



# Impact of Rapid Molecular Diagnostic Testing of Respiratory Viruses on Outcomes of Adults Hospitalized with Respiratory Illness: a Multicenter Quasi-experimental Study

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**ABSTRACT** A standard multiplex PCR offers comprehensive testing for respiratory viruses. However, it has traditionally been performed in a referral laboratory with a lengthy turnaround time, which can reduce patient flow through the hospital. We aimed to determine whether the introduction of a rapid PCR, but with limited targets (Cepheid Xpert Flu/RSV XC), was associated with improved outcomes for adults hospitalized with respiratory illness. A controlled quasi-experimental study was conducted across three hospitals in New South Wales, Australia. Intervention groups received standard multiplex PCR during the preimplementation, July to December 2016 ( $n = 953$ ), and rapid PCR during the postimplementation, July to December 2017 ( $n = 1,209$ ). Control groups (preimplementation,  $n = 937$ , and postimplementation,  $n = 1,102$ ) were randomly selected from adults hospitalized with respiratory illness during the same periods. The outcomes were hospital length of stay (LOS) and microbiology test utilization (blood culture, urine culture, sputum culture, and respiratory bacterial and virus serologies). The introduction of rapid PCR was associated with a nonsignificant 8.9-h reduction in median LOS (95% confidence interval [CI],  $-21.5$  h to  $3.7$  h;  $P = 0.17$ ) for all patients and a significant 21.5-h reduction in median LOS (95% CI,  $-36.8$  h to  $-6.2$  h;  $P < 0.01$ ) among patients with positive test results in an adjusted difference-in-differences analysis. For patients receiving test results before disposition, rapid PCR use was associated with a significant reduction in LOS, irrespective of test results. Compared with standard PCR testing, rapid PCR use was significantly associated with fewer blood culture (adjusted odds ratio [aOR], 0.67; 95% CI, 0.5 to 0.82;  $P < 0.001$ ), sputum culture (aOR, 0.56; 95% CI, 0.47 to 0.68,  $P < 0.001$ ), bacterial serology (aOR, 0.44; 95% CI, 0.35 to 0.55,  $P < 0.001$ ) and viral serology (aOR, 0.42; 95% CI, 0.33 to 0.53,  $P < 0.001$ ) tests, but not with fewer urine culture tests (aOR, 0.94; 95% CI, 0.78 to 1.12,  $P = 0.48$ ). Rapid PCR testing of adults hospitalized with respiratory illnesses can deliver benefits to patients and reduce resource utilization. Future research should consider a formal economic analysis and assess its potential impacts on clinical decision making.

**KEYWORDS** influenza, molecular diagnostic, rapid PCR, respiratory viruses

Influenza and other respiratory viruses, such as a respiratory syncytial virus (RSV), rhinovirus, coronavirus, human metapneumovirus, and parainfluenza viruses, cause significant morbidity and mortality in Australia and worldwide (1–3). Acute respiratory viral infections are estimated to cause 5.8 million deaths worldwide (4), and each year, up to 650,000 deaths are related to respiratory diseases from seasonal influenza (5). In

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Australia, influenza-like illness is a common presentation among working-age adults and is responsible for decreased productivity and lost working days (3).

The laboratory diagnosis of respiratory viruses has evolved considerably in recent years with the development of novel molecular methods (6). Molecular methods use nucleic acid amplification techniques, such as reverse transcription PCR (PCR), and are considered effective diagnostic tests to confirm respiratory viral infections (7). A standard multiplex PCR test can detect several respiratory viruses with very high sensitivity and specificity in a single test (8). While it offers comprehensive testing for respiratory viruses, multiplex PCR requires specialized laboratory facilities and expertise (9), and it has traditionally been performed in a referral laboratory with a lengthy test turnaround time (TAT), which can reduce patient flow through the hospital.

New rapid PCR-based tests have recently been developed, including the Cepheid Xpert assays (Cepheid, Sunnyvale, CA) (10). Rapid viral diagnosis at hospital-based laboratories can improve the timeliness of care in hospital and has the potential for reducing resource utilization (11). The use of rapid PCR was introduced across hospitals in New South Wales (NSW), Australia, in July 2017 (N. Wabe, L. Li, R. Lindeman, R. Yimsung, M.R. Dahm, S. McLennan, K. Clezy, J.I Westbrook, and A. Georgiou, submitted for publication). It is an easy-to-use diagnostic test for accurate detection and differentiation of influenza A/B and RSV. Existing international evidence has shown an association between the implementation of rapid viral diagnosis and reductions in hospital length of stay (LOS) (12) and other laboratory test ordering practices (13–15). However, the impact of the introduction of a rapid PCR has not yet been investigated in Australia.

In this study, we compared outcomes of patients tested for influenza A/B and RSV using a rapid PCR (Cepheid Xpert Flu/RSV XC) at the local hospital laboratory during the first six months following its introduction (July to December 2017), with patients tested at a referral laboratory using a standard multiplex PCR (Allplex respiratory panels; Seegene) during the corresponding period in 2016. The objective was to determine whether the introduction of a rapid PCR testing for influenza A/B and RSV for adults hospitalized with respiratory illness was associated with a reduction in hospital LOS and a change in microbiology test utilization compared with those associated with standard multiplex PCR testing.

## MATERIALS AND METHODS

**Study setting.** The study was conducted across three major tertiary teaching hospitals in New South Wales, Australia, each offering a comprehensive range of inpatient and community services, hospital A (>65,000 admissions per year), hospital B (>50,000 admissions per year), and hospital C (>28,000 admissions per year). Ethics approval was granted by the Human Research Ethics Committee of the South Eastern Sydney Local Health District (approval HREC/16/POWH/412).

**Study design and population.** We conducted a controlled, nonrandomized, quasi-experimental study. A patient selection flow chart is presented in Fig. 1. The study periods were between July and December 2016 (preimplementation) and between July and December 2017 (postimplementation). The inclusion criteria were patients (aged >18 years) admitted with a respiratory illness [International Classification of Diseases version 10 Australian Modification (ICD-10-AM), J00-J99] during the study period and having at least one laboratory test result.

The intervention groups included patients tested for respiratory viruses using the standard multiplex PCR during the preimplementation (standard PCR group) and patients tested for influenza or RSV using the rapid PCR during the postimplementation (rapid PCR group). The standard PCR used in this study consisted of Allplex respiratory panels 1, 2, and 3 (Seegene Inc., Seoul, Republic of Korea) (16). It was available as a referral test at a large referral laboratory located at hospital B. All three hospitals sent samples to this laboratory to conduct a standard multiplex PCR. Standard multiplex PCR testing was generally batched. During the influenza season, it was performed 1 to 2 times a day (per demand) on weekdays and once a day on weekends. Outside the influenza season, the test was performed once every day. The multiplex PCR assay was offered as three panels that can detect up to 16 viruses with influenza A subtyping. Panel 1 includes influenza A (subtypes H1, H1pdm09, and H3), influenza B, and RSV (types A and B); panel 2 includes adenovirus, metapneumovirus, enterovirus, and parainfluenza virus (types 1, 2, 3, and 4); and panel 3 includes bocavirus (types 1, 2, 3, and 4), coronavirus (types OC43, 229E, and NL63), and rhinovirus (types A, B, and C) (16). In this study, all patients in the standard PCR group received panel 1 and, depending on whether a specific request for other viruses was made, panels 2 and or 3 were also ordered.

The rapid PCR used in this study was a Cepheid Xpert Flu/RSV XC assay (Cepheid, Sunnyvale, CA). The Cepheid Xpert Flu/RSV XC assay demonstrated a high sensitivity and specificity for detection of influenza

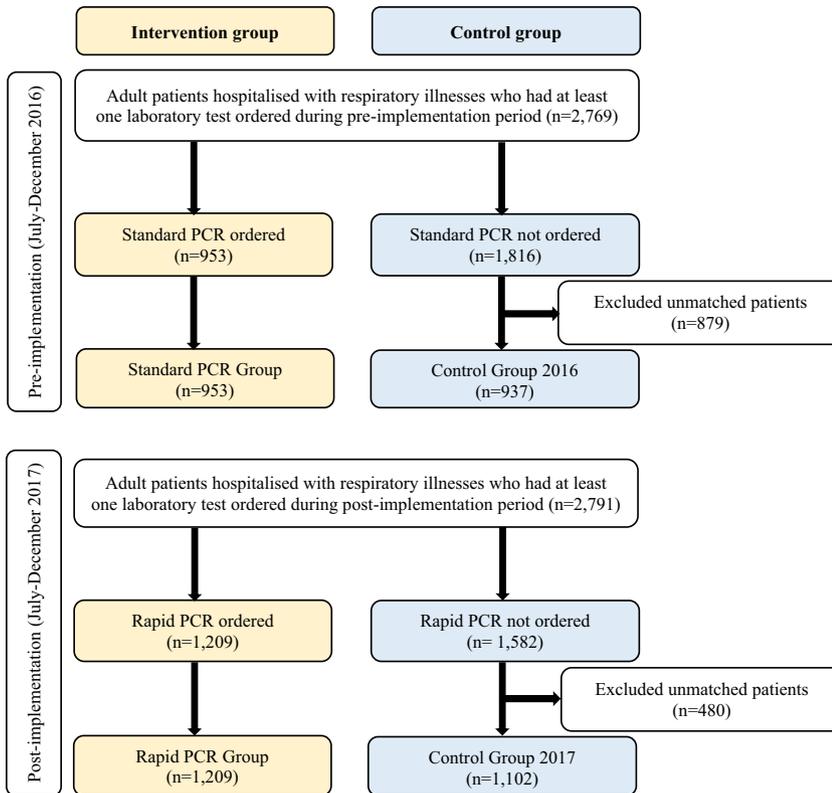


FIG 1 Patient selection flow chart.

A, influenza B, and RSV (17). However, unlike the standard multiplex PCR, Cepheid Xpert Flu/RSV XC cannot discriminate among influenza A virus subtypes. It is a hospital laboratory-based test. Hospitals A and B had an onsite laboratory to perform the rapid testing. Hospital C sent samples to hospital A because of its proximity to this hospital. During the influenza season, rapid PCR testing was available 24 h a day, 7 days a week, and outside the influenza season, the test was available on demand with approval from the pathologist or registrar on duty.

The control groups were selected from the population that fulfilled the above inclusion criteria but were not tested with standard PCR or rapid PCR. They were selected randomly from the population after matching during the respective study periods on study hospital, age quartile, mode of separation from the hospital, and gender. The purpose of matching was to balance the distribution of baseline characteristics across groups. The diagnosis of influenza among patients in the control groups was based on clinical assessment. Relevant demographic, clinical characteristics, and laboratory test data were obtained by linking the admitted patient and laboratory information system data sets. Detailed information on the linkage process has been described elsewhere (18).

**Outcome measures.** The primary outcome was hospital LOS. Hospital LOS was calculated by subtracting the admission date/time from the hospital separation date/time. As secondary outcomes, we compared utilization of commonly ordered microbiology tests, including blood culture; urine microscopy, culture, and sensitivity (urine MC&S); sputum culture; respiratory bacterial serology (e.g., *Mycoplasma pneumoniae* IgM Antibody [Ab], *Bordetella pertussis* toxin IgA Ab and *Pneumocystis jirovecii* immunofluorescence assay [IF], and *Hemophilus influenzae* B Ab), and respiratory virus serology (e.g., *Parainfluenza* Ab complement fixation test [CFT], *Adenovirus* CFT and RSV Ab CFT).

**Statistical analysis.** Descriptive statistics, including medians with interquartile ranges (IQR), were calculated. Demographic and clinical characteristics of the groups were compared using Pearson's chi-square test for categorical variables and Mann-Whitney tests for continuous variables.

The impact of a rapid PCR on hospital LOS was assessed using a median regression. As LOS data were highly positively skewed, commonly used approaches such as ordinary least-squares regression, which models the conditional mean of the outcome variable, were not appropriate (19). Median regression is robust to extreme values and therefore well suited for modeling LOS (20). A difference-in-differences (DID) analysis was conducted to estimate the effect of a rapid PCR on the LOS by comparing the median change over time in the LOS for the intervention group, compared to the median change over time for the control group.

Comparison of other microbiology test utilization (e.g., blood culture, yes/no) between intervention and control groups was performed using binary logistic regression. Analyses were adjusted for relevant demographic and clinical characteristics.

For both primary and secondary outcomes, a subgroup analysis was conducted by test results. For the primary outcome, a further subgroup analysis was conducted by limiting the analysis to patients

whose test results were returned during their stay in the hospital (i.e., excluding patients discharged before test results became available). All *P* values were 2-tailed, and a *P* value of <0.05 was considered statistically significant. Analyses were conducted using Stata version 15 (StataCorp LP, College Station, TX).

## RESULTS

**Demographic and clinical characteristics.** The intervention group included 2,162 patients (953 preimplementation [standard PCR group] and 1,209 postimplementation [rapid PCR group]). The control group consisted of 2,039 patients (937 preimplementation [control group 2016] and 1,102 postimplementation [control group 2017]) (Fig. 1). All patients in the standard PCR group received panel 1 with or without other panels; 64.1% (*n* = 611) received panel 1 only; 32.2% (*n* = 307) received all 3 panels; 2.3% (*n* = 22) received panels 1 and 3; and 1.4% (*n* = 13) received panels 1 and 2. Postimplementation, 20.2% (*n* = 244) of patients in the rapid PCR group were ordered at least one standard multiplex PCR panel in addition to the rapid PCR itself. Of the 244 patients, 40.9% (*n* = 100) received all 3 panels; 43.4% (*n* = 106) received panels 2 and 3; 12.3% (*n* = 30) received panel 1; and 3.3% (*n* = 8) received panels 1 and 3 (*n* = 3), panels 1 and 2 (*n* = 2), panel 2 (*n* = 2), or panel 3 (*n* = 1).

The distributions of patient demographic and clinical characteristics in the intervention and control groups were similar in terms of gender, age, study hospital, weekday of admission, and mode of separation from the hospital. However, the distributions of the source of referral, intensive care admission status, and principal diagnosis were different across groups (Table 1).

Influenza and pneumonia were the most common principal diagnosis in all groups. However, intervention groups had higher proportions of admissions due to influenza and pneumonia compared to those of the control groups (*P* < 0.01). Similarly, intensive care unit (ICU) admissions were higher in intervention groups than those in the control groups. Over 80% of patients were subsequently discharged home, and in-hospital death occurred in 3.6 to 7.7% of patients (Table 1).

For all intervention patients admitted to hospital through emergency departments (ED), respiratory virus testing (both standard and rapid PCR) was ordered while patients were in the ED. The tests were ordered a median of 3.8 h and 3.2 h before separation from the ED for standard PCR and rapid PCR groups, respectively. The median ED LOS of the two groups (prior to hospital admission) were roughly comparable, namely, 6.8 h for standard PCR and 7.2 h for the rapid PCR.

Twenty-two percent of patients receiving the standard PCR and 36.4% in the rapid PCR group were positive for at least one respiratory virus. The overall median TAT was significantly shorter for rapid PCR compared to standard multiplex PCR (2.3 h versus 27.4 h, *P* < 0.01). Significantly more patients were discharged from the hospital before test results became available in the standard PCR group than in the rapid PCR group (18.9% versus 2.2%; *P* < 0.01) (Table 2). Of these discharged patients, the results of 19.4% (*n* = 35) in the standard PCR and 11.1% (*n* = 3) in the rapid PCR groups eventually came back positive for at least one respiratory virus.

**Primary outcome.** Table 3 presents the results of median regression for the main and subgroup analyses. In the adjusted analyses, we controlled for multiple factors potentially affecting LOS (Table S1). For the control group, there was no significant difference in the median LOS between controls 2016 and 2017 (75.3 h versus 78.9 h, *P* = 0.68). For the intervention group, an analysis that included all patients, regardless of test result outcome, showed no significant difference in the median LOS. Similarly, for patients with negative results, there was no significant difference in the median LOS. However, for patients with positive results, the median LOS decreased significantly, by 21.2 h (95% CI, -38.0 to -4.5; *P* < 0.01) in the within-intervention-groups analysis and by 21.5 h (95% CI, -36.8 to -6.2; *P* < 0.01) in the DID analysis following the introduction of rapid PCR.

When the analysis was limited to patients whose test results were returned during their stay in the hospital, rapid PCR was associated with a significant decrease in the median LOS, irrespective of test result. Adjusted analysis showed a significant reduction

**TABLE 1** Comparison of demographic and clinical characteristics of intervention and control groups

Variable	Data for:				Intervention vs control groups ( <i>P</i> value) <sup>e</sup>	
	Intervention group		Control group		Standard PCR vs control 2016	Rapid PCR vs control 2017
	Standard PCR ( <i>n</i> = 953)	Rapid PCR ( <i>n</i> = 1,209)	2016 ( <i>n</i> = 937)	2017 ( <i>n</i> = 1,102)		
Female, <i>n</i> (%)	480 (50.4)	619 (51.2)	473 (50.5)	565 (51.3)	0.96	0.97
Age (yrs), median (IQR)	75 (62–84)	77 (65–86)	75 (62–84)	76 (64–84)	0.97	0.11
Hospital, <i>n</i> (%)					0.94	0.58
A	434 (45.5)	672 (55.6)	431 (46.0)	589 (53.5)		
B	340 (35.7)	205 (17.0)	327 (34.9)	198 (18.0)		
C	179 (18.8)	332 (27.5)	179 (19.1)	315 (28.6)		
Time of day of hospital admission, <i>n</i> (%)					<0.01	0.95
7 a.m.–7 p.m.	598 (62.8)	866 (71.6)	657 (70.1)	788 (71.5)		
7 p.m.–7 a.m.	355 (37.2)	343 (28.4)	280 (29.9)	314 (28.5)		
Day of week of admission, <i>n</i> (%)					0.99	0.24
Weekdays	700 (73.5)	920 (76.1)	688 (73.4)	815 (74.0)		
Weekends	253 (26.5)	289 (23.9)	249 (26.6)	287 (26.0)		
Source of referral, <i>n</i> (%)					<0.01	<0.01
Emergency department	926 (97.2)	1,196 (98.9)	874 (93.3)	1,007 (91.4)		
Other <sup>d</sup>	27 (2.8)	13 (1.1)	63 (6.7)	95 (8.6)		
Intensive care admission, <i>n</i> (%)	110 (11.5)	113 (11.0)	41 (4.4)	52 (4.7)	<0.01	<0.01
Mode of separation, <i>n</i> (%)					0.68	0.09
Discharged by hospital	805 (84.5)	972 (80.4)	805 (85.9)	924 (83.9)		
Transferred/discharged at own risk	110 (11.5)	171 (14.1)	98 (10.5)	125 (11.3)		
Died in hospital	38 (4.0)	66 (5.5)	34 (3.6)	53 (4.8)		
Charlson comorbidity index, median (IQR)	1 (0–2)	1 (0–2)	1 (0–2)	1 (0–2)	0.01	0.13
Principal diagnosis, <i>n</i> (%)					<0.01	<0.01
Influenza and pneumonia	527 (55.3)	758 (62.7)	276 (29.5)	358 (32.5)		
Chronic lower respiratory diseases	229 (24.0)	258 (21.3)	293 (31.3)	307 (27.9)		
Acute upper respiratory infections	53 (5.6)	45 (3.7)	75 (8.0)	95 (8.6)		
Other acute lower respiratory infections <sup>b</sup>	70 (7.4)	51 (4.2)	64 (6.8)	62 (5.6)		
Lung diseases due to external agents <sup>c</sup>	36 (3.8)	43 (3.6)	97 (10.4)	91 (8.3)		
Other respiratory diseases <sup>d</sup>	38 (4.0)	54 (4.5)	132 (14.1)	189 (17.2)		

<sup>a</sup>Medical practitioner, outpatient, day procedure center, community health.

<sup>b</sup>ICD-10-AM: J20 to J22.

<sup>c</sup>ICD-10-AM: J60 to J70.

<sup>d</sup>ICD-10-AM: J30 to J39; J85 to J84; J85 to J86; J90 to J94, and J95 to J99.

<sup>e</sup>Comparisons were made using Pearson's chi-square test for categorical variables and the Mann-Whitney test for continuous variables, and the *P* value shows the distribution of a given variable across groups.

in the LOS both in the within-intervention-groups analysis (median, –22.8 h; 95% CI, –33.1 to –12.5; *P* < 0.01) and in the DID analysis (median, –25.5 h; 95% CI, –38.1 to –12.9; *P* < 0.01) following the introduction of rapid PCR. For patients with positive test results, the median LOS decreased by 27.9 h (95% CI, –45.2 to –10.5; *P* < 0.01) in the within-intervention-groups analysis and by 33.4 h (95% CI, –49.5 to –17.3; *P* < 0.01) in the DID analysis following the introduction of rapid PCR (Table 3). Of patients in the rapid PCR group, the median LOS of patients who received an additional standard multiplex PCR panel was almost 1 day longer than that of those not requiring the multiplex PCR.

**Secondary outcome.** Figure 2 compares utilization of five common microbiology tests by intervention and control groups during the postimplementation versus baseline. Overall, after adjusting for confounding variables, patients in the rapid PCR group had significantly fewer blood culture, sputum culture, and respiratory bacterial and viral serology tests compared to standard PCR (Fig. 2A). For positive test results, patients in the rapid PCR group had significantly fewer sputum culture, and respiratory bacterial

**TABLE 2** Test results and timeliness of care among intervention groups

Variable	Results for:	
	Standard PCR ( <i>n</i> = 953)	Rapid PCR ( <i>n</i> = 1,209)
Positive test result to at least one virus, <i>n</i> (%)	210 (22.0)	440 (36.4)
Virus detected, <i>n</i> (%)		
Influenza A/B	24 (1.3)	405 (17.7)
Respiratory syncytial virus	38 (2.0)	36 (1.6)
Rhinovirus	70 (7.4)	
Coronavirus	36 (3.8)	
Human metapneumovirus	26 (2.7)	
Parainfluenza virus	22 (2.3)	
Adenovirus/enterovirus/bocavirus <sup>a</sup>	7 (0.4)	
Test TAT (h), median (IQR) <sup>b</sup>		
Hospital A	29.8 (25.2–40.6)	1.7 (1.4–2.4)
Hospital B	23.9 (17.4–28.5)	2.6 (1.9–3.9)
Hospital C	29.2 (24.7–41.4)	3.7 (2.9–5.8)
Overall	27.4 (23.0–36.8)	2.3 (1.6–3.7)
Time to sample (h) received at the lab from admission, median (IQR)	5.7 (3.2–17.4)	3.9 (2.1–7.3)
Time to test result from admission, median (IQR)	40.3 (28.3–52.6)	7.5 (4.5–16.7)
Discharged before a test result became available, <i>n</i> (%)	180 (18.9)	27 (2.2)

<sup>a</sup>Adenovirus = 3, bocavirus = 3, and enterovirus = 1.

<sup>b</sup>TAT, turnaround time; IQR, interquartile range.

and viral serology tests but not fewer blood culture and urine MC&S tests compared to those in the standard PCR group (Fig. 2B). For negative test results, with the exception of urine MC&S, there were significantly fewer other tests ordered for the rapid PCR group compared to those in the standard PCR group (Fig. 2C). In the control group, with the exception of blood culture, there was no difference in ordering practices over time (Fig. 2C).

## DISCUSSION

In this controlled, quasi-experimental study, we investigated the impact of rapid PCR testing of influenza A/B and RSV across three hospitals on hospital LOS and related microbiology test utilization. We found that the introduction of rapid PCR was associated with significant reductions in the hospital LOS among patients with positive test results and in the use of other common microbiology tests, including blood culture, sputum culture, and respiratory bacterial and viral serology tests, compared with those for patients in the preimplementation period who received standard multiplex PCR testing.

In our study, the impact of rapid PCR on hospital LOS was statistically significant among patients with positive test results. This finding is unsurprising, given that patients can leave the hospital earlier if a bacterial infection has been ruled out. For patients requiring antiviral therapy, treatment can be started more quickly in patients with positive results, which may expedite their discharge (21), whereas patients with negative results are more likely to stay in the hospital, as they may require further investigation and management. This was consistent with results of a previous U.S. study (11). Rappo et al. retrospectively assessed the impact of early diagnosis of respiratory infections using rapid PCR (FilmArray) on the outcomes of over 337 adult patients over two flu seasons at a tertiary care center in New York, NY, USA, and found a significant reduction in hospital LOS among influenza-positive patients with rapid PCR (11).

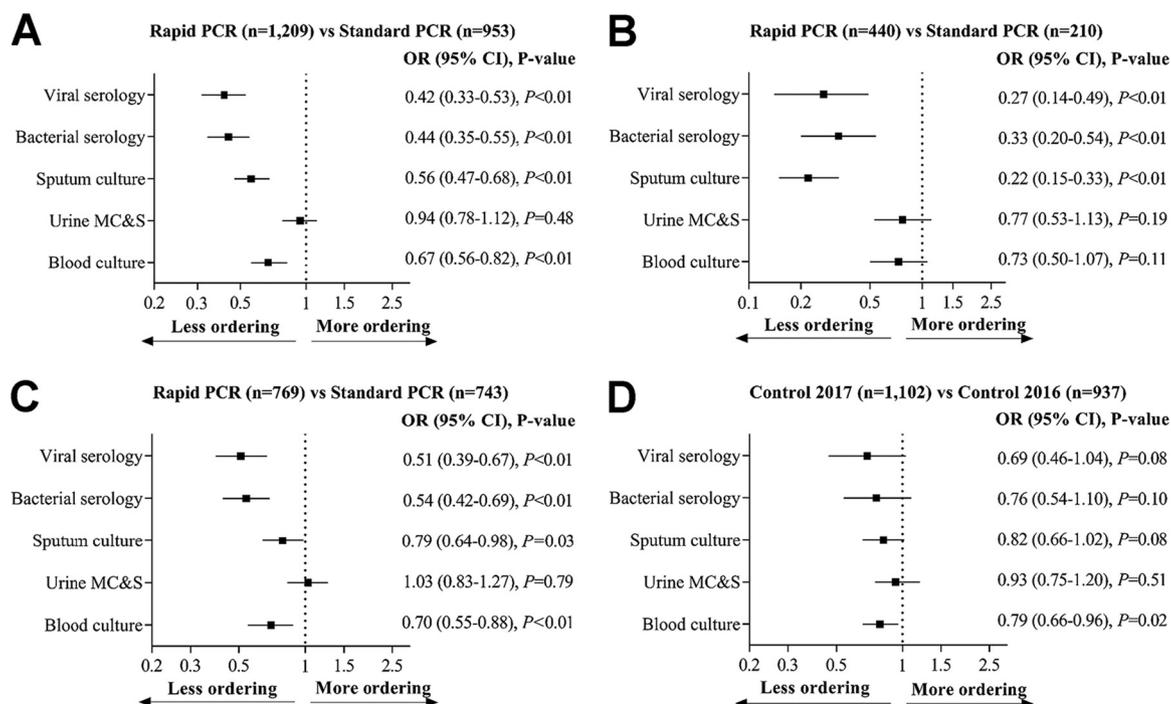
In the analysis that included all patients regardless of test results, there was no significant difference in LOS between rapid PCR and standard PCR groups. This was because more patients in the standard PCR group (18.9% versus 2.2%) were discharged from the hospital even before the test results were available. These patients had a

**TABLE 3** Median regression analysis showing changes in the median LOS following the introduction of rapid PCR

Variable	No. of patients	Median LOS (h [95% CI])	Change in LOS within the intervention groups (pre to post) <sup>a</sup>					
			Unadjusted			Adjusted <sup>b</sup>		
			Coefficient (95% CI)	P value	Coefficient (95% CI)	P value	Adjusted DID analysis <sup>b</sup> Coefficient (95% CI)	P value
<b>Intervention group</b>								
<i>Regardless of test results</i>								
Standard PCR	953	100.9 (96.2 to 109.9)	Reference group		Reference group		Reference group	
Rapid PCR	1,209	101.8 (97.1 to 112.8)	0.9 (-7.9 to 9.6)	0.85	-6.5 (-16.0 to 2.9)	0.18	-8.9 (-21.5 to 3.7)	0.17
<i>Positive test results</i>								
Standard PCR	210	98.6 (94.5 to 116.9)	Reference group		Reference group		Reference group	
Rapid PCR	440	92.9 (81.5 to 98.6)	-5.5 (-19.9 to 8.8)	0.45	-21.2 (-38.0 to -4.5)	0.01	-21.5 (-36.8 to -6.2)	<0.01
<i>Negative test results</i>								
Standard PCR	743	101.4 (96.0 to 113.0)	Reference group		Reference group		Reference group	
Rapid PCR	769	116.2 (103.4 to 122.6)	14.8 (2.4 to 27.1)	0.02	1.4 (-10.8 to 13.5)	0.83	-1.2 (-14.9 to 12.4)	0.86
Subgroup analysis including only patients whose test results were returned during their stay in the hospital								
<i>Regardless of test results</i>								
Standard PCR	773	126.0 (117.9 to 139.0)	Reference group		Reference group		Reference group	
Rapid PCR	1,182	104.0 (98.3 to 114.8)	-22.2 (-32.5 to -11.9)	<0.01	-22.8 (-33.1 to -12.5)	<0.01	-25.5 (-38.1 to -12.9)	<0.01
<i>Positive test results</i>								
Standard PCR	175	117.5 (100.5 to 142.8)	Reference group		Reference group		Reference group	
Rapid PCR	437	93.2 (82.2 to 98.6)	-24.3 (-42.0 to -6.6)	<0.01	-27.9 (-45.2 to -10.5)	<0.01	-33.4 (-49.5 to -17.3)	<0.01
<i>Negative test results</i>								
Standard PCR	598	129.7 (121.2 to 142.7)	Reference group		Reference group		Reference group	
Rapid PCR	745	117.9 (109.6 to 125.2)	-11.9 (-25.9 to 2.2)	0.09	-17.3 (-30.4 to -4.3)	<0.01	-19.6 (-33.7 to -5.5)	<0.01
<b>Control group</b>								
2016	937	75.3 (71.0 to 80.5)	Reference group		Reference group		Reference group	
2017	1,102	78.9 (73.0 to 87.3)	3.2 (-5.6 to 12.1)	0.47	1.6 (-6.0 to 9.2)	0.68	Not applicable	

<sup>a</sup> A negative coefficient indicates a median decrease in the LOS compared to the reference group.

<sup>b</sup> Adjusted for age, study hospital, the source of referral, intensive care admission status, mode of separation, Charlson comorbidity index and type of principal diagnosis. DID (difference-in-difference) = ΔLOS (h) in intervention group - ΔLOS (h) in control group.



**FIG 2** Common microbiology test utilization. (A) Regardless of test results, (B) positive test results, (C) negative test results, and (D) control groups. The middle square shows the odds ratio (OR), and the horizontal line shows the 95% confidence interval (CI) of the OR. The inclusion of “1” in the 95% CI indicates a nonsignificant difference in the ordering practices between groups. All analyses were adjusted for baseline variables, including age, study hospital, the source of referral, intensive care admission status, mode of separation, Charlson comorbidity index, and type of principal diagnosis. MC&S, microscopy, culture, and sensitivity.

shorter LOS compared with that of patients who were discharged after test results were available, confounding the effect of rapid PCR on hospital LOS. When the analysis was limited to patients whose test results were returned during their stay in the hospital, rapid PCR use was associated with a significant reduction in the median LOS, irrespective of test results.

In a recent large UK study, the use of a molecular point-of-care-test (POCT) resulted in a reduction in hospital LOS regardless of test results, although the effect was more noticeable among patients with positive test results (22). Brendish et al. conducted a pragmatic randomized controlled trial enrolling 720 adult patients (aged  $\geq 18$  years) at a large teaching hospital in the United Kingdom (the ResPOC trial). That study assessed the impact of molecular POCT testing for respiratory viruses compared with referral laboratory testing and found a significant reduction in hospital LOS by 1.1 days in the patients receiving POCT. In a subgroup analysis, patients with positive rapid PCR results had a hospital stay that was an average of 1.7 days shorter than that of patients with negative results (22).

In contrast to our findings and the results of the ResPOC trial (22) and Rappo et al. (11), the use of rapid PCR did not decrease hospital LOS in a study by Andrews et al. (21). The authors conducted a quasi-randomized trial to assess the impact of the FilmArray respiratory panel as a POCT compared to laboratory-based PCR, serology, or culture testing on a range of outcomes, including hospital LOS. The study enrolled 545 patients aged  $\geq 16$  years and presenting with influenza-like illness or other respiratory tract infections at an Acute NHS Hospital Trust in London, United Kingdom. Interestingly, the authors found an 11% increase in hospital LOS in the rapid PCR group, although it was not statistically significant. The authors attributed the lack of impact on LOS to a delay in initiating FilmArray testing. The median time to test result from admission was 19 h in their study. This was substantially longer than ours (the median time to test result from admission was 7.5 h), and this may explain the differences between our findings and those of Andrews et al. (21).

The limitation of the rapid PCR (Cepheid Xpert Flu/RSV XC) used in our study is its inability to diagnose other respiratory viruses. While it can accurately detect influenza A/B and RSV, other clinically relevant respiratory viruses, such as rhinovirus, coronavirus, metapneumovirus, and parainfluenza viruses, cannot be detected by our current rapid PCR method. This means that patients can still be ordered standard multiplex PCR following the results of rapid PCR if other respiratory viruses are suspected. Our analysis showed that 20% of the patients in the rapid PCR group also received multiplex PCR, and their median LOS was almost 1 day longer than that of those not requiring the multiplex PCR. Other studies implemented multiplex PCR respiratory panel (i.e., our version of the referral laboratory-based standard PCR) as a POCT test (11, 21, 22), avoiding the need for further respiratory viral investigations. The introduction of a broader multiplex PCR as a POCT in the future may offer better outcomes for patients.

Our results show that the utilization of blood culture, sputum culture, and respiratory bacterial and virus serology tests was significantly lower among patients tested with rapid PCR. To the best of our knowledge, no previous studies have studied the impact of PCR-based rapid tests on other laboratory test utilization in hospitalized adults. Previous studies that assessed other test ordering practices were conducted using antigen-based POCT in ED (23). Poehling et al. (13) and Blaschke et al. (14) reported significantly fewer blood culture tests when POCT was used, similar to our findings. Other studies have reported fewer biochemistry tests, including full blood count (14, 15), C-reactive protein (15), and urinalysis (14, 15), when POCT was used.

Our findings have some important implications. Reducing hospital LOS has large economic benefits. Cost savings for hospitals can be achieved through avoidance of unnecessary hospital resource utilization, including that of supplementary laboratory and imaging testing (24, 25). In a Spanish study by Soto et al., rapid PCR decreased the isolation time of hospitalized patients by 23.7 h, which resulted in a cost reduction of \$70 per hospitalized patient (24). A half-day reduction in hospital LOS in one U.S. study was associated with an estimated annual savings of \$500 to 900 million (26). Rapid diagnostic testing can have safety implications too. Reducing LOS in hospital leads to a reduction in hospital-acquired infections, including the disruption of *Clostridium difficile* transmission (27). According to a large U.S. study, reducing LOS by about 1 day lowers the risk of 30-day readmission through a reduction in infection risk (28).

The strengths of this study include its relatively large sample size, its utilization of real-world routinely collected clinical practice data, and multiple-center inclusion of three teaching hospitals, which enhances the generalizability of our findings. Unlike several prior studies (11, 12, 14, 29), we applied a quasi-experimental, controlled study design that allowed us to account for secular trends or sudden changes over time (30). Moreover, we used a number of strategies to reduce the potential impact of confounding variables. We used matching on key baseline variables to identify comparable control groups. To reduce seasonal effects, a similar time frame for both groups (i.e., July to December) was selected. To decrease the potential for selection bias, all hospitalized adults tested for influenza and RSV during the study period were included. Finally, we conducted within-intervention-groups and DID analyses, adjusting for differences in several baseline patient and clinical characteristics.

There are some limitations to consider when interpreting the findings of the current study. Given that this study was not a randomized controlled trial, our findings should not be interpreted as a cause and effect relationship. Due to a lack of randomization, there is always a risk of unidentified or hidden confounders in a quasi-experimental study (31). While our results indicated that the use of rapid PCR was associated with fewer other microbiology test utilizations, because of lack of data on regarding the time when the clinicians ordered the tests, our study did not examine whether the larger number of tests ordered with the standard PCR was associated with orders after results were returned to the patient. As we did not have access to treatment data, we were unable to assess the potential impact of rapid PCR testing on clinical decision making, including whether it influenced the decision to initiate antiviral therapy following positive results or to deescalate its use if patients were already using the medication

and the test result returned negative. Although our current study result suggests economic benefits of rapid PCR testing, a formal health economic study is required.

**Conclusion.** The introduction of rapid PCR testing of influenza and RSV viruses for hospitalized adults was associated with a significant reduction in hospital LOS for patients with positive results and a significant reduction in microbiology test use compared with those for patients who received standard multiplex PCR testing. These findings suggest there may be economic and patient outcome benefits from this intervention that should be tested in a future cost-effectiveness study.

## SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at <https://doi.org/10.1128/JCM.01727-18>.

**SUPPLEMENTAL FILE 1**, PDF file, 0.1 MB.

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