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## Estimating the Potential Effects of a Vaccine Program against an Emerging Influenza Pandemic, United States

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### Abstract

**Background**—Human illness from influenza A(H7N9) was identified March 2013, and candidate vaccine viruses were soon developed. To understand factors that may impact influenza vaccination programs, we developed a model to evaluate hospitalizations and deaths averted considering various scenarios.

**Methods**—We utilized a model incorporating epidemic curves with clinical attack rates of 20% or 30% in a single wave of illness, case hospitalization ratios of 0.5% or 4.2%, and case fatality ratios of 0.08% or 0.53%. We considered scenarios that achieved 80% vaccination coverage, various starts of vaccination programs (16 or 8 weeks before, the same week of, or 8 or 16 weeks after start of pandemic), an administration rate of 10 or 30 million doses/week (the latter rate is an untested assumption), and two levels of vaccine effectiveness (two doses of vaccine required; either 62% or 80% effective for persons aged <60, and either 43% or 60% effective for persons aged 60).

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**Results**—The start date of vaccination campaigns most influenced impact; from 141,000 – 2,200,000 hospitalizations and 11,000 – 281,000 deaths were averted when campaigns started before a pandemic and <100 – 1,300,000 hospitalizations and 0 – 165,000 deaths were averted for programs beginning the same time as or after the introduction of the pandemic virus. The rate of vaccine administration and vaccine effectiveness did not influence campaign impact as much as timing of start of campaign.

**Conclusions**—Our findings suggest that efforts to improve the timeliness of vaccine production will provide the greatest impacts for future pandemic vaccination programs.

## Keywords

Influenza; influenza A (H7N9); influenza vaccine; mathematical modeling; pandemic

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## Introduction

Four influenza pandemics have occurred since the beginning of the 20th century and have ranged widely in transmissibility and clinical severity [1, 2]. On March 29, 2013, the Chinese Center for Disease Control and Prevention (China CDC) confirmed three human infections with an avian influenza A(H7N9) virus not previously reported in humans [3]. The pandemic potential of this virus was unknown. However, the high case fatality rate among humans ill from infection with this H7N9 virus (44 of 135 cases) [4, 5], the increased affinity of H7N9 for human-receptor-binding when compared to avian influenza A(H5N1) [6, 7], and the lack of pre-existing immunity among humans to H7N9 viruses [7, 8] raised concerns about the potential for substantial impact on human health if H7N9 were to develop the ability to transmit efficiently among humans. As a precautionary measure, US CDC and other partners began development of H7N9 vaccine candidate viruses [9].

The potential impact of a pandemic influenza vaccination program can vary widely based on a number of factors, including the size, speed, and number of waves of the pandemic outbreak, the number of doses administered, the timing of the vaccination program relative to the spread of the novel influenza virus, and the vaccine effectiveness (VE) [10]. To help public health officials and policy makers evaluate the impact of a hypothetical vaccination program against a future influenza pandemic, we developed a spreadsheet-based model that allowed quick exploration of the number of hospitalizations and deaths averted in the United States under various vaccination scenarios.

## Methods

We adapted a spreadsheet model (Excel, Microsoft Corp., Redmond, Washington, USA) that was originally created to estimate the effects of a vaccine program against influenza A(H1N1)pdm09 [10]. The model user enters an epidemic curve (the number of persons becoming ill by time) and other variables that define the impact of both the pandemic and the vaccination campaign. These variables include the timing of the vaccination program relative to the introduction of cases into the United States, the number of doses administered per week and the allocation by age group, the clinical attack rate, and the ratios of health outcomes to the number of cases (e.g., the case hospitalization and case fatality ratios)

(Table 1). We adjusted calculations to account for individuals who were naturally immunized through infection but who may still be vaccinated.

## Calculation Overview

To estimate the number of infections prevented by the vaccination program, we took [the number of persons fully vaccinated two weeks prior to the current week in the model]  $\times$  [the probability of not having been previously infected with influenza before being fully vaccinated and having developed immunity]  $\times$  [probability of becoming infected with influenza after being fully vaccinated and having developed immunity]  $\times$  [VE] [10]. We utilized standardized epidemic curves, using 20% and 30% clinical attack rates in one wave of illnesses and different levels of clinical severity and assumed that the pandemic began with 100 persons initially infected (Table 1) [11].

For our model, we assumed that two doses of vaccine administered three weeks apart would be needed to be fully effective, based on data indicating that previous H5 and H7 influenza vaccines have low immunogenicity [12–14]. We further assumed that, during a pandemic with moderate or high mortality, demand for vaccine would be such that 80% of the U.S. population would receive two doses of vaccine. We prioritized persons returning for their second dose of vaccine over persons who were receiving their first dose. We also assumed that vaccine was allocated in a pattern similar to the doses administered among four age groups (persons 6 months–9 years of age, persons 10–19 years, persons 20–59 years, and persons  $\geq$  60 years) during the 2012–13 influenza season [15] (Table 1). Once 80% of an age-group was fully vaccinated, we assumed vaccination would end in that age group. Remaining vaccine would then be allocated to other age groups until they reached 80% coverage. We allowed for a 2-week delay in protection against the virus after administration of the second dose of the vaccine [16].

We ran multiple scenarios to explore the effects of the quantity of the doses administered, the timing of the vaccine program, and the VE of the first and second doses. We first assumed that the program would administer either 10 million doses per week (approximating the maximum number of doses administered per week during seasonal influenza programs [17]) or 30 million doses per week; the latter has yet to be achieved during seasonal influenza vaccination programs. To explore the effects of timing of the vaccine program, we modeled programs starting at 5 different time points, separated by 8-week intervals: 16 weeks before, 8 weeks before, the same week as, 8 weeks after, and 16 weeks after the first cases of the novel influenza virus were introduced into the United States. We also assumed that one dose of vaccine was 0% effective for all age groups and two doses of vaccine were 62% effective in protecting against subclinical and clinical cases, hospitalizations, and deaths for persons aged  $<$ 60 years old and 43% for persons aged  $\geq$  60 years (Table 1). These values were based on the VE of the monovalent, inactivated, unadjuvanted influenza A(H1N1)pdm09 vaccine [10]. For the second scenario, we assumed a high VE due to the use of higher concentrations of hemagglutinin antigen [18] or the addition of an adjuvant to the vaccine [19]. In the high VE scenario, we assumed some VE with one dose (40% VE for persons  $<$ 60 years old and 30% for persons  $\geq$  60s) and higher VE

with two doses (80% effective for persons <60 years old and 60% for persons ≥60 years old) (Table 1).

To calculate the number of vaccine-associated averted outcomes (hospitalizations and deaths), we assumed 50% of infected cases were symptomatic and either 1.05% of symptomatic cases were hospitalized and 0.084% of symptomatic cases would die (low severity scenario) or 4.2% of symptomatic cases were hospitalized and 0.53% of symptomatic cases would die (high severity scenario) (Table 1). We adjusted the risk of hospitalization and death by age group (Table 1). The values for hospitalizations and deaths were based on estimates predicted for a pandemic with high clinical severity, and the adjustments for age were based on historic pandemics [11, 20].

## Results

### Base case Scenario

For the scenario with a cumulative clinical attack rate of 20%, without any other intensive interventions, the simulated pandemic peaked in the United States 20 weeks after the introduction of the first 100 cases and resulted in 127,000,000 infections, 63,500,000 clinical cases, and 669,000 hospitalizations and 54,000 deaths in the low severity scenario or 2,700,000 hospitalizations and 336,000 deaths in the high severity scenario (Figures 1, 2, and 3). For the scenario with a cumulative clinical attack rate of 30%, the simulated epidemic peaked 12 weeks after the start and resulted in 188,000,000 infections, 94,000,000 clinical cases, and 1,070,000 hospitalizations and 86,000 deaths in the low severity scenario or 4,300,000 hospitalizations and 538,000 deaths in the high severity scenario (Figure 1). Vaccination programs distributing 10 million doses per week would take 54 weeks to achieve 80% coverage of a two-dose vaccine series among all age groups, while programs distributing 30 million doses per week would take 21 weeks (Figure 1).

**Effects of a program beginning before (8 or 16 weeks) the introduction of influenza infections—**For an influenza pandemic with a 20% overall cumulative attack rate and high clinical severity, we estimated that a vaccination program beginning 8 weeks before the pandemic started in the United States and that administered 10 million doses of vaccine with the moderate VE per week could avert 568,000 hospitalizations and 71,000 deaths (21% reduction relative to no vaccine). Starting the vaccination program 16 weeks before the pandemic started would avert 777,000 hospitalizations and 97,000 deaths (29% reduction) (Tables 2 and 3; Figures 2 and 3). A vaccine program administering 30 million doses per week that started 8 or 16 weeks before the pandemic would avert 1,000,000 hospitalizations and 130,000 deaths (39% reduction) (Tables 2 and 3; Figures 2 and 3). Assuming that the VE would be 80% and 60% for persons <60 years and ≥60 years respectively (compared to base assumption of 62% and 43% for persons <60 years and ≥60 years respectively) would further reduce hospitalizations and deaths by at least an additional 10% relative to no vaccine for both the 10 and 30 million administration scenarios (Tables 2 and 3).

For an influenza pandemic with a 30% overall cumulative attack rate and high clinical severity, we estimated that a vaccination program beginning 8 weeks before the pandemic

started in the United States that administered 10 million doses of vaccine with the moderate VE per week could avert 570,000 hospitalizations and 71,000 deaths (13% reduction relative to no vaccine). Starting the vaccination program 16 weeks before the pandemic started would avert 924,000 hospitalizations and 116,000 deaths (21% reduction) (Tables 2 and 3; Figures 2 and 3). A vaccine program administering 30 million doses per week that started 8 weeks before the pandemic would avert over 1,400,000 hospitalizations and 180,000 deaths (33% reduction) while one that started 16 weeks before the pandemic would avert 1,700,000 hospitalizations and 207,000 deaths (38% reduction) (Tables 2 and 3; Figures 2 and 3). Using the high VE would further reduce hospitalizations and deaths relative to no vaccine for both the 10 and 30 million administration scenarios by at least an additional 7% (Tables 2 and 3).

**Effects of a program beginning the same week as the introduction of influenza infections**—For an influenza pandemic with a 20% overall cumulative attack rate and high clinical severity, we estimated that a vaccination program beginning the same week as the pandemic started in the United States that administered 10 million doses of vaccine with the moderate VE per week could avert 375,000 hospitalizations and 47,000 deaths (14% reduction). A vaccine program administering 30 million doses per week that started the same week as the pandemic would avert 916,000 hospitalizations and 114,000 deaths (34% reduction) (Tables 2 and 3; Figures 2 and 3). Using the high VE would further reduce hospitalizations and deaths by at least an additional 7% relative to no vaccine for both the 10 and 30 million administration scenarios (Tables 2 and 3).

For an influenza pandemic with a 30% overall cumulative attack rate and high severity scenario, we estimated that a vaccination program beginning the same week as the pandemic started in the United States that administered 10 million doses of vaccine with the moderate VE per week could avert 260,000 hospitalizations and 32,000 deaths (6% reduction) (Tables 2 and 3; Figures 2 and 3). A vaccine program administering 30 million doses per week would avert 777,000 hospitalizations and 97,000 deaths (18% reduction) (Tables 2 and 3; Figures 2 and 3). Using the high VE would reduce hospitalizations and deaths by at least an additional 4% relative to no vaccine for both the 10 and 30 million administration scenarios (Tables 2 and 3).

**Effects of a program beginning after (8 or 16 weeks) the introduction of influenza infections**—For an influenza pandemic with a 20% overall cumulative attack rate and high clinical severity, we estimated that a vaccination program beginning 8 weeks after the pandemic started in the United States that administered 10 million doses of vaccine with the moderate VE per week could avert 183,000 hospitalizations and 23,000 deaths (6.8% reduction relative to no vaccine). Beginning the vaccination program 16 weeks after the pandemic started would avert 17,000 hospitalizations and 2,000 deaths (0.6% reduction) (Tables 2 and 3; Figures 2 and 3). A vaccine program administering 30 million doses per week that started 8 weeks after the pandemic would avert over 538,000 hospitalizations and 67,000 deaths (20% reduction) while one that started 16 weeks after the pandemic would avert 50,000 hospitalizations and 6,000 deaths (1.9% reduction) (Tables 2 and 3; Figures 2 and 3). Using the high VE would reduce hospitalizations and deaths by at least an additional

1% relative to no vaccine for both the 10 and 30 million administration scenarios (Tables 2 and 3).

For an influenza pandemic with a 30% overall cumulative attack rate and high clinical severity, we estimated that no vaccination program that began 8 or 16 weeks after the pandemic started in the United States would avert more than 36,000 hospitalizations and 4,600 deaths (<1% reduction), regardless of whether 10 million or 30 million doses of vaccine per week with the moderate VE were administered (Tables 2 and 3; Figures 2 and 3). Using the high VE would reduce hospitalizations and deaths by no more than 5% relative to no vaccine for the vaccination programs beginning 8 weeks after the pandemic started in the United States. For the high VE scenarios starting 16 weeks after the pandemic started in the United States, no additional reductions in hospitalizations or deaths were observed (Tables 2 and 3).

## Discussion

In our analysis, the clinical attack rate and case-hospitalization-and-fatality ratios had the greatest impact on the number of severe outcomes averted in the United States while the vaccination program factor with the greatest impact was the timing of the start of vaccination relative to the start of a pandemic. For example, under the 30% clinical attack rate and 30 million doses per week scenario, a vaccination program starting 16 weeks before the start of the pandemic in the United States results in a 38% reduction in hospitalizations and deaths. Delaying the start of vaccination to the same week as the pandemic starts in the United States drops the reductions to 18%. The number of vaccine doses administered each week is also very important. Decreasing the doses administered to 10 million per week causes the impact of vaccination in the above two scenarios to decline to 21% and 6% reductions, respectively. The assumptions related to VE of the first and second doses were relatively less important.

This study highlights several key components to pandemic influenza preparedness, especially for a severe pandemic, including the importance of ensuring readiness to initiate large-scale vaccination programs as early as possible and ideally before the introduction of a novel influenza virus into the United States [15, 21]. Factors that may impact vaccine dose availability include how soon we develop an appropriate vaccine virus candidate, growth characteristics of vaccine virus candidates, influenza vaccine production capacity, efficiency of vaccine allocation and distribution, and vaccine administration capacity. Increased investment and research in vaccine production technologies, including the use of cell derived recombinant proteins [22], virus-like particles [23], or adjuvants (by conserving the use of hemagglutinin antigen) have the potential to increase the speed with which the number of vaccine doses can be produced.

Also important is the need to identify ways to invest in improvements that will notably increase the capacity to administer large number of doses of pandemic influenza vaccine. Currently, the peak administration rates for seasonal influenza in the United States are between 5–12 million doses per week. CDC is working with state and local health officials and vaccine providers to identify means to enhance vaccination administration capabilities.

In addition to exercising large-scale mass vaccination clinics, this includes increasing partnerships with non-traditional vaccine providers, such as pharmacies, supermarket chains, and other community vaccine providers, including diverse health, faith, and community based-organizations that reach vulnerable, at-risk, hard to reach, and minority populations [24, 25].

We evaluated a wide range of vaccine program initiation times relative to disease introduction in the United States in this paper. While this information cannot be known in advance, beginning vaccination 16 weeks prior to the introduction of disease in the United States might be possible if the pandemic virus was identified, a stockpiled influenza vaccine were available and appropriate for use, officials were prepared to administer vaccine, and the decision to vaccinate was made at least 28 weeks before the establishment of the virus in the United States (based on current estimates of 12 weeks to fill and finish and begin distribution of stockpiled pandemic vaccine) [26]. This timeline is dependent on robust novel influenza virus surveillance that can identify influenza viruses with pandemic potential before widespread transmission has occurred. During the 2009 H1N1 pandemic, however, the first cases were identified in the United States after widespread transmission had already occurred, and no stockpiled vaccine or vaccine candidate seed viruses were available. In this example, the first doses of vaccine became available 26 weeks after identification of the first case in the United States and 8 weeks after the start of the main wave of pandemic illness in the fall of 2009 [27]. This situation is demonstrated by the scenarios beginning 8 or 16 weeks after the introduction of the virus into the United States [28].

This study has several limitations. Most important, because infections with influenza A (H7N9) so far have been rare [5], the modeled number of pandemic-related hospitalizations and deaths, and the numbers of such that would be prevented by a vaccination program, can only be considered as illustrative and are not based on the current epidemiology of H7N9 or other novel influenza virus illnesses. These results, therefore, should not be interpreted as a prediction of the impact of a widespread outbreak of H7N9 or any other novel influenza A virus with pandemic potential. Additionally, we did not account for the effects of other interventions (e.g., non-pharmaceutical interventions such as mass gathering cancellations or school closures), the seasonality of when a novel virus might be introduced into the United States, or “waves of illness,” which are thought to have occurred in three modern pandemics [1, 2]. These factors could slow the course of the pandemic and thereby increase the amount of time to initiate and complete a vaccine program, increasing the number of hospitalizations and deaths averted. For ease of estimation, we also did not account for any adverse events associated with vaccination or for the indirect effects of vaccination (e.g. herd immunity). Theoretically, accounting for indirect effects would likely increase the number of hospitalizations and deaths averted for those vaccination programs assumed to start before the pandemic; this effect would likely be lower for those programs assumed to start the same time as or after the pandemic. Another important assumption is that 80% of the population would want to be vaccinated. This is distinctly different from recent seasonal influenza coverage estimates of approximately 45% [17]. We do not know the precise correlation between severity of an influenza pandemic and public demand for vaccination but 80% coverage may be an overestimate. Another potential limitation is that no data are available

on the VE of a possible H7N9 vaccine. Thus, we based our estimates of VE of either an unadjuvanted influenza vaccine, using data from the 2009 H1N1 pandemic, or a hypothetical vaccine with high VE based on data from adjuvanted 2009 H1N1 vaccine VE estimates. Limited data indicate that H7 vaccines have lower immunogenicity than seasonal influenza vaccines, which may result in lower VE [29, 30]. The population coverage or the effectiveness of a H7N9 or other future pandemic vaccine may be lower than what is assumed here, leading to a smaller number of averted outcomes.

Historically, influenza pandemics have been largely unpredictable events, and it is likely that the set of assumptions used in this study will vary from the actual events seen in the next pandemic, even if influenza A (H7N9) is the virus involved. However, the finding that variations in the timing of vaccination administration yield the greatest effect on the reduction in hospitalizations and deaths than do variations in rate of vaccine administration or effectiveness would likely remain consistent. Continued research and investment in work that improves the timeliness of vaccine production and administration will have the greatest benefits in the event of another influenza pandemic.

## References

1. Cox NJ, Subbarao K. Global epidemiology of influenza: past and present. *Annu Rev Med.* 2000; 51:407–21. [PubMed: 10774473]
2. Jhung MA, Swerdlow D, Olsen SJ, et al. Epidemiology of 2009 pandemic influenza A (H1N1) in the United States. *Clin Infect Dis.* 2011; 52 (Suppl 1):S13–26. [PubMed: 21342884]
3. Gao R, Cao B, Hu Y, et al. Human infection with a novel avian-origin influenza A (H7N9) virus. *N Engl J Med.* 2013; 368(20):1888–97. [PubMed: 23577628]
4. Gao HN, Lu HZ, Cao B, et al. Clinical findings in 111 cases of influenza A (H7N9) virus infection. *N Engl J Med.* 2013; 368(24):2277–85. [PubMed: 23697469]
5. World Health Organization. [Accessed 8/19/2013] Human infection with avian influenza A(H7N9) virus – update. Available at: [http://www.who.int/csr/don/2013\\_08\\_11/en/index.html](http://www.who.int/csr/don/2013_08_11/en/index.html)
6. Xiong X, Martin SR, Haire LF, et al. Receptor binding by an H7N9 influenza virus from humans. *Nature.* 2013; 499(7459):496–9. [PubMed: 23787694]
7. Zhou J, Wang D, Gao R, et al. Biological features of novel avian influenza A (H7N9) virus. *Nature.* 2013; 499(7459):500–3. [PubMed: 23823727]
8. Boni MF, Chau NV, Dong N, et al. Population-Level Antibody Estimates to Novel Influenza A/ H7N9. *J Infect Dis.* 2013; 208(4):554–8. [PubMed: 23687225]
9. Centers for Disease Control and Prevention. Emergence of avian influenza A(H7N9) virus causing severe human illness - China, February–April 2013. *MMWR Morb Mortal Wkly Rep.* 2013; 62(18): 366–71. [PubMed: 23657113]
10. Borse RH, Shrestha SS, Fiore AE, et al. Effects of Vaccine Program against Pandemic Influenza A(H1N1) Virus, United States, 2009–2010. *Emerg Infect Dis.* 2013; 19(3)
11. Meltzer MI, Gambhir M, Atkins CY, Swerdlow D. Introduction and standardizing scenarios to assess the needs to respond to an influenza pandemic. *Clin Infect Dis.* 2014 this supplement.
12. Sambhara S, Poland GA. H5N1 Avian Influenza: Preventive and Therapeutic Strategies Against a Pandemic. *Annu Rev Med.* 2010; 61(1):187–98. [PubMed: 20059335]
13. Couch RB, Patel SM, Wade-Bowers CL, Nino D. A randomized clinical trial of an inactivated avian influenza A (H7N7) vaccine. *PLoS One.* 2012; 7(12):e49704. [PubMed: 23239968]
14. Cox RJ, Madhun AS, Hauge S, et al. A phase I clinical trial of a PER. C6 cell grown influenza H7 virus vaccine Vaccine. 2009; 27(13):1889–97. [PubMed: 19368768]

15. Centers for Disease Control and Prevention. [Accessed 8/2/2013] Influenza Vaccination Coverage: How Well Did We Do in 2012–13. Available at: <http://www.cdc.gov/flu/fluview/1213season.htm>
16. Vega V, Hancock K, Schiffer J, et al. Sensitivity and specificity of serologic assays for detection of human infection with 2009 pandemic H1N1 virus in U.S. populations. *J Clin Microbiol.* 2011; 49(6):2210–5. [PubMed: 21471339]
17. Centers for Disease Control and Prevention. [Accessed 9/16/2013] FluVaxView Influenza Vaccination Coverage. Available at: <http://www.cdc.gov/flu/fluview/interactive.htm>
18. Falsey AR, Treanor JJ, Tornieporth N, Capellan J, Gorse GJ. Randomized, double-blind controlled phase 3 trial comparing the immunogenicity of high-dose and standard-dose influenza vaccine in adults 65 years of age and older. *J Infect Dis.* 2009; 200(2):172–80. [PubMed: 19508159]
19. Puig-Barbera J, Arnedo-Pena A, Pardo-Serrano F, et al. Effectiveness of seasonal 2008–2009, 2009–2010 and pandemic vaccines, to prevent influenza hospitalizations during the autumn 2009 influenza pandemic wave in Castellon, Spain. A test-negative, hospital-based, case-control study. *Vaccine.* 2010; 28(47):7460–7. [PubMed: 20875486]
20. Reed C, Biggerstaff M, Finelli L, et al. Novel framework for assessing epidemiologic effects of influenza epidemics and pandemics. *Emerg Infect Dis.* 2013; 19(1):85–91. [PubMed: 23260039]
21. Centers for Disease Control and Prevention. Final estimates for 2009–10 Seasonal Influenza and Influenza A (H1N1) 2009 Monovalent Vaccination Coverage – United States, August 2009 through May, 2010. [Accessed August 20, 2013]
22. Treanor JJ, El Sahly H, King J, et al. Protective efficacy of a trivalent recombinant hemagglutinin protein vaccine (FluBlok(R)) against influenza in healthy adults: a randomized, placebo-controlled trial. *Vaccine.* 2011; 29(44):7733–9. [PubMed: 21835220]
23. Lopez-Macias C. Virus-like particle (VLP)-based vaccines for pandemic influenza: performance of a VLP vaccine during the 2009 influenza pandemic. *Hum Vaccin Immunother.* 2012; 8(3):411–4. [PubMed: 22330956]
24. Goad JA, Taitel MS, Fensterheim LE, Cannon AE. Vaccinations administered during off-clinic hours at a national community pharmacy: implications for increasing patient access and convenience. *Ann Fam Med.* 2013; 11(5):429–36. [PubMed: 24019274]
25. Interfaith Health Program. [Accessed 2/5/2014] Public Health and Faith Community Partnerships: Model Practices to Increase Influenza Prevention Among Hard to Reach Populations. Available at: <http://ihpemory.org/ihp-programs/public-health-and-faith-community-partnerships/>
26. U.S. Department of Health & Human Services. [Accessed 2/18/2014] News Release. Available at: <http://www.phe.gov/Preparedness/news/Pages/fill-finish-130930.aspx>
27. Brammer L, Blanton L, Epperson S, et al. Surveillance for influenza during the 2009 influenza A (H1N1) pandemic-United States, April 2009–March 2010. *Clin Infect Dis.* 2011; 52 (Suppl 1):S27–35. [PubMed: 21342896]
28. President’s Council of Advisors on Science and Technology. [Accessed September 19, 2013] Report to the President on Reengineering the Influenza Vaccine Production Enterprise to Meet the Challenges of Pandemic Influenza. Available at: <http://www.whitehouse.gov/sites/default/files/microsites/ostp/PCAST-Influenza-Vaccinology-Report.pdf>
29. De Groot AS, Ardito M, Terry F, et al. Low immunogenicity predicted for emerging avian-origin H7N9: Implication for influenza vaccine design. *Hum Vaccin Immunother.* 2013; 9(5)
30. Fries LF, Smith GE, Glenn GM. A recombinant viruslike particle influenza A (H7N9) vaccine. *N Engl J Med.* 2013; 369(26):2564–6. [PubMed: 24224560]

**Key points**

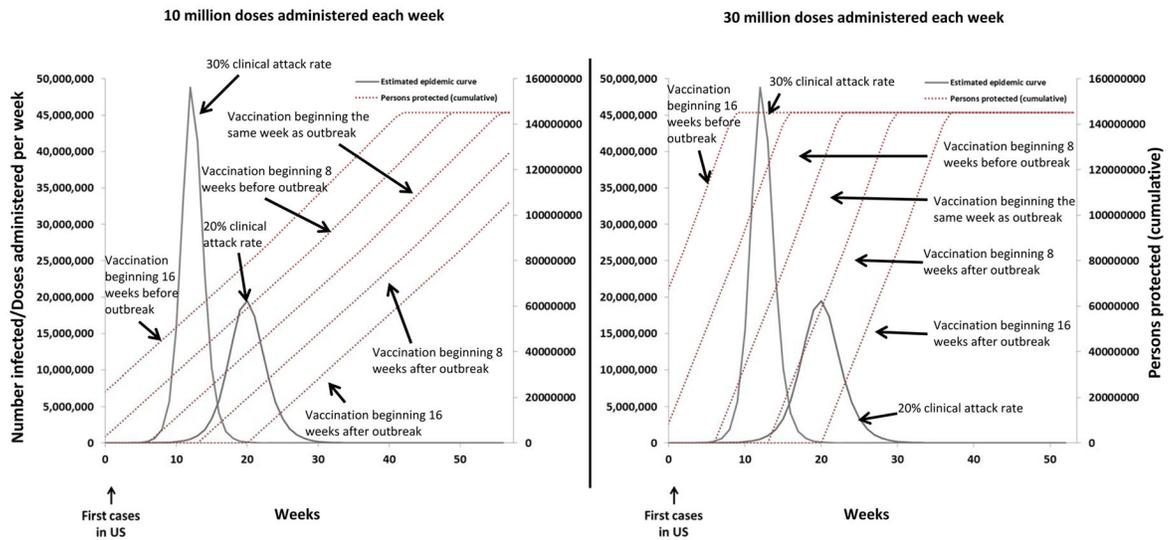
After comparing the impact of factors that could affect a vaccination program, we found that the early administration of a vaccine with a high vaccine effectiveness and a high rate of dose administration would prevent the most hospitalizations and deaths.

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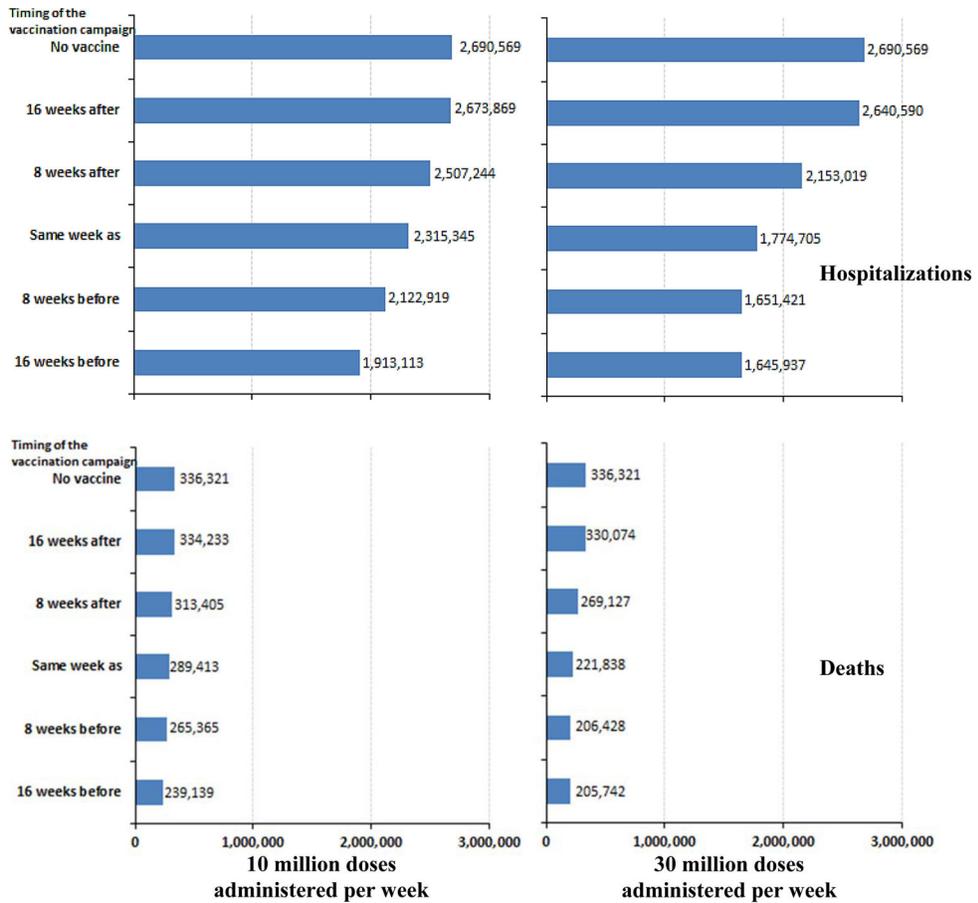
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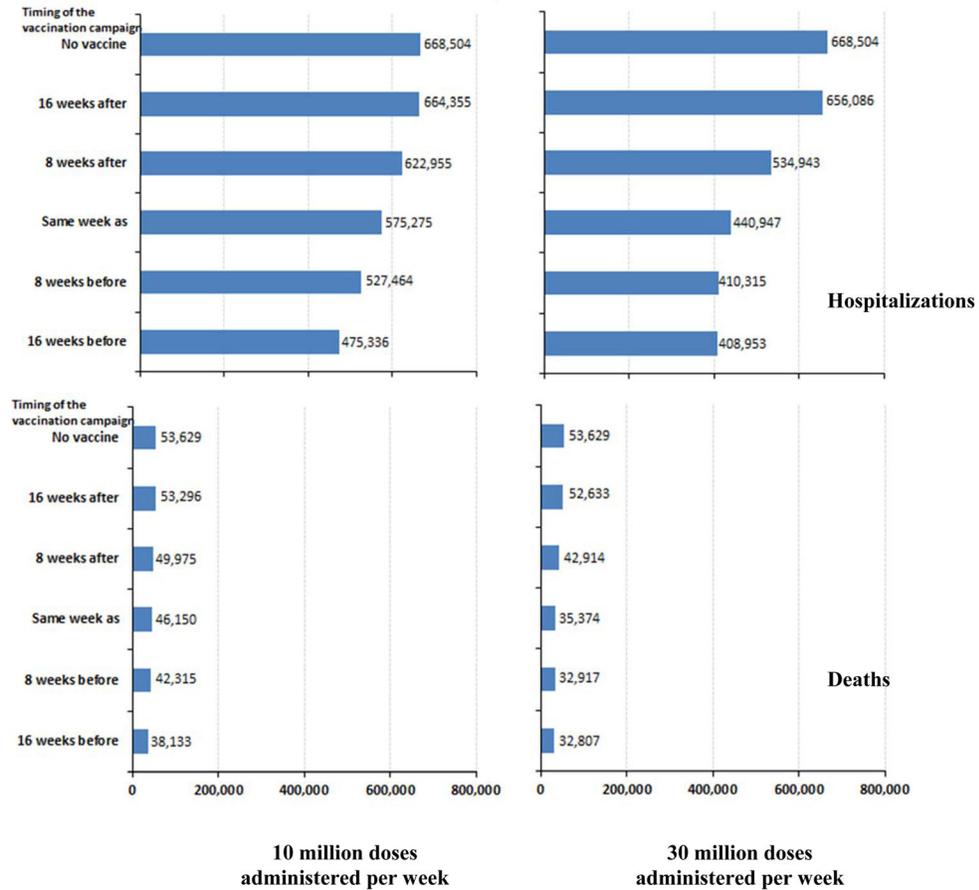
**Figure 1.**

The estimated epidemic curve without vaccination and the cumulative number of persons protected by an influenza vaccination program with the following assumptions: An overall clinical attack rate of the influenza pandemic of 20% or 30%; administered 10 million (left) or 30 million (right) vaccine doses; vaccination programs that begin 8 or 16 weeks before, the same week, or 8 or 16 weeks after the first cases of a novel influenza virus occur in the United States; and the vaccine effectiveness equivalent to the H1N1pmd09-monovalent-vaccine\*.

\*2009 H1N1-like vaccine effectiveness: Two doses of vaccine administered three weeks apart required to be fully effective (62% for persons <60 years old and 43% for persons ≥ 60) in protecting against subclinical and clinical cases, hospitalizations, and deaths. We assumed one dose of vaccine to be 0% effective for all age groups.



**Figure 2.** Number of hospitalizations (top) and deaths (bottom) if the overall clinical attack rate of the influenza pandemic is 20% and the overall case fatality ratio is 0.53% (high severity scenario), 10 million doses (left) or 30 million doses (right) of vaccine are administered each week, the vaccination program begins 16 weeks after, 8 weeks after, the same week as, 8 weeks before, and 16 weeks before the first cases of a novel influenza virus occur in the United States, and the efficacy is “H1N1pmd09-monovalent-vaccine”-like\*.  
 \*2009 H1N1-like” vaccine effectiveness: Two doses of vaccine administered three weeks apart required to be fully effective (62% for persons <60 years old and 43% for persons ≥ 60) in protecting against subclinical and clinical cases, hospitalizations, and deaths. We assumed one dose of vaccine to be 0% effective for all age groups.



**Figure 3.** Number of hospitalizations (top) and deaths (bottom) if the overall clinical attack rate of the influenza pandemic is 20% and the overall case fatality ratio is 0.084% (low severity scenario), 10 million doses (left) or 30 million doses (right) of vaccine are administered each week, the vaccination program begins 16 weeks after, 8 weeks after, the same week as, 8 weeks before, and 16 weeks before the first cases of a novel influenza virus occur in the United States, and the efficacy is “H1N1pmd09-monovalent-vaccine”-like\*. \*2009 H1N1-like” vaccine effectiveness: Two doses of vaccine administered three weeks apart required to be fully effective (62% for persons <60 years old and 43% for persons 60) in protecting against subclinical and clinical cases, hospitalizations, and deaths. We assumed one dose of vaccine to be 0% effective for all age groups.

**Table 1**

List of input values used to calculate the number of hospitalizations and deaths prevented due to a national vaccination program against an influenza pandemic, United States.

Input	Value		Reference
Number of initial cases	100		[11]
Cumulative attack rate, %	20 or 30		[11]
Case hospitalization ratio, %	Low	High	[11, 20]
All ages	1.05	4.2	
0–19 years	0.15	0.6	
20–59 years	0.45	1.8	
60+ years	6.96	28	
Case fatality ratio, %	Low	High	[11, 20]
All ages	0.084	0.53	
0–19 years	0.012	0.075	
20–59 years	0.036	0.225	
60+ years	0.558	3.5	
Vaccine Effectiveness, %			[10]
Monovalent H1N1pdm09-like			
1 <sup>st</sup> dose	All ages: 0		
2 <sup>nd</sup> dose	<60 years: 62 60 years: 43		Assumption
High vaccine effectiveness			
1 <sup>st</sup> dose by age group	<60 years: 40 60 years: 30		
2 <sup>nd</sup> dose by age group	<60 years: 80 60 years: 60		[16]
Delay in protection against the virus after the administration of the vaccine doses	2 weeks		
Cumulative vaccine coverage, all ages, %	80		Assumption
Start of vaccination program relative to the first cases of pandemic influenza in the United States	16 weeks before 8 weeks before The same week 8 weeks after 16 weeks after		Assumption
Vaccination administration, in millions per week*	10; 30		[17]; Assumption
Distribution of available doses by age group, %**	6 months–9 years: 20 10–19 years: 15 20–59 years: 35 60 years: 30		Assumption

\* We prioritized persons returning for their second doses of vaccine over persons who were receiving their first doses.

\*\* Once 80% of an age-group was fully vaccinated, we assumed vaccination would end in that age group. Remaining vaccine would then be allocated to other age groups until they all reached 80% coverage.

**Table 2**

Number and percent of hospitalizations averted relative to the number of hospitalizations without vaccination, by number of vaccine doses administered per week (10 million or 30 million), cumulative clinical attack rate (20% vs. 30%), clinical severity (high vs. low), start of vaccination relative to cases (from 16 weeks before to 16 weeks after), and vaccine efficacy (moderate vs. high).

Timing of the vaccination program relative to the start of cases in the U.S.	Number of hospitalizations averted in high <sup>d</sup> (H) or low <sup>e</sup> (L) severity scenarios															
	% hospitalizations averted relative to no vaccine						30 million doses administered per week									
	10 million doses administered per week		20%		30%		20%		30%		20%					
	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>c</sup>	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>c</sup>	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>c</sup>	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>c</sup>	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>c</sup>	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>c</sup>				
16 weeks before	777,456 (H) 193,168 (L)	1,105,624 (H) 274,706 (L)	924,056 (H) 229,153 (L)	1,362,815 (H) 338,608 (L)	1,044,632 (H) 259,551 (L)	1,417,695 (H) 352,243 (L)	1,656,461 (H) 411,568 (L)	2,252,778 (H) 559,730 (L)	29%	41%	21%	32%	39%	53%	38%	52%
8 weeks before	567,650 (H) 141,040 (L)	836,312 (H) 207,792 (L)	569,629 (H) 141,531 (L)	875,376 (H) 217,498 (L)	1,039,148 (H) 258,189 (L)	1,413,622 (H) 351,231 (L)	1,441,707 (H) 358,209 (L)	2,066,515 (H) 513,451 (L)	21%	31%	13%	20%	39%	53%	33%	48%
Same week as	375,224 (H) 93,229 (L)	572,394 (H) 142,218 (L)	259,566 (H) 64,492 (L)	448,871 (H) 111,527 (L)	915,864 (H) 227,557 (L)	1,303,629 (H) 323,902 (L)	776,967 (H) 193,047 (L)	1,326,099 (H) 329,485 (L)	14%	21%	6.0%	10%	34%	48%	18%	31%
8 weeks after	183,325 (H) 45,549 (L)	308,448 (H) 76,638 (L)	12,132 (H) 3014 (L)	68,936 (H) 17,128 (L)	537,550 (H) 133,561 (L)	877,110 (H) 217,929 (L)	36,394 (H) 9,043 (L)	206,782 (H) 51,377 (L)	6.8%	11%	0.3%	1.6%	20%	33%	0.8%	4.8%
16 weeks after	16,700 (H) 4,149 (L)	52,206 (H) 12,971 (L)	5 (H) 1 (L)	79 (H) 20 (L)	49,979 (H) 12,418 (L)	156,030 (H) 38,768 (L)	16 (H) 4 (L)	236 (H) 59 (L)	0.6%	1.9%	0%	0%	1.9%	5.8%	0%	0%

<sup>a</sup> Overall case hospitalization ratio is 4.23% (high) or 1.05% (low).

<sup>b</sup> Two doses of vaccine administered three weeks apart required to be fully effective (62% for persons <60 years old and 43% for persons 60) in protecting against subclinical and clinical cases, hospitalizations, and deaths. We assumed one dose of vaccine to be 0% effective for all age groups.

<sup>c</sup> Two doses of vaccine administered three weeks apart required to be to fully effective (80% for persons <60 years old and 60% for persons 60) in protecting against subclinical and clinical cases, hospitalizations, and deaths. We assumed one dose of vaccine is 40% effective for persons <60 years old and 30% for persons 60s.

Number and percent of deaths averted relative to the number of hospitalizations without vaccination, by number of vaccine doses administered per week (10 million or 30 million), cumulative clinical attack rate (20% vs. 30%), clinical severity (high vs. low), start of vaccination relative to cases (from 16 weeks before to 16 weeks after), and vaccine efficacy (moderate vs. high).

**Table 3**

Timing of the vaccination program relative to the start of cases in the U.S.	Number of deaths averted in high <sup>a</sup> (H) or low <sup>a</sup> (L) severity scenarios										
	% deaths averted relative to no vaccine					30 million doses administered per week					
	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>**</sup>	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>**</sup>	2009 H1N1-like efficacy <sup>b</sup>	Cumulative clinical attack rate of 20%	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>**</sup>	2009 H1N1-like efficacy <sup>b</sup>	High efficacy <sup>**</sup>	Cumulative clinical attack rate of 30%
16 weeks before	97,182 (H)	138,203 (H)	115,507 (H)	170,352 (H)	130,579 (H)	177,212 (H)	207,058 (H)	281,597 (H)	207,058 (H)	33,017 (L)	44,902 (L)
	15,496 (L)	22,037 (L)	18,418 (L)	27,164 (L)	20,822 (L)	28,258 (L)	33,017 (L)	44,902 (L)	33,017 (L)	38%	52%
8 weeks before	70,956 (H)	104,539 (H)	71,204 (H)	109,422 (H)	129,893 (H)	176,703 (H)	180,213 (H)	258,314 (H)	180,213 (H)	28,736 (L)	41,190 (L)
	11,314 (L)	16,669 (L)	11,354 (L)	17,448 (L)	20,712 (L)	28,176 (L)	28,736 (L)	41,190 (L)	28,736 (L)	33%	48%
Same week as	46,903 (H)	71,549 (H)	32,446 (H)	56,109 (H)	114,483 (H)	162,954 (H)	97,121 (H)	165,762 (H)	97,121 (H)	15,487 (L)	26,432 (L)
	7,479 (L)	11,409 (L)	5,174 (L)	8,947 (L)	18,255 (L)	25,984 (L)	15,487 (L)	26,432 (L)	15,487 (L)	18%	31%
8 weeks after	22,916 (H)	38,556 (H)	1,516 (H)	8,617 (H)	67,194 (H)	109,639 (H)	4,549 (H)	25,848 (H)	4,549 (H)	725 (L)	4,122 (L)
	3,654 (L)	6,148 (L)	242 (L)	1,374 (L)	10,715 (L)	17,483 (L)	725 (L)	4,122 (L)	725 (L)	0.8%	4.8%
16 weeks after	2,088 (H)	6,526 (H)	1 (H)	10 (H)	6,247 (H)	19,504 (H)	2 (H)	30 (H)	6,247 (H)	0 (L)	5 (L)
	333 (L)	1,041 (L)	0 (L)	2 (L)	996 (L)	3,110 (L)	0 (L)	5 (L)	996 (L)	0%	0%
	0.6%	1.9%	0%	0%	1.9%	5.8%	0%	0%	5.8%	0%	0%

<sup>a</sup> Overall case hospitalization ratio is 0.53% (high) or 0.084% (low).

<sup>b</sup> Two doses of vaccine administered three weeks apart required to be fully effective (62% for persons <60 years old and 43% for persons 60) in protecting against subclinical and clinical cases, hospitalizations, and deaths. We assumed one dose of vaccine to be 0% effective for all age groups.

<sup>c</sup> Two doses of vaccine administered three weeks apart required to be fully effective (80% for persons <60 years old and 60% for persons 60) in protecting against subclinical and clinical cases, hospitalizations, and deaths. We assumed one dose of vaccine is 40% effective for persons <60 years old and 30% for persons 60s.