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**Supplemental Material for Seasonality of respiratory viruses causing hospitalizations for acute respiratory infections in children in Nha Trang, Vietnam**

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**Highlights**

- Hospitalizations for respiratory viruses are seasonal in Vietnam
- Respiratory syncytial virus peaks in the late summer months, and influenza A in April to June
- No clear seasonality is seen for human rhinovirus
- Human parainfluenza 3 and human rhinovirus are positively associated with dew point
- This work can inform the timing of influenza and RSV vaccination and the judicious use of antibiotics in Vietnam

# Seasonality of respiratory viruses causing hospitalizations for acute respiratory infections in children in Nha Trang, Vietnam

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## Abstract

### Background

Acute respiratory infections (ARIs) are the most common causes of death in children under 5 years of age. While the etiology of most pneumonia and ARI episodes is undiagnosed, a broad range of ARI-causing viruses circulate widely in South East Asia. However, the patterns and drivers of the seasonal transmission dynamics are largely unknown. Here we identify the seasonal patterns of multiple circulating viruses associated with hospitalizations for ARIs in Nha Trang, Vietnam.

### Methods

Hospital based enhanced surveillance of childhood ARI is ongoing at Khanh Hoa General Hospital in Nha Trang. RT-PCR was performed to detect 13 respiratory viruses in nasopharyngeal

samples from enrolled patients. Seasonal patterns of childhood ARI hospital admissions of various viruses were assessed, as well as their association with rainfall, temperature, and dew point.

### **Results**

Respiratory syncytial virus peaks in the late summer months, and influenza A in April to June. We find significant associations between detection of human parainfluenza 3 and human rhinovirus with the month's mean dew point. Using a cross-wavelet transform we find a significant out-of-phase relationship between human parainfluenza 3 and temperature and dew point.

### **Conclusions**

Our results are important for understanding the temporal risk associated with circulating pathogens in Southern Central Vietnam. Specifically, our results can inform timing of routing seasonal influenza vaccination and for when observed respiratory illness is likely viral, leading to judicious use of antibiotics in the region.

## 1 Introduction

2 Acute respiratory infections (ARIs) in South East Asia cause substantial morbidity and mortality,  
3 especially in children under 5 years of age [1]. Pneumonia continues to be the number one cause  
4 of under 5 death despite effective treatments [2]. A substantial contributor to this is the largely  
5 unknown etiology of most pneumoniae, with both viral and bacterial origin. The patterns of  
6 pneumonia and other ARI hospitalizations serves as a proxy for determining the transmission  
7 dynamics of viruses and bacteria contributing to these hospitalizations, and is of key importance  
8 in understanding and limiting burden of this childhood killer.

9 In addition to informing on the relative likelihood of the potential viral etiology of a pneumonia  
10 case based on seasonally circulating pathogens, knowledge of seasonal influenza epidemiological  
11 dynamics can aid in informing optimal timing for vaccination efforts [3, 4] and judicious use of  
12 antivirals [5, 6]. Identification of low incidence seasons would provide a target window for vaccina-  
13 tion, with the hope of maximizing population immunity before the onset of the influenza season.  
14 Similarly, when respiratory syncytial virus (RSV) vaccination becomes available, knowledge of its  
15 seasonality will be useful to maximize benefit [7, 8], and in the potential administration of pas-  
16 sive immunoprophylaxis with palivizumab to reduce the number of severe outcomes associated  
17 with RSV infection among high-risk infants [9]. Finally, knowledge of seasonal patterns of virus  
18 circulation can inform the clinical use of antibiotics, again limiting use when viral circulation is  
19 traditionally high to minimize antibiotic resistance [10, 11, 12, 13].

20 Previous work has identified the potential etiology of pneumonia and other ARIs in South-  
21 ern Central Vietnam [14, 15]. Adenovirus (AoV), bocavirus (HBoV), coronavirus (CoV), human  
22 metapneumovirus (HMPV), human parainfluenza 1–4 viruses (HPIV1–4), human rhinovirus (HRV),  
23 Influenza A and B, and respiratory syncytial virus (RSV) all contribute to the disease burden and  
24 circulate widely. Information on viral transmission dynamics across seasons in Vietnam is relatively  
25 under-explored. Do et al. found seasonality of RSV infections and slight seasonality of HMPV in-  
26 fections in Ho Chi Minh City, but no seasonality of influenza [14]. Several studies have similarly  
27 found high variability in Influenza A & B incidence over the year [16, 17, 18]. Yoshida et al. found  
28 RSV occurring in the hot months, influenza A in the cool months, and year-round detection of  
29 HRV in Nha Trang [15]. Few other studies have examined seasonality of these common viruses

30 across South East Asia.

31 Here we examine the seasonal trends of hospitalizations and circulation of multiple viruses in  
32 Nha Trang, Vietnam. Using enhanced hospital based surveillance of childhood ARI we identify  
33 seasonal patterns in hospitalizations as a proxy for transmission and explore the relationship of  
34 hospitalizations associated with virus detection with rainfall, temperature, and dew point, to try  
35 and identify contributing factors to observed seasonality.

## 36 **Methods**

### 37 **Study site**

38 The study site is Nha Trang, central Vietnam, where the study population has been described  
39 previously [19, 15, 20]. The hospital based enhanced surveillance of childhood ARI is ongoing. We  
40 analyze data from January 29, 2007 to April 26, 2012 at Khanh Hoa General Hospital (KHGH)  
41 which is the only tertiary care facility located in Khanh Hoa Province. According to the field site  
42 census survey in July 2006, the study catchment area encompassing the 16 non-touristic of the 27  
43 communes in Nha Trang city, had 198,729 residents including 13,952 children less than 5 years of  
44 age. An ARI case was defined as any child presenting to KHGH with cough or/and difficulty in  
45 breathing. Before study enrollment, informed consent was obtained from parents of children who  
46 presented with ARI and lived in the study catchment area. Clinical and demographic information,  
47 chest radiographs (CXR), laboratory data, and nasopharyngeal (NP) samples were collected from  
48 all enrolled patients. KHGH is the only hospital in Nha Trang, Khanh Hoa province and the only  
49 one accessible for residents of the catchment area. Hence for incidence calculations we assume  
50 that all children with ARI are eligible to be hospitalized and enrolled into the study and use the  
51 population of the catchment area as denominator. Acute respiratory infection patients with normal  
52 CXR were categorized as upper respiratory tract infection (URTI). Patients with abnormal CXR  
53 were categorized as lower respiratory tract infection (LRTI).

54 NP samples were collected at the time of admission and viral nucleic acid was extracted using  
55 QIA viral RNA minikit (QIAGEN Inc., Valencia, CA). Four multiplex-PCR assays (1: influenza  
56 A, influenza B, RSV, hMPV; 2: PIV-1, -2, -3, and -4; 3: rhinovirus, coronavirus 229E, coronavirus  
57 OC43; 4: adenovirus and bocavirus) were performed to detect 13 respiratory viruses in each NP

58 sample. A second confirmatory-PCR was performed for samples positive on the initial PCR test.  
59 Samples positive for both PCR assays were defined as positive. Reverse transcription-PCR (RT-  
60 PCR) assays were performed using one-step RT-PCR kit from QIAGEN. For the multiplex PCR  
61 and hemi-nested PCR assays, TaqDNA polymerase (Promega, San Luis Obispo, CA) was used as  
62 previously described [15]. Positive templates were used in each assay for quality control.

### 63 **Weather data**

64 Three weather variables – rainfall (inches), temperature ( $F^{\circ}$ ), and dew point (the temperature to  
65 which air must be cooled in order to reach saturation with water) – were collected from the Nha  
66 Trang Station (ID: 488770) reported by the US National Oceanic and Atmospheric Administra-  
67 tion [21]. We considered monthly averages of all weather variables as well as both the weather  
68 variable on the day of admission ( $t_0$ ) as well as averaged over the previous 7 days ( $t_{-7}$  to  $t_{-1}$ )  
69 assuming up to a week incubation period for the viral infections (see supplementary material) [22].  
70 The weather in Nha Trang central Vietnam is warm throughout the year (between 20 and 30  $C^{\circ}$ ).  
71 In terms of temperature, December to February months are cooler (referred to here as the winter  
72 months) while June to August months are hottest months (referred to here as the summer months).  
73 September, October, November are the wettest months.

### 74 **Statistical analysis**

75 For each PCR+ for virus a series of log-link Poisson models were fit to assess respective seasonality  
76 with calendar month as the main predictor, log-commune population size as an offset term, monthly  
77 averaged rainfall, temperature, and dew point, and calendar year as adjusting variables. The out-  
78 come was monthly aggregate cases, with resulting coefficients as incidence rate ratios as compared  
79 to January. We excluded hospitalizations with more than one virus detected in the nasopharynx  
80 to adjust for a potential bias through inclusion of cases who by virtue of being co-infected may  
81 otherwise have been asymptomatic [23]. This approach underestimates the true incidence of NP  
82 carriage among ARI cases but allows estimation of seasonal patterns that are not biased by other  
83 circulating viruses; although co-circulation of bacteria was not accounted for. We assessed the  
84 numbers and variety of viruses in ARI hospitalizations using binomial proportion tests for each  
85 virus.

## 86 Cross-wavelet transform

87 To examine the relationship between monthly average rain, temperature, and dew point and inci-  
88 dence hospitalized childhood ARI infections, we estimated the cross-wavelet transform between the  
89 z-standardized time series (we subtracted the mean of the time series and divided by the standard  
90 deviation) of weather and viral detections [24]. The cross-wavelet transform identifies regions of  
91 high power in phase-space and identifies the relative phases of each time series, i.e., in-phase or  
92 out-of-phase [25]. The wavelet transform can be thought of a Fourier transformation over time  
93 that can identify what is the dominant frequency composing a time series as the signal changes  
94 in time. The cross-wavelet transform allows us to compare how two time signals co-vary: we can  
95 identify if the presence of a particular frequency at a given time in the time series of hospitalizations  
96 corresponds to the presence of that same frequency at the same time in a weather covariate [26].  
97 Additionally, we can identify the magnitude by which weather precedes or follows hospitalizations  
98 through the phase angle of the two time series. Finally, we can identify the statistical significance of  
99 the identified constituent frequencies over time by comparing the observed frequencies to a red-noise  
100 process.

## 101 Sensitivity analyses

102 Sensitivity analyses, presented in the supplementary materials, were performed as follows: 1) case  
103 counts of less than 70 per virus over the whole study period were deemed too low for robust  
104 statistical inference; 2) alternative Poisson regression models where the reference category is July;  
105 3) logistic regression models were formulated as an alternative to the Poisson regressions above with  
106 detection of a virus by PCR (yes/no) as the outcome, with month as the main predictor, adjusted  
107 for weather, commune of residence, age, sex, smoking indoors, socioeconomic status (SES), and  
108 calendar year, with weather variables on the day of admission ( $t_0$ ) as well as averaged over the  
109 previous 7 days ( $t_{-7}$  to  $t_{-1}$ ); and 4) additional wavelet analyses of viral isolations not presented in  
110 the main text.



## 111 **Results**

112 The study enrolled 3431 children between 2007 to 2013. Among those, 374 (11%) had multiple  
113 viruses detected in their NP swabs, for 59 presence of viruses in the NP was not determined, and  
114 were excluded from the analyses, thus the total study population was 2998. Among all cases with  
115 a virus detected, HRV, RSV, and Influenza A were the most frequently detected viruses, with 569  
116 (33.5% of all viral detections), 455 (26.8%), and 282 (16.6%) detections, respectively (Table 1 and  
117 Figure 1). Counts of bocavirus (HBoV), coronavirus (CoV), and human parainfluenza 1, 2, and 4  
118 viruses (HPIV1–4) were less than 70 and are reported in the supplementary material.

## 119 **Seasonality**

120 Strong seasonality, as defined by at least three consecutive months with a consistently higher or  
121 lower incidence than expected (IRR or OR greater or less than 1, respectively) and at least one of  
122 those statistically significantly different from the baseline, was observed for Influenza A, RSV, and  
123 the presence of any virus (Figures 2 and 3). RSV peaked in July through November, with August  
124 seeing a 15.69 (95% confidence interval [CI]: 3.05, 80.56) times higher risk of identifying RSV as  
125 the sole viral agent from the nasopharynx of a childhood case as compared to January. Influenza  
126 A peaked in May with an IRR of 6.28 (95% CI: 2.2, 17.89) as compared to January. Estimates  
127 of odds ratios from the supplemental logistic regression are qualitatively similar to the Poisson  
128 regression, save for HPIV3, which exhibits a consistent yet non-significant peak in the cool months  
129 in the primary analysis which becomes significant in the supplemental analysis (see supplemental  
130 material).

## 131 **Association with weather**

132 Weather patterns over the study period were similar to patterns before and after the study period  
133 (supplementary material, Figure S3). Monthly average rainfall (in inches), temperature (1° F),  
134 and dew point (1° F) correlated with the seasonality of some of our endpoints. Figure 4 shows  
135 the incidence rate ratios for the three weather effects from the seasonally-adjusted models. Over-  
136 all hospitalizations for ARI were negatively associated with temperature (IRR 0.92 per 1 degree  
137 increase, 95% CI: 0.87, 0.97,  $p = 0.003$ ) and positively associated with dew point (IRR 1.08 per 1

138 degree increase, 95% CI: 1.04, 1.13,  $p < 0.001$ ). Of the few other significant effects, Influenza A  
139 and HRV had a negative associations with temperature (IRR 0.85, 95% CI: 0.75, 0.9,  $p = 0.0116$ ,  
140 and IRR 0.86, 95% CI: 0.79, 0.94,  $p < 0.001$ , respectively), and HPIV3 and HRV were positively  
141 associated with dew point, with IRRs 1.26 (95% CI: 1.04, 1.52,  $p = 0.0164$ ) and 1.13 (95% CI: 1.05,  
142 1.21,  $p = 0.0009$ ), respectively. Logistic regression found that RSV was positively associated with  
143 the previous week's rain, with an odds ratio of of 1.90 (95% CI: 1.21, 2.99,  $p = 0.0053$ ). Previous  
144 week's temperature was marginally associated with RSV (OR: 1.14 (95% CI: 0.98, 1.33,  $p = 0.08$ )  
145 (see supplementary material).

146 Figure 5 shows the cross-wavelet transform of all hospitalizations and the three weather vari-  
147 ables in the month of admission. Significant bands of high power around 1 year can be seen for  
148 temperature and dew point. This indicates that hospitalizations and temperature and dew point  
149 share variability at yearly frequencies over the study period, and the phase relationship indicates  
150 that changes in weather slightly precede changes in hospitalizations (arrows point about  $45^\circ$  down).  
151 Figure 6 shows the cross-wavelet transform of RSV and the three weather variables with similar  
152 significance bands around 1 year for temperature and dew point. The phase indicates temperature  
153 and dew point lead RSV incidence. While not significant, RSV was found to be leading rainfall  
154 by  $90^\circ$  (3 months) at the one year period band (see supplementary material). Significant bands of  
155 power were seen between temperature and dew point with HPIV3 with an indicated phase relation-  
156 ship of nearly completely out-of-phase (Figure 7). Similar patterns were seen (1 year significant  
157 bands between temperature and dew point and virus) for ADV, HBoV, CoV, HPIV1,2&4, HRV,  
158 and influenza A, though the phase differences varied across these viruses; HMPV and influenza B  
159 had bands lying outside the cone of influence (ie, not statistically significant; see supplementary  
160 material). These results in general indicate strong associations between weather covariates and  
161 viral hospitalizations at a yearly timescale.

## 162 Discussion

163 Here we have identified seasonal trends of several common respiratory viruses in hospitalized chil-  
164 dren in Nha Trang, Vietnam. By fitting a series of statistical models to the observed data, we allow  
165 the data to identify salient features contributing to the seasonality of these viruses. We evaluated

166 seasonal patterns and associations with weather of hospitalizations for several respiratory viruses  
167 using three lines of evidence: 1) Poisson regression examining the relative incidence across months  
168 of virus detections adjusted for weather covariates, 2) cross-wavelet transforms of hospitalizations  
169 with viral detections, and 3) a sensitivity analysis with a logistic regression model finding odds  
170 ratio of hospitalizations with viral detections and weather variables.

171 Any viral detection showed distinct seasonality with peaks in May through September, a nega-  
172 tive association with temperature, positive association with dew point, and cross-wavelets indicating  
173 temperature and dew point leading viral detection. Of commonly detected viruses, RSV, Influenza  
174 A, and HPIV3 had significant seasonality. RSV peaked in July through December, was positively  
175 associated with the week's previous average rainfall. Cross-wavelets showed temperature and dew  
176 point to lead RSV, rain was found to non-significantly fall behind RSV at 1 year frequencies, and  
177 precede RSV at shorter ( $< 100$  day) frequencies. Finally, HPIV3 while not significant, had peaks  
178 in January and February, was positively associated with dew point, and was completely out of  
179 phase with temperature and dew points in cross-wavelet analyses. These results contribute to the  
180 growing body of knowledge on the epidemiology of respiratory pathogens in South East Asia and  
181 Southern Central Vietnam.

182 Using a cross-wavelet transform we evaluated the time-dependence of virus hospitalizations  
183 with the weather covariates. We found strong yearly associations with RSV over the study period  
184 with temperature and dew point in phase with hospitalizations. This seasonality is opposite to  
185 observed seasonality in temperate climates, where RSV typically peaks in winter [27, 28]. Recent  
186 reviews of RSV seasonality in tropical regions highlights the uncertainty in the effects of weather  
187 on RSV transmission. Studies in Brazil [29], Hawaii [30], India [31], Kenya [32], and Malaysia [33,  
188 34, 35], have shown negative associations between temperature and RSV, while studies in Hong  
189 Kong [36], Mexico [37], Singapore [33], and Taiwan [38] have shown positive associations. We find  
190 strongly positive associations between temperature and RSV hospitalizations with a slight lead of  
191 temperature on RSV. This is evident both from the cross-wavelet transform and the regression  
192 results indicating a positive effect of the previous week's temperature on RSV.

193 There is less uncertainty in the role of rainfall on RSV transmission in tropical areas, where  
194 the majority of work indicates that RSV generally occurs during rainy seasons [28]. Colombia [39],  
195 The Gambia [40], Hong Kong [36], Kenya [32], Malaysia [34], and Papua New Guinea [41] all show

196 positive RSV associations with rainfall. Omer et al. [42] showed significant positive associations  
197 between rainfall and temperature in the previous 8 days and RSV incidence in Lombok, Indonesia.  
198 We find similar associations, with the mean rainfall and temperature over the previous 7 days  
199 having odds ratios for RSV of 1.98 and 1.23, respectively. This association is plausible as the  
200 incubation period of RSV is estimated to be between 4 and 5 days [22]. Detailed contact tracing  
201 studies, coupled with climatological data could refine this association.

202 Somewhat surprisingly, the cross-wavelet transform of RSV and rain showed no significant  
203 association, though areas of high power were observed in the 1-year and 6-month bands, with phase  
204 indicating RSV leading weather. However, none of the viruses studied here showed appreciable  
205 associations with rain in the cross-wavelet transform, possibly indicating the dominance of other  
206 weather effects (temperature and dew point) on virus hospitalizations.

207 As with previous studies examining HPIV incidence, we found a predominance of HPIV3 (72  
208 detections) compared to 41, 13, and 3 for HPIV1, 2, and 4, respectively. Typical seasonality for  
209 HPIV3 is the spring and early summer months in the temperate regions [43, 44], and has little to  
210 no observed seasonality in the tropics and subtropics [33, 45]. We find evidence for winter peaks in  
211 HPIV3 hospitalizations when employing both the Poisson regression model as well as the logistic  
212 regression model to estimate odds ratios, though the peaks were not statistically significant when  
213 controlling for weather.

214 The cross-wavelet transforms reveals HPIV3 to vary significantly at 1 year periodicity with  
215 temperature and dew point across the study. It also shows that HPIV3 hospitalizations are nearly  
216 completely out of phase with temperature and dew point throughout the study period. HPIV3 has  
217 been associated with low temperature and low relative humidity [46] though there is in general a  
218 paucity of data on the transmission routes of HPIV3 [47, 48]. Future work could explore in more  
219 depth the epidemiological relationship between HPIV and weather variables.

220 This study is not without limitations. First, while we have nearly 6 years of data, this is  
221 still a relatively short period to assess long-term seasonal trends, or to increase confidence in the  
222 estimates of seasonal patterns. However, the length of the analyzed time series is similar to other  
223 studies examining seasonal trends in viral respiratory pathogens in the tropics [33, 44], and gives  
224 indications for areas of future study. Second, we excluded individuals with more than one virus  
225 detected. Examination of changes in the seasonality of other viruses and coinfection over this

226 period is worthy of study and is outside the scope of this paper. Third, this study used hospital-  
227 based surveillance, necessarily presenting the most ill children. We take as an assumption that  
228 hospitalizations are a fraction of all transmission and severity of illness is not related to weather.  
229 Finally, this study examines the influence of weather on viral hospitalizations and does not address  
230 other drivers of seasonal patterns of transmission, such as school closures [49], differences in other  
231 social behavior such as contact rates [50, 51], susceptible recruitment through births [52], or possible  
232 seasonal changes in host immune responses [53]. Future work examining these drivers in this setting  
233 is necessary.

234 Limitations aside, our study adds to the body of literature on seasonality of common respiratory  
235 patterns in tropical regions and will be of use when consideration of the epidemiology of these  
236 pathogens is necessary. For example the timing of influenza peaks in the mid-spring months (April,  
237 May, June) would indicate routine vaccination in the winter months would be of biggest impact.  
238 Similarly, knowing RSV peaks in the late summer/early fall when RCP is at its lowest may help  
239 in limiting unnecessary antibiotic [54] or antiviral use [6]. In Vietnam and most of Asia, most  
240 antibiotics are acquired from a pharmacist without a formal prescription [10]. This fact makes  
241 results like those presented here of high importance to public health decision-makers to inform  
242 pharmacists of seasonality of respiratory infection etiologies and urge judicious prescribing practices.

243 Additional future work could include examination of the effects of contact clustering [55], co-  
244 infection [5, 56], and asymptomatic carriers [23] on transmission of the examined viruses all of  
245 which may be influenced by weather.

## 246 Contributions

247 Study design: BMA, SF, HH, LMY; Data collection: LMY, LNM, VDT; Data analysis: BMA,  
248 SF; Writing first draft: BMA, Writing subsequent drafts: BMA, SF, GR, LMY; Contributed  
249 intellectually: all authors

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257 or preparation of the manuscript.

### 258 **Ethical approval**

259 The study was approved by institutional review boards in the National Institute of Hygiene and  
260 Epidemiology, Vietnam and the Institute of Tropical Medicine, Nagasaki University, Japan.

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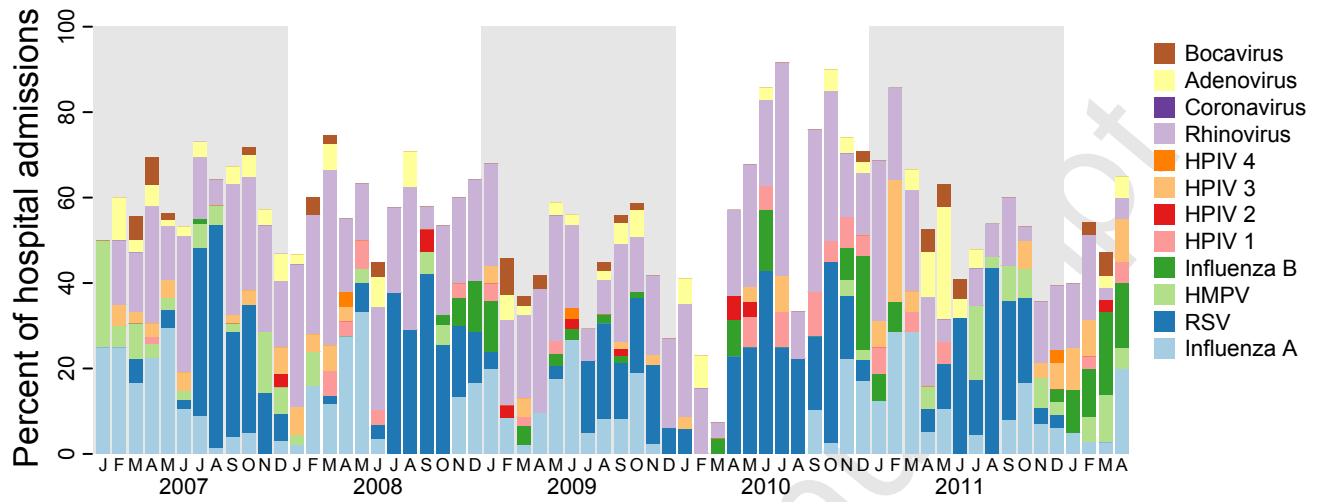
425 **Figures**

Figure 1: **Monthly hospitalizations by virus in Nha Trang, Vietnam.** Figure shows monthly detections of virus as percentages of all enrolled ARI hospitalizations.

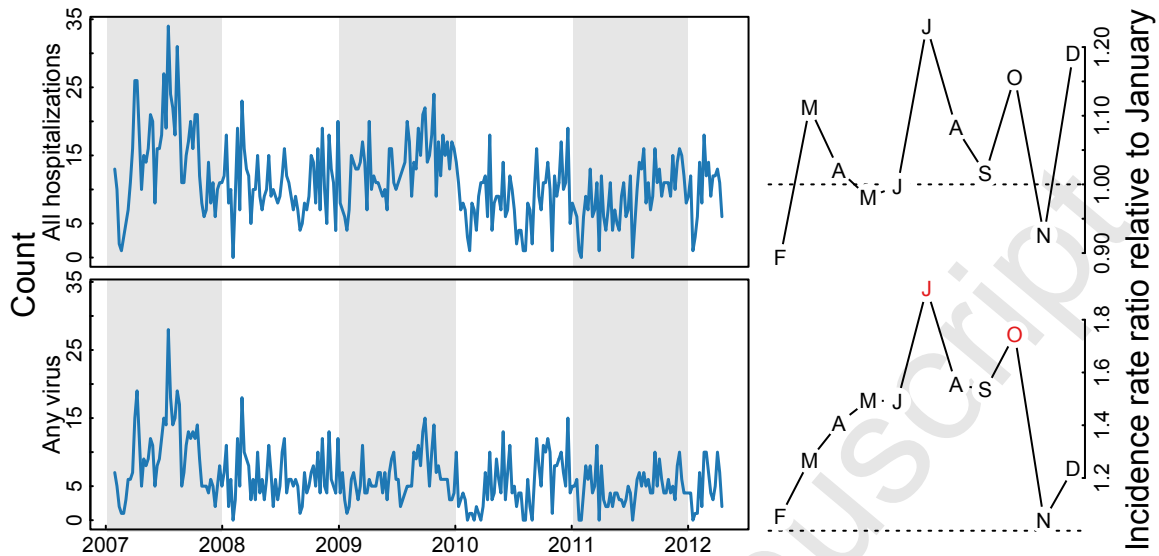


Figure 2: **Seasonality of hospitalizations and any viral detection in Nha Trang, Vietnam.** Figure shows weekly counts of all hospitalizations (top row) and any virus detection (bottom row). Right-hand column shows model-adjusted incidence rate ratios for month of year as compared to January. Red months indicates statistically significant deviations from January.

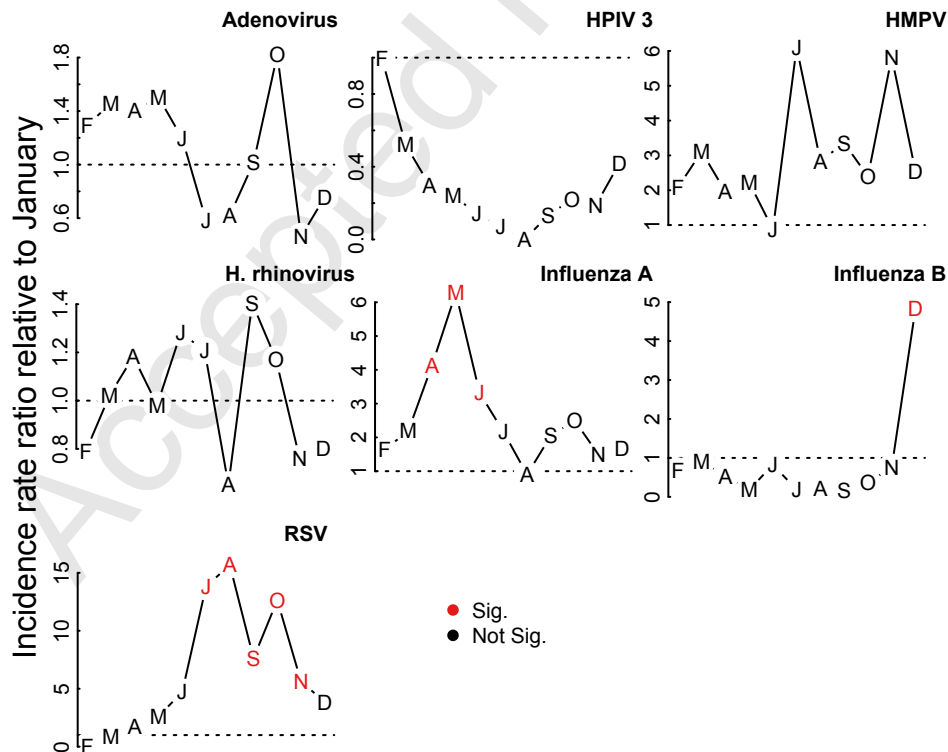


Figure 3: **Seasonality of common respiratory viruses in Nha Trang, Vietnam.** Figure shows model-adjusted incidence rate ratios for month of year as compared to January. Red months indicates statistically significant deviations from January.

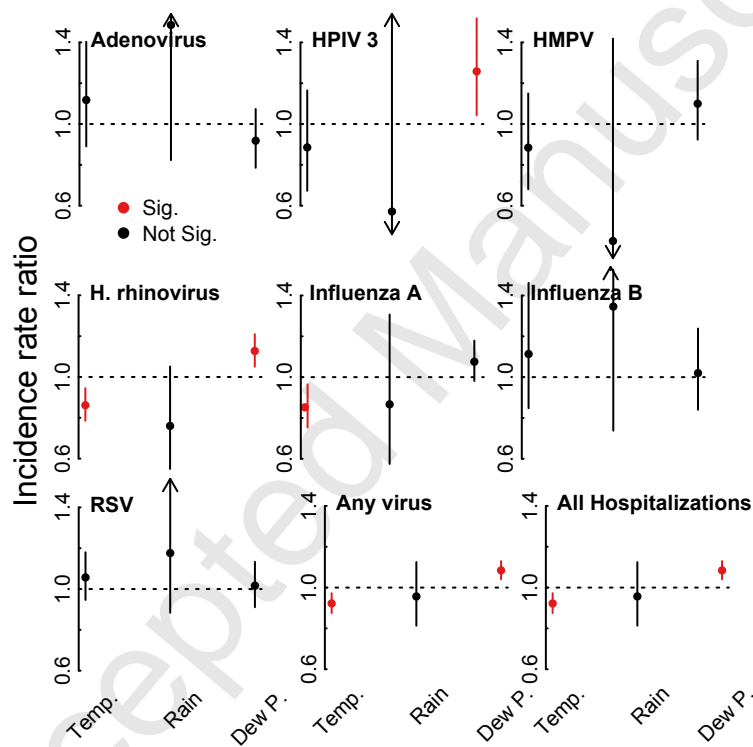


Figure 4: **Weather effects.** Figure shows model-adjusted incidence rate ratios for the main three weather effects: monthly mean rainfall (in inches), temperature ( $1^{\circ}$  F), and dew point ( $1^{\circ}$  F).

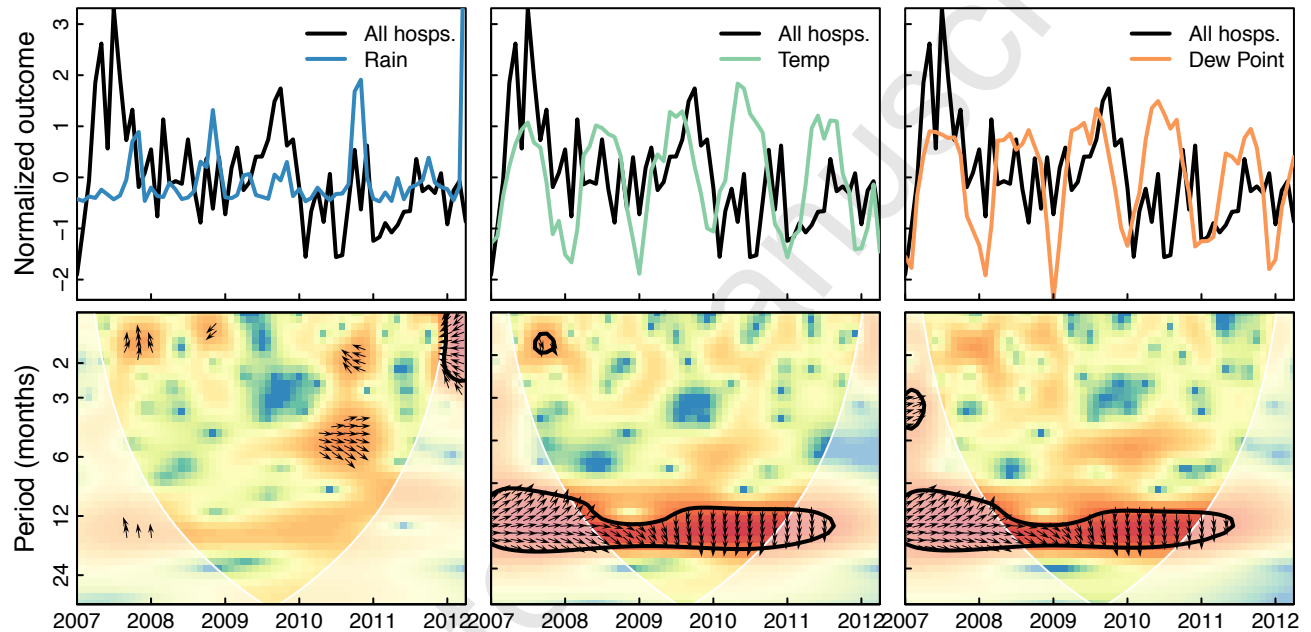


Figure 5: **Cross-wavelet transform of the z-standardized all hospitalization counts and weather time series.** Figure shows the cross-wavelet of all hospitalization counts and rainfall, temperature, and dew point. Colors indicate increasing cross-wavelet power (strength of coherence between the time series) blue to red. The 5% significance level against red noise is shown as a thick contour and the cone of influence (within which the wavelets are not influenced by the edges of the time series) is shown in white shading. The relative phase relationship is shown as arrows (with in-phase pointing right, out-of-phase pointing left, and weather leading hospitalizations by  $90^\circ$  pointing straight down).

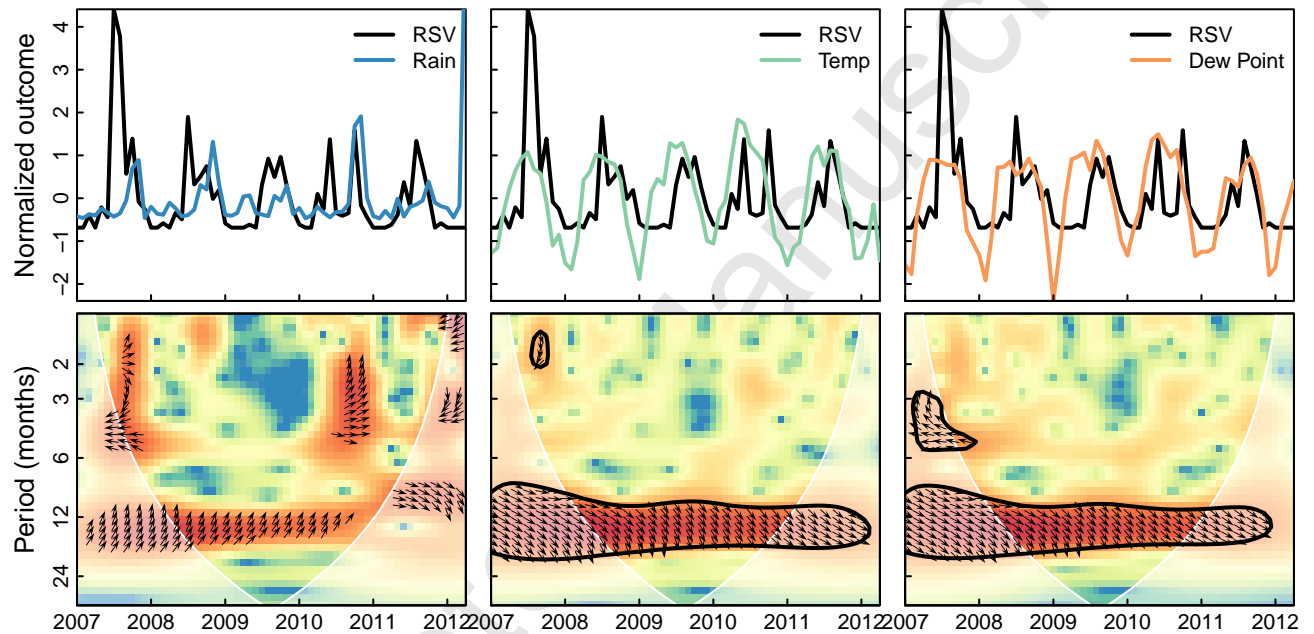


Figure 6: **Cross-wavelet transform of the z-standardized RSV and weather time series.** Figure shows the cross-wavelet of RSV and rainfall, temperature, and dew point. Colors indicate increasing cross-wavelet power (strength of coherence between the time series) blue to red. The 5% significance level against red noise is shown as a thick contour and the cone of influence (within which the wavelets are not influenced by the edges of the time series) is shown in white shading. The relative phase relationship is shown as arrows (with in-phase pointing right, out-of-phase pointing left, and weather leading RSV by  $90^\circ$  pointing straight down).

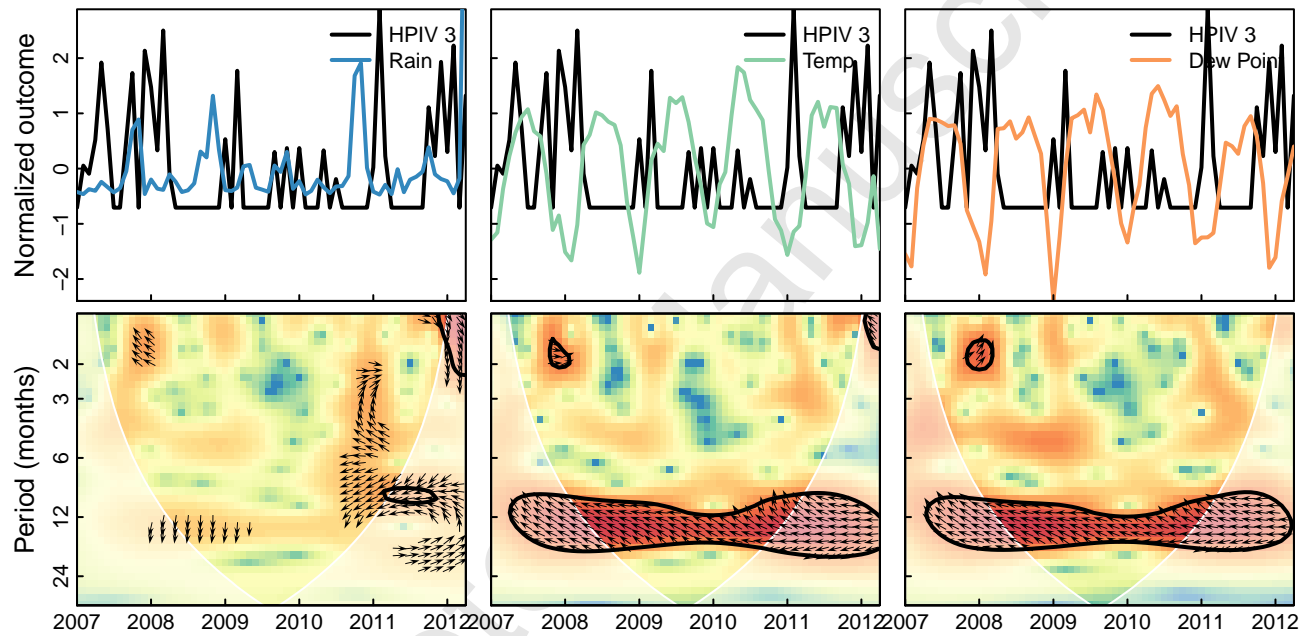


Figure 7: **Cross-wavelet transform of the z-standardized HPIV3 and weather time series.** Figure shows the cross-wavelet of HPIV3 and rainfall, temperature, and dew point. Colors indicate increasing cross-wavelet power (strength of coherence between the time series) blue to red. The 5% significance level against red noise is shown as a thick contour and the cone of influence (within which the wavelets are not influenced by the edges of the time series) is shown in white shading. The relative phase relationship is shown as arrows (with in-phase pointing right, out-of-phase pointing left, and weather leading HPIV3 by  $90^\circ$  pointing straight down).



|           | ADV       | HMPV      | HPIV3     | HRV         | Influenza A | Influenza B | RSV         | Any         |
|-----------|-----------|-----------|-----------|-------------|-------------|-------------|-------------|-------------|
| 2007      | 21        | 28        | 19        | 137         | 84          | 1           | 149         | 451         |
| 2008      | 15        | 10        | 10        | 134         | 51          | 8           | 85          | 330         |
| 2009      | 20        | 1         | 9         | 129         | 74          | 10          | 68          | 333         |
| 2010      | 12        | 2         | 4         | 95          | 25          | 24          | 84          | 271         |
| 2011      | 13        | 24        | 17        | 61          | 37          | 8           | 69          | 235         |
| 2012      | 6         | 7         | 12        | 13          | 11          | 19          | 0           | 76          |
| Total (%) | 87 (5.1%) | 72 (4.2%) | 71 (4.2%) | 569 (33.5%) | 282 (16.6%) | 70 (4.1%)   | 455 (26.8%) | 1696 (100%) |

Table 1: **Annual hospitalizations by virus.** Table shows counts of hospitalizations for adenovirus (ADV), human metapneumovirus (HMPV), human parainfluenza virus 3 (HPIV3), influenza A, influenza B, rhinovirus (HRV), and respiratory syncytial virus (RSV). Viruses with less than 70 cases were excluded from this table () and are presented in the supplement. Counts of all virus detections (including those presented in the supplement are in the column ‘any virus’).