Types and Predilection of Human Avian Influenza Virus among Patients at Abubakar Tafawa Balewa Teaching Hospital Bauchi, Nigeria

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Abstract

Background: Nigeria suffered waves of Highly Pathogenic Avian Influenza (HPAI) outbreaks that peaked twice in February 2006 and February 2007. The outbreaks affected 3,057 commercial and rural household farms causing 1.3 million of the country’s poultry destroyed.

Aim: The burden of Influenza is likely to be underestimated and because of the unstable nature of the virus it allows for the mapping of this virus to specific region, this study therefore seeks to identify the types, prevalence, strain, predilection of the organism so as to monitor the occurrence of influenza in this part of the country with the aim of providing a base line level of intervention of influenza virus and find the candidate virus for vaccine selection and production.

Methods: This study is a cross sectional survey, our target populations were all pediatric and adult patients admitted at the pediatrics and internal medicine department of ATBUTH who has met the criteria for presence of cough, fever (> 37°C), nasal congestion and dyspnoea. Samples were collected and analyzed using the polymerase chain reaction (PCR), for the presence and sub typing of influenza viruses in respiratory specimens.

Results: Overall 5% of the samples collected tested positive for human influenza type A and B. During average epidemics, overall attack rates are estimated to be 10 - 20%, but in certain susceptible populations such as schoolchildren or nursing home residents, attack rates of 40 - 50% may occur. 60% of the positive results were among the female samples whereas 40% from the male samples.

Keywords: Human Influenza Virus; Prevalence; Types; Subtypes and Predilections

Introduction

Nigeria suffered waves of Highly Pathogenic Avian Influenza (HPAI) outbreaks that peaked twice in February 2006 and February 2007. The outbreaks affected 3,057 commercial and rural household farms causing 1.3 million of the country’s 160 million poultry to be destroyed at the cost of $5.4M paid in compensation by the Government of Nigeria (Federal Department of Livestock [FDL], 2008) [1,2].

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The unstable nature of the virus strain calls for mapping of HPAI to specific state with the aim of utilizing specific regional or state level approach in prevention and management of HPAI.

Influenza virus particularly type A infect large number of warm blooded animals, including wild birds, domestic birds, pigs, horses and humans influenza virus can switch host to form new lineage in novel hosts, the most significant of this is the emergence of antigenically novel influenza A in humans leading to pandemics [3]. Pandemics occur when a new avian influenza strain acquires the ability to infect people and to spread easily person to person. This can occur in two ways: Reassortment due to exchange of seasonal and avian influenza genes in a person or pig infected with both strains and Mutation, when an avian strain becomes more transmissible through adaptive mutation of the virus during human avian influenza infection [4].

Influenza viruses are sub typed according to surface glycoprotein: hemagglutinin (HA) and neuraminidase (NA) Currently, there are 16 hemagglutinin (H1 to H16) and 9 neuraminidases (N1 to N9), about 144 possible sub-types have been identified [5,6].

Hemagglutinin attaches the virus to the surface of the host cell which enables the virus to replicate while Neuraminidase lets the newly replicated viruses out of the cell to infect more cells. The Influenza viruses are divided into three main types: influenza A, B, and C, the A viruses infect birds and other animals, as well as humans, it is the source of seasonal influenza epidemics and all pandemics while B and C viruses infect humans only and are not associated with pandemics [5]. Human influenza originates as avian (birds) influenza, typically from migratory water bird to domestic birds, swine and then human infection occurs as a result of consumption or contact with these domestic animals [6]. Human Influenza under goes mutation through "Antigenic Drift and Shift "which enhances its pathogenicity [6]. Antigenic Drift - Point mutations in HA or NA generally results in relatively small changes in virus, these are small changes in the genes of influenza viruses that happened continually over time as the virus replicates while antigenic Shift is as a result of reassortment of gene segments leading to novel and potentially pandemic strains. Influenza viruses mutate by antigenic drift most times, antigenic shift happens only occasionally [6,7]. Influenza viruses spreads easily from person to person through coughing and sneezing and transmitted by inhaling respiratory aerosols containing the virus, produced when infected person talks, coughs, or sneezes and by touching an infected person or an item contaminated with the virus and then touching the eyes, nose, or mouth [8]. The incubation period is usually 3 to 7 days depends on the strain of virus, dose of inoculums, age and immune status of bird, management and environmental factors plays a significant role in the infectivity of the virus [8]. Major clinical presentation include Sudden onset of fever, headache, muscle aches, severe weakness, respiratory symptoms, e.g., cough, sore throat, difficulty breathing [7,8].

Pandemics are less predictable, not always in winter, great variations occurs in mortality, severity of illness, and pattern of illness or age most severely affected, rapid surge in number of cases over brief period of time can occur, often measured in weeks and it tend to occur in waves of 6 - 8 weeks, subsequent waves may be more or less severe, more than one wave of influenza is likely, gaps between the waves may be weeks or months, a subsequent wave can be worse than the first [9].

Objective

The burden of Influenza is likely to be under estimated, this study therefore seeks to identify the types, prevalence, strain and predilection of the organism so as to monitor the occurrence of influenza in this part of the country with the aim of providing a base line level of intervention of influenza virus admitted in this hospital, this will also help give an insight into the candidate virus for vaccine selection and production, similarly it will provide a foundation for detecting outbreaks and pandemics or emergence of a novel strain of influenza so as to create an early warning system to trigger a rapid public health response and advocate for a targeted approach in terms of vaccine procurement and distribution.

Methods

This study is a cross sectional survey, our target populations were all pediatric and adult patients admitted at the pediatrics and internal medicine department of ATBUTH who has met the criteria for presence of cough, fever (> 37°C), nasal congestion and dyspnoea.

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Samples were collected between 8.00am and 8.00pm, 50 samples were collected for each of the 6 age bracket 0-12 months, 13 months - 5yr, 6 - 25yr, 26 - 45yr, 46 - 65yr and > 65yr. Samples were analyzed using the polymerase chain reaction (PCR) at a national influenza reference laboratory (NIRL) Abuja Nigeria. Samples were analyzed for the presence and sub typing of influenza viruses in respiratory specimens which did not require invitro isolation. Here, primers and probes were used for typing and sub typing. Because the influenza genome consists of single-stranded ribonucleic acids (RNA), complimentary deoxyribonucleic acids (cDNA) were synthesized using the reverse-transcriptase (rt) enzyme prior to the PCR reaction. Therefore, any copy of the viral RNA specimens that tested positive for a novel or new subtype of influenza virus were sent to the CDC Atlanta for confirmation specimens were sent with the original sample in case of possible contamination. The results of this technique were obtained within 24 - 72 hrs. Appropriate controls that identify poor-quality of samples e.g. extraction were utilized to checkmate or avoid false positive results, the most common cause of false-positive results is contamination, this was mitigated by the use of real-time rt-PCR operating in a contained system. Specimens were kept at -70°C and were transporting to the National reference lab in cold chain. Human Influenza positivity was considered as the positive outcome variable whereas positive contact with animal or human cases were considered as the exposure variables for this study, possible confounders were presence of other respiratory tract infections e.g Pneumonias, Tuberculosis, respiratory syncytial virus, laryngitis and sinusitis.

Sample size was determined by the number of samples collected for nasal and oral swabs within a period of 12 months.

Data management and analysis: Results is presented as frequency of Human Influenza virus and its subtypes for both sex and age groups. The result were stored on secured excel data sheet and was analyzed using a statistical package for social science (SPSS) version 21.

Means (±SD) were used to describe continuous variables and proportions were used for categorical data. Two-tailed student’s t-test was used for comparisons of group means. When comparing groups of subjects, the chi-squared (X²) test was applied to determine the significance of the differences observed.

Ethical Approval

This was obtained from the ethical committee of the ATBUTH where the study was conducted.

Results

There were 300 samples collected for this study. 140 (46%) of the samples were from female patients, while 160 (53%) were samples collected from male patient. The age groups are as stated in Table 1 below.

<table>
<thead>
<tr>
<th>Age group</th>
<th>Frequency- n (%)</th>
<th>Positive results/age Grp</th>
<th>Age related Flu Types</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 - 12mo</td>
<td>50 (16.7%)</td>
<td>0</td>
<td>Flu A 9, Flu B 6</td>
</tr>
<tr>
<td>13mo - 5y</td>
<td>50 (16.7%)</td>
<td>15 (F11, M4)</td>
<td>Flu A 3, Flu B 0</td>
</tr>
<tr>
<td>6 - 25y</td>
<td>50 (16.7%)</td>
<td>3 (2F1M)</td>
<td>Flu A1, Flu B 1</td>
</tr>
<tr>
<td>26 - 45y</td>
<td>50 (16.7%)</td>
<td>2 (1 F 1M)</td>
<td></td>
</tr>
<tr>
<td>46 - 65y</td>
<td>50 (16.7%)</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>&gt; 65y</td>
<td>50 (16.7%)</td>
<td>0</td>
<td></td>
</tr>
</tbody>
</table>

Table 1: Frequency of Influenza virus based on age group.

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Table 2: Frequency of Human Influenza by sex.

<table>
<thead>
<tr>
<th>Sex</th>
<th>Frequency</th>
<th>Confirmed cases</th>
<th>Not confirmed</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>160</td>
<td>Flu A 4, Flu B 1</td>
<td>155</td>
</tr>
<tr>
<td>Female</td>
<td>140</td>
<td>Flu A 9, Flu B 6</td>
<td>125</td>
</tr>
<tr>
<td>Total</td>
<td>300</td>
<td>Flu A 13, Flu B 7</td>
<td>280</td>
</tr>
</tbody>
</table>

Figure 1: Frequency of Human Influenza %.

Figure 2: Frequency of Flu A and Flu B among subjects.

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Discussion

This study is a cross-sectional study that was conducted to determine the frequency, types, and predilection of human influenza virus among patients admitted with severe acute respiratory infection (SARI) in a tertiary hospital in Nigeria.

Overall, 6.7% of the samples collected tested positive for human influenza type A and B. During average epidemics, overall attack rates are estimated to be 10 - 20%, but in certain susceptible populations such as schoolchildren or nursing home residents, attack rates of 40 - 50% may occur [11]. 5% of the positive result was among the female samples whereas 1.7% from the male samples. Similar study also shows female predilection [12] even though some studies have shown higher mortality among male children with respiratory tract viral infection including avian influenza [13]. Among the positive sample about 65% were positive for Human Influenza type A (Flu A), whereas 35% where positive for Human Influenza type B (Flu B).

In Nigeria of the 2803 specimens tested, 217 (7.7%) were positive for influenza viruses, 167 (8%) were from subjects with influenza-like illness (ILI), 17 (5%) were from subjects with severe acute respiratory illness (SARI) and 33 were from subjects with unclassified condition. During the pre-pandemic period, subtype H3N2 was found to be the dominant circulating influenza A virus subtype while during the 2009 pandemic, influenza A virus subtype H1N1 replaced H3N2 as the dominant circulating virus [1,2]. Among persons with ILI, A/H1N1 pdm09 was most frequently found in children aged 5 - 17 years, whereas among subjects with SARI, it was most frequently found in persons aged 65 years. The percentage of specimens that tested positive for influenza virus peaked at 18.9% in 2010 and majority were A/H1N1 pdm09 [1].

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In sub-Saharan Africa, only Burkina-Faso, Cameroon, Ethiopia, Kenya, Madagascar, Rwanda and Mauritius reported low influenza activity with influenza A (H3N2) being predominant in DR Congo, Kenya, Mauritius and influenza A (H1N1) pdm09 in Ivory Coast, with influenza B being predominant in Niger, and Uganda. In Ethiopia, Ghana, Tanzania, Madagascar and Togo influenza A and influenza B have been identified in similar proportions in 2012 [14].

In tropical Asia, Influenza activity in this region is decreasing or at low or undetectable levels. Although influenza B remains the most commonly detected influenza type in the region, a high proportion of influenza A (H1N1) pdm09 has been detected in India and Bangladesh in the past. In southern China, of the 1,074 specimens that tested positive for influenza. Of the positive specimens, 67% were influenza A (H3N2), 17% were influenza A unsubtyped, and only 17% were influenza B, indicating an increase in the proportion of A (H3N2) cases in this region [14].

There were more cases of human influenza among the age group 1 - 5 years representing 5% of total samples collected compared to 1% for age group 6 - 25 years and 0.7% for age group 26 - 45 years, the incidence were found to be less or absent from 45 years and above. Studies conducted during both pandemic years and interpandemic periods demonstrate that age-specific attack rates are often highest among schoolchildren [11]. Family studies conducted in Houston and documented age-specific attack rates during various epidemics during the 1970s and 1980s demonstrated high rates of infection in school-age children and the importance of schoolchildren as vehicles of infection within families [15,16].

About 33% of patient with influenza are asymptomatic [8,17]. Influenza reaches peak prevalence in winter and because of the variability of winter period in northern and southern hemisphere, there are actually 2 flu seasons each year [18]. Flu occur seasonally rather than uniformly throughout the year. One possible explanation is that, because people are indoors more often during the winter, they are in close contact more often, and this promotes transmission from person to person, another factor is that cold temperatures lead to drier air, which may dehydrate mucus, preventing the body from effectively expelling virus particles. The virus also survives longer on surfaces at colder temperatures and aerosol transmission of the virus is highest in cold environments (less than 5°C) with low relative humidity [19].

Conclusion

The burden of Influenza is likely to be under estimated in this part of the country, Avian influenza infection even though has less prevalence, it is not an uncommon infection in Nigeria and northeastern region, however concerted effort has not been made in the past to identify and type these virus, and determine its predilections, it is worth of note that this study has revealed that avian influenza type A is commoner than B and C and it has much predilection to female and under 5 age group, with seasonality in occurrence. More research is needed to subtype and identify novel types during outbreaks of avian influenza.

Acknowledgement

We wish to sincerely thank the National Influenza Surveillance Unit and reference laboratory of Nigeria for given us such opportunity to conduct this research, the laboratory unit has help analyze all our samples, also we recognize the effort of the entire team of this research in this hospital who have worked round the clock to produce this peace of article. We also wish to thank the entire management of Abubakar Tafawa Balewa University Teaching Hospital for given us the enabling environment to carry out this research.

Disclosures

This research is a hospital based research and neither the author nor any of the corresponding authors has been paid in any form or collected any resource in any form in order to conduct this research.

Signed:

Dr Jacob A Dunga

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Bibliography


