Seasonality of acute kidney injury in a tertiary hospital academic center: an observational cohort study

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Abstract

Background: The aim of our study was to describe seasonal trends of acute kidney injury (AKI) and its relationship with weather conditions in a hospitalized population.

Methods: We retrospectively collected demographic (age, sex), clinical (ICD-9-CM codes of diagnosis discharge) and laboratory data (creatinine values) from the inpatient population admitted to Fondazione Policlinico Universitario A. Gemelli IRCCS between January 2010 and December 2014 with inclusion of all patients ≥ 18 years with at least two values available for creatinine. The outcome of interest was AKI development, defined according to creatinine kinetics criteria. The exposures of interest were the months and seasons of the year; air temperature and humidity level were also evaluated. Log-binomial regression models adjusted for age, sex, eGFR, comorbidities, Charlson/Deyo index score, year of hospitalization were used to estimate risk ratios (RR) and 95% confidential intervals (CI).

Results: A total of 64,610 patients met the inclusion criteria. AKI occurred in 2864 (4.4%) hospital admissions. After full adjustment, winter period was associated with increased risk of AKI (RR 1.16, 95% CI 1.05, 1.29, p = 0.003). Lower air temperature and higher humidity level were associated with risk of AKI, however in multivariable-adjusted models only higher humidity level showed a significant and independent association.

Conclusions: AKI is one of the most common complications of hospitalized populations with a defined seasonal pattern and a significant increase in incidence during wintertime; weather conditions, particularly higher humidity level, are independent predictors of AKI and could partially justify the observed seasonal variations.

Keywords: Acute kidney injury, Epidemiological study, Seasonality, Weather conditions

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Background

Acute kidney injury (AKI) is a complex clinical syndrome characterized by an abrupt reduction of renal function [1]. It is a common and severe complication in hospitalized populations. The reported incidence is quite varied especially since it is strongly influenced by definition and study population with an incidence ranging from 7 to 57% [2–6] and with a significant burden on patient morbidity and mortality [7–9]. As reported by clinical and epidemiological studies, several diseases [10–14] show a defined seasonal variation. “Winter peaks” have been observed in cardiovascular [13], respiratory [14] and infectious [10] diseases.

Understanding the impact of seasonal patterns of diseases could have a role in clinical practice and preventive care. However, despite the importance of AKI, to date only a Japanese retrospective study on an inpatient population has investigated seasonal variations of AKI describing a peak of incidence during wintertime [15]. Although the observed pattern may be in part related to seasonality of underlying illnesses associated with kidney injury, as reported in the aforementioned study, a significant relationship between AKI and wintertime persisted even after adjustment for potential confounders (demographic factors, comorbidities and medications). It may be reasonable to speculate that even environmental factors such as weather conditions might partially influence AKI occurrence.

Therefore, the aim of our study was to describe trend and seasonality of AKI in an Italian cohort of hospitalized patients and to investigate the possible relationship between AKI and weather conditions.

Methods

Study population

We performed a retrospective observational cohort study on the inpatient population admitted to Fondazione Polyclinico Universitario A. Gemelli IRCCS (Rome, Italy), a tertiary level hospital serving more than 1 million people in Rome collecting patients all around the city, between January 1, 2010 and December 31, 2014. We included only adult patients (18 years or older) with at least two measurements available for serum creatinine (Cr) during hospital stay. Patients with End Stage Kidney Disease (ESKD) were excluded. For patients with multiple hospital admissions, only the first one was considered.

Data collection

All clinical, demographic and laboratory data were extracted from the electronic hospital database. We exported the following variables for each hospital admission: age, sex, ICD-9-CM codes of primary and secondary discharge diagnoses, Cr, date of hospital admission. Using the Italian meteorological historical records [16] we extracted the following weather variables, at city level, during the same period: average monthly air temperature and relative humidity levels.

Definitions

AKI diagnosis and severity were defined according to creatinine kinetics (CrK) criteria [17] as an absolute increase of Cr concentration during hospitalization. To do this, we calculated the difference between each Cr and the previous measured value during hospitalization. According to CrK criteria, we defined AKI as an absolute increase in Cr of 0.3 mg/dL over 24 h or a 0.5 mg/dL increase over 48 h.

ESKD was identified according to ICD-9-CM codes in primary and secondary discharge diagnoses.

Estimated glomerular filtration rate (eGFR) was evaluated in all patients at hospital admission by the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) equation [18].

The seasons were classified according to the month of the year as spring (March, April, May), summer (June, July, August), autumn (September, October, November) and winter (December, January, February).

ICD-9-CM codes in primary and secondary discharge diagnoses were used to identify comorbidities (cardiovascular diseases, endocrine/metabolic diseases, malignancies, genito/urinary diseases, infectious diseases, respiratory diseases). Charlson/Deyo comorbidity index score [19] was calculated for each patient using ICD-9-CM codes in primary and secondary discharge diagnoses.

The outcome of interest was AKI development (reported as categorical variable [yes/no] in the regression models). The exposures of interest were the months/seasons of the year; secondly, we evaluated air temperature and relative humidity level as exposures. Covariates assessed to control for confounding were age, sex, baseline eGFR, comorbidities, Charlson/Deyo index score, year of hospitalization.

Statistical analysis

Continuous variables were reported as means and standard deviations (SD) or medians and interquartile ranges (IQR) as appropriate. Categorical variables were described using numbers and percentages (%). Student t-test and chi-square ($\chi^2$) test were used to compare continuous and categorical distributions in the descriptive analysis between two groups; two-way ANOVA was performed for multiple groups comparisons (three or more groups).
Log-binomial regression models adjusted for all covariates were used to estimate risk ratios (RRs) and 95% confidential intervals (CIs) of months/seasons (June and summer were selected as the reference groups because it contained the smallest proportion of patients with AKI among hospitalized patients). Because AKI occurrence vary largely across different age and sex groups, and hence the seasonal pattern of AKI in different groups may vary, stratified analysis by sex (male vs female) and age (more or less 60 years old) were performed.

Multivariable regression models were subsequently fitted to explore the association between weather conditions (analyzed as continuous and categorical variable grouped in quartiles) and AKI. Three models were built: Model 1 represents unadjusted RRs; Model 2 was adjusted for all covariates (age, sex, eGFR, comorbidities, Charlson/Deyo score, year of hospital admission); Model 3 was adjusted for relative humidity and mean temperature in addition to factors included in Model 2. Ultimately, mediation analysis was performed to evaluate the "mediator role" of weather conditions on AKI seasonality and estimate the average causal mediation effect (ACME) as measurement of the amount of the indirect (or mediate) effect or as the amount of the reduction of the causal variable effect (direct effect, [season period]) on the outcome of interest. Confidence intervals were calculated with bootstrapping approach and a high number (500) of simulations. We use the linear regression fit with least squares and the binomial regression for the mediator and outcome models, respectively.

Statistical analyses were performed using R version 3.4.4 (Free software Foundation, California). A p-value < 0.05 was considered as statistically significant.

Results

Descriptive measurements of study population

Overall, 64,610 out of 201,304 patients met our inclusion criteria (Fig. 1). AKI occurred in 2864 (4.4%) patients. Patients with AKI had higher comorbidity index score (0.8 [1.1] vs 0.3 [0.8], p<0.001) with an increase prevalence of cardiovascular diseases (1747 [61.0%] vs 23,208 [37.6%], p<0.001), endocrine/metabolic diseases (702 [24.5%] vs 10,680 [17.3%], p<0.001), genitourinary disorders (782 [27.3%] vs 6876 [11.1%], p<0.001), infectious diseases (269 [9.4%] vs 3007 [4.9%, p<0.001], respiratory diseases (698 [24.4%] vs 7580 [12.3%], p<0.001). AKI was also more commonly observed in older individuals (70.7 [13.9] years vs 60.2 [18.0] years, p<0.001) and males (1771 [61.8%] vs 28,149 [45.6%, p<0.001) (Table 1).

Fig. 1 Flowchart of the study
Overall 17,213 patients were admitted during spring period, 14,674 during summer period, 16,330 in autumn and 16,393 in winter. As reported in Table 2, patients hospitalized during spring and winter period showed a higher comorbidity index score and a higher prevalence in cardiovascular diseases. In patients admitted during winter period we observed an higher prevalence in respiratory diseases. Summertime was instead characterized by a higher prevalence in infectious diseases.

### Association between seasonality and AKI

AKI was more commonly observed during wintertime [788 (4.8%)] compared with other seasons [spring 776 (4.5%), summer 597 (4.1%), autumn 703 (4.3%)] (Table 3). As reported in Table 3, the winter period was significantly associated with higher risk of AKI (RR 1.18, 95% CI 1.07, 1.31, \(p=0.002\)) (Table 3). This association was confirmed even after correction for potential confounders (RR 1.15, 95% CI 1.04, 1.38, \(p=0.006\)). No significant relationships were observed after comparing other seasons to each other (Supplementary Table 1). Stratified analysis, by age and sex, demonstrated a significant and independent increase in AKI risk during wintertime only in male subjects older than 60 years (Supplementary Table 2).

The risk of AKI was significantly higher in January (RR 1.28, 95% CI 1.08, 1.57, \(p=0.004\)) [Fig. 2]) than in the rest of the year.

### Table 1  Descriptive characteristics of study population stratified by AKI

|              | No AKI | AKI   | \(p\) value |
|--------------|--------|-------|-------------|
| \(n\)        | 61,746 | 2864  |             |
| Age, years, mean (SD) | 60.2 (18.0) | 70.7 (13.9) | <0.001     |
| Sex, Males, n (%) | 28,149 (45.6) | 1771 (61.8) | <0.001     |
| Charlson/Deyo score, n (%) |                  | <0.001     |
| 0            | 48,693 (78.9) | 1662 (58.0) |             |
| 1            | 9190 (14.9) | 624 (21.8) |             |
| 2            | 2199 (3.6) | 319 (11.1) |             |
| > 2          | 1664 (2.7) | 259 (9.0) |             |
| Charlson/Deyo score, mean (SD) | 0.3 (0.8) | 0.8 (1.1) | <0.001     |
| Comorbidity, n (%) |        |       |             |
| Cardiovascular | 23,208 (37.6) | 1747 (61.0) | <0.001     |
| Gastrointestinal | 9487 (15.4) | 410 (14.3) | 0.134      |
| Malignancies  | 19,471 (31.5) | 758 (26.5) | <0.001     |
| Endocrine/Metabolic | 10,680 (17.3) | 702 (24.5) | <0.001     |
| Genito/Urinary | 6876 (11.1) | 782 (27.3) | <0.001     |
| Infectious    | 3007 (4.9) | 269 (9.4) | <0.001     |
| Respiratory   | 7580 (12.3) | 698 (24.4) | <0.001     |
| eGFR, ml/min/1.73m², median (IQR) | 82.0 (35.0) | 50.0 (43.0) | <0.001     |

\(\text{eGFR at hospital admission}\)

### Table 2  Descriptive characteristics of study population stratified by season

|              | Spring | Summer | Autumn | Winter | \(p\) value |
|--------------|--------|--------|--------|--------|-------------|
| \(n\)        | 17,213 | 14,674 | 16,330 | 16,393 |             |
| Age, years, mean (SD) | 60.8 (17.8) | 60.7 (18.2) | 60.1 (18.0) | 61.0 (17.9) | <0.001     |
| Sex, Males, n (%) | 8005 (46.5) | 6831 (46.6) | 7475 (45.8) | 7609 (46.4) | 0.463     |
| Charlson/Deyo score, n (%) |                  | <0.001     |
| 0            | 13,332 (77.5) | 11,556 (78.8) | 12,870 (78.8) | 12,597 (76.8) |             |
| 1            | 2631 (15.3) | 2190 (14.9) | 2579 (14.6) | 2614 (15.9) |             |
| 2            | 701 (4.1) | 538 (3.7) | 624 (3.8) | 655 (4.0) |             |
| > 2          | 549 (3.2) | 390 (2.7) | 457 (2.8) | 527 (3.2) |             |
| Charlson/Deyo score, mean (SD) | 0.4 (0.9) | 0.3 (0.8) | 0.3 (0.8) | 0.4 (0.9) | <0.001     |
| Comorbidity, n (%) |        |       |       |       |             |
| Cardiovascular | 6780 (39.4) | 5506 (37.5) | 6135 (37.6) | 6534 (39.9) | <0.001     |
| Gastrointestinal | 2675 (15.5) | 2232 (15.2) | 2480 (15.2) | 2510 (15.3) | 0.8     |
| Malignancies  | 5365 (31.2) | 4536 (30.9) | 5160 (31.6) | 5168 (31.5) | 0.526     |
| Endocrine/Metabolic | 3122 (18.1) | 2533 (17.3) | 2805 (17.2) | 2922 (17.8) | 0.067     |
| Genito/Urinary | 2011 (11.7) | 1733 (11.8) | 1992 (12.2) | 1922 (11.7) | 0.453     |
| Infectious    | 876 (5.1) | 816 (5.6) | 778 (4.8) | 806 (4.9) | 0.01     |
| Respiratory   | 2241 (13.0) | 1790 (12.2) | 1945 (11.9) | 2302 (14.0) | <0.001     |
| eGFR, ml/min/1.73m², median (IQR) | 81.0 (37.0) | 79.0 (37.0) | 81.0 (36.0) | 82.0 (36.0) | <0.001     |

\(\text{eGFR at hospital admission}\)
### Table 3 Association between AKI and seasons

|       | Spring | Summer | Autumn | Winter |
|-------|--------|--------|--------|--------|
| n     | 17,213 | 14,674 | 16,330 | 16,393 |
| AKI   | 776 (4.5) | 597 (4.1) | 703 (4.3) | 788 (4.8) |
| RR (95% CI) | 1.11 (1.00, 1.23) | 1.00 (Reference) | 1.06 (0.95, 1.18) | p=0.300 | 1.18 (1.07, 1.31) | p=0.002 |
| RR (95% CI)* | 1.09 (0.99, 1.21) | p=0.100 | 1.00 (Reference) | 1.07 (0.97, 1.19) | p=0.200 | 1.16 (1.05, 1.29) | p=0.003 |

*Adjusted for age, sex, eGFR comorbidities, Charlson/Deyo score, year of hospital admission

### AKI and weather conditions
As depicted in Fig. 3 an association between air temperature and relative humidity level can be describe. Progressive increase in the average monthly relative humidity level and a progressive decrease in the average monthly air temperature was associated with increased AKI occurrence (Fig. 3).

Multivariable regression model demonstrate a significant and association between air temperature and relative humidity level with the risk of AKI (RR 1.16, 95% CI 1.05, 1.28, p=0.004, in the highest quartile of humidity level compared with the lowest; RR 1.15, 95% CI 1.04, 1.28, p=0.008, in the lowest quartile of air temperature compared with the highest; p-value for trend=0.003; Table 4).

However, in multivariable model fully adjusted for weather factors (Model 3, Table 4), only humidity level (RR 1.15, 95% CI 1.01, 1.82, p=0.038, in the highest quartile of humidity level compared with the lowest) was independently associated with a higher risk of AKI.

In order to evaluate the “complete mediation effect” played by humidity we performed a mediation analysis (after controlling for correlation between humidity level and winter period, β=9.29, p<0.001) that confirmed the significant mediator effect (or indirect effect) of humidity level (ACME average, point estimate 0.0026, 95% CI 0.0005, 0.0100, p=0.028) with an estimate mediated proportion of 0.52 (as ratio between the indirect “mediated” effect and the total effect) on the relationship between AKI and winter period.

### Discussion
Our study describes the relationship between AKI and seasonality and between AKI and weather conditions, air temperature and humidity level.

![Fig. 2 Multivariable-adjusted risk ratios (95% confidence of interval) of AKI associated with month of the year; adjusted for age, sex, comorbidities, eGFR, year of hospital admission, Charlson/Deyo score; June used as reference](image)
The incidence of AKI increases during wintertime; weather conditions, such as air temperature and relative humidity level are associated with AKI; humidity level is an independent predictor of AKI occurrence and could partly explain and mediate the observed seasonal variations.

Seasonality patterns in human illnesses are a well-recognized phenomenon since Hippocrates [20]. In particular, variations in the seasonal patterns of cardiovascular diseases, infectious diseases, respiratory diseases have been already described [10–14]. Most studies reported “winter peaks” in cardiovascular [13], respiratory [14] and infectious [10] related hospitalizations and mortality. A complex interaction between individual susceptibility and environmental factors could justify such relationship.

Recently, a Japanese study [15] was the first to describe a seasonality pattern of AKI. The authors, analyzing a historical cohort of hospitalized patients, documented an increase in incidence rates of AKI during winter period. As reported, an increase in cardiovascular and pulmonary diseases during wintertime, might partly explain such association. Although not investigated, authors concluded that weather conditions should also be considered.

In our study AKI occurred in 4.4% (2864/64,610) of hospitalized population. Cardiovascular diseases were the most common disorders associated with kidney injury. As reported in Table 3 and Fig. 2, and similarly to the aforementioned Japanese study, we observed a “winter peak” in kidney injury. Although this relationship could partly be justified by an increase incidence in common diseases generally associated AKI during wintertime (we found an increased prevalence in AKI patients of respiratory and cardiovascular diseases in winter, [Table 2]), an independent association persisted even after adjustment for all covariates.

In order to better understand this independent seasonal variation, we investigated the possible relationship between kidney injury and weather conditions. We
found that higher relative humidity level and lower air temperature were associated with risk of AKI development. However, after multivariable adjustment, only humidity level showed an independent relationship with kidney injury. Furthermore, a "mediate role of humidity level on the relationship between winter period and AKI emerges from the analysis. Even if mediation analysis does not imply causal relationships (unless it is based on experimental design) a significant mediation of humidity level on the relationship between winter period and AKI development.

Some limitations in our study must be considered. Ours is a monocentric study with a retrospective design; furthermore, we used ICD-9-CM codes for comorbidities assessment and for risk adjustment. On the other hand, we determined the Charlson/Deyo comorbidity index, a valid score based on ICD-9-CM diagnosis codes. Furthermore, we were unable to define the representativeness of our hospital for database limitations.

Ours is the first study describing AKI seasonality in a European cohort and the first one demonstrating an association between weather conditions (humidity levels) and AKI development.

**Conclusions**

We conclude that AKI has a seasonal pattern and a significant increase in incidence during wintertime; the seasonality is not explained by common clinical factors. On the other hand, weather conditions, particularly humidity level, are associated with the observed seasonal variations. It remains to be understood if such an association is independent or due to other unknown variables. The climate and its influence on body physiology could represent a new element to be considered and investigated in AKI epidemiology.

### Table 4 Association between weather conditions and AKI

| Humidity level (%) | No. of patients | No. of events | Model 1 | | Model 2 | | Model 3 |
|--------------------|----------------|---------------|---------|---------|---------|---------|---------|
| - Q1: < 63.0       | 16,777         | 656 (4.5)     | 1.00    | <0.001  | 1.00    | 0.003   | 1.00    | 0.064   |
| - Q2: 63-68.2      | 15,643         | 711 (4.5)     | 1.16 (1.05, 1.29) | p = 0.005 | 1.17 (1.06, 1.30) | p = 0.002 | 1.17 (1.05, 1.31) | p = 0.006 |
| - Q3: 68.2-72.8    | 16,497         | 774 (4.7)     | 1.20 (1.08, 1.33) | p < 0.001 | 1.20 (1.08, 1.32) | p < 0.001 | 1.18 (1.03, 1.35) | p = 0.016 |
| - Q4: > 72.8       | 15,693         | 723 (4.6)     | 1.18 (1.06, 1.31) | p = 0.002 | 1.16 (1.05, 1.28) | p = 0.004 | 1.15 (1.01, 1.32) | p = 0.038 |

| Air temperature (°C) | No. of patients | No. of events | Model 1 | | Model 2 | | Model 3 |
|-----------------------|----------------|---------------|---------|---------|---------|---------|---------|
| - Q1: < 12.1          | 16,476         | 787 (4.8)     | 1.19 (1.07, 1.32) | p = 0.002 | 1.15 (1.04, 1.28) | 0.009 | 1.03 (0.89, 1.19) | 0.675 |
| - Q2: 12.1-16.9       | 16,603         | 742 (4.5)     | 1.11 (1.00, 1.24) | p = 0.056 | 1.08 (0.98, 1.20) | p = 0.136 | 0.99 (0.87, 1.13) | 0.867 |
| - Q3: 16.9-23.8       | 17,424         | 767 (4.4)     | 1.09 (0.98, 1.22) | p = 0.100 | 1.10 (0.99, 1.22) | 0.081 | 1.01 (0.89, 1.14) | 0.901 |
| - Q4: > 23.8          | 14,107         | 568 (4.0)     | 1.00     | (Reference) | 1.00     | (Reference) | 1.00     | (Reference) |

Model 1: unadjusted
Model 2: Multivariable regression model adjusted for age, sex, eGFR, comorbidities, Charlson/Deyo score, year of hospital admission
Model 3: Fully adjusted model, multivariable regression model adjusted for relative humidity and mean temperature in addition to factors included in Model 2
Supplementary Information

The online version contains supplementary material available at https://doi.org/10.1186/s12940-021-00691-5.

Additional file 1: Table S1. Association between AKI and seasons.

Supplementary Table 2. Association between AKI and seasons, stratified analysis by age and sex

Abbreviations

AKI: Acute kidney injury; Cr: Serum creatinine; ESKD: End Stage Kidney Disease; ICD-9-CM: International Classification of Diseases, Ninth Revision, Clinical Modification; CrK: Creatinine kinetics; eGFR: Estimated glomerular filtration rate; CKD-EPI: Chronic Kidney Disease Epidemiology Collaboration

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Authors' contributions

G.G., P.M.F. and G.L. contributed to the research idea and study design. G.L. contributed to data acquisition. G.G., P.M.F. and G.L. contributed to data analysis/interpretation. P.M.F. and G.L. contributed to statistical analysis. G.L. drafted the paper. G.G., P.M.F., A.N., N.P. and M.B. contributed to supervision. Each author contributed important intellectual content during manuscript drafting for the overall work. The authors read and approved the final manuscript.

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Availability of data and materials

The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate

The ethics committee of Fondazione Policlinico Universitario A. Gemelli IRCCS approved the research protocol (Prot. number 34327/18 ID 2210).

Consent for publication

All of the authors have reviewed and approved the manuscript for publication.

Competing interests

The authors declare that they have no competing interests.

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