

Impact of Repeated Vaccination on Vaccine Effectiveness Against Influenza A(H3N2) and B During 8 Seasons

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Background. Recent studies suggest that influenza vaccination in the previous season may influence the effectiveness of current-season vaccination, but this has not been assessed in a single population over multiple years.

Methods. Patients presenting with acute respiratory illness were prospectively enrolled during the 2004–2005 through 2012–2013 influenza seasons. Respiratory swabs were tested for influenza and vaccination dates obtained from a validated registry. Vaccination status was determined for the current, previous, and prior 5 seasons. Vaccine effectiveness (VE) was calculated for participants aged ≥ 9 years using logistic regression models with an interaction term for vaccination history.

Results. There were 7315 enrollments during 8 seasons; 1056 (14%) and 650 (9%) were positive for influenza A (H3N2) and B, respectively. Vaccination during current only, previous only, or both seasons yielded similar protection against H3N2 (adjusted VE range, 31%–36%) and B (52%–66%). In the analysis using 5 years of historical vaccination data, current season VE against H3N2 was significantly higher among vaccinated individuals with no prior vaccination history (65%; 95% confidence interval [CI], 36%–80%) compared with vaccinated individuals with a frequent vaccination history (24%; 95% CI, 3%–41%; $P = .01$). VE against B was 75% (95% CI, 50%–87%) and 48% (95% CI, 29%–62%), respectively ($P = .05$). Similar findings were observed when analysis was restricted to adults 18–49 years.

Conclusions. Current- and previous-season vaccination generated similar levels of protection, and vaccine-induced protection was greatest for individuals not vaccinated during the prior 5 years. Additional studies are needed to understand the long-term effects of annual vaccination.

Keywords. influenza; vaccine effectiveness.

The Advisory Committee on Immunization Practices (ACIP) has recommended annual influenza vaccinations for all persons aged ≥ 6 months since 2010 [1].

Although annual vaccination has been recommended for adults aged ≥ 65 years and certain high-risk groups for decades, the impact of prior vaccination history on current-season vaccine effectiveness (VE) is unclear. Despite previous debates on the relevance of repeated vaccination [2, 3], few influenza VE studies have considered the effect of vaccinations received in prior seasons. In the 1980s, a multiseason randomized, placebo-controlled trial among healthy adults found no consistent differences in efficacy of primary vs repeated vaccination against serologically confirmed influenza [4, 5]. Recent observational studies have suggested that VE may be influenced by prior season vaccination [6–12]. However, these studies were limited to 1 or 2 seasons,

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historical vaccination data were limited, confounders related to repeated vaccination behavior may have gone unmeasured, and findings were inconsistent.

Given the universal recommendation for annual influenza re-vaccination in the United States, a better understanding of the relationship between prior vaccination history and current season VE is needed. Annual VE studies have been conducted in a Wisconsin community cohort since 2004–2005, and 5 years of vaccination history is available for most study participants. This provided an opportunity to explore the effects of prior vaccination during multiple seasons and to evaluate the relationship between propensity to be vaccinated and current-season vaccine effectiveness.

METHODS

Study Population and Enrollment

From 2004–2005 through 2012–2013, observational studies of influenza VE were conducted in a defined community cohort using consistent enrollment and laboratory methods [7, 13–17]. Community-dwelling residents of a 14-zip-code area around Marshfield, Wisconsin, with ≥ 12 months of continuous residency were eligible for enrollment each season. From 2004–2005 through 2006–2007, the cohort was restricted to individuals for whom vaccination was recommended based on age or the presence of a high-risk medical condition as recommended by ACIP [18–20]. In 2007–2008 and all subsequent seasons, the cohort was expanded to include all individuals aged ≥ 6 months living in the community.

During each influenza season, eligible individuals were actively recruited during a clinical encounter for acute respiratory illness (ARI). Participants completed an interview to assess illness onset and symptoms. Sample collection methods varied by season, but either a nasal, nasopharyngeal, or a combined shallow nasal swab and oropharyngeal swab was obtained from all participants for influenza testing.

Swab samples were tested for influenza (type and subtype) using real-time reverse transcription polymerase chain reaction and viral cultures at the Marshfield Clinic Research Foundation as previously described [13]. Culture alone was performed on samples collected in 2004–2005.

We restricted this analysis to enrolled persons aged ≥ 9 years at the beginning of each season (1 August). We excluded children aged 6 months through 8 years because the criteria to be considered fully vaccinated varied during the study period and were based on prior vaccination history. Most individuals were enrolled no more than once per season. For individuals with multiple enrollments in 1 season for distinct illness episodes, the analysis included either the first enrollment (if all were negative) or the first enrollment associated with a positive influenza test; 94.3% of all enrollment visits were included based on these criteria.

Study procedures were approved by the Marshfield Clinic Institutional Review Board. Informed consent was obtained from all adults and parents/guardians of children.

Vaccination History

Influenza vaccination history was obtained from a real-time, Internet-based immunization registry used by all vaccination providers serving the population (<http://www.recin.org>). Validation studies during 2006–2007 and 2007–2008 [21], and 2011–2012 (unpublished) seasons demonstrated that the registry captured 95% of all influenza vaccinations received by study participants.

Vaccination status was determined for the current and previous seasons. Persons were considered vaccinated during the current season if they had received seasonal vaccine ≥ 14 days prior to illness onset. Previous season vaccination status was determined based on vaccination received 1 season prior (August–July). Five-year vaccination history was determined for individuals with ≥ 5 years of continuous residency in the community prior to study enrollment. Five-year vaccination history (excluding the current season) was classified into 3 categories: frequent vaccinees (vaccinated in 4–5 of the past 5 seasons), infrequent vaccinees (vaccinated in 1–3 of the past 5 seasons), and nonvaccinees (no vaccinations in the past 5 seasons). Persons who received live attenuated influenza vaccine were excluded.

Estimation of Vaccine Effectiveness

Vaccine effectiveness was assessed using a test-negative case-control study design [22–25]. Cases were persons with medically attended ARI that was laboratory confirmed as influenza. Controls were persons with ARI who tested negative for influenza. Two models were considered. The first model included exposure variables for current-season vaccination and previous-season vaccination, and an interaction term for current- and previous-season vaccination [6]. Using this model, VE was estimated for all combinations of vaccine exposure in the current and previous season: (1) vaccinated in both the current and previous season, (2) vaccinated in the current season only, (3) vaccinated in the previous season only, and (4) not vaccinated in either current or previous season (reference group).

The second model included exposure variables for current-season vaccination and 5-year vaccination history, and an interaction term for current-season vaccination and 5-year vaccination history. Using this model, VE was estimated for all combinations of vaccine exposure in the current season and 5-year vaccination history: (1) current-season vaccination and frequent vaccinee, (2) current-season vaccination and infrequent vaccinee, (3) current-season vaccination and nonvaccinee, (4) no current-season vaccination and frequent vaccinee, (5) no current-season vaccination and infrequent vaccinee, and (6) no current-season vaccination and nonvaccinee (reference group).

Table 1. Characteristics of Enrolled Participants by Case (Influenza A[H3N2] and B Virus Infection) Status, Age ≥ 9 years

Characteristic	A(H3N2) ^a			B ^b		
	H3N2 Cases (n = 1056) No. (%)	Influenza-Negative Controls (n = 3238) No. (%)	P Value ^c	B Cases (n = 650) No. (%)	Influenza-Negative Controls (n = 4293) No. (%)	P Value ^c
Influenza season			<.001			<.001
2004–2005	108 (10)	314 (10)		8 (1)	314 (7)	
2005–2006	17 (2)	181 (6)		14 (2)	181 (4)	
2006–2007	16 (2)	280 (9)		5 (1)	280 (7)	
2007–2008	469 (44)	619 (19)		169 (26)	619 (14)	
2008–2009	0	0		238 (37)	1055 (25)	
2010–2011	58 (5)	847 (26)		13 (2)	847 (20)	
2011–2012	100 (9)	501 (15)		12 (2)	501 (12)	
2012–2013	288 (27)	496 (15)		191 (29)	496 (12)	
Age			.2			<.001
9–19 y	210 (20)	664 (21)		286 (44)	979 (23)	
20–49 y	445 (42)	1246 (38)		232 (36)	1717 (40)	
50–59 y	175 (17)	524 (16)		61 (9)	657 (15)	
60–69 y	96 (9)	374 (12)		46 (7)	429 (10)	
70–79 y	84 (8)	269 (8)		21 (3)	325 (8)	
≥ 80 y	46 (4)	161 (5)		4 (1)	186 (4)	
Male sex	463 (44)	1217 (38)	<.001	315 (48)	1605 (37)	<.001
High-risk condition ^d						
Any	410 (39)	1522 (47)	<.001	148 (23)	1831 (43)	<.001
Cardiovascular	153 (14)	576 (18)	.01	40 (6)	684 (16)	<.001
Diabetes	94 (9)	385 (12)	.007	35 (5)	455 (11)	<.001
Pulmonary	164 (16)	715 (22)	<.001	67 (10)	858 (20)	<.001
Other high-risk	176 (17)	670 (21)	.004	61 (9)	804 (19)	<.001
Interval from onset to enrollment			<.001			<.001
<3 d	493 (47)	1086 (34)		233 (36)	1491 (35)	
3–4 d	390 (37)	1126 (35)		284 (44)	1488 (35)	
5–7 d	173 (16)	1026 (32)		133 (20)	1314 (31)	
Influenza diagnosis ^e						
Previous season	8 (1)	77 (2)	.001	6 (1)	134 (3)	.002
Prior 5 seasons	50 (5)	288 (9)	<.001	37 (6)	395 (9)	.003

^a Excludes 2008–2009 and 2009–2010 seasons because H3N2 viruses were not identified in the study population.

^b Excludes 2009–2010 season because type B viruses were not identified in the study population.

^c P values were calculated with χ^2 test.

^d Presence of ≥ 1 medical record–documented high-risk code in prior year, as defined by the Advisory Committee on Immunization Practices guidance for conditions that increase risk for complications from influenza.

^e Presence of *International Classification of Diseases, Ninth Revision* code for influenza infection in medical record.

Vaccine effectiveness was estimated as $100\% \times (1 - \text{odds ratio})$ using logistic regression models for each level of interaction. Like other VE studies using the test-negative approach [13, 14, 16], we adjusted for age (using a cubic B-spline with 3 equally spaced knots), high-risk medical conditions, interval from illness onset to specimen collection (<3 days, 3–4 days, and 5–7 days), season, and sex. In addition, any influenza diagnosis documented in the medical record in the previous season and prior 5 seasons was included in the adjusted model for examining the previous-season and 5-year vaccination history, respectively.

High-risk conditions were classified into the following groups: cardiovascular, diabetes, pulmonary, and other. The latter included cancer, kidney disease, liver disease, blood disorders, immunosuppressive disorders, metabolic disorders, and neurological/musculoskeletal disorders (list of *International Classification of Diseases* codes available on request).

We estimated VE separately for influenza A(H3N2) and B infections. Only seasons with H3N2 (2004–2005 through 2007–2008, and 2010–2011 through 2012–2013) or B infections (2004–2005 through 2008–2009 and 2010–2011 through

Table 2. Characteristics of Enrolled Participants by Vaccination History, Age ≥ 9 Years

Characteristic	Current- and Previous-Season Vaccination Status				Current-Season Vaccination Status and 5-Year Vaccination History ^a					
	Vaccinated in Both Current and Previous Season	Vaccinated in Current Season Only	Vaccinated in Previous Season Only	Not Vaccinated in Either Current or Previous Season	Current-Season Vaccination and Frequent Vaccinee	Current-Season Vaccination and Infrequent Vaccinee	Current-Season Vaccination and Nonvaccinee	No Current-Season Vaccination and Frequent Vaccinee	No Current-Season Vaccination and Infrequent Vaccinee	No Current-Season Vaccination and Nonvaccinee
Age										
9–19 y	308 (17)	178 (10)	149 (8)	1211 (66)	151 (10)	169 (11)	76 (5)	42 (3)	312 (21)	758 (50)
20–49 y	759 (26)	326 (11)	292 (10)	1557 (53)	347 (17)	327 (16)	95 (5)	69 (3)	344 (17)	813 (41)
50–59 y	486 (45)	100 (9)	98 (9)	392 (36)	317 (33)	180 (19)	24 (2)	51 (5)	121 (13)	271 (28)
60–69 y	392 (58)	63 (9)	61 (9)	160 (24)	285 (46)	116 (19)	14 (2)	29 (5)	67 (11)	110 (18)
70–79 y	381 (76)	31 (6)	30 (6)	57 (11)	318 (69)	56 (12)	4 (1)	21 (5)	23 (5)	36 (8)
≥ 80 y	217 (76)	19 (7)	17 (6)	31 (11)	190 (75)	20 (8)	2 (1)	14 (6)	9 (4)	19 (7)
Sex										
Female	1666 (38)	470 (11)	434 (10)	1855 (42)	1055 (31)	552 (16)	128 (4)	141 (4)	523 (15)	1039 (30)
Male	877 (30)	247 (9)	213 (7)	1553 (54)	553 (23)	316 (13)	87 (4)	85 (4)	353 (15)	968 (41)
High-risk condition^b										
Any	1519 (53)	296 (10)	268 (9)	760 (27)	1058 (44)	427 (18)	70 (3)	128 (5)	280 (12)	427 (18)
Cardiovascular	635 (61)	93 (9)	86 (8)	233 (22)	471 (52)	146 (16)	25 (3)	50 (6)	75 (8)	139 (15)
Diabetes	463 (67)	64 (9)	65 (9)	102 (15)	324 (54)	118 (20)	13 (2)	38 (6)	50 (8)	57 (10)
Pulmonary	676 (52)	143 (11)	118 (9)	355 (27)	444 (42)	205 (19)	32 (3)	54 (5)	134 (13)	190 (18)
Other	743 (59)	126 (10)	106 (8)	277 (22)	532 (50)	187 (18)	26 (2)	54 (5)	99 (9)	166 (16)
None	1024 (23)	421 (9)	379 (8)	2648 (59)	550 (16)	441 (13)	145 (4)	98 (3)	596 (17)	1580 (46)
Influenza diagnosis in previous season^c										
Yes	46 (26)	25 (14)	19 (11)	85 (49)	27 (23)	16 (14)	7 (6)	5 (4)	20 (17)	40 (35)
No	2497 (35)	692 (10)	628 (9)	3323 (47)	1581 (28)	852 (15)	208 (4)	221 (4)	856 (15)	1967 (35)
Influenza diagnosis in prior 5 seasons^c										
Yes	207 (35)	75 (13)	57 (10)	252 (43)	125 (28)	79 (18)	23 (5)	18 (4)	71 (16)	127 (29)
No	2336 (35)	642 (10)	590 (9)	3156 (47)	1483 (28)	789 (15)	192 (4)	208 (4)	805 (15)	1880 (35)
Received monovalent pandemic vaccine^d										
Yes	696 (63)	120 (11)	132 (12)	155 (14)	482 (53)	182 (20)	25 (3)	59 (6)	95 (10)	72 (8)
No	655 (24)	254 (9)	253 (9)	1560 (57)	388 (18)	278 (13)	86 (4)	83 (4)	402 (19)	932 (43)

Data are presented as No. (row %).

^a Among persons with ≥ 5 years of continuous residency.

^b Presence of ≥ 1 medical record–documented high-risk code in prior year, as defined by the Advisory Committee on Immunization Practices guidance for conditions that increase risk for complications from influenza.

^c Presence of *International Classification of Diseases, Ninth Revision* code for influenza infection in medical record.

^d Among participants enrolled during or after the 2009–2010 season.

2012–2013) in the community were included. We separately estimated VE against H3N2 for each of the 2 seasons (2007–2008 and 2012–2013) that accounted for the majority of H3N2 cases over the study period. The H3N2 component of the vaccine in 2007–2008 was the same as the previous season (2006–2007). In contrast, the 2012–2013 vaccine included a new H3N2 strain compared with the previous year. There were too few cases of seasonal H1N1 infection during the study period to estimate VE for H1N1. The 2009–2010 season was excluded because monovalent vaccine was not available before the local pandemic wave, and seasonal viruses were absent locally. We conducted sensitivity analyses restricting to certain age groups (9–49 years, 18–49 years, and ≥ 50 years), as repeated vaccination is common in older adults and unmeasured confounding is a particular concern [26]. We also performed an analysis restricted to participants swabbed within 4 days of illness onset because of potential misclassification bias with delayed sample collection.

Analyses were performed using SAS software (version 9.3; SAS Institute, Cary, North Carolina).

RESULTS

The analysis included 7315 persons aged ≥ 9 years with medically attended ARI during 8 influenza seasons: 1056 (14%) had influenza A(H3N2), 650 (9%) had influenza B, and 3 were coinfecting with H3N2 and B. The median annual number of cases was 79 (range, 0–469) for H3N2 and 13.5 (range, 5–238) for influenza B. The median age was 27 and 40 years, respectively, for influenza B and H3N2 cases. The age distribution

of controls was similar to that of H3N2 cases. Cases were more likely to be male, were less likely to have a high-risk condition, had a shorter interval from illness onset to sample collection, and were less likely to have had a prior influenza diagnosis compared with controls (Table 1).

Influenza vaccination history varied with age, sex, and high-risk condition (Table 2). Unvaccinated persons were more likely to be younger and male. Older adults and persons with a high-risk condition were more likely to be vaccinated during both the current and previous seasons, and they were more likely to be frequent vaccinees during the prior 5 years (Supplementary Table 1).

Vaccine Effectiveness Against Influenza A(H3N2)

VE estimates against medically attended H3N2 infection was similar for vaccination in both current and previous season, current season only, and previous season only, but confidence intervals (CIs) were wide. The adjusted VE was 31% (95% CI, 16%–43%), 30% (95% CI, 8%–47%), and 36% (95% CI, 14%–52%), respectively, for each of these groups (Figure 1).

The analysis including 5-year vaccination frequency was limited to 5800 individuals with at least 5 years of residency. Current-season VE against H3N2 was significantly higher (65%; 95% CI, 36%–80%) among nonvaccinees (ie, those vaccinated in the current season but not during the prior 5 years) compared with frequent vaccinees (24%; 95% CI, 3%–41%; $P = .01$). Current-season VE was 35% (95% CI, 13%–51%) among infrequent vaccinees.

The precision of VE estimates for the 2007–2008 and 2012–2013 seasons was lower, but we observed similar patterns

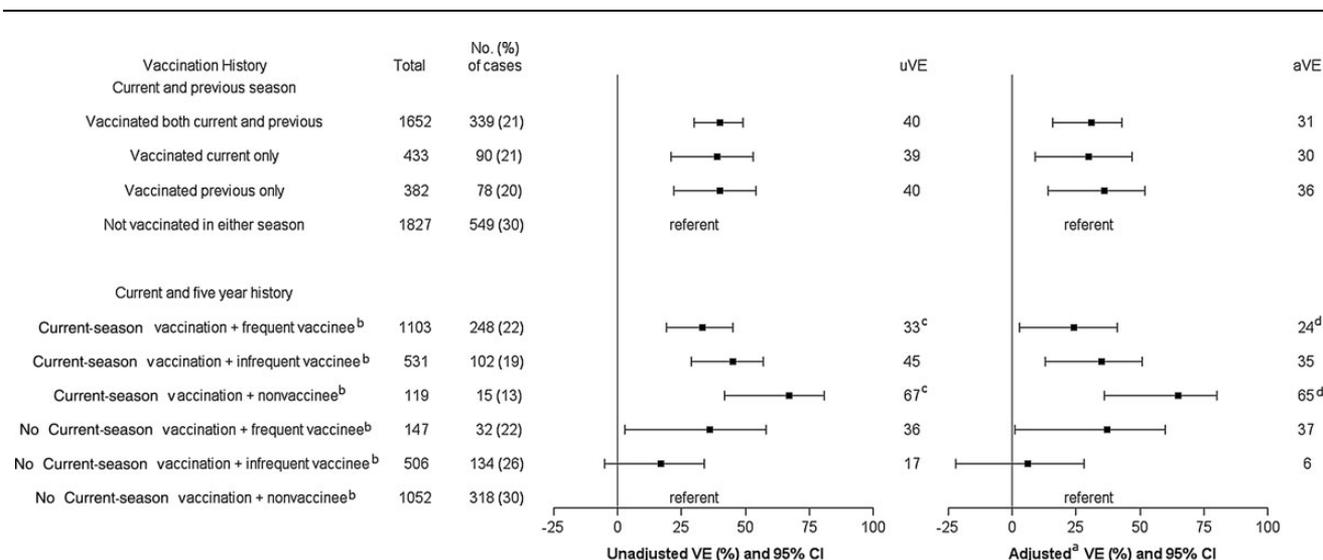


Figure 1. Unadjusted and adjusted vaccine effectiveness (VE) of various vaccination histories against influenza A(H3N2) virus infection–associated medically attended acute respiratory illness, age ≥ 9 years. ^aAdjusted for age, sex, high-risk conditions, season, interval (days) from onset to sample collection, and influenza diagnosis code in prior seasons. ^bAmong persons with ≥ 5 years of continuous residency. ^c $P = .01$ for comparison between the 2 groups. ^d $P = .01$ for comparison between the 2 groups. Abbreviations: aVE, adjusted vaccine effectiveness; CI, confidence interval; uVE, unadjusted vaccine effectiveness.

compared with the 8-season analysis (Supplementary Figures 1 and 2). VE for vaccination in both seasons (current and previous) and in the current season only was higher during 2007–2008 compared with 2012–2013, but differences between seasons were not statistically significant. Data for 5-year vaccination history were limited in some groups, but current-season VE was lowest among frequent vaccinees, consistent with overall findings, although confidence intervals were wide and included zero for some groups.

Vaccine Effectiveness Against Influenza B

For protection against medically attended influenza B infections, VE was similar for vaccination in both current and previous season, current season only, and previous season only. The adjusted VE was 52% (95% CI, 38%–62%), 66% (95% CI, 51%–77%), and 64% (95% CI, 45%–76%), respectively, for each of these groups (Figure 2). In the analysis of 5-year vaccination history, adjusted VE against B was significantly higher ($P = .05$) among nonvaccinees with current-season vaccination (75%; 95% CI, 50%–87%) compared with frequent vaccinees with current-season vaccination (48%; 95% CI, 29%–62%).

There was evidence of residual protection among frequent and infrequent vaccinees who did not receive current-season vaccine. The adjusted VE was 50% (95% CI, 10%–73%) and 27% (95% CI, 4%–45%) for these groups, respectively.

Sensitivity Analyses

VE estimates for H3N2 and B were similar when the analysis was restricted to individuals aged 9–49 years and 18–49 years, but

sample size was smaller and VE differences between groups were not significant (Tables 3 and 4). VE estimates were also similar when restricting to participants who were swabbed within 4 days of illness onset. The absolute VE difference was $\leq 8\%$, with a few exceptions. However, CIs were wide (data not shown).

DISCUSSION

With our longitudinal influenza vaccination records, we were able to examine the independent and combined effectiveness of current-season vaccination and prior vaccination history over multiple seasons. Current-season vaccination was effective against medically attended H3N2 and B virus-associated illness regardless of prior vaccination history.

The observed protection among persons vaccinated during the previous season and not the current season suggests that influenza vaccines received >1 year earlier may provide some residual protection. A case-control study of influenza VE in pregnant women during the 2010–2011 and 2011–2012 seasons found patterns similar to our study with evidence of residual protection from previous-season vaccination [10]. VE results from the 2010–2011 season in Canada were also consistent with residual protection against A(H1N1)pdm09 among individuals who were previously vaccinated with an adjuvanted monovalent vaccine [9]. In our cohort, residual protection against H3N2 viruses occurred despite the fact that the predominant H3N2 viruses were antigenically distinct from the previous season H3N2 vaccine component in 5 of 7 seasons that were included in the H3N2 analysis. Similarly, the predominant

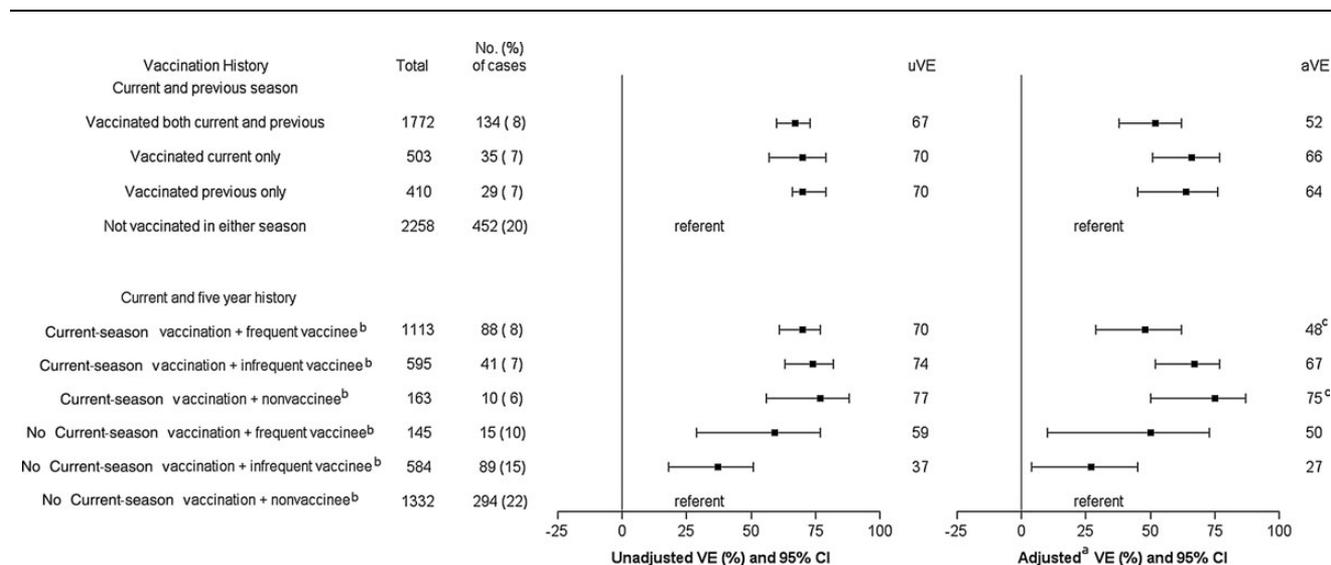


Figure 2. Unadjusted and adjusted vaccine effectiveness (VE) of various vaccination histories against influenza B virus infection–associated medically attended acute respiratory illness, age ≥ 9 years. ^aAdjusted for age, sex, high-risk conditions, season, interval (days) from onset to sample collection, and influenza diagnosis code in prior seasons. ^bAmong persons with ≥ 5 years of continuous residency. ^c $P = .05$ for comparison between the 2 groups. Abbreviations: aVE, adjusted vaccine effectiveness; CI, confidence interval; uVE, unadjusted vaccine effectiveness.

Table 3. Unadjusted and Adjusted Vaccine Effectiveness Estimates Against Influenza A(H3N2) Virus Infection–Associated Medically Attended Acute Respiratory Illness Showing Effect of Individual Covariates

Vaccination Status	Current and Previous Season			Current-Season Vaccination Status and 5-Year Vaccination History ^a				
	Vaccinated Both	Vaccinated Current Season Only	Vaccinated Previous Season Only	Current-Season Vaccination + Frequent Vaccinee	Current-Season Vaccination + Infrequent Vaccinee	Current-Season Vaccination + Nonvaccinee	No Current-Season Vaccination + Frequent Vaccinee	No Current-Season Vaccination + Infrequent Vaccinee
Age ≥9 y								
Unadjusted	40 (30–49)	39 (21–53)	40 (22–54)	33 (19–45)	45 (29–57)	67 (42–81)	36 (3–58)	17 (–5 to 34)
Adjusted for age	42 (31–51)	40 (23–53)	41 (23–55)	34 (18–47)	46 (30–58)	67 (42–81)	37 (4–58)	17 (–6 to 34)
Adjusted for sex	39 (28–47)	38 (20–52)	39 (20–53)	31 (17–43)	44 (28–57)	67 (42–81)	35 (1–57)	16 (–7 to 34)
Adjusted for season	32 (19–42)	32 (11–48)	36 (16–52)	25 (7–39)	36 (16–51)	65 (38–81)	36 (1–59)	10 (–16 to 30)
Adjusted for high-risk condition	35 (24–45)	36 (18–51)	38 (19–53)	27 (11–41)	42 (25–55)	66 (40–80)	32 (–4 to 55)	15 (–8 to 33)
Adjusted for interval from onset to sample collection	36 (25–46)	38 (20–52)	39 (20–54)	29 (13–41)	44 (27–56)	68 (44–82)	33 (–2 to 56)	15 (–8 to 33)
Adjusted for prior influenza history ^b	40 (30–49)	38 (20–52)	40 (22–54)	33 (19–45)	45 (29–57)	66 (41–81)	35 (2–57)	17 (–6 to 34)
Fully adjusted ^c	31 (16–43)	30 (8–47)	36 (14–52)	24 (3–41)	35 (13–51)	65 (36–80)	36 (1–60)	6 (–22 to 28)
Age 9–49 y								
Unadjusted	45 (31–56)	49 (29–63)	46 (25–62)	37 (14–54)	54 (34–67)	74 (47–87)	49 (4–73)	10 (–18 to 32)
Fully adjusted ^c	31 (11–47)	43 (19–61)	41 (15–59)	17 (–19 to 42)	48 (23–65)	70 (36–86)	41 (–19 to 71)	–3 (–40 to 24)
Age 18–49 y								
Unadjusted	49 (33–61)	37 (11–56)	55 (33–70)	42 (16–60)	55 (32–70)	66 (26–84)	69 (25–87)	16 (–20 to 41)
Fully adjusted ^c	38 (15–54)	37 (5–57)	53 (26–70)	26 (–15 to 52)	53 (25–70)	69 (28–87)	68 (17–88)	3 (–45 to 36)
Age ≥50 y								
Unadjusted	30 (10–46)	16 (–28 to 44)	23 (–23 to 52)	29 (4–48)	34 (3–55)	45 (–39 to 78)	22 (–38 to 55)	34 (–7 to 59)
Fully adjusted ^c	28 (1–47)	10 (–44 to 43)	29 (–20 to 57)	28 (–3 to 50)	20 (–25 to 48)	58 (–14 to 84)	37 (–19 to 66)	32 (–16 to 60)

^a Among persons with ≥5 years of continuous residency.

^b Presence of *International Classification of Diseases, Ninth Revision* code for influenza infection in medical record for the previous season in model of vaccine exposure for the current and previous season and prior 5 seasons in model of vaccine exposure for current and 5-year vaccination history.

^c Adjusted for age, sex, high-risk conditions, season, interval (days) from onset to sample collection, and influenza diagnosis code in prior seasons.

Table 4. Unadjusted and Adjusted Vaccine Effectiveness Estimates Against Influenza B Virus Infection–Associated Medically Attended Acute Respiratory Illness Showing Effect of Individual Covariates

Vaccination Status	Current and Previous Season			Current-Season Vaccination Status and 5-Year Vaccination History ^a				
	Vaccinated Both	Vaccinated Current Season Only	Vaccinated Previous Season Only	Current-Season Vaccination + Frequent Vaccinee	Current-Season Vaccination + Infrequent Vaccinee	Current-Season Vaccination + Nonvaccinee	No Current-Season Vaccination + Frequent Vaccinee	No Current-Season Vaccination + Infrequent Vaccinee
Age ≥9 y								
Unadjusted	67 (60–73)	70 (57–80)	70 (55–79)	70 (61–77)	74 (63–82)	77 (56–88)	59 (29–77)	37 (18–51)
Adjusted for age	59 (49–67)	69 (56–79)	67 (51–78)	58 (44–68)	72 (60–80)	79 (58–89)	54 (20–74)	38 (19–52)
Adjusted for sex	66 (58–73)	69 (56–78)	68 (53–79)	69 (59–76)	73 (62–81)	77 (55–88)	58 (27–76)	35 (16–50)
Adjusted for season	62 (53–69)	68 (54–78)	66 (50–78)	63 (52–72)	70 (58–79)	76 (53–88)	57 (24–76)	30 (8–47)
Adjusted for any high-risk condition	59 (50–67)	67 (53–77)	67 (50–77)	60 (48–69)	69 (57–78)	75 (53–87)	50 (13–71)	33 (13–48)
Adjusted for interval from onset to sample collection	67 (59–73)	70 (57–79)	70 (55–79)	69 (59–76)	74 (63–81)	77 (55–88)	59 (28–76)	36 (17–51)
Adjusted for prior influenza history ^b	68 (60–74)	70 (57–79)	70 (55–79)	69 (61–76)	74 (63–81)	77 (55–88)	59 (29–76)	36 (17–51)
Fully adjusted ^c	52 (38–62)	66 (51–77)	64 (45–76)	48 (29–62)	67 (53–77)	75 (50–87)	50 (10–73)	27 (4–45)
Age 9–49 y								
Unadjusted	54 (41–65)	64 (47–76)	67 (49–79)	50 (30–65)	62 (45–74)	75 (51–88)	40 (–14 to 68)	34 (12–51)
Fully adjusted ^c	42 (23–57)	61 (41–74)	60 (35–75)	27 (–9 to 51)	56 (33–71)	70 (38–86)	27 (–45 to 64)	21 (–9 to 43)
Age 18–49 y								
Unadjusted	55 (36–68)	71 (47–84)	61 (32–78)	39 (5–62)	70 (46–83)	74 (28–91)	40 (–43 to 75)	33 (–4 to 57)
Fully adjusted ^c	54 (32–69)	70 (44–84)	60 (27–78)
Age ≥50 y								
Unadjusted	71 (58–80)	85 (59–95)	72 (35–88)	74 (60–83)	89 (74–95)	86 (–4 to 98)	79 (31–94)	44 (–1 to 69)

^a Among persons with ≥5 years of continuous residency.

^b Presence of *International Classification of Diseases, Ninth Revision* code for influenza infection in medical record for the previous season in model of vaccine exposure for the current and previous season and prior 5 seasons in model of vaccine exposure for current and 5-year vaccination history.

^c Adjusted for age, sex, high-risk conditions, season, interval (days) from onset to sample collection, and influenza diagnosis code in prior seasons.

circulating B lineage differed from the previous season vaccine lineage in 5 of 7 seasons, and both lineages circulated widely in 1 season (2012–2013). This residual protection may be conferred by immune responses elicited by previous vaccines that cross-react with current-season virus antigens. Consistent with this idea, several studies have shown evidence that seasonal influenza vaccination can induce antibodies and/or T cells capable of cross-reacting with antigenically distinct viruses [27–31].

Previous epidemiologic studies on clinical effectiveness of current-season influenza vaccination among persons with and without prior history of influenza vaccination have yielded inconsistent results [2, 3]. A placebo-controlled randomized clinical trial from 1984 to 1988 among healthy adults found no consistent differences in efficacy of primary vs repeated vaccination [4, 5]. However, the trial identified few virus-confirmed influenza cases and had limited power to detect differences between groups. More importantly, serologic endpoints were used to identify influenza cases in addition to virus culture. Use of serologic endpoints can cause bias in studies of influenza VE because vaccinated individuals are less likely to seroconvert after influenza infection compared with unvaccinated subjects [32, 33]. Although randomized clinical trials are less susceptible to confounding compared with observational studies, the negative results from this trial should be interpreted with caution due to these limitations.

Vaccine interference may be occurring if protection is lower in individuals who were vaccinated in both the current and previous season compared with those vaccinated in the current season only. Our analysis did not suggest any evidence of vaccine interference when we considered previous-season vaccination. However, the analysis using 5 years of historical vaccination data suggested a significant difference in current-season VE among frequent vaccinees compared with nonvaccinees. Interference due to repeated prior vaccination is one possible explanation for this difference, but unmeasured confounding could also account for the observed differences. The potential immunologic mechanisms for vaccine interference are not well understood. Contributing factors may include “original antigenic sin,” immune exhaustion, and/or antigenic drift of circulating influenza viruses with respect to vaccine antigens. The potential role of original antigenic sin is particularly intriguing as it postulates that exposure to influenza antigens can preferentially expand preexisting memory responses to historical virus antigens at the expense of *de novo* responses to the current vaccine or infecting strain [34–36]. In most people, vaccination appears to boost preexisting memory responses against antigenically related, previously circulating strains [28, 37]. Reviewing the available evidence, authors of a commentary suggested that the potential for original antigenic sin is greatest when “sequential viruses are of intermediate antigenic relatedness; when they are antigenically complex; and when the sequential exposure

intervals are long . . .” [34]. The evidence of original antigenic sin is strongest for sequential natural influenza infections. It is unclear whether repeated influenza vaccination generates effects of similar magnitude.

This study had several limitations. Unmeasured confounding may have influenced VE estimates, as people who choose to get vaccinated every year may have different characteristics and susceptibility to influenza compared with those who do not seek vaccination every year. Although we adjusted for prior influenza using diagnosis codes, prior influenza infection history could be a source of unmeasured confounding as not all illnesses result in medical attention or documentation. However, we found similar trends for the effect of prior vaccination even when analysis was limited to younger persons who were less likely to be frequent vaccinees or to have prior infections. Also, the test-negative design has been shown to yield VE results that were virtually identical to those obtained using the gold standard clinical trial analysis in a recent reanalysis of influenza vaccine clinical trial data [25]. In addition, several other studies have indicated that it is a valid method for estimating VE under a wide range of assumptions [22–24]. Nonetheless, this study examined VE against medically attended influenza and results should not necessarily be interpreted as VE against influenza infection in general.

Despite data from 8 seasons, the sample size in some groups was limited. Furthermore, we were unable to examine VE patterns for each season to determine if patterns were consistent across all seasons. Although our findings represent aggregate effects across multiple seasons, it is apparent from the 2 largest H3N2 seasons that current-season VE is modified by prior vaccination history, irrespective of season or whether there was or was not a change in the vaccine component. We were not able to evaluate timing of past vaccinations for patients with an infrequent vaccination history. However, most patients in this group were vaccinated at least once in the most recent 2 seasons, so results for this group may reflect more recent vaccination history. Finally, this analysis was restricted to inactivated influenza vaccine, and it is unknown whether similar results would be observed with live attenuated influenza vaccine.

In conclusion, we found that vaccination provided protection against medically attended influenza infection, regardless of prior vaccination history. This is consistent with randomized clinical trials of influenza vaccine efficacy [38]. However, this study raises relevant questions about the potential interference of repeated annual influenza vaccination and possible residual protection from previous season vaccination that have not been considered in most trials. Further observational studies that include simultaneous assessment of immune response, both to vaccine antigens and previously circulating viruses, and clinical protection would be helpful. In addition, we believe the results of this study and others support the need for another

multiseason randomized clinical trial. Although a randomized trial cannot be conducted in the United States where influenza vaccination is universally recommended, it may be feasible in countries that do not currently recommend influenza vaccination for healthy adults. This would provide valuable new information about the impact of repeated annual influenza vaccination, and it will help guide future vaccine policy recommendations.

Supplementary Data

Supplementary materials are available at *Clinical Infectious Diseases* online (<http://cid.oxfordjournals.org>). Supplementary materials consist of data provided by the author that are published to benefit the reader. The posted materials are not copyedited. The contents of all supplementary data are the sole responsibility of the authors. Questions or messages regarding errors should be addressed to the author.

Notes

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