

**West Nile Virus infection in Northern Italy: case-crossover study on the short-term effect of climatic parameters.**

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**Short title:** West Nile Virus and climatic parameters in Italy

**Competing interests statement:** The authors have no competing financial interests to declare

**Abstract:**

**Background:** Changes in climatic conditions are hypothesized to play a role in the increasing number of West Nile Virus (WNV) outbreaks observed in Europe in recent years.

**Objectives:** We aimed to investigate the association between WNV infection and climatic parameters recorded in the 8 weeks before the diagnosis in Northern Italy.

**Methods:** We collected epidemiological data about new infected cases for the period 2010-2015 from the European Center for Disease Control and Prevention (ECDC) and meteorological data from 25 stations throughout the study area. Analyses were performed using a conditional Poisson regression with a time-stratified case-crossover design, specifically modified to account for seasonal variations. Exposures included weekly average of maximum temperatures, weekly average of mean temperatures, weekly average of minimum temperatures and weekly total precipitation.

**Results:** We found an association between incidence of WNV infection and temperatures recorded 5-6 weeks before diagnosis (Incidence Rate Ratio (IRR) for 1°C increase in maximum temperatures at lag 6: 1.11; 95% CI 1.01-1.20). Increased weekly total precipitation, recorded 1-4 weeks before diagnosis, were associated with higher incidence of WNV infection, particularly for precipitation recorded 2 weeks before diagnosis (IRR for 5 mm increase of cumulative precipitation at lag 2: 1.16; 95% CI 1.08-1.25).

**Conclusions:** Increased precipitation and temperatures might have a lagged direct effect on the incidence of WNV infection. Climatic parameters may be useful for detecting areas and periods of the year potentially characterized by a higher incidence of WNV infection.

**Key Words:** West Nile Virus, Temperatures, Precipitations, Lag-distributed Models, Case-crossover

## 1        **1. Introduction**

2        West Nile Virus (WNV) is a globally distributed RNA virus of *Flaviviridae* family (Campbell  
3        et al. 2002). It is maintained in nature through an enzootic cycle. Adult mosquitoes, generally  
4        of *Culex* genus, represent primary bridge vectors, while susceptible bird species play the role  
5        of amplification hosts (Chancey et al. 2015). Humans usually develop infection after being  
6        bitten by an infected mosquito. Infection in humans is generally asymptomatic, but 20% of  
7        infected subjects can develop a febrile syndrome, known as West Nile Fever (WNF), and less  
8        than 1% of infected subjects can develop a West Nile Neuroinvasive Disease (WNND)  
9        characterized by encephalitis or meningitis symptoms (David and Abraham 2016).

10       In recent years, several outbreaks of WNV infection have been recorded in many European and  
11       Mediterranean countries (Rizzoli et al. 2015). Infected migratory birds are responsible for the  
12       introduction of the virus in new areas, while native mosquitoes feeding behaviour, presence of  
13       susceptible endemic birds and local environmental conditions are essential for persistence and  
14       amplification of the virus in new areas (Reisen and K. 2013, Rizzoli et al. 2015). Climatic and  
15       meteorological conditions have been suggested as important factors for virus transmission in  
16       newly affected areas (Paz 2015a; Paz et al. 2013). High extrinsic temperatures are associated  
17       with virus replication and the growth rate of the vector population (Gubler et al. 2001). Levels  
18       of precipitation are also believed to play an important role in pathogen/vector ecology: some  
19       studies reported that vector replication and activity are positively associated with heavy rainfall  
20       and other studies reported that mosquitoes' abundance is associated with drought periods (Nile  
21       et al. 2009, Paz 2015).

22       In Italy, the WNV was isolated for the first time in 1998 in 14 equine cases and the first human  
23       case was identified in 2008. Since then, human cases of WNV infection have been repeatedly  
24       notified, and now the virus is considered endemic in Italy (Rizzo et al. 2016). Concurrently  
25       the number of provinces set in Northern Italy affected by WNV circulation has increased during  
26       the study period (3 provinces in 2010 vs 16 in 2015). Thus, Italy can be considered as an  
27       example of area that is facing the process of endemization of an emerging pathogen.

28       The purpose of this study is to evaluate the short-term effects of air temperatures and  
29       precipitation on the incidence of WNV infection to understand the role of climatic parameters  
30       in the spread of WNV infection in an area, such as Northern Italy, where the process of  
31       endemization has recently started.

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34        **2. Methods**

35        **2.1 Data collection and elaboration**

36        Epidemiological data were obtained from the European Center for Disease Control and  
37        Prevention (ECDC). In our study, WNV cases are subjects resident in Northern Italy who,  
38        during the period 2010-2015, met the European criteria for probable or confirmed case of WNV  
39        infection (European Commission Decision 2008/426/E). Cases are confirmed if at least one  
40        following laboratory criterion is present: isolation of WNV from blood or Cerebrospinal Fluid  
41        (CSF), detection of WNV nucleic acid in blood or CSF, WNV specific IgM in CSF, WNV IgM  
42        high titer and subsequent detection of WNV IgG. Cases are considered probable in presence of  
43        stable and elevated virus specific serum antibody titer in association with one clinical criterion  
44        (fever, meningitis or encephalitis) or evidence of an epidemiological link that proves  
45        animal/human to human transmission. Thus, notified cases recorded by ECDC are a  
46        heterogeneous population and include: WNV positive blood donors, cases of WNF and cases  
47        of WNND. For each case, the ECDC provides information on the year, the week and the  
48        geographical province of diagnosis.

49        Meteorological data were obtained from the Regional Environmental Protection Agency  
50        (ARPA) for each province that reported at least one case of infection between 2010 and 2015.  
51        We used the information recorded by the land-based meteorological stations set in the capital  
52        of each province. Meteorological data included minimum, mean, maximum daily temperatures,  
53        and daily precipitation. On the daily data of temperatures and precipitation a quality control  
54        was carried out to exclude the possibility of measurement error (Fortin et al 2017; Acquotta  
55        et al, 2016; Zandonadi et al, 2016). In order to conform meteorological data to epidemiological  
56        data, we calculated the weekly average of the minimum, mean and maximum temperatures, as  
57        well as, the weekly total precipitation. We considered missing all weeks with at least one  
58        missing daily information (information missing on weekly scale: 4.4% for maximum  
59        temperatures, 6.4 % for mean temperatures, 5.1% for minimum temperatures and 6.1% for total  
60        precipitation).

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## 65 2.2 Study design

66 To estimate the association between climatic parameters and WNV infection, we used a case-  
67 crossover design, which is a special case-control design where every case serves as its own  
68 control and originally developed to study the acute effect of transient exposures on the risk of  
69 rapid onset events (Maclure and Mittleman 2000). For each case, exposures occurring during  
70 the period prior to the event (known as “hazard period”) are compared to exposures at  
71 comparable control periods (known as “reference periods”) (Janes et al. 2005a; Janes et al.  
72 2005b, Levy et al. 2001). In our study, control periods were identified according to a time-  
73 stratified sampling scheme, which uses fixed and relatively short time strata (e.g. calendar  
74 month) to match case and control periods (e.g. calendar week). Time-stratified case-crossover  
75 design has been repeatedly applied in environmental studies as it can control for long time  
76 trends (e.g. variability from year to year) and seasonality (variability from month to month)  
77 and can provide results equivalent to time series regression (Bateson and Schwartz 1999;  
78 Navidi 1998; Lu and Zeger 2007). We further modified the original time-stratified approach  
79 with the inclusion of a b-spline function of time to control for residual temporal variation within  
80 strata, given the strong seasonality of WNV infection (Whitaker et al. 2007).

81 After observing the 2010-2015 cumulative epidemic curve, we firstly defined the transmission  
82 period of WNV, identifying the time interval going from the 27<sup>th</sup> to the 46<sup>th</sup> weeks of each year  
83 (length of 20 weeks). We secondly divided the identified period into 5 strata, each of 4 weeks  
84 length. For each week in which at least one human WNV case was reported (case period), we  
85 selected the other 3 weeks of the stratum as control periods. Exposure to meteorological  
86 variables, recorded in the capital of the province, were attributed to each case on the basis of  
87 the province in which her/his diagnosis was made.

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## 89 2.3 Statistical analysis

90 The analysis was performed using conditional Poisson regression (Armstrong et al. 2014).  
91 Since weather effects on infectious disease risk may be delayed (lag-effect), we studied the  
92 incidence of WNV infection in relation to meteorological data recorded during the 8 weeks  
93 prior to the diagnosis. Therefore, we implemented a conditional Poisson regression in the  
94 context of lag-distributed models, which are suitable to explore the delayed effect of an  
95 exposure. Specifically, we used distributed lag non-linear models (DNLM), two-dimensional  
96 models developed to explore exposure-lag-response relationships along both the dimensions of  
97 exposure and lag (Gasparrini et al. 2010; Imai et al. 2015). These models use a cross-basis

98 function, derived through a special tensor product of two independent functions, in order to  
99 analyze the exposure-response relationship and lag-response effect jointly. In our study, the  
100 effect of climatic parameters was modelled with a linear function, while the lag effect was  
101 modelled through a cubic basis spline with 4 degrees of freedom (df). The selection of the  
102 proper spline function for the lag-effect was based on the Akaike Information Criterion (AIC).  
103 We began the distributed lag models at lag 1 (the week before the week of diagnosis),  
104 hypothesizing that, since that WNV incubation period lasts 0-7 days (Rudolph et al. 2014), the  
105 risk should be null at lag 0 (week of diagnosis). The estimates can be plotted using a three-  
106 dimensional graph to show the Incidence Rate Ratio (IRR) along both exposure and lag  
107 dimension. Since the effect of climatic parameters was modelled as linear we estimated, for  
108 each lag, the IRR for an increase of 1 °C for the weekly average of minimum, mean and  
109 maximum temperatures and an increase of 5mm for the weekly total precipitation. The lag-  
110 specific IRR was derived by exponentiating the estimated regression coefficient, namely the  
111 variation in log-rate, for a unit increase of each climatic parameter for all specific lag (lag 1-  
112 8). In addition, we estimated the overall cumulative effect, that is the sum of each specific lag  
113 contribution over the whole lag period and can be interpreted as the overall risk. To control  
114 further for residual seasonal confounding, we included a cubic basis spline function with 5 df  
115 of the week number of the year, able to capture the seasonal pattern of the case distribution  
116 observed during the transmission period.

117 In addition, during summer holidays people are more likely to move out from their area of  
118 residence for leisure reasons. Thus, change of geographical location between the case and the  
119 control period would violate an assumption of the case-crossover design and possibly introduce  
120 bias. The potential impact of this source of bias was assessed in a sensitivity analysis in which  
121 we adjusted for holiday periods, defined as the two weeks around the 15<sup>th</sup> of August.

122 The software used to compute analysis is R, version 3.5.0 (R Development Core Team 2018).  
123 The packages used for statistical analysis are “splines” “dlnm” and “gnm”.

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### 3. Results

129 In total, 213 cases were diagnosed during the study period in Northern Italy and included in  
130 the case-crossover analysis. During 2010-2015 period, 25 provinces of Northern Italy out of  
131 42 (60%) reported human cases of WNV infection. Figure 1 shows the average of crude  
132 incidences of WNV infection per 1,000,000 inhabitants in each province over the 6-year period.  
133 Distribution of cases by week of the year (Fig 2) shows that the WNV infection has a seasonal  
134 pattern in Italy, with all cases being notified during the summer/autumn period. All human  
135 cases occurred between the 28<sup>th</sup> and 44<sup>th</sup> week of the year with a peak at the end of August (36<sup>th</sup>  
136 week). This pattern has suggested the inclusion of the spline function of time to further adjust  
137 seasonal confounding.

138 Results, both crude and adjusted for seasonality, conducted on climatic parameters recorded up  
139 to 8 weeks prior to the diagnosis in relation to the risk of WNV infection are shown in Figure  
140 3 and Table 1. The three-dimensional plots, show the entire surface of the adjusted IRRs in  
141 relation to maximum temperatures/precipitation at all lags considered (Figure 3a). Figure 3b  
142 shows the estimated effect of a unit increase in maximum temperatures and precipitation over  
143 the 8-week lag (continuous line: adjusted IRR, dashed line: crude IRR). Crude and adjusted  
144 lag-specific estimates for a unit increase in temperatures/precipitation are reported in Table 1.  
145 We found that the weekly average of maximum temperatures might affect the risk of WNV  
146 infection after 5 and 6 weeks (Fig 3). As shown in Table 1, the highest effect on WNV incidence  
147 was observed considering maximum temperatures recorded in the 6<sup>th</sup> week prior to diagnosis  
148 (adjusted IRR for 1°C increase in maximum temperatures at lag 6: 1.11; 95% CI 1.01-1.20).  
149 However, we did not find evidence of a positive overall cumulative effect for 1°C increase in  
150 maximum temperatures on WNV infection risk in the following weeks (Table 1). Weekly  
151 average of mean and minimum temperatures was not associated with the risk of WNV infection  
152 at any lag (Table 1). Weekly total precipitation recorded at lag 1-4 resulted positively  
153 associated with the risk of WNV infection (Fig 2b). As reported in Table 1, the maximum effect  
154 of precipitation was found with the precipitation recorded two weeks before diagnosis (lag 2)  
155 (adjusted IRR for 5 mm increase of weekly total precipitation at lag 2: 1.16; 95% CI 1.08-1.25).  
156 We found that 5 mm increase in weekly total precipitation was associated with a positive  
157 overall cumulative effect in the following 8 weeks: adjusted overall risk of 1.62 (95% CI 1.03-  
158 2.56). Lastly, when we adjusted for summer holidays in sensitivity analyses results were not  
159 affected more than marginally (results not shown).

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**Table 1**  
Risk of WNV infection in relation to unit increase<sup>a</sup> in temperature and precipitation.

<b>1°C increase in weekly average of maximum temperature</b>				
<b>Lag (Weeks)</b>	<b>IRR1<sup>b</sup></b>	<b>95% CI</b>	<b>IRR2</b>	<b>95% CI</b>
1	0.95	0.88-1.03	0.91	0.81-1.01
2	1.00	0.95-1.03	0.93	0.83-1.04
3	1.04	1.00-1.09	0.98	0.88-1.10
4	1.09	1.05-1.14	1.04	0.95-1.15
5	1.13	1.08-1.17	1.09	1.00-1.19
6	1.13	1.08-1.18	1.11	1.01-1.20
7	1.09	1.04-1.14	1.06	0.98-1.15
8	0.99	0.91-1.08	0.94	0.84-1.04
<b>Cumulative effect</b>	<b>1.48</b>	<b>1.22-1.80</b>	<b>1.03</b>	<b>0.56-1.87</b>
<b>1°C increase in weekly average of mean temperature</b>				
<b>Lag (Weeks)</b>	<b>IRR1</b>	<b>95% CI</b>	<b>IRR2</b>	<b>95% CI</b>
1	0.95	0.86-1.05	0.88	0.77-1.01
2	1.00	0.96-1.04	0.90	0.79-1.03
3	1.05	1.00-1.11	0.95	0.83-1.09
4	1.10	1.05-1.15	1.02	0.90-1.15
5	1.13	1.08-1.18	1.08	0.97-1.20
6	1.13	1.08-1.19	1.09	0.99-1.21
7	1.09	1.03-1.15	1.04	0.94-1.15
8	1.00	0.91-1.12	0.91	0.79-1.04
<b>Cumulative effect</b>	<b>1.53</b>	<b>1.23-1.92</b>	<b>0.86</b>	<b>0.41-1.80</b>
<b>1°C increase in weekly average of minimum temperature</b>				
<b>Lag (Weeks)</b>	<b>IRR1</b>	<b>95% CI</b>	<b>IRR2</b>	<b>95% CI</b>
1	0.96	0.86-1.07	0.91	0.80-1.05
2	1.01	0.96-1.06	0.90	0.79-1.03
3	1.06	1.00-1.12	0.93	0.81-1.07
4	1.10	1.05-1.15	0.98	0.86-1.12
5	1.12	1.08-1.17	1.03	0.92-1.15
6	1.12	1.07-1.18	1.04	0.93-1.17
7	1.09	1.03-1.16	1.00	0.89-1.12
8	1.02	0.92-1.15	0.88	0.75-1.02
<b>Cumulative effect</b>	<b>1.60</b>	<b>1.24-2.07</b>	<b>0.71</b>	<b>0.32-1.56</b>
<b>5 mm increase in weekly total precipitation</b>				
<b>Lag (Weeks)</b>	<b>IRR1</b>	<b>95% CI</b>	<b>IRR2</b>	<b>95% CI</b>
1	1.02	0.97-1.08	1.12	1.06-1.20
2	1.05	1.00-1.10	1.16	1.08-1.25
3	1.03	0.98-1.09	1.15	1.06-1.24
4	1.00	0.95-1.05	1.10	1.02-1.19
5	0.95	0.90-1.01	1.04	0.97-1.12
6	0.92	0.87-0.97	0.99	0.92-1.07
7	0.91	0.86-0.96	0.97	0.90-1.03
8	0.94	0.88-0.99	0.98	0.92-1.05
<b>Cumulative effect</b>	<b>0.82</b>	<b>0.57-1.14</b>	<b>1.62</b>	<b>1.03-2.56</b>

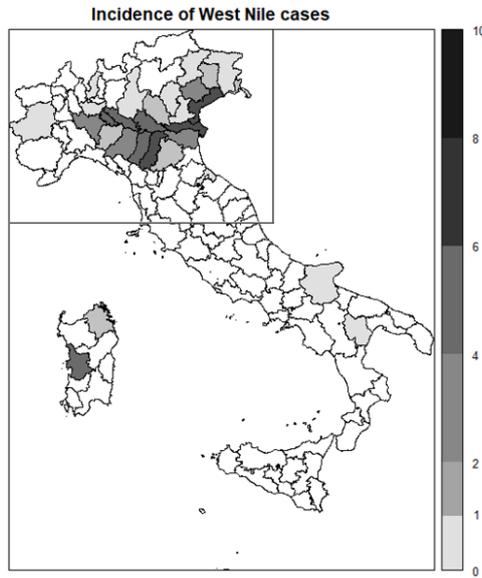
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<sup>a</sup> Estimates for a unit increase are derived by exponentiating the estimated regression coefficient, namely the variation in log-rate, for a unit increase of meteorological variables. Estimates for *n*-fold unit increase is obtainable by raising the estimate to the *n*-power

<sup>b</sup> IRR1: Crude Incidence Rate Ratio; IRR2: Incidence Rate Ratio adjusted for seasonality; CI: Confidence Interval

**Figure 1**

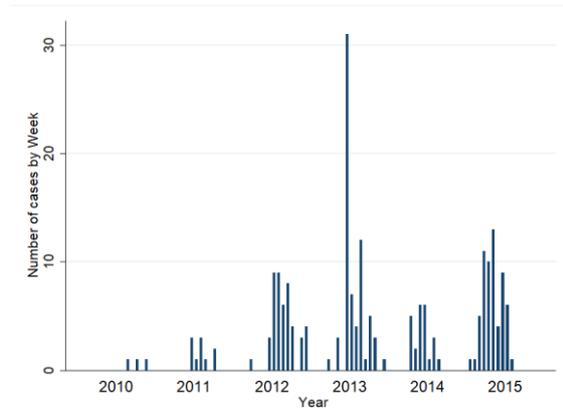
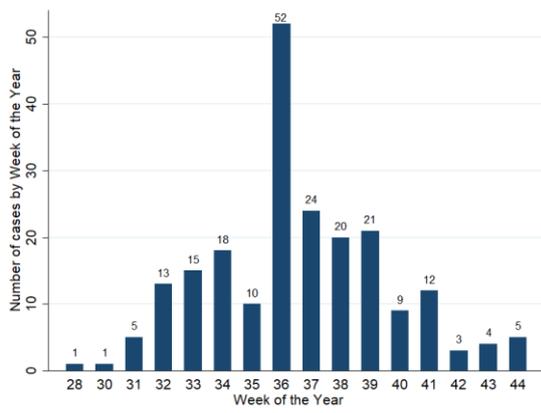
Average of crude incidences of WNV infection per 1,000,000 person-years in Italian provinces during the study period. Framed area corresponds to the study area.



**Figure 2**

Total number of WNV infection cases observed in Northern Italy during the study period (2010-2015) by week of the year (left) and by week and year (right)

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**Figure 3**

**Fig. 3a (left)** IRR2 (adjusted for seasonality) of WNV infection by weekly average of maximum temperatures (°C) and weekly total precipitation (mm), using a natural cubic spline–linear effect DLNM with 4 df basis cubic spline for lag and linear effect for exposure.

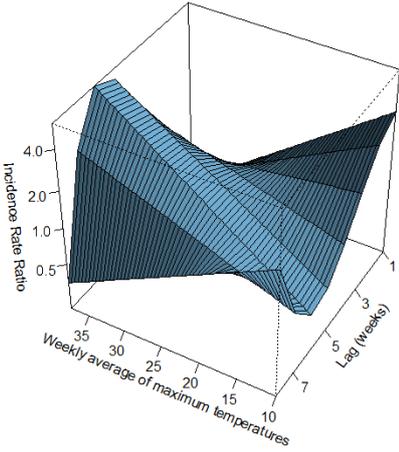
**Fig. 3b (right)** The estimated IRR2 (adjusted for seasonality) and 95% confidence intervals in unit increase of weekly average of maximum/minimum temperature (1 °C) and of weekly total precipitation (5mm) over 8 weeks of lag. Dashed line: IRR1 (not adjusted for seasonality)

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**Figure 3**

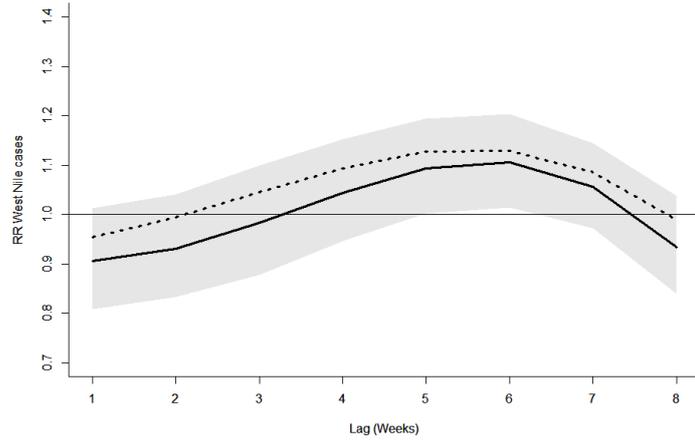
**Figure 3a**

*3D Graph of effect of Max T*

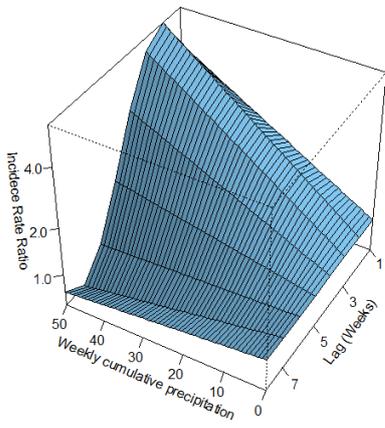


**Figure 3b**

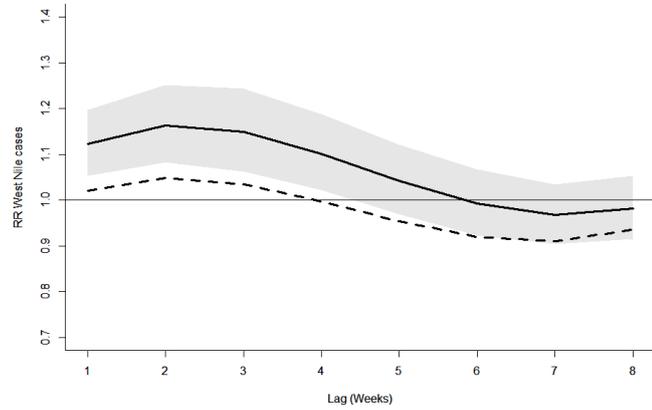
*Lag effect of 1°C increase in weekly average of Max T*



*3D Graph of effect of Tot Prec*



*Lag effect of 5mm increase in weekly Tot Prec*



170 **4. Discussion**

171 Our study revealed that cases in Northern Italy are notified between July and October, with a  
172 peak at the end of August. The transmission season is similar to the activity period (May-  
173 November) of mosquito *Culex Pipiens*, the main WNV vector in Italy (Bisanzio et al. 2011).

174  
175 Our study is, to our knowledge, the first to assess the lag-effect of meteorological exposures  
176 and risk of WNV infection in Italy, including all incident cases diagnosed in Northern Italy  
177 between 2010 and 2015. Methodologically, the main strength of this study is the application of  
178 DLNMs in the context of a time stratified case-crossover design in order to explore delayed  
179 effects of exposures. We further included in the model a seasonal term (namely a spline  
180 function of time) to enhance the study validity, as it has been shown that in presence of a strong  
181 seasonal pattern of exposures and outcomes, time-stratified case-crossover studies might still  
182 be biased by residual seasonal confounding (Whitaker et al. 2007). Since we were interested in  
183 evaluating the short-term effect of the weekly variation of climatic parameters on the incidence  
184 of WNV infection from here onwards we will discuss only results adjusted for seasonality.

185  
186 We found evidence of association, despite no overall cumulative effect, between maximum  
187 temperatures recorded in the 5<sup>th</sup> and 6<sup>th</sup> weeks prior to diagnosis (lags 5 and 6) and the  
188 incidence of WNV infection. Several studies have evaluated the effect of the temperatures on  
189 WNV ecology and transmission among mosquitoes, birds and humans in different areas  
190 worldwide (Gubler 2007; Paz 2015a; Paz and Semenza 2013), and many of them showed that  
191 temperatures may play an important role in the virus transmission cycle. However, only few  
192 studies have assessed the risk of WNV infection in humans in relation to temperatures with  
193 the specific aim of evaluating the lag effect. One correlation study conducted in Israel,  
194 Greece, Romania and Russia analyzed human cases of WNV infection notified during the  
195 summer of 2010 in relation to temperature anomalies, namely temperatures recorded in 2010  
196 compared with the perennial weekly average of 1981–2010. This study found an association  
197 between WNV cases and temperature at lag 0-1 (weeks) in Israel and Greece and at lag 3-4  
198 (weeks) in Romania and Russia (Paz et al. 2013). One US study, a bidirectional case-  
199 crossover, not adjusted for seasonality, analyzed all incident cases of WNV infection notified  
200 between 2001 and 2005 (n= 16.298) in relation to the temperatures recorded in the 4 previous  
201 weeks, finding associations of similar strength for each lag (0-4 weeks) (Nile et al. 2009).

202 The lag of 5-6 weeks observed in our study might be explained by the complexity of the  
203 host/pathogen ecology. However, our study was not designed to assess the underlying  
204 mechanisms through which temperatures and precipitation may affect WNV infection, thus we  
205 can only speculate on the effects of climate parameters on vector and virus ecology.

206 It has been observed that the air temperature can augment virus replication rate and lead to  
207 higher viremia level in mosquito population (Reisen et al. 2006). Higher temperatures have  
208 been also shown to impact the vector transmission rate, by shortening the extrinsic incubation  
209 period (namely “the time from ingestion of an infectious bloodmeal until a mosquito is capable  
210 of transmitting virus infection to a susceptible organism”) (Reisen 1989, Reisen et al. 2006).  
211 In addition, elevated temperatures can cause an expansion of the absolute number of  
212 mosquitoes and affect their feeding behaviours (Bisanzio et al. 2011; Conte et al. 2015). Thus,  
213 higher temperatures are believed to first impact the virus transmission in the enzootic cycle  
214 among mosquitoes and birds (Kilpatrick et al. 2008; Reisen et al. 2006) and, second, to affect  
215 the expansion of the proportion of infective mosquitoes, on which depend the human infection.  
216 The aforementioned pathways intrinsically imply a latency of the effect that, in addition to an  
217 incubation period of 0-7 days of human infection (Rudolph et al. 2014), might explain the  
218 overall latency of 5-6 weeks observed between increased temperatures and higher incidence of  
219 WNV infection cases.

220 However, it is noteworthy that the whole lag pattern presents negative point estimates at lag 1-  
221 2 and that the overall cumulative effect estimate is close to zero. For these reasons we cannot  
222 exclude that our findings of association between increased maximum temperatures and  
223 incidence of WNV infection at lag 5-6 might be due to chance.

224

225 Our results revealed an association between WNV infection and total precipitation recorded  
226 between the 1 and 4 weeks prior the diagnosis (lag 1-4). Levels of precipitations are believed  
227 to affect the patterns and the transmission of WNV (Paz 2015). However, findings about the  
228 relationship between precipitation and incidence of WNV cases are contradictory. Some  
229 studies reported that above-average precipitation can lead to higher risk of WNV outbreaks by  
230 expanding mosquitoes (Di Sabatino et al. 2014; Nile et al. 2009). On the contrary, other studies  
231 found that drought periods can induce outbreaks favoring the bird-to-bird viral transmission by  
232 facilitating the concentration of avian species in the few existing pools (Shaman et al. 2005). It  
233 is plausible that the response to precipitation might change over different geographical areas,  
234 depending on the differences in the characteristics of the local environment and in the ecology  
235 of vectors (Shaman et al. 2002, Paz 2015). Our results of associations between WNV infection

236 cases and increased precipitation at lag 1-4 (weeks) can be due to the close relationship between  
237 aquatic environment and mosquito proliferation. Intermediate stages of Culex mosquitoes, such  
238 as larvae, are water dependent, and therefore, precipitation might be important, especially in  
239 drought periods such as summer, to create and maintain water pools that are necessary for the  
240 development of mosquitoes. Accordingly, an observational study reported that the WNV  
241 outbreak recorded in 2010 in central Macedonia, Greece, was preceded by unusually  
242 precipitation (Danis et al 2011).

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244 Our study has three main limitations. First, we had information on the week but not on the day  
245 of diagnosis. Thus, we could not date back the exposure history starting from the day of  
246 symptoms onset, but only from the week preceding the week of the diagnosis. However, our  
247 study aligns with most of environmental studies conducted on infectious diseases, as typically  
248 surveillance systems for communicable diseases notify cases on a weekly scale. Second, since  
249 we had no information about the municipality but only about the province of residence of the  
250 cases, we linked each case to the meteorological station of the capital of its province in order  
251 to obtain data on the corresponding environmental exposures. This linkage might have  
252 introduced some non-negligible degree of exposure misclassification. However, since in case-  
253 crossover analysis the same subject is used both as case and as its own control, misclassification  
254 is likely to be non-directional, which would likely lead to conservative estimates. Third, the  
255 reason of the diagnosis (asymptomatic subjects: WNV positive blood; symptomatic subjects:  
256 West Nile Fever or West Nile Neuroinvasive Disease) was not available at the individual level.  
257 Asymptomatic subjects, such as blood donors, can be diagnosed during the incubation period,  
258 and therefore the lag-effect of environmental exposures might be different between  
259 asymptomatic and symptomatic groups. However, WNV infection cases diagnosed among the  
260 blood donors represent a minority of cases identified through the surveillance system. For  
261 instance, only 13 out of 61 cases (21% of the total) observed in Italy in 2015 were blood donors  
262 (ISS, 2015).

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270 **5. Conclusions**

271 In conclusion, our results suggest that high temperatures might be associated with the incidence  
272 of WNV infection after a lag of 5-6 weeks, while heavy precipitation after a lag of 2-3 weeks.  
273 These results strengthen the evidence that the WNV is a climate-sensitive disease in an area  
274 where the process of endemization has recently started and underline that climatic parameters  
275 might be useful for detecting areas and periods of the year potentially characterized by a higher  
276 incidence of WNV infection

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