Association between winter season and risk of death from cardiovascular diseases: a study in more than half a million inpatients in Beijing, China

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Background: Seasonal associations of cardiovascular mortality have been noted in most populations of European origin years ago, but are not well evaluated in Asian populations recently.

Methods: Utilizing the electronic Hospitalization Summary Reports (HSRs) from 32 top-ranked hospitals in Beijing, China, we evaluated the association between winter season and the risk of cardiovascular death among hospitalized individuals. General additive models and logistic regression models were adjusted for confounding factors.

Results: Older patients who were admitted to the hospital in the winter months (January, February, November and December) had a death risk that was increased by approximately 30% to 50% (P < 0.01) over those who were admitted in May. However, younger patients did not seem to experience the same seasonal variations in death risk. The excess winter deaths among older patients were associated with ischemic heart disease (RR = 1.22; 95% CI 1.13 to 1.31), pulmonary heart disease (RR = 1.42; 95% CI 1.10 to 1.83), cardiac arrhythmias (RR = 1.67; 95% CI 1.36 to 2.05), heart failure (RR = 1.30; 95% CI 1.09 to 1.54), ischemic stroke (RR = 1.30; 95% CI 1.17 to 1.43), and other cerebrovascular diseases (RR = 1.78; 95% CI 1.40 to 2.25). The risks of mortality were higher in winter months than in the month of May, regardless of the presence or absence of respiratory disease.

Conclusions: Winter season was associated with a substantially increased risk of cardiovascular death among older Chinese cardiovascular inpatients.

Keywords: Cardiovascular disease, Winter, Seasonality, Older adults, Mortality, Asian population

Background
Excess winter mortality from cardiovascular diseases has been an active research area since 1960s [1-9]. Some studies have concluded that the increase in winter deaths results from cold indoor and outdoor temperatures, while others posited that the increased winter mortality may be associated with epidemics of respiratory syncytial virus and influenza [10-15]. Studies are difficult because cardiovascular and respiratory diseases are codependent disorders and often coexist in one patient. Prior studies of seasonal variation used death certification data, which did not record disease diagnoses but recorded a primary cause of death [13-15].

Many previous studies reporting seasonal variations of cardiovascular mortality were conducted in Europe [1,4,5] and North America [2,3,6], while a few were conducted in Asia [7-9]. In those countries, excess seasonal cardiovascular mortality varies widely from 5% to 30% [16]. Moreover, excess winter mortality is reported to be relatively lower in Scandinavian countries than in the UK [17,18]. In addition to cold temperatures, the variations among countries may be related to differences in socioeconomic factors, healthcare systems, or individual lifestyles [19]. However, most previous studies of seasonal variations and health were conducted years ago, and there is little current information [1-9]. More importantly, existing studies focused on either...
overall or some specific cardiovascular or cerebrovascular deaths without accounting for other circulatory diseases such as pulmonary heart disease and rheumatic heart disease [1-9].

There has been a dramatic increase in the incidence of cardiovascular disease in China over recent decades [20]. Cardiovascular diseases have become the largest cause of hospitalization and mortality in China, and are expected to remain the dominant trend [20]. Due to the lack of high quality data, the association between winter season and deaths from cardiovascular disease has not been well evaluated among the Chinese population (20% of the world population). This study used data from 626,950 cardiovascular inpatients from 32 top-ranked general hospitals in Beijing to investigate whether associations between winter season and cardiovascular deaths were modified by age and cardiovascular subtypes and to determine the fraction attributable to respiratory diseases.

**Methods**

**Data source and study subjects**

Data were obtained from electronic Hospitalization Summary Report (HSR) in 32 top-ranked (Grade 3A) hospitals in Beijing from Jan 1, 2006 to Dec 31, 2010. Hospitals in China are ranked by the existing hospital hierarchical management approach. For both public and private hospitals, the rank is assessed according to medical services and management, quality and safety of clinical care, and technical level and efficiency. There are three grades and ten classes, and the first class hospitals are designated Grade 3A, which is the highest rank except the national special hospitals in China.

To fulfill the administrative requirements of the China Ministry of Health, every hospital in Beijing must submit an electronic HSR to a centralized health information system of the Beijing Municipal Health Bureau (full document on the official website: http://www.bjhb.gov.cn/zwfwq/ztlj/drg/ppt/201110/t20111008_41256.htm). The standard summary report is documented with basic demographics, dates of admission and discharge, pre- and post-hospitalization diagnoses, discharge status, treatments, and financial costs. This study is considered exempt since it used data collected for administrative purpose without any personal identifiers.

The HSR diagnosis was coded according to the International Classification of Diseases, 10th Revision, Clinical Modification (ICD-10-CM). The HSR contains a maximum of eight discharge diagnoses with the first diagnosis listed designated as the principal diagnosis, while the rest diagnoses were supplementary. Cardiovascular inpatients were identified if the first-listed diagnoses were cardiovascular diseases (ICD-10-CM codes I00-I99), rheumatic heart disease (codes I00-I09), hypertension (codes I10-I15), angina pectoris (codes I20), acute ischemic heart disease (codes I21-I24), chronic ischemic heart disease (code I25), pulmonary heart disease (codes I26-I28), cardiac arrhythmias (codes I44-I49), heart failure (code I50), other heart disease (codes I30-I42, I51), hemorrhagic stroke (codes I60-I62), ischemic stroke (code I63), other cerebrovascular diseases (codes I64-I62), diseases of arteries and arterioles (codes I70-I79), or other circulatory diseases (codes I80-I89, I95-I99). For this study, nonparametric estimation which requires enough sample size was used to predict the daily average mortality rate. Those aged < 30 yrs were excluded from this study for small sample size. Therefore, the final analytic sample size was 626,950. To distinguish individuals with or without respiratory diseases, cardiovascular inpatients were considered to have a respiratory disease if at least one of the supplementary diagnostic codes was J00-J47.

**Statistical analysis**

Mortality rates for cardiovascular patients were calculated for each different group such that the numerator was the number of deaths for calendar-month, the winter and summer seasons, and disease subtypes. The Generalized Addictive Model (GAM), an extension of linear regression models, was used to describe average daily variations in mortality from 2006 to 2010 [21]. It models the daily average mortality rate (on a logit scale) with unspecified smooth function of admission date \( \lambda(t) \), which is estimated by cubic smoothing spline. The knots of the spline located at the beginning and end of each season as well as the first and last day of the five years. Using log-binomial regression models and the mortality in May as the reference, we estimated gender-, year-, and hospital-adjusted rate ratios (RRs) and 95% confidence intervals (CIs) for the other months. Analyses were performed separately for older (≥ 65 years) and younger patients (30 to < 65 years), since there is considerable heterogeneity in seasonal patterns of mortality among these two populations. To define seasons for further analysis, we grouped data of the four months of November, December, January, and February as the winter season, and the four months of May, June, July, and August as the summer season based on the results of the approximation in the risk of death between months (see Table 1). We then analyzed the associations between the winter season and the risk of cardiovascular death among older inpatients. Similar analyses were performed separately among patients with and without respiratory disease. All analyses were carried out using STATA 12.0 (StataCorp LP, College Station, Texas).

**Results**

Descriptive characteristics of the study sample were shown in Table 2. Compared with younger inpatients, older inpatients had a statistically significantly higher mortality rate...
(4.0% vs. 1.2%, p < 0.001), longer length of hospital stay (13 days vs. 10 days, P < 0.001), and more supplementary diagnoses (5 diagnoses vs. 3 diagnoses, P < 0.001), but a lower rate of surgery (55.4% vs. 70.6%, P < 0.001) and a lower cost per stay (USD 2,436 vs. USD 2,583, P < 0.001).

Mortality due to cardiovascular disease was presented in Figure 1. Among older patients, the mortality rate peaked during winter months and was lowest in the summer months between 2006 and 2010. Mortality rates trended downwards during both the winter (6.2% to 4.3%) and summer (3.9% to 3.2%) over the 5 years studied. The cyclical seasonal mortality pattern found among older inpatients was not observed among younger inpatients. Similar seasonal trends and overall downward trends existed for both older and younger inpatients with heart disease and cerebrovascular disease (data not shown).

Table 1 shows the adjusted risks of cardiovascular death associated with hospitalization months among older and younger inpatients. Older patients who were hospitalized in the winter months had a 30% to 50% increased risk of death (P < 0.01) than those who were hospitalized in

| Month   | ≥ 65 years (n = 308,952) | 30 to < 65 years (n = 317,998) |
|---------|--------------------------|--------------------------------|
| January | 23,912 5.0 1.49 (1.36, 1.63) < 0.01 | 23,244 1.3 1.06 (0.91, 1.24) 0.43 |
| February| 23,054 4.6 1.39 (1.27, 1.52) < 0.01 | 22,266 1.3 1.02 (0.87, 1.20) 0.82 |
| March   | 28,587 3.5 1.06 (0.96, 1.16) 0.24 | 30,463 1.0 0.86 (0.74, 1.01) 0.07 |
| April   | 27,745 3.9 1.18 (1.08, 1.29) < 0.01 | 27,770 1.2 1.01 (0.87, 1.18) 0.91 |
| May†    | 26,758 3.3 Reference | 27,248 1.2 Reference |
| June    | 24,555 3.5 1.03 (0.94, 1.14) 0.51 | 25,270 1.1 0.92 (0.78, 1.08) 0.29 |
| July    | 23,689 3.7 1.10 (1.00, 1.21) 0.05 | 25,376 1.1 0.89 (0.76, 1.04) 0.15 |
| August  | 23,695 3.7 1.09 (0.99, 1.19) 0.09 | 25,120 1.2 0.92 (0.78, 1.08) 0.30 |
| September| 24,213 4.0 1.19 (1.08, 1.30) < 0.01 | 24,395 1.3 1.02 (0.88, 1.19) 0.79 |
| October | 28,006 4.2 1.26 (1.15, 1.37) < 0.01 | 28,149 1.3 1.05 (0.90, 1.22) 0.53 |
| November| 28,027 4.3 1.29 (1.18, 1.41) < 0.01 | 30,110 1.3 1.06 (0.91, 1.23) 0.45 |
| December| 26,711 4.5 1.35 (1.24, 1.48) < 0.01 | 28,587 1.2 0.99 (0.85, 1.15) 0.86 |

*The rate ratios (RRs) were adjusted for gender, year, and hospital.
†Mortality rates in May were taken as the reference levels for comparison.

Table 2 Characteristics of cardiovascular inpatients by two age groups

| Year | ≥ 65 years (n = 308,952) | 30 to < 65 years (n = 317,998) | Total (n = 626,950) |
|------|--------------------------|--------------------------------|------------------|
|      | No. (%) | No. (%) | No. (%) |
| 2006 | 53,960 (17.5) | 50,114 (15.8) | 104,074 (16.6) |
| 2007 | 57,873 (18.7) | 55,898 (17.6) | 113,771 (18.2) |
| 2008 | 58,909 (19.1) | 60,511 (19.0) | 119,420 (19.1) |
| 2009 | 66,031 (21.4) | 71,523 (22.5) | 137,554 (21.9) |
| 2010 | 72,179 (23.4) | 79,952 (25.1) | 152,131 (24.3) |

| Gender | ≥ 65 years (n = 308,952) | 30 to < 65 years (n = 317,998) | Total (n = 626,950) |
|--------|--------------------------|--------------------------------|------------------|
|        | No. (%) | No. (%) | No. (%) |
| Male   | 172,130 (55.7) | 215,099 (67.6) | 387,229 (61.8) |
| Female | 136,822 (44.3) | 102,899 (32.4) | 239,721 (38.2) |

*Numbers in brackets are the 25th and 75th percentile.
May (the month with the lowest death rate). Compared with older inpatients, however, the risks of death associated with the winter months were not statistically significant among younger inpatients. Separate analyses of deaths from cardiovascular and cerebrovascular diseases were similar (results not shown).

To evaluate which cardiovascular diseases were associated with the increased winter deaths among older inpatients (Table 3), we examined winter and summer hospitalization groups. The rates of death in winter and summer were highest for hospitalized patients diagnosed with hemorrhagic stroke (17% and 15.5%), acute ischemic heart disease (13.6% and 11.9%), and pulmonary heart disease (11.1% and 8.2%). In addition, a significantly increased mortality risk was associated with the winter season among those patients diagnosed with ischemic heart disease (RR = 1.22; 95% CI 1.13 to 1.31), pulmonary heart disease (RR = 1.42; 95% CI 1.10 to 1.83), cardiac arrhythmias (RR = 1.67; 95% CI 1.36 to 2.05), heart failure (RR = 1.30; 95% CI 1.09 to 1.54), ischemic stroke

Table 3 Associations between winter season and deaths among older cardiovascular inpatients

| Cardiovascular disease subtypes | Winter season (n = 101,704) | Summer season (n = 98,748) | RR (95% CI) | P-value |
|--------------------------------|-----------------------------|-----------------------------|-------------|---------|
| Ischemic heart disease (I20-I25) | 1,826/42,761 4.3 | 1,358/38,946 3.5 | 1.22 (1.13, 1.31) | < 0.01 |
| Angina pectoris (I20) | 145/19,367 0.8 | 113/17,653 0.6 | 1.16 (0.90, 1.48) | 0.25 |
| Acute ischemic heart disease (I21-I24) | 1,274/9,342 13.6 | 942/7,893 11.9 | 1.14 (1.04,1.25) | 0.01 |
| Chronic ischemic heart disease (I25) | 407/14,052 2.9 | 303/13,400 2.3 | 1.28 (1.10, 1.49) | < 0.01 |
| Pulmonary heart disease (I26-I28) | 194/1,753 11.1 | 121/1,477 8.2 | 1.42 (1.10, 1.83) | < 0.01 |
| Cardiac arrhythmias (I44-I49) | 248/7,221 3.4 | 160/7,737 2.1 | 1.67 (1.36, 2.05) | < 0.01 |
| Heart failure (I50) | 372/4,454 8.4 | 243/3,755 6.5 | 1.30 (1.09, 1.54) | < 0.01 |
| Cerebrovascular disease (I60-I69) | 1,691/27,829 6.1 | 1,328/28,975 4.6 | 1.35 (1.25, 1.45) | < 0.01 |
| Hemorrhagic stroke (I60-I62) | 575/3,378 17.0 | 439/2,825 15.5 | 1.12 (0.98, 1.29) | 0.10 |
| Ischemic stroke (I63) | 924/18,647 5.0 | 768/19,721 3.9 | 1.30 (1.17, 1.43) | < 0.01 |
| Other cerebrovascular diseases (I64-I69) | 192/5,804 3.3 | 121/6,429 1.9 | 1.78 (1.40, 2.25) | < 0.01 |
| Rheumatic heart disease (I00-I09) | 63/979 6.4 | 57/1,032 5.5 | 1.17 (0.80, 1.71) | 0.42 |
| Hypertensive diseases (I10-I15) | 35/8,389 0.4 | 34/7,889 0.4 | 1.07 (0.66, 1.73) | 0.78 |
| Other heart disease (I30-I42, I51, I52) | 73/1,587 4.6 | 72/1,648 4.4 | 1.03 (0.74, 1.45) | 0.85 |
| Diseases of arteries and arterioles (I70-I79) | 139/3,969 3.5 | 114/4,024 2.8 | 1.22 (0.94, 1.57) | 0.13 |
| Other circulatory diseases (I80-I89, I95-I99) | 21/2,762 0.8 | 16/3,214 0.5 | 1.47 (0.76, 2.86) | 0.26 |

1Winter season consisted of January, February, November, and December, while summer season consisted of May, June, July, and August. Mortality rate in summer season was taken as the reference levels.

2The rate ratios (RRs) were adjusted for gender, year, and hospital.
(RR = 1.30; 95% CI 1.17 to 1.43), and other cerebrovascular diseases (RR = 1.78; 95% CI 1.40 to 2.25). However, there was no positive association between winter season and deaths from hemorrhagic stroke.

Table 4 listed the risks of mortality associated with hospitalization months for older cardiovascular inpatients with and without respiratory disease. The mortality risks were higher in the winter months than in the month of May, regardless of whether they had a respiratory disease. However, winter increases in deaths were nearly 20% higher among patients diagnosed with respiratory diseases (40% to 60%) than those not diagnosed with respiratory diseases (20% to 40%), and the monthly mortality rates among the former were also about twice that of the latter. The analytic results of cardiovascular subtypes after excluding patients with respiratory diseases were similar to those with the influence of respiratory disease, except for hemorrhagic stroke, which also showed increases in winter deaths (RR = 1.18; 95% CI 1.01 to 1.37; results not shown).

Discussion

Our study demonstrated that overall cardiovascular mortality increased by approximately 30% to 50% during winter among cardiovascular inpatients aged ≥ 65 yrs in Beijing, China. The winter increase in mortality was seen among those with diverse subtypes of cardiovascular diseases, including ischemic heart diseases, pulmonary heart disease, cardiac arrhythmias, heart failure, and stroke. It should be emphasized that our estimates were independent of the impact of concurrent diagnosis of respiratory diseases. Results for younger patients, however, did not show a statistically significant elevated risk of cardiovascular deaths during the winter months.

Despite improved health care quality in recent years, mortality continues to peak during the winter months among older cardiovascular inpatients. Although the reason for this peak is not fully understood, the winter increase in mortality may be reduced or prevented by adequate clinical interventions since recognized triggers such as cold temperatures and respiratory infections can be effectively remediated during hospitalization.

Winter cardiovascular mortality among older inpatients is higher in Beijing than in other regions of the world. A previous study reported that excess winter mortality from coronary events was about 10% in 21 countries [22]. In addition, studies in the U.S. and in Canada also found that the difference in winter-summer mortality was about 10% for myocardial infarction, 20% for stroke, and 17% for sudden cardiac death [23,24]. The magnitude of excess winter mortality from cardiovascular disease was also found to be 28% and 48% higher in Norway and Ireland, respectively [17]. The winter peak in mortality from coronary heart disease was noted to diminish by about 2% per year from 1937 until around 1970 in the U.S., however, the trend reversed until 1991 [25]. The winter-summer difference has fluctuated in different countries, and reported data were generally based on statistics collected in the 1980s and the 1990s [22-25].

Our results showed the overall risk of death increased by 10% or more if cardiovascular inpatients aged ≥ 65 yrs

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**Table 4 Risks of mortality associated with hospitalization month among older cardiovascular inpatients with or without respiratory diseases in Beijing, China**

| Month    | Cardiovascular inpatients with respiratory diseases* (n = 50,451) | Cardiovascular inpatients without respiratory diseases (n = 258,501) |
|----------|-----------------------------------------------------------------|-------------------------------------------------------------------|
|          | Death/No. | Death rate (%) | RR (95% CI)† | P-value | Death/No. | Death rate (%) | RR (95% CI)† | P-value |
| January  | 355/4492  | 7.9           | 1.54 (1.28, 1.85) | < 0.01 | 843/19420 | 4.3           | 1.44 (1.30, 1.60) | < 0.01 |
| February | 311/3865  | 8.1           | 1.59 (1.32, 1.92) | < 0.01 | 757/19189 | 3.9           | 1.32 (1.19, 1.47) | < 0.01 |
| March    | 266/4583  | 5.8           | 1.14 (0.94, 1.37) | 0.20  | 743/24004 | 3.1           | 1.03 (0.93, 1.15) | 0.55  |
| April    | 302/4437  | 6.8           | 1.33 (1.11, 1.61) | < 0.01 | 787/23308 | 3.4           | 1.13 (1.02, 1.25) | 0.02  |
| May†     | 210/4089  | 5.1           | Reference       |       | 680/22669 | 3.0           | Reference       |       |
| June     | 211/3661  | 5.8           | 1.07 (0.87, 1.31) | 0.54  | 636/20894 | 3.0           | 1.03 (0.92, 1.15) | 0.61  |
| July     | 233/3736  | 6.2           | 1.20 (0.99, 1.47) | 0.07  | 652/19953 | 3.3           | 1.07 (0.96, 1.19) | 0.21  |
| August   | 242/3873  | 6.3           | 1.16 (0.95, 1.41) | 0.14  | 639/19822 | 3.2           | 1.06 (0.95, 1.19) | 0.27  |
| September| 241/3864  | 6.2           | 1.22 (1.01, 1.49) | 0.04  | 733/20349 | 3.6           | 1.18 (1.06, 1.31) | < 0.01 |
| October  | 309/4490  | 6.9           | 1.34 (1.11, 1.62) | < 0.01 | 863/23516 | 3.7           | 1.23 (1.11, 1.36) | < 0.01 |
| November | 337/4662  | 7.2           | 1.43 (1.19, 1.72) | < 0.01 | 861/23365 | 3.7           | 1.24 (1.12, 1.38) | < 0.01 |
| December | 332/4699  | 7.1           | 1.34 (1.11, 1.61) | < 0.01 | 866/22012 | 3.9           | 1.34 (1.21, 1.49) | < 0.01 |

* Cardiovascular inpatients with respiratory diseases were those with a secondary diagnosis as acute upper respiratory infections, influenza and pneumonia, other acute lower respiratory infections, other diseases of upper respiratory tract, chronic lower respiratory diseases, and those with primary diagnosis or secondary diagnosis as pulmonary heart disease.
† The rate ratios (RRs) were adjusted for gender, year, and hospital.
‡ Mortality rates in May were taken as the reference levels.
were also diagnosed with a respiratory disease. However, the respiratory disease factor seemed not to contribute a large effect on the excess winter mortality, because patients diagnosed with both diseases accounted for a small fraction (about 25%) of all cardiovascular patients, and the risks of mortality were one-fold higher than those with a single cardiovascular disease in all months. Our finding is consistent with previous studies that influenza vaccination prevented only a small fraction of hospitalizations (20%) for cardiovascular diseases [26].

We also found that most of the cardiovascular disease subtypes were associated with a winter increase in mortality except for angina pectoris, rheumatic heart disease, hypertension, and arteries and arterioles diseases. For those cardiovascular subtypes with a positive association, however, the magnitudes of excess winter mortality were different, and a high difference was found for cardiac arrhythmias with the adjusted rate ratios being 1.67 (95% CI 1.36 to 2.05). The high mortality risk of cardiac arrhythmias may be associated with some abnormal activity in the cardiovascular system during winter.

The winter season may act as a surrogate variable representing other causes that ultimately lead to death. Excess winter cardiovascular mortality is principally driven by cold temperatures, and the temperature hypothesis is plausible biologically [27]. Previous studies showed that cold temperatures could increase systemic vascular resistance and fibrinogen levels, which further induced an increase in blood pressure, thrombus formation, and also enhanced fibrinolytic activity, platelet adheriveness, and lipid levels [28-30]. Beijing is located in Northern China, with a typical cold and dry winter. Springs and autumns are both of relatively short duration. The public heating system lasts from November to March, with fairly dry and filthy air due to limited ventilation. Due to the cold winter climate, residents in Beijing tend to stay indoors in a climate-controlled condition. The large differences in the indoor/outdoor temperatures as well as long stay in indoor environment with poor ventilation may induce or worsen the onset of vascular conditions. Because of reduced physiologic reserves, older patients may be more vulnerable to winter stress [23]. No visible seasonal fluctuations of mortality among young patients may be due to the existence of different reasons for suffering cardiovascular diseases.

Young adults suffer cardiovascular diseases because of their adverse health behaviors and job stress. Aside from cold temperatures, however, other factors, including holiday effects also increase the cardiovascular risk during the winter [27-31]. An increase in cardiovascular deaths in winter months among older patients may be associated with two important holidays - the New Year and the Chinese Spring Festival - and may have been caused by hospital factors rather than by individual holiday behaviors. During the holiday season, on-duty staffing and care may be reduced.

Our data from the 32 top-ranked hospitals in Beijing allowed us to examine seasonal associations with multiple disease endpoints and the large sample size allows statistical analysis with reliable results once potential referral biases are accounted for. The reliability of our results relates to the high quality of the HSR data. The HSR data were required by the Beijing Municipal Health Bureau for computing allocations of financial resources and evaluating hospital performance. Hospitals included in this study have a high reputation for quality in all aspects of healthcare, including diagnosis, treatment, hospital management, coding, and electronic medical record systems in China. Only inclusion of top-ranked hospitals in this study can help with eliminating quality of care as a strong potential confounding factor. However, results from this study can be only applicable to top-ranked hospitals and are not representative for all hospitals. The quality of care for other hospitals may not be as well as these top-ranked hospitals and thus can possibly have a more notably seasonal variation of daily average mortality rate.

Our study has some limitations. Despite quality controls, the HSR data are still subject to measurement errors, which include errors due to incomplete or inaccurate information recorded on summary reports as well as processing errors. It is also possible that misclassification of diseases in coding may have occurred, especially for some subtypes of cardiovascular diseases. However, the misclassification of coding, if it exists, should occur randomly and should not depend upon the seasons. The HSR data are restricted to hospitalized patients; therefore, we could not analyze cardiovascular patients who were not hospitalized—those who were not admitted to the hospital, those who died before being treated, or those who died after being discharged from the hospital. Thus, the amplitudes of excess winter mortality in our study could have been underestimated. The HSR data are abstracts describing an individual’s hospitalization, and lack clinical details that may influence the risks of death in relation to the winter season. In this study, we assumed that the standards of admission and treatment for cardiovascular diseases are the same throughout the year, and that the impact of the absence of particular clinical details is small. Finally, although we have adjusted for several confounding factors, it is possible that those residual confounding effects of other unidentified factors remain and that these could distort the positive association found between cardiovascular deaths and the winter season.

Our study has important clinical implications. We suggest that hospitals and physicians evaluate their care for older cardiovascular patients, particularly those with high-risk conditions such as respiratory infection and cardiac arrhythmias, during the winter season. The plan of care for these patients should include controlling the temperature of the wards, early identification of high-risk
patients, use of preventive therapies including vascular and anti-infection treatments, sufficient health care staffing, use of adequate monitoring devices and complete discharge instructions to prevent re-admission. Clinical guidelines related to the care of older patients during the winter season may be necessary to ensure that physicians and other hospital staff are aware of the increase in cardiovascular deaths among older individuals during the winter season.

**Conclusions**

Overall cardiovascular mortality increased by approximately 30% to 50% during winter among cardiovascular inpatients aged ≥ 65 yrs, while the excess winter mortality was not observed among younger cardiovascular inpatients in Beijing, China. The winter increase in mortality was seen among those with diverse subtypes of cardiovascular diseases, including ischemic heart diseases, pulmonary heart disease, cardiac arrhythmias, heart failure, and stroke. It should be emphasized that our estimates were independent of the impact of concurrent diagnosis of respiratory diseases.

**Competing interests**

All authors declare that they have no competing interests.

**Authors’ contributions**

BX, HL, GPY and LZ designed the research, analyzed the data, drafted the manuscript and interpreted the results. All authors have contributed to, carried out the data management and study design. JL contributed to draft the manuscript and interpreted the results. All authors have contributed to, seen and approved the manuscript.

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