mixed-effects linear regression, for control gulls that remained alive for the duration of the study; shaded area represents the 95% confidence intervals. The same statistical model predicted an average constant loss of 26 grams for H5N1-inoculated gulls ($P = 0.029$), and an average constant loss of 26 grams for gulls that died during the study ($P = 0.047$)—these estimates are not shown in graph.

Table 3 Histological lesions (HE) and presence of antigen (IHC) in tissues of H5N1-inoculated gulls during the course of the experiment

Tissue	Number of birds positive/number of birds examined per day											
	2 dpi		4 dpi		5 dpi		6 dpi		7 dpi		12 dpi	
	HE ¹	IHC ²	HE	IHC	HE	IHC	HE	IHC	HE	IHC	HE	IHC
Thymus	0/4	2/4	0/5	0/5	0/2	0/2	0/2	0/2	0/1	0/1	0/2	0/2
Adrenal	1/4	1/4	0/5	0/5	0/2	0/2	1/2	1/2	0/1	0/1	0/2	0/2
Lung/Trachea	0/4	3/4	1/5	0/5	1/2	0/2	1/2	0/2	0/1	0/1	0/2	0/2
Air sacs	4/4	4/4	5/5	5/5	2/2	1/2	1/2	1/2	0/1	0/1	1/2	1/2
Pancreas	0/4	0/4	2/5	2/5	1/2	1/2	2/2	2/2	1/1	1/1	1/2	0/2
CNS	0/4	0/4	5/5	5/5	2/2	2/2	2/2	2/2	1/1	1/1	1/2	0/2

¹HE: presence of histological lesions by examination of hematoxylin-and-eosin-stained tissue sections.

²IHC: presence of viral antigen by examination of tissue sections stained by immunohistochemical technique.

of inflammatory cells, consisting of variable proportions of lymphocytes, plasma cells, macrophages, and heterophils. The cerebellum of one bird (#49) and brainstem of another one (#43) had similar lesions as the cerebrum. Virus antigen expression was observed in epithelial cells

(#42, 49) and unspecified inflammatory cells (#40, 41, 42, 43, 49) of the air sacs, in acinar cells (#40, 49) of the pancreas (Figure 3D), and in neurons, glial cells, and ependymal cells of the brain, localized to lesions in these tissues (Figure 3F). There also was virus antigen

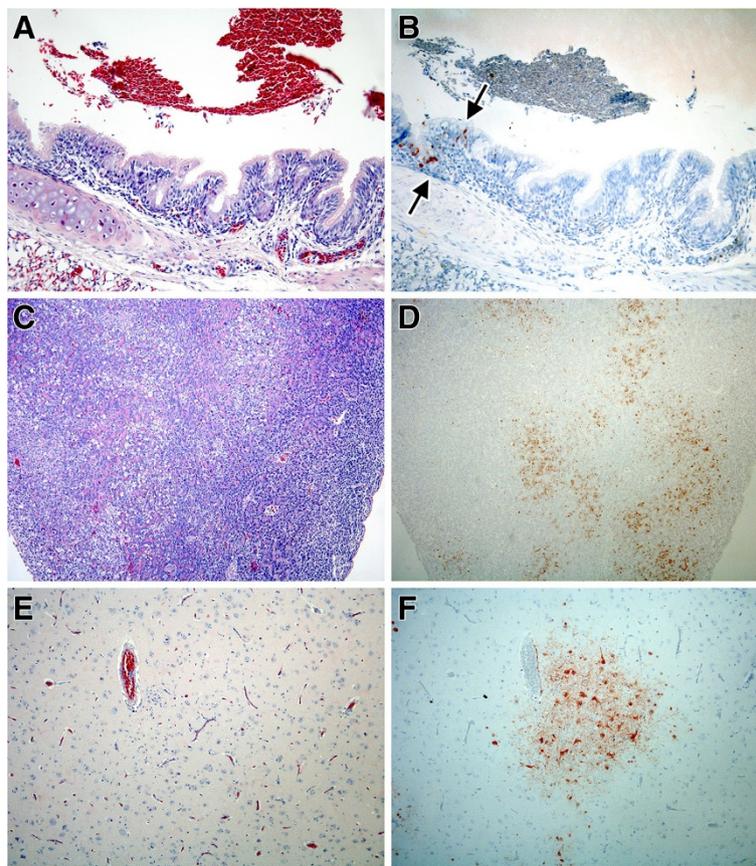


Figure 3 Histopathology and antigen expression in selected tissues from infected gulls. By immunohistochemistry, influenza virus antigen expression is visible as a red staining. (A, B): Bronchus of 2 dpi infected gull without histologic lesions (A), and scarce bronchial epithelial cells showing positivity on the immuno-stained consecutive section (arrows) (B). (C, D): Pancreas of 4 dpi infected gull showing multiple foci of lytic necrosis (C), and intense positivity associate to necrotic foci in the immune-stained consecutive section. (E, F): Cerebral cortex showing necrotic neurons, and satellitosis (E), and intense positivity in neurons and associated glial cells in the immune-stained consecutive section. HE stain (A, C, E) and immunoperoxidase counterstained with hematoxylin (B, D, F). All the pictures are at 10x power fields.

expression in ganglion cells of the peripheral nervous system in adrenal gland (#43) in absence of histological lesions.

On day 5, airsacculitis and air sac granulomas were seen in both birds (#46, 50) and mild heterophilic tracheitis in one of them (#50). The myocardium of both birds showed scattered foci of mixed inflammatory infiltrate. The pancreas of one bird (#50) showed multifocal pancreatitis. The CNS of both birds, similarly to day 4 dpi, presented multifocal mixed encephalitis; however in one of them (#46) no lesions were observed in cerebellum. Viral antigen expression was observed in few inflammatory cells in air sacs and pancreatic acinar cells in bird #50, in nuclei of heart myocytes of bird #46, and in neurons, glial cells and ependymocytes in both birds (#46, 50), localized to lesions in these tissues. There also was virus antigen expression in ganglion cells of the peripheral nervous system in adrenal gland (#46) in absence of histological lesions.

On day 6, the air sacs presented scattered granulomas in one bird (#44), and mild tracheitis and multifocal adrenocortical necrosis could be seen in the other one (#45). The pancreas of both birds (#44, 45) showed multifocal pancreatitis. The CNS of both birds presented multifocal mixed encephalitis, similarly to the previous days; however, in one of them (#44), no lesions were observed in cerebellum. Viral antigen expression was observed in few inflammatory cells associated to air sac, adrenocortical cells, pancreas acinar cells, neurons, glial cells and ependymocytes, localized to lesions in these tissues.

On day 7 the only bird evaluated (#48) showed mild tracheal epithelial hyperplasia, and some granulomas in lungs and air sacs associated with *Aspergillus*-like hyphae. Kidney presented focal interstitial nephritis, tubular casts and gout and liver showed isolated hepatocyte death, activation of Kupffer cells. In the large and small intestine we observed a mild, diffuse and generalized lymphocytic enteritis; and the presence of a heterophilic granuloma in the cloaca. The pancreas showed multifocal pancreatitis. In the CNS, cerebrum and brainstem presented multifocal mixed encephalitis, while cerebellum showed no relevant lesions except choroiditis in the fourth ventricle. Viral antigen expression was observed in pancreatic acinar cells, neurons, glial cells and ependymocytes, localized to lesions in these tissues. There also was virus antigen expression sporadically in Kupffer cells in absence of histological lesions.

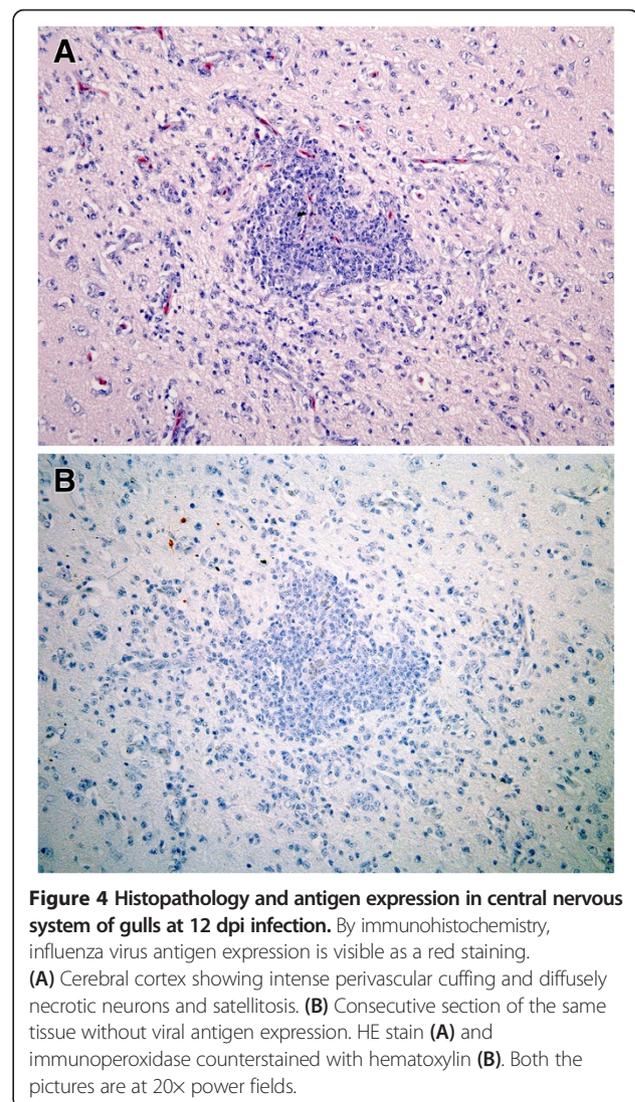
On day 12, one infected bird (#51) showed airsacculitis and multiple granulomas consisting of a large central necrotic area containing bacterial colonies and a wide ring of inflammatory cells (histiocytes and heterophils). The other one (#47) showed mild scattered multifocal lymphocytic infiltrate in myocardium and pancreas; and

severe inflammation in the CNS. Cerebrum, cerebellum and brainstem presented a severe multifocal mixed encephalitis (Figure 4A) and a severe lymphocytic choroiditis in lateral ventricles. Weak vestigial viral antigen expression was observed only in bird #51, located in epithelial cells of air sacs and in inflammatory cells of one of the air sac granulomas. Bird #47 showed no positivity at any organ or tissue (Figure 4B).

Tissues of sham-inoculated birds were examined according to the same methods as those of H5N1-inoculated birds. Histological lesions and viral antigen expression were not observed in sham-inoculated birds on days 3 and 12 dpi.

Virus titration in selected tissues

Virus was isolated from tissues of all H5N1-inoculated birds euthanized or found dead between 2 and 7 dpi, but not at 12 dpi (Additional file 1). Virus was isolated from



multiple organ and systems: respiratory, digestive, urinary, and lymphoid systems of all birds; brain in all but four birds; skeletal muscle in four birds; and skin in none of the birds. At 2 dpi, the highest viral titres ($> 10^7$ TCID₅₀ per g tissue) in each bird were observed in respiratory tract: bronchus or air sac. At 4 dpi, the highest viral titre in two of four birds was detected in the brain ($\geq 10^7$ TCID₅₀ per g tissue) whereas viral titres in respiratory tract tissues (trachea, bronchus lung, air sac) were lower than at 2 dpi (range: $10^{2.7}$ - $10^{6.4}$ TCID₅₀ per g tissue). From 5 to 7 dpi the highest viral titres were detected in brain and pancreas, and all the birds on those days showed a titre $\geq 10^7$ TCID₅₀ per g tissue in one or both of these organs. Virus was not isolated in the tissues coming from sham-inoculated birds.

Discussion

Our study shows that black-headed gulls are highly susceptible to disease from HPAIV A/turkey/Turkey 1/2005 (H5N1). All inoculated birds were productively infected and shed infectious virus until 7 dpi from the pharynx and 6 dpi from the cloaca, at maximal titres of more than 10^4 TCID₅₀ per mL. They developed systemic disease with a high morbidity and mortality rate and conspicuous nervous signs. The mortality was high and surviving animals showed mild nervous signs until the last day of the experiment. Infected birds presented pharyngeal and cloacal excretion detected by RT-PCR until the last day of the experiment, and viral isolation could be done until 6 dpi and 7 dpi, respectively, from pharyngeal and cloacal swabs (Table 2; Figure 1A, B).

The results from these black-headed gulls can be compared directly with an earlier study on six wild duck species [10], because the virus, inoculation dose, and route of inoculation used were identical. The black-headed gulls had a slightly higher pattern of virus excretion to that of mallards (*Anas platyrhynchos*), Eurasian pochards (*Aythya ferina*), and tufted ducks (*A. fuligula*), which had a maximum titre of about 10^3 TCID₅₀ per mL and excreted up to 6 dpi. The black-headed gulls also were more susceptible to disease from HPAIV H5N1 than any of the six duck species studied. For the most susceptible duck species, the tufted duck, clinical signs were severe in 3 of 7 (43%) and mild in 4 of 7 (57%). In contrast, clinical signs were severe in 14 of 16 (87.5%) of H5N1-inoculated black-headed gulls and mild in 2 of 16 (12.5%). Together, these results indicate that black-headed gulls are more likely to act as sentinels of the presence of HPAIV H5N1 in the environment than as long-distance carriers of the virus to new geographical areas. Older black-headed gulls may be more resistant to disease from HPAIV H5N1 infection, since an age-related association with dissemination and clinical outcome in Peking ducks has been observed following infection with H5N1 HPAIV [18].

In our black-headed gulls, the clinical picture—showing exclusively neurological signs—was similar to that described for laughing gulls [6] and herring gulls [5]. However, none of our black-headed gulls showed total recovery, while some herring gulls recovered completely after experimental infection using two different HPAIV H5N1 isolates (A/Whooper Swan/Mongolia/244/05 and

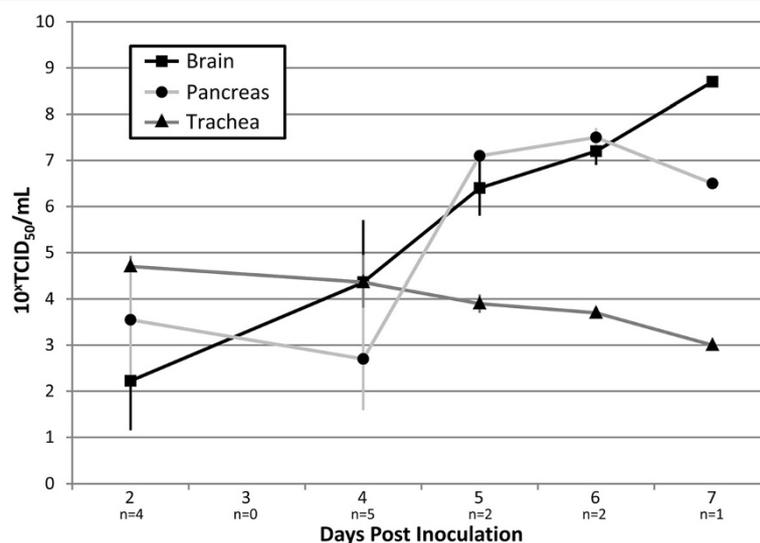


Figure 5 Comparison of viral loads in selected tissues during the course of the experiment. The viral loads are expressed as $10^x \times$ median tissue culture infectious dose per mL (TCID₅₀/mL), among trachea, brain, and pancreas in H5N1-inoculated black-headed gulls from day 2 to 7 post inoculation. The sample sizes (n) indicate the number of black-headed gulls that contributed to each mean value, with their corresponding standard errors (vertical bars).

A/Duck Meat/Anyang/01), and one laughing gull, infected with HPAIV H5N1 (A/Whooper Swan/Mongolia/244/05 H5N1) recovered without clinical signs. In these species, clinically healthy birds showed pharyngeal viral excretion detected by viral isolation, but no longer than 10 dpi for laughing gulls [6] and 5 dpi for herring gulls [5]. Although comparison between our study and these studies is difficult because of differences in virus strains, inoculation dose, and route of inoculation, it suggests that black-headed gulls are more susceptible to disease from HPAIV H5N1 than either herring gulls or laughing gulls and show a similar level of virus excretion as laughing gulls. In the case of common gulls, it is not clear whether they are resistant to the infection with (A/Gull/Chany/P/06) H5N1 or this isolate is not pathogenic for gulls [7].

The absence of gross lesions in our black-headed gulls, despite abundant virus replication and associated histological lesions in multiple organs, is remarkable, but corresponds to results of other experimental HPAIV H5N1 infections in wild birds, where only a small proportion of infected individuals shows gross lesions [10]. Specifically, we did not observe multifocal pancreatic necrosis, the most common gross lesion in most wild birds, or multi-organ haemorrhage, as is seen specifically in swans [19]. We did observe a clear decrease of the weight in the H5N1-inoculated birds in comparison with sham-inoculated ones, and also between the birds that died and birds that survived within the inoculated group. These findings correspond with severe anorexia, leading to rapid loss in body weight.

The location and severity of microscopic lesions generally corresponded to the distribution and intensity of the immunostaining and to the viral load in each evaluated organ or tissue. At 2 dpi, viral antigen expression and viral titres were higher in organs of the respiratory tract than in other organ systems, although viral antigen was expressed in some other organs (thymus, adrenal gland). This suggests that the viremic period is reached very soon after virus inoculation. At 4 dpi, the respiratory tract presented a similar degree of microscopic lesions, viral antigen expression and viral titres as at 2 dpi. Moreover, necrotizing pancreatitis and neuronal necrosis in the brain were present, as described in other gull species (6, 5). Viral titres in pancreas and brain were similar to those respiratory tract tissues. From 4 dpi onwards, the severity of microscopic lesions, viral antigen expression and viral titres were increased in brain and pancreas and decreased in respiratory tract tissues. Interestingly, while the two birds euthanatized on day 12 dpi both showed neurological signs, and one of them showed microscopic brain lesions, no virus could be detected in the brain or any other tissue by virus isolation or immunohistochemistry in either bird (Table 3 and Additional file 1). As a general trend in this HPAIV H5N1 infection

of black-headed gulls, we can assert that the virus is mainly in the respiratory tract on the first days after inoculation and then concentrates more in pancreas and CNS from 4 dpi onwards (Figure 5).

Additional file

Additional file 1: Distribution of infectious virus in tissues of H5N1-inoculated gulls during the course of the experiment.

Competing interests

TK is part-time advisor and AO is part-time chief scientific officer of Viroclinics Biosciences B.V. The other authors declare that they have no competing interests.

Authors' contributions

Conceived and designed the study: AR, AO, TK. Performed the study: AR, GvA, MvdB, LL, TK. Analysed the data: AR, MvdB, RV, TK. Wrote the manuscript: AR, MvdB, RV, TK. All authors read and approved the final manuscript.

Acknowledgements

This work was funded by the Dutch Ministry of Economic Affairs (Impulse Veterinary Avian Influenza Research in the Netherlands program). AR sabbatical stay at the Department of Virology in the Erasmus Medical Center was supported by a grant of the Spanish Government (PR-2009-0145). We thank P. van Run for technical assistance and F. van der Panne for figure preparation.

Author details

¹CRESA and Departament de Sanitat i Anatomia Animals, Universitat Autònoma de Barcelona, Barcelona, Spain. ²Department of Virosciences, Erasmus Medical Center, Rotterdam, The Netherlands. ³Department of Health Management, Atlantic Veterinary College, University of Prince Edward Island, Charlottetown, Canada.

Received: 4 March 2014 Accepted: 30 July 2014

Published online: 19 August 2014

References

1. Webster RG, Bean WJ, Gorman OT, Chambers TM, Kawaoka Y: **Evolution and ecology of influenza A viruses.** *Microbiol Revs* 1992, **56**:152–179.
2. Ellis TM, Bousfield RB, Bissett LA, Dyrting KC, Luk GSM, Tsim ST, Sturm-Ramirez K, Webster RG, Guan Y, Peiris JSM: **Investigation of outbreaks of highly pathogenic H5N1 avian influenza in waterfowl and wild birds in Hong Kong in late 2002.** *Avian Pathol* 2004, **33**:492–505.
3. Sturm-Ramirez KM, Ellis T, Bousfield B, Bissett L, Dyrting K, Rehag JE, Poon L, Guan Y, Peiris M, Webster RG: **Reemerging H5N1 influenza viruses in Hong Kong in 2002 are highly pathogenic to ducks.** *J Virol* 2004, **78**:4892–4901.
4. Liu J, Xiao H, Lei F, Zhu Q, Qin K, Zhang XW, Zhang XL, Zhao D, Wang G, Feng Y, Ma J, Liu W, Wang J, Gao GF: **Highly pathogenic H5N1 influenza virus infection in migratory birds.** *Science* 2005, **309**:1206.
5. Brown JD, Stallknecht DE, Swayne DE: **Experimental infections of herring gulls (*Larus argentatus*) with H5N1 highly pathogenic avian influenza viruses by intranasal inoculation of virus and ingestion of virus-infected chicken meat.** *Avian Pathol* 2008, **37**:393–397.
6. Brown JD, Stallknecht DE, Beck JR, Suarez DL, Swayne DE: **Susceptibility of North American ducks and gulls to H5N1 highly pathogenic avian influenza viruses.** *Emerg Infect Dis* 2006, **12**:1663–1670.
7. Zaykovskaya AV, Sharshov KA, Sherstkov EA, Yurlov AK, Shestopalov AM: **Experimental infection caused by influenza A (H5N1) virus in common gull (*Larus canus*).** *Vopr Virusol* 2012, **57**:43–45 (in Russian).
8. Piersma T, Wiersma P: **Familia Charadriidae.** In *Handbook of the Birds of the World*, Volume 3. Edited by del Hoyo J, Elliott A, Sargatal J. Barcelona, Spain: Lynx Edicions; 1996:615–616.
9. European Food Safety Authority (EFSA): **Scientific Statement on Migratory Birds and their Role in the Spread of Highly Pathogenic Avian Influenza.** 2006.

10. Keawcharoen J, van Riel D, van Amerongen G, Bestebroer T, Beyer WE, van Lavieren R, Osterhaus ADME, Fouchier RA, Kuiken T: **Wild ducks as long-distance vectors of highly pathogenic avian influenza virus (H5N1).** *Emerg Infect Dis* 2008, **14**:600–607.
11. Löndt BZ, Nunez A, Banks J, Nili H, Johnson LK, Alexander DJ: **Pathogenesis of highly pathogenic avian influenza A/turkey/Turkey/1/2005 H5N1 in Pekin ducks (*Anas platyrhynchos*) infected experimentally.** *Avian Pathol* 2008, **37**:619–627.
12. Rimmelzwaan GF, Kuiken T, van Amerongen G, Bestebroer TM, Fouchier RA, Osterhaus AD: **Pathogenesis of influenza A (H5N1) virus infection in a primate model.** *J Virol* 2001, **75**:6687–6691.
13. Ward C: **Design and performance testing of quantitative real time PCR assays for influenza A and B viral load measurement.** *J Clin Virol* 2004, **29**:179–188.
14. Rimmelzwaan GF, Baars M, Claas EC, Osterhaus AD: **Comparison of RNA hybridization, hemagglutination assay, titration of infectious virus and immunofluorescence as methods for monitoring influenza virus replication in vitro.** *J Virol Methods* 1998, **74**:57–66.
15. Wilcoxon F: **Individual comparisons by ranking methods.** *Biometrics* 1945, **1**:80–83.
16. Royston P, Sauerbrei W: *Multivariable Model-building, a Pragmatic Approach to Regression Analysis Based on Fractional Polynomials for Modelling Continuous Variables.* New Jersey, USA: John Wiley & Sons Ltd; 2008.
17. Dohoo I, Martin W, Stryhn H: *Veterinary Epidemiologic Research.* Canada: AVC Incorporated Charlottetown; 2009.
18. Löndt BZ, Núñez A, Banks J, Alexander DJ, Russell C, Richard- Löndt AC, Brown IH: **The effect of age on the pathogenesis of a highly pathogenic avian influenza (HPAI) H5N1 virus in Pekin ducks (*Anas platyrhynchos*) infected experimentally.** *Influenza Other Respir Viruses* 2010, **4**:17–25.
19. Reperant LA, Osterhaus ADME, Kuiken T: **Influenza virus infections.** In *Infectious Diseases of Wild Mammals and Birds in Europe.* Edited by Gavier-Widen D, Duff JP, Meredith A. Chichester, U.K: Wiley-Blackwell; 2012:37–58.

doi:10.1186/s13567-014-0084-9

Cite this article as: Ramis *et al.*: Experimental infection of highly pathogenic avian influenza virus H5N1 in black-headed gulls (*Chroicocephalus ridibundus*). *Veterinary Research* 2014 **45**:84.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

Submit your manuscript at
www.biomedcentral.com/submit

 **BioMed Central**