Phenotypic Differences in Virulence and Immune Response in Closely Related Clinical Isolates of Influenza A 2009 H1N1 Pandemic Viruses in Mice

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Abstract

To capture the possible genotypic and phenotypic differences of the 2009 influenza A virus H1N1 pandemic (H1N1pdm) strains circulating in adult hospitalized patients, we isolated and sequenced nine H1N1pdm viruses from patients hospitalized during 2009–2010 with severe influenza pneumonia in Kentucky. Each viral isolate was characterized in mice along with two additional H1N1 pandemic strains and one seasonal strain to assess replication and virulence. All isolates showed similar levels of replication in nasal turbinates and lung, but varied in their ability to cause morbidity. Further differences were identified in cytokine and chemokine responses. IL-6 and KC were expressed early in mice infected with strains associated with higher virulence. Strains that showed lower pathogenicity in mice had greater IFN-α, MIG, and IL-10 responses. A principal component analysis (PCA) of the cytokine and chemokine profiles revealed 4 immune response phenotypes that correlated with the severity of disease. A/KY/180/10, which showed the greatest virulence with a rapid onset of disease progression, was compared in additional studies with A/KY/136/09, which showed low virulence in mice. Analyses comparing a low (KY/136) versus a high (KY/180) virulent isolate showed a significant difference in the kinetics of infection within the lower respiratory tract and immune responses. Notably by 4 DPI, virus titers within the lung, bronchoalveolar lavage fluid (BALF), and cells within the BAL (BALc) revealed that the KY/136 replicated in BALC, while KY/180 replication persisted in lungs and BALC. In summary, our studies suggest four phenotypic groups based on immune responses that result in different virulence outcomes in H1N1pdm isolates with a high degree of genetic similarity. In vitro studies with two of these isolates suggested that the more virulent isolate, KY/180, replicates productively in macrophages and this may be a key determinant in tipping the response toward a more severe disease progression.

Introduction

The 2009 pandemic H1N1 influenza virus (H1N1pdm) arose through reassortment of two preexisting swine influenza viruses, a Eurasian avian-like virus and a North American triple reassortant virus [1–3]. The risk factors associated with human cases of H1N1pdm mirrored those of seasonal influenza [4]. As observed with seasonal influenza, the most common underlying chronic conditions among hospitalized patients were respiratory disease, asthma, cardiac disease, and diabetes [4–9]. However, in contrast to seasonal influenza, a greater proportion of severe and fatal cases had a pre-existing chronic illness. A second notable difference was the age distribution of hospitalized and severe cases. Children less than 17 years old had the greatest rates of hospitalization per capita and adults over 64 had the greatest rates of mortality per capita. Retrospective clinical studies focused on surveillance of H1N1pdm sequences present in ICU admissions suggest that pre-existing medical conditions may be a more important factor in severity rather than particular viral variants [10]. However, this does not explain why a greater proportion of persons with pre-existing medical conditions had more severe disease than typically observed for seasonal influenza. Further, previously unreported comorbidities such as morbid obesity have been widely suggested for H1N1pdm for increased risk for admission to the ICU and death [11]. At this time, it is difficult to rule out the contribution of viral variants to the resulting illness observed with the various comorbidities. Nonetheless, the course of illness and the progression to more severe disease are most likely due to the combined interplay of the individual’s health, the intrinsic phenotype of the infecting viral variant and the treatment regime.

In contrast to seasonal influenza viruses, the H1N1pdm viruses replicate well and show greater pathogenicity with viral antigen in the bronchiolar epithelia and the alveolus by day 3 post-infection.
In the BALB/c mouse, A/California/04/09 and other H1N1pdm viruses show lethality, but only at the highest dose of $10^5$ plaque forming units (PFU) [2,12]. Studies of infection of BALB/c with several different 2009 H1N1pdm virus isolates show high virus titers in nasal turbinates (NT) and lung tissues by 3 DPI. Virus titers show a slight decrease by 6 DPI in lung and NT although the decrease varies among strains, too. Proinflammatory cytokines and chemokines were elevated for most mice in whole lung specimens at 3 DPI for KC, IL-6, IL-12 (p40), G-CSF, M-CSF, MCP-1, MIG, MIP-1b, and LIF. By 6 DPI, IL-10 was present, albeit low, whereas some of the cytokines were reduced (e.g., MIP-2, IL-6), but most remained elevated [2,12]. The levels of cytokines and chemokines in humans or mice do not suggest hypercytokinemia common to H5N1 and 1918 viruses [13–15].

The overall genetic distance among H1N1pdm isolates remained low with 7 distinct clades in the first wave of the pandemic, while in the second wave a single viral lineage dominated [16]. Molecular surveillance, while important, will not confirm potential phenotypic differences dominated [16]. Molecular surveillance, while important, will not confirm potential phenotypic differences. Pandemic, while in the second wave a single viral lineage remained low with 7 distinct clades in the first wave of the pandemic. These data suggest four distinct immunological phenotypes that correlate with potential variability in the virulence of 2009 H1N1 influenza A viruses [13–15].

Potential phenotypic differences may be related to infection with certain hospitalized patients. Studies of influenza H1N1pdm viruses from adult patients hospitalized in Kentucky during the second pandemic wave, September 2009 and April 2010. Four of the nine patients died and all of the patients for whom data was available had an underlying chronic condition. This group of clinical isolates, with high genetic similarity, was characterized for virus load, virulence, and host immune response in mice. Immune responses in the lungs suggest four distinct immunological phenotypes that correlate with the observed mortality in mice. This study underscores the potential pathogenicity and immunogenicity of circulating strains. Further, such studies have potential to reveal the efficacy of treatment regimes.

Herein we report the isolation, sequence, and characterization of nine H1N1pdm influenza A viruses from adult patients hospitalized in Kentucky during the second pandemic wave, September 2009 and April 2010. Four of the nine patients died and all of the patients for whom data was available had an underlying chronic condition. This group of clinical isolates, with high genetic similarity, was characterized for virus load, virulence, and host immune response in mice. Immune responses in the lungs suggest four distinct immunological phenotypes that correlate with the observed mortality in mice. This study underscores the potential pathogenicity and immunogenicity of circulating strains. Further, such studies have potential to reveal the efficacy of treatment regimes.

Results

Isolation and Sequence Analyses of H1N1pdm Isolates from Hospitalized Patients

H1N1pdm strains were isolated from nasopharyngeal swab samples obtained from nine de-identified patients enrolled in the Severe Influenza Pneumonia Surveillance (SIPS) project, a clinical study of hospitalized patients with severe community-acquired pneumonia in Kentucky from December 2008–December 2011 (Table 1). Hospitals included in the SIPS project included two rural areas of Kentucky, one in the east and one in the west and one in an urban area (Louisville). The Midwest is active in agriculture and one of the highest livestock producers of hogs and pigs. The age of the patients ranged from 31 to 58 and included 5 females and 4 males. Many of those patients had underlying comorbidities commonly associated with severe influenza disease such as obesity, diabetes and chronic obstructive pulmonary disorder (COPD) (Table S1). Four of the nine patients died.

Viruses were isolated by passage through both Madin-Darby canine kidney (MDCK) cells and eggs and virus isolates were designated according to a tracking number that contains no patient identifier information. Herein, isolates will be referred to by the tracking number and virus isolation method (e.g., A/Kentucky/180/2010 egg-passaged isolate will be referred to as “KY/180E”). Viruses were amplified and the virus titers were determined by 50% tissue culture infectious dose (TCID$_{50}$), plaque forming units (PFU) and hemagglutination assays in MDCK cells (data not shown). Viral titers measured in TCID$_{50}$ and PFU assays were similar for MDCK-adapted viruses and egg-adapted isolates (e.g., median titer $\sim 10^5$). Some viruses (A/KY/96/09, A/KY/99/09, A/KY/108/09, A/KY/180/10) showed a 10–100 fold higher titers in MDCK cell culture than in egg, but some (A/KY/80/09, KY/110/09, KY/136/09) did not. Egg-adapted viruses that were not recovered included A/KY/104/09 and A/KY/190/10. Hemagglutination titers were generally lower from egg-adapted viruses and showed variability among isolates (data not shown).

Universal and gene-specific primers were designed and used to amplify and sequence full-length cDNAs of all viral segments from each isolate. The amino acid sequences encoded by each gene were deduced and compared. All of the isolates have stop codons in PB1-F2 at positions 12, 58, and 88 (data not shown) [18]. Amino acids that showed polymorphisms are listed for each viral protein for the majority of genes (Table S2). The most amino acid changes were observed in HA1 and acidic polymerase (PA) with 18 and 9 mutations collectively across the 7–8 the KY isolates (not all isolates were completely sequenced). Neuraminidase and the two basic polymerase subunits (NA, PB1 and PB2) each had 4–5 mutations while the nonstructural protein 1 (NS1) and HA2 showed 3 mutations collectively. In contrast, matrix (M1), nucleoprotein (NP) had only 1 mutation and there were none in M2, NS2 or PA-X. Of the mutations only a few have been previously studied functionally to our knowledge. For example, in the HA1/HA2 of KY/180E we noted changes in D222G, S83P, S183P, and E374K. E374K was identified as a vaccine escape mutation [19]. The S183P and the Q293H mutations in HA1 (found in KY/180E and KY/96E, respectively) have been reported by others [20–23], and have been associated with alterations in receptor binding and increases in disease severity in humans [24,25].

Differential Progression of Virus Replication and Disease in Mice Infected with H1N1pdm Strains

To make an initial assessment of the levels of replication and virulence potential of each of the clinical H1N1pdm isolates, groups of six DBA2 mice were intranasally-infected and monitored daily for clinical signs and body weight. In addition to the nine clinical isolates, we included two pandemic strains (A/CA/07/09 and A/NY/18/09) and a seasonal H1N1 strain (A/BN/59/07). On days three and six, three mice each were euthanized and analyzed for viral load and antibody titers (Table 2). All mice infected with influenza strains showed weight loss (Table 2). Mice infected with KY/96M, KY/80M, KY/180E, KY/90E, and NY/18E showed the greatest weight loss in this short study (Table 2). Mice infected with the seasonal influenza strain BN/59E showed the least weight loss (about 9%). All other mice infected with H1N1pdm influenza isolates (KY/136M, KY/99M, KY/136E, KY/110M, CA/07E, KY/108M, KY/110M, KY/104M, and KY/108E) showed weight loss averages ranging from 13% to 26%, respectively.

The mortality rates for mock- and virus-infected groups were expressed as a percentage of lethal infections (Table 2). BN/59, CA/07, KY/110 and KY/136 showed no lethality and low morbidity; whereas KY/190 and KY/180 showed the greatest
lethality and substantial weight loss. Both KY/80 and NY/18 showed lethality and weight loss. On 3 and 6 days post-infection (DPI), three animals were sacrificed and the lung tissues were collected for virus titration analysis (TCID\textsubscript{50} assay of lung tissue homogenate using MDCK cells). Virus titers of lungs from each mouse-infected group showed relatively little difference between the various isolates on 3 and 6 DPI (p>0.05 using pairwise Wilcoxon Rank-Sum test without correction for multiple comparisons, Table 2). Mice infected with egg-adapted viruses showed a higher mortality than several of the corresponding MDCK isolates (e.g., see KY/96, KY/99, KY/108). However, both egg and MDCK-adapted KY/80 and KY/180 showed lethality in mice.

The influenza-specific serum IgG antibody responses in mice were measured at 6 DPI by ELISA using antigen prepared from inactivated NY/18 virus (Table 2) and reciprocal endpoints were determined. Strains KY/80E, KY/180, and KY/190 had no detectable IgG antibody by 6 DPI. The seasonal strain and all other pandemic viruses produced detectable IgG antibody; the highest endpoint titers were collected from BN/59, KY/108E, NY/18E, KY/99M, KY/136, and KY/110M. In some cases, these numbers reflect a small numbers of animals remaining at 6 DPI.

### Cytokine and Chemokine Responses in H1N1pdm Isolates Show Four Distinct Phenotypic Profiles

Mouse cytokine and chemokine panels were employed to gain a broad overview of the immune response at 3 and 6 DPI in the lungs and sera of mice infected with each of the viral isolates (Figures 1 and 2). No significant difference was seen between egg- and MDCK-passaged isolates; therefore, the combined data are presented in these two figures. There was an overall increase in proinflammatory cytokines (TNF\textsubscript{a}, IL-1\textbeta, IL-6 and KC) albeit different levels were noted. The levels of TNF\textsubscript{a} and IL-1\textbeta were greater in the lungs on day 3 in viral isolates with greater lethality (e.g., KY/180, KY/190) as compared to those with no lethality (e.g., KY/136, p<0.05). These same cytokines were higher in the mice with no lethality on day 6 as compared to those showing lethality. A similar temporal pattern was noted with some chemokines (e.g., RANTES, Figure 2). Interferon-gamma (IFN\textgamma) and IL-10, an anti-inflammatory cytokine, were present on day 6 in mice infected with all but the most lethal strains, KY/180, KY/190 (Figure 1, p<0.05).

With the exception of isolate KY/110 and KY/104, the chemokines CCL2 (MCP-1), CCL3 (MIP1\textalpha), CXCL9 (MIG), and CXCL10 (IP-10) displayed similar levels in all isolates on 3 DPI followed by a relative increase in animals infected with nonlethal versus a decrease in animals infected with lethal isolates at 6 DPI (Figure 2 and Table 2). IFN\textgamma stimulates both IP-10 and MIG, and as expected the IFN\textgamma levels were low in lethal isolates as compared to nonlethal isolates at 6 DPI. Cytokines and chemokines that showed limited response in mice with any viral infection included Eotaxin, GM-CSF, IL-1\textalpha, M-CSF, IL-2, IL-3, IL-4, IL-5, IL-7, IL-9, IL-12p40, IL-13, IL-15, IL-17, MIP2, LIX, MIP1\textbeta, RANTES, IL-12p70, and VEGF (data not shown).

A principal components analysis (PCA) was performed using 11 cytokine/chemokine measurements from the lungs of mice infected with each virus (Figure S1). The cytokines and chemokines included in the analysis were selected because they were determined to be the most significantly different between isolates on 3 and 6 DPI by generalized linear model fitness testing (data not shown). The first two components explained 72% of the variance (Figure S1) and the third component explained an additional 10% (data not shown). CCL3, TNF\textalpha, IL-10, IL1\textbeta, and IFN\textgamma were all highly correlated with the first component dimension (>80%), meaning these cytokines were principally important in differentiating the isolates. CXCL10, KC and G-CSF were similarly highly correlated with the second component (>70%), and CXCL9 was highly correlated with the third component dimension (83%, data not shown). The remaining variables, IL-6 and CCL2, were not highly correlated with the components used in this model, although they were significantly different between isolates on 3 and 6 DPI. The mean for each isolate on 3 and 6 DPI are plotted according to the coordinates of the first two components in Figure 3 (and Figure S1).

The method used for computing this PCA is similar to k-means cluster analysis, and therefore we expected to observe the isolates to group according to their cytokine/chemokine signatures. To better visualize these clusters, we connected each isolates’ 3 DPI coordinate to its 6 DPI coordinate with an arrow (Figure 3). According to this method of using a “time-resolved” PCA, the isolates were observed to cluster clearly into four groups representing various trajectories of disease.

The first three groups of isolates could be differentiated by their immunogenicity and low to moderate lethality. The first group

### Table 1. General Patient Data for Nasal Swabs Used for Virus Isolation.

| Sample ID/Locality | Patient Age | Patient Sex | Hospital Admission Date | Nasal Swab Date | LOS \(^{(D)}\) (Days) | Comorbidity | Mortality |
|-------------------|-------------|-------------|--------------------------|----------------|-----------------------|-------------|----------|
| 80/3              | 31/F        |             | 9/30/2009                | 9/30/2009      | 10                     | COPD, MRSA  | Died     |
| 96/2              | 51/M        |             | 10/24/2009               | 10/26/2009     | 4                      | BMI = 123.6 | Survived |
| 99/3              | 35/M        |             | 10/28/2009               | 10/29/2009     | 3                      | ND          | Survived |
| 104/2             | 58/M        |             | 10/30/2009               | 11/4/2009      | 2                      | ND          | Died     |
| 108/6             | 57/F        |             | 11/2/2009                | 11/3/2009      | 19                     | ND          | Survived |
| 110/6             | 54/F        |             | 11/3/2009                | 11/4/2009      | 3                      | COPD, diabetes, renal disease, BMI = 79.7 | Survived |
| 136/3             | 46/M        |             | 12/10/2009               | 12/11/2009     | 8                      | Diabetes, MRSA | Survived |
| 180/2             | 53/M        |             | 3/24/2010                | 4/1/2010       | 19                     | COPD, renal disease | Died     |
| 190/2             | 55/F        |             | 4/10/2010                | 4/15/2010      | 12                     | ND          | Died     |

\*locality, 2 = western Kentucky (KY); 3 = eastern KY; 6 = Louisville metro; ⁹LOS, Length of stay in hospital; MRSA, methicillin-Resistant Staphylococcus aureus; COPD, chronic obstructive pulmonary disease; BMI- body mass index; ND, none determined.

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Phenotypic Differences of H1N1pdm Isolates
includes isolates with a low lethality in mice, and was exemplified by infection with the seasonal strain BN/59, and 2009 pandemic strains CA/07, KY/80, and KY/136. The second group (KY/96, KY/99, and KY/104) included those strains with moderate lethality in the mouse model, and was similar to the first group in course of disease, but was characterized by increased inflammatory cytokines (e.g., IL-6, KC-like, and G-CSF) and chemokines (e.g., CXCL9). In contrast to the first group, these had a higher level of the anti-inflammatory cytokine IL-10 later in disease, as well as increased IFNγ at 6 DPI. The third group of influenza strains (KY/108 and KY/110) was the most immunogenic compared to the other isolates, and differed from the two groups above in that they showed the highest levels of inflammatory cytokines late in infection (e.g., IL-6, TNF-α, IL-1β) but were similar to Group 2 in terms of lethality. They showed the highest level of chemoattractant chemokines, particularly CCL2, CCL3, and CXCL10, and produced the highest levels of IL-10 and IFNγ detected in this study.

Finally, the fourth group of isolates, consisting only of KY/180, KY/190, and NY/10, were the most lethal of the viruses screened in mice. Their proinflammatory cytokines were elevated throughout the course, but not different from 3 DPI in Group 1 isolates. Most notably, these isolates failed to produce any IFNγ or IL-10 by 6 DPI. In Figure 3, the trajectory of these isolates point in an opposite direction from all other isolates, perhaps indicating a different course of disease. In summary (see Table S5), the immune responses of the virus isolates clustered similar to other phenotypic markers of virulence and lethality (Table 2).

### Table 2. Summary of viral titers, lethality of H1N1pdm isolates in DBA2 mice.

| Inoculum | DPI | Virus titer (Lung) | Lethality |
|----------|-----|--------------------|-----------|
| Mock     | 3   | <1.5 *             | 0.0       |
|          | 6   | <1.5 *             | 0.0       |
| KY/80M   | 3   | 7.2±0.5            | 0.0       |
|          | 6   | 6.4±0.9            | 33.3      |
| KY/80E   | 3   | 7.6±0.1            | 0.0       |
|          | 6   | 7.0±0.3            | 33.3      |
| KY/96M   | 3   | 7.0±0.5            | 0.0       |
|          | 6   | 7.2±0.5            | 0.0       |
| KY/96E   | 3   | 7.3±0.4            | 0.0       |
|          | 6   | 7.4±0.4            | 100.0     |
| KY/99M   | 3   | 6.4±0.2            | 0.0       |
|          | 6   | 5.8±0.7            | 0.0       |
| KY/99E   | 3   | 7.4±0.9            | 0.0       |
|          | 6   | 7.3±0.3            | 33.3      |
| KY/104M  | 3   | 6.6±0.9            | 0.0       |
|          | 6   | 6.9±0.5            | 33.3      |
| KY/108M  | 3   | 7.5±0.6            | 0.0       |
|          | 6   | 6.6±0.4            | 0.0       |
| KY/108E  | 3   | 6.5±0.5            | 0.0       |
|          | 6   | 6.4±0.8            | 33.3      |
| KY/110M  | 3   | 5.9±0.4            | 0.0       |
|          | 6   | 6.2±0.9            | 0.0       |
| KY/110E  | 3   | 5.9±0.2            | 0.0       |
|          | 6   | 6.8±0.7            | 0.0       |
| KY/136M  | 3   | 5.3±0.3            | 0.0       |
|          | 6   | 5.8±0.9            | 0.0       |
| KY/136E  | 3   | 5.8±0.4            | 0.0       |
|          | 6   | 5.2±0.5            | 0.0       |
| KY/180M  | 3   | 6.8±0.3            | 0.0       |
|          | 6   | 5.7±0.7            | 100.0     |
| KY/180E  | 3   | 6.9±0.4            | 33.3      |
|          | 6   | 6.4±0.1            | 66.7      |
| KY/190M  | 3   | 6.8±0.5            | 0.0       |
|          | 6   | 6.3±0.3            | 100.0     |
| NY/18E   | 3   | 6.9±0.6            | 0.0       |
|          | 6   | 6.8±1.0            | 33.3      |
| CA/07E   | 3   | 7.2±0.6            | 0.0       |
|          | 6   | 6.9±0.8            | 0.0       |
| BN/59E   | 3   | 6.7±0.0            | 0.0       |
|          | 6   | 5.9±0.1            | 0.0       |

Legend: Virus titer (log10 TCID50/mL at a limit of detection = 10^1.5 TCID50/mL). doi:10.1371/journal.pone.0056602.t002

In-depth Comparison of the Temporal Progression of Survival, Viral Load and Immune Responses of Two Clinical Isolates, KY/136E and KY/180E, with Low and High Virulence, Respectively

KY/136E and KY/180E were selected from the nine H1N1pdm clinical isolates for further characterization as representatives of lower and higher virulence based on the apparent disease in mice. These isolates had similar concentrations of lung cytokines and chemokines on 3 and 6 DPI compared to other isolates and clustered together in the PCA. However, the time-resolved PCA revealed that the isolates differed in the progression of immune responses in the course of infection. To gain additional insight, we first assessed dose response to infection in DBA/2 mice. Mice were infected with 10^2, 10^3 or 10^5 TCID50 of KY/136E or KY/136E and examined daily for clinical signs. Each day, mice were weighed and data from each dose group are presented in Figure 4A (KY/136E) and Figure 4B (KY/180E). The Kaplan-Meier survival curves of mice infected with KY/136E or KY/136E (Figure 4C and 4D) confirmed the observations reported above in terms of the general lethality of each virus, although KY/136E did show lethality in 1–2 mice on 10 DPI (Figure 4C). Mice infected with KY/180E succumbed to infection starting at 3 DPI at the high dose and on 6 DPI for the middle dose (Figure 4D). Animals were humanely euthanized upon showing a moribund state or upon a 25% loss in body weight as described in the materials and methods.

The levels of infectious virus present in the upper and lower respiratory tracts, nasal turbinates (NT) and lung, respectively, were measured at 1, 3 and 5 DPI at the three different doses (10^5, 10^6 or 10^7 PFU per mouse) of KY/180E or KY/136E (Table S3). As expected, levels of virus were greatest in mice with higher doses of infection. Mice infected with KY/180E persisted at higher levels of virus over the 5 day period as compared to KY/136E which had approximately 2–3 logs lower virus in the lungs and had virus levels that decreased from 3 to 5 DPI. The infectious dose (ID50) and lethal dose (LD50) were <10^5 TCID50/mouse and 10^1.2 TCID50 per mouse, respectively, for KY/180E (data not shown).

To further dissect the immune responses of these two viruses, multiplex cytokine/chemokine bead arrays were employed to provide insight into the temporal patterns for the key cytokine and chemokine profiles noted in the broad survey of all the...
isolate. Mice (5 per group/per day) were infected with each of the isolates and humanely euthanized on 1, 3, and 5 DPI (Figure 5 and 6). There was an overall increase in proinflammatory cytokines (TNFα, IL-1β, IL-6 and KC) in the lungs of all mice, however, there were much higher levels of all cytokines in lungs of mice infected with KY/180E. Further the responses were much earlier and correlated with the high levels of infection noted in the lung. In contrast with KY/136E-infected mice showed a gradual progression in immune responses, however, the overall responses on 6 DPI of KY/136E remained low. Interferon-alpha (IFNα) and IL-12p70 levels were higher in the lungs of KY/180E-infected mice as compared to KY/136E-infected mice (Figure S2). Virus-infected mice that recovered from KY/136E infection showed a high IL-10 response by 6 DPI whereas KY/180E-infected mice showed no response (Figures 1 and 5, p<0.05). As expected, and also shown in Figure 1, the IFNγ levels were low in the lethal KY/180E isolate as compared to the robust response observed in the...
nonlethal KY/136E isolates at 6 DPI (no statistically significant difference).

Mice infected with the middle dose of KY/180E that survived after 6 DPI, and produced high concentrations of IFNγ, eventually became moribund with all but one euthanized by Day 10. These data suggest that the immune response differences between the two isolates are due to the high levels of infection in the lung by KY/180E. The levels of cytokines and chemokines in the lung support this observation (Figures S3–S6), showing a dose response for both isolates on 1, 3, and 5 DPI in mouse lung homogenate.

Notably, a dose of $10^2$ pfu/mouse of KY/180E showed a progressive increase in chemotactic chemokines to 5DPI whereas a higher dose ($10^5$ PFU/mouse) began declining in concentration of these analytes after 3DPI (Figure S3). Similar patterns were seen for proinflammatory cytokines in the lung (Figure S5) and cytokines involved in adaptive immunity initiation, IFNγ and IL12p70 (Figure S6). The endpoint titer IgG responses of the mice for the low and middle doses were similar for both viruses (Figure 7; all significantly different from control using Student's $t$-test, $p<0.05$). Further, the HI titers were also very similar except

Figure 2. Chemokine levels in mice infected with pandemic and seasonal H1N1 influenza viruses. Six to eight week old DBA/2 mice were infected intranasally with $10^5$ TCID$_{50}$ with a seasonal virus isolate (BN/59), Kentucky (KY/80, KY/136, KY/96, KY/99, KY/104, KY/108, KY/108, KY/180, KY/190) or other H1N1 pandemic isolates (CA/07, NY/18) from 2009. Chemokine levels were measured at 3 and 6 DPI as described in the materials and methods and presented as mean +/- SEM (n = 6 per group when possible).

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for the lower dose, which was about 2-fold higher in mice infected with KY/180E (Figure 7).

The leukocyte chemoattractant chemokines CCL2, CXCL10, KC (CXCL8-like), and G-CSF were greater and earlier in the lungs of mice infected with high doses of KY/180E than with a similar dose of KY/136E (Figure 6, p<0.05). As expected, cytopsins of BAL revealed that KY/180E contained more neutrophils and monocytes on Day 4 compared to KY/136E (Figure 8A,B, p<0.05). Additionally, we observed a higher proportion of macrophages in the lungs of the low dose group of KY/136E compared to the higher dose group (p<0.05, Figure 8C). Despite there being a greater number of infiltrating leukocytes in the lungs of KY/180E-infected mice, these mice had a delayed clearance of virus in the lung.

Given the apparent clearance of virus from the lungs of KY/136E (Table S4), we were interested in whether the apparent differences in lethality of the two viruses could be due to the site of virus replication. We tested virus titers from bronchoalveolar lavage fluid (BALf), from the cellular pellet of the BALc (BALc) of the mice, and from the homogenized lung after lavage using TCID_{50} assay. Samples were collected on 3 and 4 DPI because we began to see decreases in viral titer in the lungs of KY/136E-infected mice after 3 DPI. In our earlier studies (Table S4), the virus titer in lung reflects lung, BALf, and BALc cells combined and we observed a nearly 3 log difference between KY/136E and KY/180E. When BAL is collected before isolation of lung tissue, we found the majority of infectious KY/136E virus present in lungs of mice was localized in the cellular fraction of the BAL (Figure 9). In contrast to KY/180E, no infectious KY/136E virus was present in the BALf by 4 DPI. Flow cytometric analysis revealed that the influenza NP-positive cells in the BAL from both isolates were also Gr1-positive (clone RB6-8C5), a marker of mouse phagocytic leukocytes, i.e., Ly6C/Ly6G positive (Figure 5D).

### Infection and Replication of H1N1pdm Viruses in Macrophage Cell Lines

Given the differences noted within the levels of virus in BALf and BALc, we used a BALB/c mouse macrophage cell line (RAW264.7) and a C57BL/6 mouse macrophage cell line (NR-9465) to test for the ability of KY/136E and KY/180E to infect and replicate in macrophage cells. Confocal microscopy at 24 hours post-infection showed that both viruses were able to infect macrophage cell lines (Figure 10). However, KY/180E was more efficient at replication, reaching 10-fold higher virus titers that persisted longer in both cell culture systems (Figure 11 and Figure S7). Although the cell culture supernatant was positive at 24 hours post-infection for both viruses in vivo, KY/180E was the only isolate detected from the cellular component in BAL (Figure 8; p<0.05 at 24 and 48 hours post-infection). These observations agree with our findings in the DBA/2 mouse model and suggest that KY/180 is more successful in vivo in replication and production of infectious virus in macrophages (Figure 8).

### Discussion

Herein we present data characterizing several isolates of H1N1pdm influenza virus taken from human patients with severe pneumonia in Kentucky, USA. Clinical surveillance from foci around the world showed that the majority of human cases of H1N1pdm were mild [26,27]. Severe disease from H1N1pdm infection was more likely to be seen in patients with pre-existing chronic illness [28]. It was initially observed that antigenic and genetic variation in circulating pdmH1N1 2009 influenza viruses was less than what is seen during seasonal human influenza A (H1N1) [29]. Similarly, the isolates listed here show relatively low variation, differing at only 43 amino acid positions. Each isolate contained from 2 to 12 unique non-synonymous mutations from the consensus sequence.

Three of the four lethal cases occurred in western Kentucky, and 1 of these lethal cases (KY/180) contained a virus isolate with the avian-like HA RBS (G222). The D222 is most commonly associated with human H1 strains while the G222 is common in avian strains. The HA 222 position, D222G/N/S/E/Y, was a common polymorphism in severe cases of influenza in human patients, which is within the HA receptor binding site, RBS [10,30–32] [33]. Additionally, the D222G has been associated with increased pathogenesis in some animal models possibly due to increased replication of the virus [30,31,34–38]. The aspartic acid functions in binding of the receptor and as a calcium antigenic site. Each of these amino acids will give rise to a specific binding affinity to the 2,6-sialic acid receptor with the D222G increasing 2,3-sialic acid receptor binding specificity [35,37,39]. However, it is clear that other amino acids constellations at positions 183, 186, 187, 216 and 224 will also influence these interactions [3], and may result in different overall outcomes. Additional amino acid signatures in closely related H1N1pdm isolates have also been implicated in differences in virulence in animal models. For example, in

**Figure 3. Principal components analysis of immune responses in lungs of mice to infection with pandemic and seasonal influenza viruses.** A principal components analysis was performed using the 14 cytokines/chemokines analytes shown to be the most significantly different across all isolates from days 3 and 6 (determined by generalized linear model fitness testing, data not shown). The data were normalized and scaled (zero mean-centered) cytokine responses after influenza infection at 3 and 6 DPI for all viruses. The ordinate and abscissa represent the first and second components from the PCA, which explain approximately 72% of the variance. Each arrow represents the mean of a virus isolate tested in mice. Arrow tails represent day 3 components and arrow heads represent day 6 components. Using this tool to visualize the immune response, the arrows depict the trajectory of disease of the various influenza isolates as tested in DBA/2 mice. Additionally, the 12 virus isolates clustered into four distinct patterns: Group 1, BN/59, CA/07, KY/80, KY/136; Group 2, KY/96, KY/99, KY/104, KY/108; Group 3, KY/108, KY/110; Group 4, KY/180, KY/190, NY/19 (See Table S5). doi:10.1371/journal.pone.0056602.g003
a study in a cynomolgus macaque model of two highly similar strains of H1N1pdm, A/Mexico/4487 and A/Mexico/4108, showed notable heterogeneity in virulence [40]. The amino acid variations responsible for the differences noted in these studies have not yet been reported. The HA1 of KY/180E also has a P183 (noted as 186 in [20]) rather than a S183 at the Sb antigenic site, which also has been noted to affect receptor specificity ([21–23]). This site, independently or together with the D222G mutation, is thought to allow binding to α-2,3 linked sialic acid residues. This substitution may allow the virus to more efficiently enter and replicate in the lower airway epithelium [23,41–43].

Other mutations observed in the HA gene of the Kentucky isolates have also been reported by others. Belser, et al. (2010) published data that showed A/CA/04/09 also has the S83P mutation in HA (as in KY/180E), although this mutation is not a known pathogenicity determinant [12]. Byushina, et al. (2010) found the S183P (as seen in KY/180E) mutation arose from serial passage of A/CA/04/09 in mice and resulted in increased pathogenicity [21]. Xu, et al. (2011) report isolates from China show similar variability and share some of our mutations in HA (S83P, T203S, and V321I; seen in KY/180E, KY/96E, and KY/80E, respectively), as well as mutations in NS1 (V123I, seen in KY/99E) and PA (V14I, seen in KY/180E) [44]. Melidou et al. have data that suggest V321I mutation in HA may be associated with increased disease severity [24]. Finally, HA1 amino acid mutation Q293H, identified in KY/96E, has been associated with increased severity of disease observed in human cases during the pandemic [24,25]. We include a summary of those mutations that correlate with virulence in humans and mice (Table S6). An understanding of these and other signatures associated with more virulent phenotypes will benefit insight into the biology of the virus, patient management and public health responses.

In total, KY/180E contains mutations that differ from KY/136E in seven amino acid sites in the HA protein, four sites in PA, two sites in NA, PB1 and PB2, and one site in each of M1, NS1, and NP. Comparison of the CA/07/09, NY/18, NL/602 isolates with the KY isolates showed additional variation among all isolates (Table S2). Specifically, CA/07/09, NY/18, NL/602 were similar to the KY consensus in M, HA2, PA, PB1 and PB2. In HA1, a Q223R occurred in NY/18 only. In NA, a D248N changed occurred in CA/07 and NL/602. In NS1, a D247N was noted for KY/110, CA/07, NY/18, and NL/602. Most additional changes were conserved hydrophobic changes such as Val to Iso or Met to Leu. No differences were observed among the KY, CA/07/09, NY/18, NL/602 isolates in M2 or NS2 (Table S2). Despite the small apparent genomic variation between H1N1pdm isolates, we and others have shown that there exists substantial variation in the course of disease, pathogenicity, and immune response in human patients, in animal models, and in primary cell culture [25,40,44–46]. As virulence determinants of influenza viruses typically involve the genes encoding the HA, NA, and polymerase proteins, we would expect changes in these genes to be of future interest for

Figure 4. Weight loss and Kaplan-Meier curve of mice infected with KY/136 and KY/180. Mean weight change (+/- SEM) after six to eight week old DBA/2 mice were infected intranasally with 10⁵, 10⁶ and 10⁷ TCID₅₀ with KY/136E (A) or KY/180E (B). Mice were examined daily for clinical signs and weighed (n = 10 mice per virus group). Kaplan-Meier Survival curves for these mice show KY/136E to have low lethality (C) compared to KY/180E (D).

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biological function [17]. Addition genome diversity that might give rise to unique phenotypes during infection may be revealed be deep sequencing or isolation of clones that sample the population of the virus [17].

To gain insight into the potential phenotypic variability inferred from the genotypic variability each virus was screened in DBA2 mice. All IAV tested so far, H1N1pdm, 1918 H1N1, and seasonal influenza A H1N1 virus, infect DBA2 with varying levels of

Figure 5. Cytokine levels in mice infected with KY/136 or KY/180. The levels of notable cytokine responses are shown for 1, 3 and 5 DPI in six to eight week old DBA/2 mice that were infected intranasally with 10^5 TCID_{50} of KY/180E or KY/136E (n = 10 mice per virus group per time point). Statistical significance was determined by day using Kruskal-Wallis test followed by pairwise Wilcoxon Rank Sum post hoc test with Holm’s adjustment for multiple comparisons. P-values <0.05 are indicated by the following method: “a” = KY/180 is significantly different from mock; “b” = KY/136 is significantly different from mock; “c” = KY/180 is significantly different from KY/136. doi:10.1371/journal.pone.0056602.g005
Figure 6. Chemokine levels in mice infected with KY/136 or KY/180. The levels of notable chemokine responses are shown for 1, 3 and 5 DPI in six to eight week old DBA/2 mice that were infected intranasally with $10^5$ TCID$_{50}$ of KY/180E or KY/136E ($n=10$ mice per virus group per time point). Statistical significance was determined by day using Kruskal-Wallis test followed by pairwise Wilcoxon Rank Sum post hoc test with Holm’s adjustment for multiple comparisons. $P$-values $<0.05$ are indicated by the following method: “a” = KY/180 is significantly different from mock; “b” = KY/136 is significantly different from mock; “c” = KY/180 is significantly different from KY/136.

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Figure 7. IgG responses of mice infected with different doses of KY/136 or KY/180. Serum from day 20 post-infection was taken from six to eight week old DBA/2 mice that were infected intranasally with 10^6, 10^7 and 10^8 TCID_{50} of KY/180E or KY/136E (n = 10 mice per virus group per time point). Influenza specific (A/NY/18/09 BPL-inactivated whole viral antigen) IgG titers were measured by ELISA and presented as average log_{2} endpoint titers (+/-SEM). No mice survived to day 20 at the high dose of KY/180. All endpoint titers for animals infected with influenza virus isolates were significantly different from mock (p < 0.05). Endpoint titers for both IgG and HI from animals infected with KY/180 were significantly different from the same dosage amount of KY/136 (p < 0.05).

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Phenotypic Differences of H1N1pdm Isolates

The high dose of KY/180. All endpoint titers for animals infected with KY/136 were significantly different from the same dosage amount of KY/136 (p < 0.05). Endpoint titers for both IgG and HI from animals infected with KY/180 were significantly different from the same dosage amount of KY/136 (p < 0.05).

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responses [64,65], which was seen with KY/180E. Neutrophils are important for assisting in clearing influenza-infected cells from the lungs directly and indirectly [66,67], but the precise mechanisms remain to be discovered [64,68,69]. Although important to the resolution of influenza virus infection, neutrophils contain cytotoxic granules that may cause severe pathology, and may contribute to morbidity and mortality [70]. An increase in neutrophils late in disease confers the pathogenicity seen in highly pathogenic strains of influenza viruses [69].

Natural Killer (NK) cells are known to be potent innate immune cells that recognize influenza hemagglutinin [71]. Classically, NK cell effector functions include release of IFNγ and direct cytotoxicity to influenza-infected cells by granule exocytosis (e.g., granzyme B and perforin). Mice infected with KY/180E have higher numbers of peripheral NK cells at 3 DPI compared to KY/136E using flow cytometry (data not shown). Consistent with this finding, CCL5 (RANTES) was increased in the lungs and serum of mice infected with KY/180E compared to KY/136E (Figure 2). NK cell deficiency in mouse models of influenza infection leads to increased pathogenicity [72,73]. Despite the increase in NK cells in KY/180E-infected mice, it has been shown that some influenza isolates may counteract this effector response, causing lysosomal degradation of the ζ chain, which is a critical component of the NK-activating receptor [74]. Additionally, DBA2 mice lack the NKG2A activating receptor that is responsible for recognizing the down-regulation of MHC-I that typically occurs with a viral infection [75]. Therefore, KY/180E may be able to exacerbate this deficiency by decreasing NK cell effector function. It has been shown that NK cells also contribute to immune pathology in response to influenza and other respiratory virus infections [76].

Finally, this study and others have shown that some strains of influenza virus are capable of infecting macrophages [77–80], and may alter their functionality in response to infection [81,82]. During influenza infection, alveolar macrophages are critical in clearing virus from the lungs and are key producers of IFNγ to stimulate the adaptive immunity [64,65,76,83–88]. Upon infection, influenza viruses may interfere with their normal function [86,89,90]. Of the two isolates tested here, mice infected with KY/136E showed clearance of virus from the lung by day 4 [86,89,90]. Our in vitro data using mouse macrophage cell lines indicates that KY/180E was able to produce infectious virus better than KY/136E. Additionally, mice infected with KY/180E had a higher number of Gr-1 positive infiltrating cells into the lungs by day 4, a delayed IL-10 production, and very low levels of

Figure 8. Cells within bronchial lavage fluid of mice infected with KY/136 or KY/180. The cells within the bronchoalveolar lavage fluid (BALf) at 4 DPI from DBA/2 mice infected with 10^5 TCID_{50} of KY/136E (A) or KY/180E (B) (n = 5 mice per virus group per time point) were affixed to slides by cytospin centrifuge and stained (Kwik-Diff). Microscopic images from three fields per slide were counted by three blinded, independent observers. (C) The average number of macrophages, non-specific monocytes, and neutrophils are presented as a percent of the total cell count (+/− SEM). There were significantly more neutrophils in the BALf of mice infected with KY/180E virus (p<0.05, indicated by asterisks). (D) Cells in the BALf were fixed, permeabilized, and stained with anti-influenza nucleoprotein (NP)-FITC antibody conjugate and analyzed by flow cytometry. There were significantly more NP-positive cells in mice infected with KY/180 at a dose of 10^6 pfu, and in mice infected with KY/136 compared to mock-infected controls (p<0.05, indicated by asterisks). 80% of NP-positive cells were Gr-1 positive (macrophage or neutrophil).

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TCID<sub>50</sub> assay on MDCK cells. Lung compartments taken from mice from the lavage fluid by centrifugation and the fluid (BALf), cellular (BAL)

Figure 9. Virus titer the in lung compartments of mice infected
with KY/136E or KY/180. Bronchoalveolar lavage was taken from six
to nine week old DBA/2 mice that were infected with 10<sup>5</sup> TCID<sub>50</sub> KY/136E or KY/180E on days 3 and 4 post-infection. Cells were separated from the lavage fluid by centrifugation and the fluid (BALf), cellular (BAL cell), and whole lung homogenate were tested separately for virus by TCID<sub>50</sub> assay on MDCK cells. Lung compartments taken from mice infected with KY/180 had statistically higher virus titers at 3 DPI than from mice infected with KY/136E (p<0.05, indicated by asterisks on the figure). It is not possible to compute statistical differences from 4 DPI, as there were only two mice infected with KY/180 in that experiment that survived to 4 DPI.

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IFN<sub>γ</sub> resulting in a deficient virus clearance. Both our in vivo and in vitro data show that KY/180E can infect macrophages, which may affect the cell's ability to release IFN<sub>γ</sub> and explain the increase pathogenicity observed with this virus compared to KY/136E. Future studies will focus on understanding the mechanism(s) involved in the virus-induced deregulation of macrophage function in vitro and in vivo. Understanding the genetic basis for the differences in the interaction between the virus and macrophages as well as other pulmonary and immune cells will contribute to understanding the course of influenza infection in patients, and assist in predicting potential virulence during future outbreaks.

Materials and Methods
Detection of Influenza in Clinical Nasopharyngeal Swab Specimens
Nasal swab samples were provided by the SIPS project. A nasopharyngeal swab was taken from all patients meeting the case definition, which was defined as a patient admitted to an intensive care unit with the physician diagnosis of community-acquired pneumonia. The University of Louisville Institutional Review Board Human Subject Protection Program Office (HSPPPO) approved this study prior to any data collection (#08.0399). Informed consent was waived because it was a retrospective chart review and the data were analyzed anonymously. The nasopharyngeal swabs were taken as part of 'standard of care' diagnosis. Detection of influenza virus was accomplished using the Respiratory Viral Panel (RVP) detection kit (Luminex Corp., Austin, TX), recently approved for in vitro diagnostics by the FDA. The RVP is a reverse-transcriptase, real-time PCR assay which is multiplexed to detect 12 viral targets in a single reaction well. Those targets include Influenza A, Influenza A-subtype H1, Influenza A-subtype H3, Influenza B, RSV-A, RSV-B, Parainfluenza 1, Parainfluenza 2, Parainfluenza 3, human metapneumovirus, Rhinovirus and Adenovirus. Nasal swabs were collected from each patient and placed in Universal Transport Media (Copan Diagnostics, Inc, Murietta, CA). Nucleic acid was extracted using the QIAamp Mini-Elute Viral Spin Kit® (Qiagen, Valencia, CA) into a final volume of 50.0 μL. A 5.0 μL aliquot was combined with 20 μL of RVP mastermix and processed through several rounds of amplification, including a single-tube multiplex reverse-transcriptase PCR, followed by a multiplex target-specific primer extension protocol, a bead-hybridization step, and finally a data acquisition step in the Luminex reader. Results are qualitative and read-outs are Positive, Negative or a No Call (equivalent).

Viruses and Virus Isolation
The viruses CA/07, NY/18, and BN/39 were kindly provided by the Centers for Disease Control and Prevention, Virus Surveillance and Diagnosis Branch, Influenza Division. Viruses from clinical cases from KY present in nasopharyngeal swab specimens were isolated and propagated in the chorioallantoic cavity of 10-day-old embryonated chicken eggs (Charles River) and MDCK cell line purchased from ATCC (CCL-34). The allantoic fluid containing infectious particles or MDCK supernate was harvested 72 h after inoculation. MDCK cells were infected in DMEM virus culture medium (DMEM containing 0.2% BSA, 1% PEN/STREP, and 2 μg/mL of Trypsin-TPKC). The infectious virus titer of the resulting seed stock was determined by TCID<sub>50</sub> and the titer calculated by Reed and Muench [91], and confirmed by plaque assay on MDCK cells. Egg passage E2 was used for the studies involving KY/180E and KY/136E reported herein.

RNA Isolation and Sequencing
To sequence each of the gene segments of each H1N1pdm isolate, we followed the method described by Inoue et al [92] with minor modifications. In brief, total RNA was isolated with MagMAX AI/ND Viral RNA isolation kit (Ambion) from each virus seed stock solutions. Five μL of RNA was used to synthesize cDNA with SuperScriptase III (Invitrogen) from the FWuni12 and RVuni13 primers based on those reported by Inoue et al [92] and the cDNAs were PCR-amplified based with Accuprime (Invitrogen). The amplified products were separated by agarose gel electrophoresis and the DNA bands corresponding to the size of each segment were purified using the Wizard SV gel Clean-Up System (Promega). The purified DNAs were used as templates for automated dyeodeoxy sequencing with BigDye 3.1 cycle sequencing kit (Applied Biosystems). To sequence the entire gene we used approximately 500 bp overlapping gene-specific primers in both directions. For the sequencing of HA gene, we used FWuni12, RVuni13, HA462_f (5’-GACTCGAGACAAAGGTTGAACGG-3’) and HA1202_r (5’-GTCAATGGCATTCTGTGTGCTC-3’) as sequencing pri-
mers. For the sequencing of NA gene, we used FWuni12, RVuni13, NA_521r (5’-TGACCAAGGAGGACTGACTCAA-3’), NA376_f (5’-CCCTTGGAATGCAGAACCTT-3’), and NA905_f (5’-CGTGGGTGTCTTTCACCCAGAA-3’) as sequencing primers. The sequences were assembled with the SeqScape™ program (Applied Biosystems). Alignments of nucleic and amino acid sequences were completed to identify polymorphisms with other H1N1pdm isolates and sequences for all isolates were submitted through the Influenza Research Database and uploaded into the GenBank (accession numbers provided in Table S4).

Ethics Statement for Animal Studies
Mice studies were approved by the University of Louisville Institutional Animal Care and Use Committee with Veterinary Medicine tasked to monitor and support all animal experiments. Research was conducted in compliance with the Animal ...
Welfare Act and other federal statutes and regulations relating to animals and experiments involving animals and adheres to principles stated in the Guide for the Care and Use of Laboratory Animals, National Research Council, 1996. The facilities where this research was conducted are fully accredited by the Association for Assessment and Accreditation of Laboratory Animal Care International.

Mice Studies

Six to nine-week-old female DBA/2 mice were purchased from Jackson Laboratories (Bar Harbor, ME) and housed in the vivarium managed by UofL at the Research Resource Center or at the Regional Biocontainment Laboratory. The mice received food and water ad libitum, and all experiments were conducted in accordance with rules of the Institutional Animal Control and Use Committee of UofL. Mice were anaesthetized by isofluorane inhalation and infected intranasally with a total volume of 30 μl at the dose as outlined in the figure legends and text. Viruses were adjusted to the dose required in PBS (pH 7.2). Animals were observed daily for morbidity and measured for weight loss. All mice showing more than 25% body weight loss were considered to have reached the experimental end point and were euthanized humanely.

Mice were euthanized on the days noted in the results and on figure legends and blood, nasal turbinates and lung tissues were collected and stored appropriately until analyses. For collection of BAL after euthanasia, the neck and thoracic area were disinfected by soaking with 70% ethanol, and the trachea exposed by dissection. An 18G catheter needle was inserted into upper trachea, the needle removed, and the outer catheter sheath moved into the lower trachea. A 3 mL syringe filled with 1 mL of DPBS was used to carefully inject DPBS into the lungs, and aspirate BAL fluid. Collected fluids were transferred to an ice-cold, sterile 15 mL conical tube on ice and repeated once more as above. BAL fluids were immediately centrifuged at 500g for 10 minutes to separate fluids and cells. After centrifugation, supernatants are separated and kept separately at −80°C freezer until used for TCID50 assay. Cells pellets were washed once by adding 2 mL ice-cold DPBS and centrifuged again at 500g for 10 min. Cells were resuspended in 1 mL DPBS and kept at −80°C until used.

Figure 10. Detection of Influenza in mouse macrophage cell lines. Mouse macrophage cell line, RAW264.7, was infected in vitro with MOI = 1 of each influenza isolate (high pathogenic isolate, KY/180, and low pathogenic isolate, KY/136) on chambered microscopy slides. At 24 hours post-infection cells were fixed, permeabilized, and stained using a FITC antibody conjugate specific for influenza A (H1N1) nucleoprotein (green). Cells were counterstained with a nuclear dye (TO-PRO3, Molecular Probes, red), and visualized on a Zeiss LSM710 confocal microscope. Both isolates tested were observed to infect macrophages. Scale bars indicate 100 microns.
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TCID$_{50}$ Assay for Viral Load in Tissues and Bronchial Lavage Fluid

Tissues were homogenized in ice-cold virus culture medium (DMEM containing 0.2% BSA, 1% PEN/Strep) within a biosafety cabinet. Tissue homogenates were clarified by centrifugation (4000g for 20 min at 4°C) prior to storage at −80°C, and later analyzed by TCID$_{50}$ or Luminex cytometric bead array (next section). For measurement of the level of infectious virus present in tissue samples, the titers of the virus were determined by TCID$_{50}$ assay by titration of the clarified tissue homogenates or BALf on MDCK cells. The limit of virus detection was typically 10$^{1.5}$ TCID$_{50}$/ml or as indicated in each figure. Virus titers were calculated by the method of Reed and Muench [91], and are expressed as the mean log$_{10}$ TCID$_{50}$ per milliliter. Tissues in which no virus was detected were given a value of 10$^{1.0}$ TCID$_{50}$/ml for calculation of the mean titer.

Quantification of Cytokine and Chemokine Levels in Lungs

Sera and lungs from mice euthanized on days as noted in figure legends and text were analyzed by using Luminex® xMAP® technology-based assay kit (Millipore) according to the manufacturer’s protocol. The final reaction plate was read with a Luminex 100 or FlexMAP 3D machine and specific concentrations were calculated from a standard using Luminex xPONENT software.

Statistics and PCA

R (version 2.13.0) base statistical package and GraphPad software package were used in the analysis of data and generation of figures. Generalized linear models were constructed using each analyte included in the initial screen of Kentucky isolates in DBA/2 mice as a response variable (cytokine/chemokine concentrations) and isolate, DPI, and egg vs. MDCK stock, were used as predictors. The stock designation of the virus was removed from the model after it was seen that it did not contribute to explain significant proportion of the variance. Log likelihood ratios were used to compare the relative fit of each model, and the top 12 models showing the highest parameter estimates for the isolates were chosen for inclusion into the PCA (i.e., the cytokines/chemokines that were the most different between isolates, controlling for DPI, selected from the best-fit models). The R package “FactoMineR” (version 1.14) was used to perform the PCA. Statistically significant differences between multiple groups was assessed using Kruskal-Wallis tests followed by post hoc tests using pairwise Wilcoxon Rank Sum tests unless otherwise noted. Multiple comparisons were adjusted according to Holm’s method and a $p$-value <0.05 is considered significant.

Cytospin and Flow Cytometry

BAL cells were centrifuged at 300g for 5 min and washed twice in PBS. Cells were resuspended in 0.4 mL PBS and loaded into Shandon Cytospin$^\text{TM}$ centrifuge funnels. The cytospin chamber slides were spun at 1000 rpm for 5 min. Slides were air dried in a BSC and then were fixed and stained with Eosin and Methylene Blue (Kwik-Diff, Thermo Scientific). Slides were inspected under a light microscope.

For flow cytometry analysis, BAL cells were washed in PBS as above and stained with antibodies specific to mouse CD3 (500A2 Pacific Blue, BD Biosciences), Gr-1 (RB6-8C5 phycoerythrin, eBioscience), and CD49b (DX5 phycoerythrin-Cy7, eBioscience). Cells were then washed and fixed in 4% paraformaldehyde for 15 minutes at room temperature, followed by permeabilization with 0.2% saponin in 2% BSA for 20 minutes. Cells were then stained intracellularly for influenza nucleoprotein using a fluorescein isothiocyanate conjugated antibody (NP-FITC, AbCam #ab20921) for 30 minutes. Cells were washed and analyzed on a FACSaria II flow cytometer (BD).

Mouse Macrophages Experiments

RAW264.7 cells (ATCC #TIB-71) were seeded onto glass chamber slides (LabTek) and allowed to rest overnight. The cells were gently washed in PBS and virus was added at 1.0 MOI. The cells were washed and fixed at 24 hours post-infection and permeabilized with 0.2% saponin in 10% PBS. Cells were stained for intracellular influenza nucleoprotein as above, and a nuclear counter stain was included for the final 10 minutes (TO-PRO3, Molecular Probes). Cells were washed, mounted under a coverslip with ProLong Antifade media (Invitrogen), and visualized using a Zeiss LSM/710 confocal microscope. Kinetics of infection were determined by infecting mouse macrophage cell lines, RAW264.7 or BEI Resources #NR4536 (NAID, NIH), in triplicate wells of a 24-well plate in DMEM supplemented with 0.2% BSA, 1% Pen/Strep, 2.5% L-glutamine, and 25 mM HEPES. Supernatants were taken and centrifuged prior to performing a TCID50 assay using MDCK cells. Virus titers were confirmed at 24 hours using a plaque assay.

Supporting Information

Figure S1 Principal Components Analysis (PCA) of mouse lung cytokine and chemokine expression after challenge with clinical influenza A (H1N1) virus isolates from Kentucky, 2009. Standardized mean values for each cytokine/chemokine are plotted from Day 3 and Day 6 post-infection (n = 3, each) against the first two principal components, accounting for 72% of the variation in the analysis. (TIF)

Figure S2 Cytokine and chemokine profiles from mouse lung homogenate. DBA/2 mice were infected with KY/180E, KY/136E (10$^5$ pfu), or mock-infected with PBS. The mice were sacrificed 1, 3, and 5 days post-infection (n = 5 mice per group-
day). Samples from moribund mice taken after Day 5 post-infection were also analyzed when available. Bars indicate mean concentration.

**Figure S3.**

**Figure S4** Dose response of chemokines in the lungs of mice infected with KY/136 or KY/180 influenza A (H1N1) virus isolates. Mice were infected with 10^5, 10^2, or 10^5 pfu of virus and samples were collected upon euthanasia on days 1, 3, or 5 post-challenge (D1, D3, and D5, respectively; n = 5 mice per dose-day for all groups except for mice infected with 10^5 pfu of KY/180 where only 2/5 mice survived to D5.

**Figure S5** Dose response of cytokines in the lungs of mice infected with KY/136 or KY/180 influenza A (H1N1) virus isolates. Mice were infected with 10^5, 10^2, or 10^5 pfu of virus and samples were collected upon euthanasia on days 1, 3, or 5 post-challenge (D1, D3, and D5, respectively; n = 5 mice per dose-day for all groups except for mice infected with 10^5 pfu of KY/180 where only 2/5 mice survived to D5.

**Figure S6** Dose response of cytokines in the lungs of mice infected with KY/136 or KY/180 influenza A (H1N1) virus isolates. Mice were infected with 10^5, 10^2, or 10^5 pfu of virus and samples were collected upon euthanasia on days 1, 3, or 5 post-challenge (D1, D3, and D5, respectively; n = 5 mice per dose-day for all groups except for mice infected with 10^5 pfu of KY/180 where only 2/5 mice survived to D5.

**Figure S7** Replication kinetics of KY/180E and KY/180E isolates in C57BL/6 mouse macrophages. The macrophage cell line, BEIR #NR-9465, was infected at 1.0 MOI (two independent experiments at n = 3 per experiment) and clarified supernatants were taken at 4, 24, 48, and 72 hours post-infection. Virus titers were measured by TCID50 assay on MDCK cells.

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