Research Article

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The Role of Ambient Temperature and Plasma Osmolarity on Clinical Outcomes of Acute Myocardial Infarction Patients during Hajj

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ABSTRACT

Background: Performance of Hajj is physically very demanding, especially if performed during the summer season. The aim of this study is to evaluate the importance of ambient temperature and dehydration, indicated by plasma osmolarity on the clinical outcomes of cardiac patients during Hajj season in 2017.

Methods: We included all patients referred to tertiary center with acute coronary syndrome during Hajj period of 2017. Plasma osmolarity was calculated using concentrations of sodium, plasma glucose, and blood urea nitrogen at admission. Patients were stratified by groups (G) of admission osmolarity, clinical outcome was compared. The primary endpoints were in-hospital mortality, length of stay, Cardiac complications (heart failure, re-infarction, arrhythmia, shock and thrombus formation), left ventricular function and readmission rate.

Result: Total of 300 patients were identified with mean age 56.2 ± 12.1 , 84% males and 97(32%) were pilgrims. They were exposed to average heat index $61.9 \pm 10.6^\circ\text{C}$. Significantly longer admissions were found in the group of higher osmolarity (G2) (≥ 295 mos/L) as compared to patients with normal osmolarity in G1 [6.7 ± 14.9 VS 4.0 ± 4.5 , $P=0.045$]. Total in-hospital death rate was 4.3% (13). Using Binary regression analysis; osmolarity Group [$p=0.009$], Pilgrim [$P=0.005$], Heat index [$P=0.005$], were independent predictor of inhospital mortality, while Heat index is the only independent predictor for MACE [$P=0.001$].

Conclusion: Plasma osmolarity and heat index significantly affect cardiac patient's outcome. These finding underscore the importance of health awareness of protection from dehydration for pilgrims during summer season.

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Abbreviation List

BMI: body mass index

CAD: coronary artery disease

CKD: chronic kidney disease

DM: Diabetes

STEMI ST: elevation MI

MACE major adverse cardiac events

LV: left ventricle

LVEF: left ventricle ejection fraction

LOS: length of stay

MPV: mean platelet volume

Introduction

Annually, millions of Muslims travel from over 180 different countries to Makkah city for Hajj; from 8 to 13 of Dhul-Hijja. The lunar calendar only has 354 days, compared with 365 days for Gregorian

calendar, this means that the Hajj is held during different periods of seasonality [1]. Starting 2016 Hajj has entered into the summer months with ambient temperature between 43°C and 48.7°C , and relative humidity of 58–87%, which increases the risk of dehydration [2]. Performance of Hajj is physically very demanding, with extraordinary physical stressors such as heat exposure, dehydration, crowding, and physical exertion, which expose the pilgrims to many health hazards. Most of pilgrims are over the age of 60 years with the cardiovascular diseases being the leading cause of death. The risk of dehydration in older adults is increased because of loss of muscle mass, decreased kidney function, physical and cognitive impairments, reduced thirst and multiple drugs use [3].

Plasma osmolarity has been shown to be a simple and valid tool for assessing dehydration [4]. It reflects the number of dissolved particles per liter of plasma. Plasma osmolarity of 275 to < 295 mOsmol/L is considered normal; 295 to 300 mOsmol/L suggests impending water loss dehydration; and > 300 mOsmol/L suggests current water loss dehydration [5].

A recent meta- analysis demonstrated increased risk of cardiovascular hospitalization 2.2% with heat wave exposure [6]. Furthermore, several recent studies have demonstrated a significant prognostic role of hyper-osmolarity in patients with acute coronary syndromes and heart failure [7-9].

King Abdullah Medical City (KAMC) is the tertiary cardiac center for cardiac catheterizations and open-heart surgeries in Makkah. In this study, we evaluated the importance of plasma osmolarity on cardiovascular complications, length of stay, and mortality in acute myocardial infarction (AMI) patients admitted during Hajj season of 2017 (1438 H). This could play a strategic role in managing cardiac patients during Hajj in the coming decade where Hajj will be in the summer time.

Material and Methods

Definitions and Outcomes

Aim is to study the relationship between plasma osmolarity and ambient temperature with cardiovascular clinical outcomes including in-hospital mortality; defined as death during hospitalization; and MACE: defined as any cardiac complication: pulmonary edema, cardiogenic shock, intubation /ventilation, cardiac arrest, readmission, LV thrombus, and LVEF <30%.

Patient Population and Statistical Methods

This is a retrospective cohort Single center study. All ST elevation myocardial infarction (STEMI) patients referred to the KAMC Cardiac Center during Hajj of 2017 (1438 H) were included. After local KAMC IRB approval, electronic medical records of included patients were reviewed. The authors attest they have no conflict of interest statement and have not received any funding nor grants.

Plasma Osmolarity which has been calculated from the formula $(2 \times (\text{Na} + \text{K})) + (\text{BUN} / 2.8) + (\text{glucose} / 18)$ [10]. Heat index (HI) calculated by the formula: $(12) \text{ HI} = 0.5 \times (\text{T} + 61.0 + [(\text{T} - 68.0) \times 1.2] + (\text{RH} \times 0.094))$ Using website; <http://www.wpc.ncep.noaa.gov/html/heatindex.shtml> (T) is the air temperature and (rh) is the relative humidity. Daily temperatures were collected

from Saudi general authority of meteorology and environment protection website; <https://www.pme.gov.sa/ar/Pages/default.aspx>.

Assumption of ($\alpha = 0.05$) 2-sided, ($\beta = 0.2$), mean difference in plasma osmolarity between dying and living patients 10 mOsmol/L. SD = 10, mortality of 5%. Using online sample size calculator; <http://www.sample-size.net/>. Based on this assumption we needed records of at least 170 patients.

Statistical Analysis

Using SPSS version 26. student's t-test and Chi square test, used in continuous and categorical variables respectively. Cox regression analysis was used for survival analysis. All tests were two sided, and a significance level of 5% was used.

Results

300 consecutive patients admitted to KAMC cardiac center with STEMI. They were divided into 2 groups according to their plasma osmolarity level, Group (Gp) I patients with plasma osmolarity level <295 mOsmol/L (n= 161), and Gp II patients with plasma osmolarity level ≥ 295 mOsmol/L (n= 139).

Base Line Characteristics and Outcomes

The mean age of study group was 56.2 ± 12.1 , 84% were males, with a mean BMI 27.9 ± 5.4 exposed to average ambient temperature on admission days as follows: maximum temperature of 42.5 ± 3.1 °C and a minimum temperature of 30.3 ± 2.5 °C resulting in Heat index (HI) of 61.9 ± 10.6 °C. They have an average plasma osmolarity of 295.1 ± 8.7 . The majority, 127 (43.2%) were referred for anterior STEMI.

The baseline risk factors, characteristics, presentation and complication among both groups are summarized in Table I. A total 13 in-hospital deaths occurred (4.3 %) while 77 (25.8 %) suffered MACE. There was no statistically significance difference between both groups as regard age, gender, BMI, smoking, HTN, being resident, or pilgrim but Group II were more frequently Diabetics (P = 0.046) and history of kidney disease (P = 0.027).

Table I: Risk Factors and Clinical Characteristics in Group I and II

	Total	Gp I (n= 161)	Gp II (n= 139)	P
Male	252 (84 %)	133 (82.6%)	119 (85.6%)	0.48
Diabetes	147 (49%)	70 (43.5%)	77 (55.4%)	0.04*
CKD	20 (6.7%)	6 (3.7%)	14 (10.2%)	0.028*
Hypertension	164 (54.7%)	84 (51.5%)	80 (58.4%)	0.32
Known CAD	72 (24%)	33 (20%)	39 (28.1%)	0.13
Smoker	99 (33 %)	53 (32.5%)	46 (33.6%)	0.49
Pilgrims	103 (34.3 %)	60 (37.3%)	43 (30.1%)	0.25
Age	56.2 ± 12.1	55.5 ± 12.8	57.1 ± 11.1	0.27
BMI	27.9 ± 5.4	28.1 ± 5.3	27.6 ± 5.5	0.42
Heat index	61.9 ± 10.6	61.3 ± 10.9	62.6 ± 10.3	0.314
MPV	10.5 ± 1.3	10.4 ± 1.1	10.6 ± 1.5	0.088
Glucose	171.4 ± 65.9	158.3 ± 50.2	186.7 ± 77.8	<0.001
Sodium	135.8 ± 4.2	133.7 ± 3.3	138.1 ± 3.8	< 0.001
Potassium	4.2 ± 0.52	4.2 ± 0.5	4.2 ± 0.5	0.97
BUN	17.5 ± 9.4	15.5 ± 7.5	19.9 ± 1.9	<0.001
Urea	6.3 ± 3.4	5.5 ± 2.7	7.1 ± 3.9	<0.001
Osmolarity	295.1 ± 8.7	289.2 ± 4.4	302.0 ± 7.3	<0.001

LVEF	43.8 ±10.1	43.5 ±9.3	44.1 ±10.9	0.65
cardiac arrest	16 (5.3 %)	7(4.3%)	9(6.6%)	0.668
LOS (days)	5.2 ±10.7	4.01 ± 4.5	6.7 ±14.9	0.045*
LV thrombus	7 (2.3%)	3(1.8%)	6(4.4%)	0.210
Readmission rate	5 (1.7%)	2(1.2%)	3(2.2%)	0.531
LVEF <30%	41 (13.7%)	19 (11.8%)	22 (15.8 %)	0.31
MACE	77 (25.8)	37(23%)	40 (29%)	0.24
In-hospital death	13 (4.3%)	3 (1.9%)	10 (7.2%)	0.023*

BMI: body mass index; CAD: coronary artery disease; CKD: chronic kidney disease; LVEF: left ventricle ejection fraction; LV: left ventricle; LOS: length of stay; MACE: major adverse cardiac events

Pilgrims Characteristics and Outcomes

However, comparing Pilgrims and residents is summarized in Table II, pilgrims were older, less frequently male gender and diabetic, they were exposed to a higher heat index 58.4± 9.7 versus 68.5± 9.02, $p < 0.01$. However, there was no significant difference regarding plasma osmolarity, or patient outcomes (mortality or MACE).

Table II: Baseline Characteristics and Outcome among Pilgrims Versus Residents

Variable	Residents (N=197)	Pilgrims (N=103)	P value
Male	175 (88.8%)	77(74.8%)	0.02*
DM	107 (54.3%)	40 (38.8%)	0.011*
CKD	12 (6.1%)	8 (7.8%)	0.58
HTN	108 (54.8%)	56 (54.4%)	0.94
Known CAD	57 (28.9%)	15 (14.6%)	0.06
Smoker	53 (32.5%)	46 (33.6%)	0.499
Anterior STEMI	120(60.9%)	53 (51.5%)	0.115
Osmolarity >295	96 (48.7%)	43 (41.7%)	0.249
Age	54.3 ±12.8	59.7 ±9.8	0.266
Heat Index	58.4± 9.7	68.5± 9.02	<0.001
BMI	28.2±5.3	27.3 ±5.3	0.181
MPV	10.4± 1.2	10.7 ±1.6	0.298
Glucose	175.8 ±63.8	163.1± 69.3	0.298
Sodium	135.7 ±3.9	135.9 ±4.6	0.755
potassium	4.2± 0.5	4.2± 0.56	0.76
BUN	17.3 ±9.5	17.8 ± 9.2	0.724
Urea	6.2 ±3.4	6.3± 3.3	0.724
osmolarity	295.0±7.7	295.3 ± 10.4	0.764
In mortality	6 (3.0%)	7 (6.8%)	0.13
LVEF <30%	25 (12.7%)	16 (15.5%)	0.49
MACE	44 (22.3%)	23 (22.3%)	0.99

BMI: body mass index; CAD: coronary artery disease; CKD: chronic kidney disease; MACE major adverse cardiac events

Relationship between Plasma Osmolarity and Hospital Course

There was no significant difference between the groups regarding cardiac complications; or any MACE. However, Group II had significantly longer length of stay 4.01 ± 4.5 day versus 6.7 ± 14.9 $p=0.045$. A total 13 in hospital death occurred (4.3 %), In hospital mortality was significantly higher in group II 10 (7.2%) versus 3 (1.9%); $p = 0.023$, Figure 1.

Figure 1: In-Hospital Mortality Rate between the Two Groups

Binary regression model was used for predicting in hospital mortality and MACE is summarized in Table III. We started with univariate model analysis with plasma osmolarity as continuous variable. Then the effect of potential demographic and clinical confounders used univariate models. The independent predictors for mortality are Pilgrim, HI, and plasma osmolarity; conversely HI, was the only independent predictor for MACE.

TABLE III: Multivariate Predictors of In-Hospital Mortality and Mace Using Binary Regression Analysis
77 MACE COCURED (25.8%0 AND 13 DEATH (4%)

		Sig.	Exp(B)	95% C.I.for EXP(B)	
				Lower	Upper
DEATH	HI	.005	.949	.915	.984
	Osmolarity GP	.009	.175	.047	.650
	Age	.143	1.029	.990	1.070
	PILGRIM	.005	.213	.072	.631
MACE	HI	.001	.974	.959	.990
	Osmolarity GP	.113	.660	.395	1.103
	Age	.056	1.017	1.000	1.035
	PILGRIM	.309	.769	.464	1.275

Discussion

To the best of our knowledge, this is the first study analysis to establish a link between heat index, osmolarity imbalance and mortality in patients admitted with STEMI.

Makkah Geography and Climate

Makkah, the holiest of Muslim cities, located in the Şirāt Mountains, inland from the Red Sea coast. All devout Muslims attempt a hajj (pilgrimage) at least once in their lifetime [11,12]. Meckkah is 909 feet (277 metres) above sea level in western Saudi Arabia between the dry beds of the Wadi Ibrāhīm and is surrounded by the Mountains. Less than 5 inches (130 mm) of rainfall occur yearly, mainly in the winter months. Ambient temperatures are high throughout the year and in summer may reach 120 °F (49 °C) resulting in a very high heat index [13,14].

Pilgrims and Risk of Heat Stress

In a questionnaire of 412 male; age 43.48 ± 13.42 years pilgrims regarding heat stress awareness in Hajj in 1436 (September 2015) 36% of pilgrims were not aware of Makkah weather; highlighting the importance pilgrims health education on heat-related health issues and coping strategies [14]. In Hajj, activity is unavoidable in high heat index conditions, this emphasizes the importance of public awareness to minimize health risks. Pre-training to provide acclimatization for at least two weeks before hand, liberal intake of fluids before thirst begins and dressing in clothes, including hats, appropriate for the climate is of utmost importance [15]. Nowadays majority of pilgrims reach Makkah by air and arrive within, a few hours of leaving home; giving them little time to acclimatize [16]. Failure to adapt to the high load of ambient temperature can result in heat stress. Exhaustion syndromes may not be accompanied by a rise in body temperature. Several variable affect the heat adaptation namely fatigue, lack of sleep, dehydration, diabetes, cardiovascular disease, lack of acclimatization, all of which are common in Hajj [16].

Our results demonstrate the heat stress is a predictor for MACE, furthermore we demonstrated that Pilgrims are exposed, on admission, to significantly (17%) higher heat index, which is an important predictor factor for in-hospital mortality and MACE.

Heat Index and Heat Stress

To our knowledge this is the first study to demonstrate the effect of heat stress, using heat index, in Saudi Arabia during Hajj season. Recently, several studies from different continents and countries have studied the association of ambient temperature on emergency room visits, admissions with acute coronary syndromes, and mortality [17-23].

In a metanalysis of 18 studies, Bhaskaran et al concluded that there is a statistically significant association between higher temperatures and risk of myocardial infarction, with the main effects up to 3 days later [24]. In 2019, Cheng et al reported a metanalysis on 54 studies from 20 countries to study the global effect of heatwave on cardiovascular and respiratory morbidity and mortality outcomes. They concluded that heatwaves increase the risk of cardiorespiratory mortalities especially in vulnerable subgroups elderly and those with history of ischemic heart disease. In 2012 Bhaskaran et al studied the relationship between hourly temperature in 11 areas in Wales and England and concluded that ambient temperature above a threshold of 20°C , each 1°C increase in temperature was associated with a 1.9% transient increase in risk of myocardial infarction in the following 1-6 hours [24,25].

The adverse physiologic effect of heat exposure was demonstrated in a study by Keatinge et al, on volunteers under controlled conditions of moving air at 41°C for six hours. This caused a 0.84°C rise of core temperature and resulted in increase in red blood cell counts, platelet counts, and blood viscosity, as well as heart rate [26]. A recent study in healthy men age > 55 years, for each 5°C increase in mean ambient temperature HDL decreased -1.76% and LDL increased by 1.74% [27].

Several studies of age-related cardiovascular heat strain have demonstrated that healthy aging is accompanied by altered cardiovascular function, limiting the extent to which older individuals can maintain stroke volume, increase cardiac output, and increase skin blood flow when exposed to heat. This is more prominent in elderly, whereby the increased cardiovascular demand is often fatal due to increased strain on an already compromised left ventricle [28,29].

Previous studies are in agreement with our current study whereby heat index; takes in account the ambient temperature and humidity to construct a scale that describes how warm the air feels. it is easily calculated from available weather website. HI on the day of admission was a predictor of in-hospital mortality and MACE. This underscores the importance of health education to the general public and Pilgrims in specific. In 2008, Noweir et al, studied the climatic heat load in Hajj locations and reported that the Wet Bulb Globe Temperature (WBGT); a measure of the heat stress, that includes temperature, humidity, wind speed, sun angle and solar radiation. They reported that WBGT were considerably high for safe heat exposure and that it exceeded the recommended comfort zone [30]. This is in agreement with our results where the average heat index is in the danger zone and was found to be a predictor of mortality and MACE in our population study [31]. Lastly, seasonal variations in blood pressure, serum lipids, and fibrinogen have been observed and may explain the relation between temperature and cardiovascular disease mortality [32]. These data and our study underscores the importance of ambient temperature and consequently cellular dehydration on prognosis.

Dehydration and Plasma Osmolarity

Dehydration is defined as water-loss, it usually is a result of insufficient fluid intake, resulting in elevation of serum osmolality [33]. Osmolarity is defined as the number of milliosmoles of the solutes per liter of solution. Calculated serum osmolality (mOsm/L); an estimation of the osmolar concentration of serum; has been recommended as an easy and simple tool to assess dehydration [33]. A study on healthy adults from 3 European counties, showed inadequate hydration status on several days per week, which may have a negative health and cognitive impact on daily life [34].

Water balance inside the body is of vital importance for patients who are critically ill, and serum osmolality plays an important role in extracellular and intracellular water distribution. Several recent studies have reported the role of hyperosmolality with increased mortality and readmission in critically ill cardiac patients, heart failure, and in acute coronary syndrome undergoing percutaneous cardiac intervention [7,35,36]. Our results reveal that hyperosmolality is associated with increased hospital mortality of patients who presented with acute STEMI.

Pathophysiology and Osmolarity

Hyperosmolality is a result of increase of its components, namely sodium and glucose each of which have been reported as risk factors for adverse outcomes [37,38]. Secondly, hyperosmolality causes redistribution of body fluids, increasing cardiac preload volume and consequently patient outcomes [35].

Hyperosmotic Stress

Several recent studies have demonstrated that hyperosmotic stress is related to several pathologies [39-43]. Dehydration, and consequent intracellular hyperosmolality, is a major challenge; as it results in disturbance of global cellular function and cell death [44]. Increased extracellular osmolality is cytotoxic, it promotes water flux out of the cell, triggering cell shrinkage, and intracellular dehydration, triggering apoptosis, or cell death [42,43]. This adversely affect protein structure and function and altered enzyme activity; it triggers oxidative stress, protein carbonylation, mitochondrial depolarization, DNA damage, and cell cycle arrest, resulting in apoptosis [45].

Hyperosmolarity and Cardiomyocyte

Exposure of the myocytes to hypertonic media causes NF- κ B (nuclear factor kappa-light-chain-enhancer of activated B cells) and caspase activation through reactive oxygen species (ROS) production. Studies have suggested that the hyperosmotic stress triggers oxidative stress and reactive oxygen species generation within the cells [40,46]. Furthermore hyperosmotic stress triggers apoptosis in cardiac myocytes through a p53-dependent manner [43]. Lastly, when exposed to hyperosmolarity cardio-myocytes exhibit an increase in Aldose reductase (AR) expression that was accompanied by AR-mediated activation of apoptotic signaling pathways [47].

Pilgrims Health Education

Whenever activity is unavoidable in high heat index conditions (as in Hajj), health education can help minimize health risks. Pre-training to provide acclimatization for at least two weeks before the event [16]. Adaptation strategies, as educating pilgrims on Makkah climate, liberal intake of fluids before thirst begins and dressing in clothes, including hats, appropriate for the climate is of utmost importance to reduce the health effects of heat stress [16]. Moreover, alerting first responders and emergency room physicians to the importance of early intravenous fluid hydration, whenever clinically possible. Physicians should be alerted to the role of plasma osmolarity, at admission, on patient outcome and the possible preventive role of intravenous infusion of normal saline or half normal saline to improve outcome. These data and our study underscores the importance of ambient temperature and consequently cellular dehydration on prognosis.

Limitations

Firstly, only in-hospital patients and in-hospital complications were recorded. Secondly, this is a retrospective study, with its inherent limitations, conducted in Makkah; hence, the present findings may not be readily extrapolated to other populations or settings. Thirdly, our analysis is restricted to patients admitted to an interventional cardiology center, which may induce selection bias: STEMI cases managed solely medically were not included. Lastly, in the present study, osmolarity was calculated rather than being measured directly, which may be different from actual osmolarity.

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References

- Ahmed QA, Arabi YM, Memish ZA (2006) Health risks at the Hajj. *Lancet* 367: 1008-1015.
- Aleeban M, Mackey TK (2016) Global Health and Visa Policy Reform to Address Dangers of Hajj during Summer Seasons. *Front Public Health* 4: 280.
- Hooper L, Bunn D, Jimoh FO, Fairweather-Tait SJ (2014) Water-loss dehydration and aging. *Mechanisms of Ageing and Development* 136: 50-88.
- Cheuvront SN, Kenefick RW, Charkoudian N, Sawka MN (2013) Physiologic basis for understanding quantitative dehydration assessment. *American Journal of Clinical Nutrition* 97: 455.
- Thomas DR, Cote TR, Lawhorne L, Levenson SA, Rubenstein LZ, et al. (2008) Understanding clinical dehydration and its treatment. *J Am Med Dir Assoc* 9: 292-301.
- Phung D, Thai PK, Guo Y, Morawska L, Rutherford S, et al. (2016) Ambient temperature and risk of cardiovascular hospitalization: An updated systematic review and meta-analysis. *Science of The Total Environment* 550: 1084-1102.
- Rohla M, Freynhofer MK, Tentzeris I, Farhan S, Wojta J, et al. (2014) Plasma osmolarity predicts clinical outcome in patients with acute coronary syndrome undergoing percutaneous coronary intervention. *Eur Heart J Acute Cardiovasc Care* 3: 84-92.
- Çiçek G, Açıkgöz SK, Altay S, Bozbay M, Yleri M (2015) Prognostic Impact of Hyperosmolarity in Patients Undergoing Primary Angioplasty for ST-Segment Elevation Myocardial Infarction. *Abant Med J* 4: 323-330.
- Briongos Figuero S, Jiménez-Mena M, Ortega Marcos J, Camino López A, Fernández Santos S, et al. (2014) Dehydration and serum hyperosmolarity as new predictors of mortality after acute coronary syndrome. *International Journal of Cardiology* 172: e472-e4.
- Charles Hu GBMD (2017) Serum Osmolality 200 <http://www.medcalc.com/osmol.html>.
- Niu S, Xu M (2019) Impact of Hajj on Global Health Security. *J Relig Health* 58: 289-302.
- Al Mayahi ZK, Ali Kabbash I (2019) Perceptions of, and Practices for Coping with, Heat Exposure among Male Arab Pilgrims to the Hajj, 1436. *Prehosp Disaster Med* 34: 161-74.
- Glubb ASAaJB (2018) Mecca: Encyclopaedia Britannica, Inc.; [updated February 07, 2018. electronic Encyclopedia]]. Available from: <https://www.britannica.com/place/Mecca>.
- Wexler RK (2002) Evaluation and treatment of heat-related illnesses. *Am Fam Physician* 65: 2307-2314.
- Khogali M (1983) Epidemiology of heat illnesses during the Makkah Pilgrimages in Saudi Arabia. *Int J Epidemiol* 12: 267-273.
- An der Heiden M, Muthers S, Niemann H, Buchholz U, Grabenhenrich L, et al. (2019) Schätzung hitzebedingter Todesfälle in Deutschland zwischen 2001 und 2015. *Bundesgesundheitsblatt - Gesundheitsforschung - Gesundheitsschutz* 62: 571-579.
- Rodrigues M, Santana P, Rocha A (2019) Effects of extreme temperatures on cerebrovascular mortality in Lisbon: a distributed lag non-linear model. *International journal of biometeorology* 63: 549-59.
- Wichmann J, Rosengren A, Sjöberg K, Barregard L, Sallsten G (2013) Association between ambient temperature and acute myocardial infarction hospitalisations in Gothenburg, Sweden: 1985-2010. *PLoS One* 8: e62059.
- Green RS, Basu R, Malig B, Broadwin R, Kim JJ, et al. (2010) The effect of temperature on hospital admissions in nine California counties. *Int J Public Health* 55: 113-121.
- Vaneckova P, Bambrick H (2013) Cause-specific hospital admissions on hot days in Sydney, Australia. *PLoS One* 8: e55459.
- Yang J, Yin P, Zhou M, Ou C-Q, Li M, et al. (2016) The effect of ambient temperature on diabetes mortality in China: A multi-city time series study. *Science of The Total Environment* 543: 75-82.
- Nastos PT, Giaouzaki KN, Kampanis NA, Matzarakis A (2013) Acute coronary syndromes related to bio-climate in a Mediterranean area. The case of Ierapetra, Crete Island, Greece. *International Journal of Environmental Health Research* 23: 76-90.
- Bhaskaran K, Hajat S, Haines A, Herrett E, Wilkinson P, et al. (2009) Effects of ambient temperature on the incidence of myocardial infarction. *Heart* 95: 1760-1769.
- Cheng J, Xu Z, Bambrick H, Prescott V, Wang N, et al. (2019) Cardiorespiratory effects of heatwaves: A systematic review and meta-analysis of global epidemiological evidence. *Environmental Research* 177: 108610.

25. Bhaskaran K, Armstrong B, Hajat S, Haines A, Wilkinson P, et al. (2012) Heat and risk of myocardial infarction: hourly level case-crossover analysis of MINAP database. *BMJ* 345: e8050.
26. Keatinge WR, Coleshaw SRK, Easton JC, Cotter F, Mattock MB, et al. (1986) Increased platelet and red cell counts, blood viscosity, and plasma cholesterol levels during heat stress, and mortality from coronary and cerebral thrombosis. *The American Journal of Medicine* 81: 795-800.
27. Halonen JJ, Zanobetti A, Sparrow D, Vokonas PS, Schwartz J (2011) Outdoor temperature is associated with serum HDL and LDL. *Environmental research* 111: 281-287.
28. Crandall CG, Wilson TE, Marving J, Vogelsang TW, Kjaer A, et al. (2008) Effects of passive heating on central blood volume and ventricular dimensions in humans. *The Journal of Physiology* 586: 293-301.
29. Kenney WL, Craighead DH, Alexander LM (2014) Heat waves, aging, and human cardiovascular health. *Med Sci Sports Exerc* 46: 1891-1899.
30. Noweir MH, Bafail AO, Jomoah IM (2008) Study of heat exposure during Hajj (pilgrimage). *Environ Monit Assess* 147: 279-295.
31. Heat Safety Tips and Resources (2019) Available from: <https://www.weather.gov/safety/heat-index>.
32. Pan WH, Li LA, Tsai MJ (1995) Temperature extremes and mortality from coronary heart disease and cerebral infarction in elderly Chinese. *Lancet* 345: 353-355.
33. Hooper L, Abdelhamid A, Ali A, Bunn DK, Jennings A, et al. (2015) Diagnostic accuracy of calculated serum osmolarity to predict dehydration in older people: adding value to pathology laboratory reports. *BMJ Open* 5: e008846.
34. Braun H, von Andrian-Werburg J, Malisova O, Athanasatou A, Kapsokefalou M, et al. (2019) Differing Water Intake and Hydration Status in Three European Countries-A Day-to-Day Analysis. *Nutrients* 11.
35. Shen Y, Cheng X, Ying M, Chang HT, Zhang W (2017) Association between serum osmolarity and mortality in patients who are critically ill: a retrospective cohort study. *BMJ Open* 7: e015729.
36. Arevalo-Lorido JC, Gomez JC, Formiga F, Conde-Martel A, Carrera-Izquierdo M, et al. (2016) High serum osmolarity at admission determines a worse outcome in patients with heart failure: Is a new target emerging? *Int J Cardiol* 221: 238-242.
37. Plakht Y, Gilutz H, Shiyovich A (2018) Sodium levels during hospitalization with acute myocardial infarction are markers of in-hospital mortality: Soroka acute myocardial infarction II (SAMI-II) project. *Clin Res Cardiol* 107: 956-964.
38. Wannamethee SG, Shaper AG, Lennon L, Papacosta O, Whincup P (2016) Mild hyponatremia, hypernatremia and incident cardiovascular disease and mortality in older men: A population-based cohort study. *Nutr Metab Cardiovasc Dis* 26: 12-19.
39. Yancey PH (2005) Organic osmolytes as compatible, metabolic and counteracting cytoprotectants in high osmolarity and other stresses. *Journal of Experimental Biology* 208: 2819-2830.
40. Reinehr R, Haussinger D (2007) Hyperosmotic activation of the CD95 system. *Methods Enzymol* 428: 145-160.
41. Criollo A, Galluzzi L, Maiuri MC, Tasdemir E, Lavandro S, et al. (2007) Mitochondrial control of cell death induced by hyperosmotic stress. *Apoptosis* 12: 3-18.
42. Alfieri RR, Petronini PG (2007) Hyperosmotic stress response: comparison with other cellular stresses. *Pflugers Arch* 454: 173-185.
43. Burg MB, Ferraris JD, Dmitrieva NI (2007) Cellular response to hyperosmotic stresses. *Physiol Rev* 87: 1441-1474.
44. Kino T, Segars JH, Chrousos GP (2010) Brx, a link between osmotic stress, inflammation and organ physiology/pathophysiology. *Expert Review of Endocrinology & Metabolism* 5: 603-614.
45. Bocker C, Thompson DC, Vasiliou V (2012) The role of hyperosmotic stress in inflammation and disease. *Biomol Concepts* 3: 345-364.
46. Burgos JI, Morell M, Mariangelo JIE, Vila Petroff M (2019) Hyperosmotic stress promotes endoplasmic reticulum stress-dependent apoptosis in adult rat cardiac myocytes. *Apoptosis* 24: 785-797.
47. Galvez AS, Ulloa JA, Chiong M, Criollo A, Eisner V, et al. (2003) Aldose reductase induced by hyperosmotic stress mediates cardiomyocyte apoptosis: differential effects of sorbitol and mannitol. *J Biol Chem* 278: 38484-38494.