Association of circulating MR-proADM with all-cause and cardiovascular mortality in the general population: Results from the KORA F4 cohort study

Christina Gar1,2,3*, Barbara Thorand3,4, Christian Herder1,3,5,6, Chaterina Sujana4,7, Margit Heier4,8, Christa Meisinger9,10, Annette Peters3,4,11, Wolfgang Koenig11,12,13, Wolfgang Rathmann3,14, Michael Roden3,5,6, Michael Stumvoll15, Haifa Maalml3,5, Thomas Meitinger11,16, Holger Then17, Jochen Seissler1,2,3, Cornelia Then1,2,3

1 Department of Medicine IV, University Hospital, LMU Munich, Germany, 2 Clinical Cooperation Group Diabetes, Ludwig-Maximilians-Universität München and Helmholtz Zentrum München, Munich, Germany, 3 German Center for Diabetes Research (DZD), München-Neuherberg, Germany, 4 Institute of Epidemiology, Helmholtz Zentrum München, German Research Center for Environmental Health (GmbH), Neuherberg, Germany, 5 Institute of Clinical Diabetology, German Diabetes Center, Leibniz Center for Diabetes Research at Heinrich Heine University Düsseldorf, Düsseldorf, Germany, 6 Division of Endocrinology and Diabetology, Medical Faculty, Heinrich Heine University Düsseldorf, Düsseldorf, Germany, 7 Institute for Medical Information Processing, Biometry, and Epidemiology (IBE), Ludwig-Maximilians-Universität, Munich, Germany, 8 KORA Study Centre, University Hospital Augsburg, Augsburg, Germany, 9 Independent Research Group Clinical Epidemiology, Helmholtz Zentrum München, German Research Center for Environmental Health (GmbH), Neuherberg, Germany, 10 Chair of Epidemiology at University Hospital Augsburg, Augsburg, Germany, 11 DZHK (German Centre for Cardiovascular Research), Partner Site Munich Heart Alliance, Munich, Germany, 12 Institute of Epidemiology and Medical Biometry, University of Ulm, Ulm, Germany, 13 Deutsches Herzzentrum München, Technische Universität München, Munich, Germany, 14 German Diabetes Center, Leibniz Institute at Heinrich Heine University Düsseldorf, Institute of Biometrics and Epidemiology, Düsseldorf, Germany, 15 Department of Medicine, University of Leipzig, Leipzig, Germany, 16 Institute of Human Genetics, Technische Universität München, Munich, Germany, 17 Freie Waldorfschule Augsburg, Augsburg, Germany

* christina.gar@med.uni-muenchen.de

Abstract

Background and aim
Despite its vasodilatory effect, adrenomedullin and its surrogate mid-regional pro-adrenomedullin (MR-proADM) have been found to be positively associated with all-cause and cardiovascular mortality. However, the underlying mechanisms thereof remain unclear and the associations were mostly shown in geriatric cohorts or in patients with chronic diseases. Therefore, we aimed to investigate the possible involvement of abdominal obesity, selected adipokines, and biomarkers of subclinical inflammation in the association of MR-proADM with mortality in a population based study cohort.

Methods
Prospective analysis of the KORA F4 study; median follow-up 9.1 (8.8–9.4) years. Complete data on MR-proADM and mortality was available for 1551 participants, aged 56.9±12.9 years (mean±SD). Correlation and regression analyses of MR-proADM with overall (BMI)
Results

MR-proADM associated with all-cause (HR (95%CI): 2.37 (1.72–3.26) and 2.31 (1.67–3.20)) and cardiovascular mortality (4.28 (2.19–8.39) and 4.44 (2.25–8.76)) after adjustment for traditional cardiovascular risk factors including BMI or waist circumference, respectively.

MR-proADM was further associated with four out of seven examined adipokines (leptin, retinol-binding protein-4, chemerin, and adiponectin) and with five out of eleven examined biomarkers of subclinical inflammation (high-sensitivity C-reactive protein, interleukin-6, myeloperoxidase, interleukin-22, and interleukin-1 receptor antagonist) after multivariable adjustment and correction for multiple testing. However, only IL-6 substantially attenuated the association of MR-proADM with all-cause mortality.

Conclusions

We found an association of MR-proADM with (abdominal) obesity, selected adipokines, and biomarkers of subclinical inflammation. However, the association of MR-proADM with mortality was independent of these parameters. Future studies should investigate the role of IL-6 and further characteristics of subclinical inflammation in the association between MR-proADM and all-cause mortality.

Introduction

Adrenomedullin (AMD) is a 52 amino acid peptide synthesized by a variety of tissues, including adrenal medulla, vascular endothelial cells, vascular smooth muscle cells and adipose tissue [1–3], and exhibits vasodilative properties [4]. In addition to its hypotensive effects, biological actions of AMD include antioxidative and antiapoptotic but also metabolic effects, such as inhibition of insulin secretion in beta cells [4]. Increased AMD levels and its surrogate marker mid-regional pro-adrenomedullin (MR-proADM) are correlated with congestive heart failure, hypertension and incident cardiovascular events [4]. During recent years, ADM and MR-proADM have been shown to be associated with cardiovascular and all-cause mortality [4–6]. Mostly, these associations have been shown in geriatric cohorts or in patients with chronic diseases [7–12]. The underlying mechanisms that link (MR-pro)ADM to mortality remain to be determined [7–11].

Because the measurement of ADM itself is challenging due to its short half-life, MR-proADM as a surrogate for ADM levels has become increasingly prominent. MR-proADM represents a more stable peptide with a longer half-life and is co-secreted together with ADM from pro-adrenomedullin, resulting in a direct association between plasma levels of ADM and MR-proADM [12].

Besides mortality, MR-proADM levels are linked to obesity [13–16] and the metabolic syndrome [17]. Given that obesity is a known risk factor for mortality [18], the association between MR-proADM and mortality might be mediated by their common association to body fat and other cardiovascular risk factors and thus be a secondary finding. One study, however,
found that after adjustment for BMI, MR-proADM remained significantly associated with all-cause mortality, and thus, MR-proADM might affect mortality via mechanisms other than simple body fat mass [19]. In this context, abdominal fat mass was the only anthropometric index to be independently linked to MR-proADM levels [16]. Further, abdominal fat, as opposed to overall body fat, might be more relevant for an increased mortality rate in the context of the metabolic syndrome [2, 4, 13, 14, 20–22]. The BMI-independent association between MR-proADM and mortality might therefore be explained by varying levels of abdominal fat at comparable BMI levels. Further, an altered secretion profile of adipokines and markers of subclinical inflammation may indicate metabolic alterations in the adipose tissue [23–25]. Subclinical inflammation [26] as well as alterations in adipokine levels, e.g. adiponectin [27], were found to be associated with an elevated mortality risk. In this respect, ADM was suggested to exert both proinflammatory and immunomodulatory effects [4, 28–30]. Consequently, immunomodulatory adipokines and biomarkers of subclinical inflammation may also confound the relationship between MR-proADM and mortality.

Therefore, the present study aimed to investigate the association of MR-proADM with all-cause and cardiovascular mortality in the KORA (Cooperative Health Research in the Region of Augsburg) F4 cohort and to further examine whether the association between MR-proADM and mortality is independent of abdominal fat, adipokines, and biomarkers of subclinical inflammation.

Materials and methods

Study cohort

The KORA F4 (2006–2008) study is a follow-up examination of the population-based KORA S4 study (1999–2001). Recruitment and eligibility criteria for the KORA studies, study design, standardized sampling methods and data collection (medical history, medication, anthropometric and blood pressure measurements) have been described elsewhere [31, 32]. The outcomes all-cause and cardiovascular mortality (ICD-9 codes 390–459 and 798) were ascertained by regularly checking the status of the participants through the population registries until 2016. Death certificates were obtained from the local health authorities. Prevalent myocardial infarction and stroke were self-reported diagnoses. The median (1st; 3rd quartile) follow-up time was 9.1 (8.8; 9.4) years.

All study participants gave written informed consent. The study was approved by the Ethics Committees of the Bavarian Medical Association in adherence to the declaration of Helsinki.

MR-proADM was measured in the first 1596 participants of the KORA F4 study (out of a total of 3080 participants). All variables necessary for the analysis of the association of MR-proADM with mortality were available in 1551 participants. Measurements of leptin and retinol-binding protein-4 (RBP-4) were available in 1549 participants; chemerin in 1055 participants; wingless-type MMTV integration site family, member 5A (Wnt-5a), secreted frizzled-related protein-5 (SFRP-5), vaspin, progranulin, omentin, and adiponectin were available in 606 participants aged ≥62 years; interleukin-6 (IL-6), tumor necrosis factor-α (TNF-α), interleukin-18 (IL-18), soluble intercellular adhesion molecule-1 (sICAM-1), myeloperoxidase (MPO), superoxide dismutase-3 (SOD-3), interleukin-22 (IL-22) and interleukin-1 receptor antagonist (IL-1RA) were available in 603 participants aged ≥62 years.

Overall obesity was defined as a BMI ≥30 kg/m². Abdominal obesity (increased waist circumference) was defined according to the International Diabetes Federation definition (waist circumference ≥94 cm in men and ≥80 cm in women) [33].

Criteria for diabetes mellitus were a validated medical diagnosis or current self-reported use of glucose-lowering agents. Participants without clinically diagnosed diabetes underwent a
standard 75 g oral glucose tolerance test. Newly diagnosed diabetes was defined according to the World Health Organization diagnostic criteria (≥ 7.0 mmol/l fasting and/or ≥ 11.1 mmol/12-h glucose) [34]. Participants with a diabetes type other than type 2 or unknown glucose tolerance status (n = 25) were excluded from the analyses.

Arterial hypertension was defined as a systolic blood pressure ≥ 140 mmHg and/or a diastolic blood pressure ≥ 90 mmHg, and/or intake of anti-hypertensive medication, given that the participants were aware of being hypertensive.

Leisure-time physical activity was assessed with two separate questions concerning leisure time sports activity in winter and in summer (cycling included). Possible answers were (i) > 2 hours, (ii) 1–2 hours, (iii) < 1 hour and (iv) none per week. Participants who had a total score < 5, obtained by summing the numbers (i)-(iv) from the winter and summer questions, were classified as ‘physically active’.

**Biochemical measurements**

Blood samples were collected after an overnight fast of at least eight hours and were kept at room temperature until centrifugation. Plasma was separated immediately, serum after 30 min. Plasma and serum samples were assayed immediately or stored at -80°C. Plasma MR-proADM was measured by sandwich fluoro-immunoassay (BRAHMS, Hennigsdorf, Berlin, Germany) on an automated BRAHMS KRYPTOR system as previously described [35]. Blood serum glucose levels were assessed using the hexokinase method (GLU Flex; Dade Behring, Marburg, Germany). High- and low-density lipoprotein (HDL and LDL) cholesterol were measured with enzymatic methods (CHOD-PAP; Dade Behring). Triglycerides were measured by an enzymatic color test (GPO-PAP method, TGL Flex; Dade Behring). Serum creatinine was determined with a modified Jaffe test (Krea Flex; Dade Behring). Insulin was measured by an electrochemiluminescence immunoassay on a Cobas e602 instrument (Roche Diagnostics GmbH, Mannheim, Germany). HbA1c was measured in hemolyzed whole blood using the cation-exchange high performance liquid chromatographic, photometric VARIANT II TURBO HbA1c Kit—2.0 assay on a VARIANT II TURBO Hemoglobin Testing System (Bio-Rad Laboratories Inc., Hercules, USA).

High-sensitivity C-reactive protein (hsCRP) was determined in plasma with a high-sensitivity latex-enhanced nephelometric assay on a BN II analyzer (Siemens, Erlangen, Germany). Serum levels of IL-6 and TNF-α were measured with Quantikine HS ELISA kits, IL-22, IL-1RA, sICAM-1, and MPO with Quantikine ELISA kits (R&D Systems, Wiesbaden, Germany) [5, 6, 36]. Plasma levels of IL-18 were determined using ELISA kits from MBL (Nagoya, Japan). Serum SOD-3 concentrations were measured with an ELISA from Cloud-Clone Corp. (Houston, TX, USA) [7]. Intra-assay coefficient of variation (CV) for hsCRP, IL-1RA, IL-22, sICAM-1, IL-6, TNF-α, IL-18, MPO, and SOD-3 were 2.7, 2.8, 5.5, 3.5, 7.2, 6.3, 7.6, 3.2, and 7.1%, respectively. Inter-assay CVs were 6.3, 7.0, 9.3, 6.4, 11.8, 14.4, 9.4, 5.6, and 7.1%, respectively. For IL-22, 332 (31%) of the sera yielded values below the limit of detection (LOD; 3.9 pg/ml). Values below LOD were assumed to be evenly distributed between 0 and LOD and were assigned a value of 0.5 × LOD. Chemerin serum concentrations were determined using a commercially available ELISA kit (Human Chemerin ELISA, Biovendor, Heidelberg, Germany) with a sensitivity of 0.1 ng/ml, an intra-assay CV of 6%, and an inter-assay CV of 7.6% [37]. Plasma concentrations of RBP-4 were measured by immunonephelometry using a BN II analyzer with an intra- and inter-assay CV of <10%. Leptin was measured by ELISA (Merckodia, Uppsala, Sweden); mean intra- and inter-assay CVs were <10% [38]. Serum levels of omentin-1 and adiponectin were measured using the Human Omentin-1 ELISA (BioVendor, Brno, Czech Republic) [39] and the Human Total Adiponectin/Acrp30 Quantikine ELISA Kit
(R&D Systems) with intra-assay CVs of 2.0% and 4.0% respectively, and inter-assay CVs of 3.8 and 8.0%, respectively. Serum vaspin concentrations were measured using a commercial enzyme-linked immunosorbent assay kit (AdipoGen, Seoul, Korea) according to the manufacturer’s protocol [37]. The assay has a sensitivity of 12 pg/ml and the intra- and inter-assay CVs were 1.3–3.8% and 3.3–9.1%, respectively. Progranulin serum concentrations were determined using a commercially available ELISA (Progranulin human ELISA Kit AdipoGen, AdipoGen Inc., Korea) [37]. SFRP5 and Wnt-5a levels were measured using the Enzyme-linked Immunosorbent Assay Kit for Secreted Frizzled Related protein 5 (SFRP5) and Wnt-5a from CloudClone (Houston, TX; previously USCN, Wuhan, China) [40]. All sera yielded levels above the LOD. Mean intra- and inter-assay CVs were 6.4 and 18.6%, respectively.

Glomerular filtration rate (eGFR) was calculated according to the Chronic Kidney Disease Epidemiology Collaboration (CKD-EPI) equation (2009) [41].

**Statistical analysis**

Characteristics of the study participants were compared between participants with BMI < 30 kg/m² (non-obese) and BMI ≥ 30 kg/m² (obese), using t-tests in case of approximately normally distributed variables. Mann-Whitney U-tests were performed for variables with skewed distributions. Binomial proportions were compared with Chi-square tests. The associations of MR-proADM with all-cause and cardiovascular mortality were examined in Cox proportional hazard models. The cross-sectional associations of MR-proADM with categorical cardiovascular risk factors were assessed with logistic regression models, whereas the associations of MR-proADM with continuous outcomes were assessed using linear regression models. Continuous variables were transformed to a Gaussian distribution by the probability integral transformation followed by an inverse transform sampling and were used in calculations per one standard deviation. The associations of MR-proADM with the respective outcomes were examined in models without (crude model) and with adjustments for covariates:

- **Model 1**: adjusted for sex and age (continuous);
- **Model 2**: adjusted for sex, age, and BMI (continuous);
- **Model 3**: adjusted for sex, age, and waist circumference (continuous);
- **Model 4**: adjusted for sex, age, BMI, diabetes, arterial hypertension, eGFR (continuous), HDL cholesterol (continuous), physical activity (active/inactive), current smoking and former smoking;
- **Model 5**: adjusted for sex, age, waist circumference, diabetes, arterial hypertension, eGFR (continuous), HDL cholesterol (continuous), physical activity (active/inactive), current smoking and former smoking.

For further analyses, models adjusted for single parameters and adjustments according to model 4 plus single adipokines and biomarkers of subclinical inflammation were calculated. LDL cholesterol was not found to be associated with MR-proADM after adjustment for sex, age, and BMI and was therefore not included in the models. Participants with missing data were excluded from the respective adjusted and crude analyses. The level of statistical significance was set at 5% (two-sided). The Bonferroni method was used to correct for multiple testing where appropriate. The calculations were performed using the statistical environment R, version 3.6.0. Figures were created using Tableau 2020.4 (Tableau Software, Seattle, WA, USA).
Results

Clinical characteristics of the study participants

Table 1 shows the clinical characteristics of the study sample. MR-proADM was higher in obese (BMI $\geq 30 \text{ kg/m}^2$) compared to non-obese (BMI $< 30 \text{ kg/m}^2$) participants. Besides the characteristics of the metabolic syndrome, obese participants also presented with a lower eGFR and were less physically active, but more likely to have quit smoking. Sex and a history of myocardial infarction or stroke did not differ between the groups. Several of the selected biomarkers (vaspin, progranulin, omentin, adiponectin, Wnt-5a, SFRP-5, sICAM-1, MPO, SOD-3, TNF-$\alpha$, IL-6, IL-18, IL-22, and IL-1RA) were available only in participants aged $\geq 62$ years.

Association of MR-proADM with cardiovascular risk factors

First, we analyzed the association between MR-proADM and selected cardiovascular risk factors (Table 2). Except for sex, all selected cardiovascular risk factors (age, BMI, waist circumference, arterial hypertension, type 2 diabetes, LDL and HDL cholesterol, former and current smoking, and physical activity) significantly associated with MR-proADM in the crude model. All of these risk factors but LDL cholesterol remained significantly associated after adjustment for sex, age, and BMI.

Association of MR-proADM with all-cause and cardiovascular mortality

MR-proADM was associated with all-cause mortality. The association was affected by age, but hardly attenuated by the other cardiovascular risk factors. In all models, MR-proADM remained independently associated with all-cause mortality ($p < 0.001$; Table 3).

Furthermore, we analyzed the association between MR-proADM and all-cause mortality stratified by overall overweight/obesity (BMI $< 30$ vs. $\geq 30 \text{ kg/m}^2$) and subgroups of abdominal obesity (waist circumference $< 80 \text{ cm}$ in women; $< 94 \text{ cm}$ in men). Irrespective of BMI and waist circumference subgroups, MR-proADM was higher in participants who died compared to survivors ($p < 0.001$; Fig 1). The association was stronger in obese compared with non-obese participants (S1 Table). The $p$ for interaction with BMI was 0.026 in the model adjusted for BMI, the $p$ for interaction waist circumference was 0.034 in the model adjusted for waist circumference.

Likewise, MR-proADM independently associated with cardiovascular mortality (HR 4.28 (95% CI: 2.19–8.39), $p < 0.001$, S2 Table) with a stronger association in obese compared to non-obese participants, although the interaction term was not significant (interaction BMI: 0.543).

Association of MR-proADM with adipokines and biomarkers of subclinical inflammation

MR-proADM associated with four out of seven adipokines (leptin, RBP-4, chemerin, and adiponectin) and five out of eleven biomarkers of subclinical inflammation (hsCRP, IL-6, IL-1RA, IL-22, and MPO) after multivariable adjustment (Model 4) and correction for multiple testing using the Bonferroni method (Fig 2 and S3 Table).

There was a trend towards a stronger association of MR-proADM with MPO, IL-22, leptin, and adiponectin in obese compared to non-obese participants (S4 Table). However, after correcting for multiple testing, only leptin ($p < 0.001$) showed a significant interaction with BMI and waist circumference (S4 Table). MR-proADM was significantly associated with IL-6 in
Table 1. Characteristics of the KOFA F4 study participants, overall and stratified by BMI.

| Characteristic                        | Total study cohort | Non-obese | Obese | p value |
|---------------------------------------|--------------------|-----------|-------|---------|
| All participants                      | N = 1551           | N = 1160  | N = 391|         |
| Male sex, n (%)                       | 762 (49)           | 574 (49)  | 189 (48)| 0.674  |
| Age (years)                           | 56.9 ± 12.9        | 55.9 ± 13.0| 59.9 ± 12.0| <0.001 |
| BMI (kg/m²)                           | 27.4 ± 4.6         | 25.4 ± 2.7| 33.6 ± 3.4| <0.001 |
| Waist circumference (cm)              | 93.4 ± 13.6        | 88 ± 10.7 | 107.7 ± 10.9| <0.001 |
| LDL cholesterol (mmol/l)              | 3.47 (2.88; 4.03)  | 3.44 (2.85; 4.01)| 3.38 (2.95; 4.13)| 0.035 |
| HDL cholesterol (mmol/l)              | 1.40 (1.16; 1.68)  | 1.47 (1.21; 1.73)| 1.27 (1.11; 1.50)| <0.001 |
| Type 2 diabetes, n (%)                | 181 (12)           | 82 (7)    | 99 (25) | <0.001 |
| HbA1c (%)                             | 5.4 (5.2; 5.7)     | 5.4 (5.1; 5.6)| 5.6 (5.3; 5.9)| <0.001 |
| HbA1c (mmol/mol)                      | 35.5 (33.3; 38.8)  | 34.4 (32.2; 37.7)| 37.7 (34.4; 41.0)| <0.001 |
| Arterial hypertension, n (%)         | 617 (40)           | 377 (32)  | 241 (61)| <0.001 |
| Systolic blood pressure (mmHg)        | 120.5 (109; 133)   | 119.0 (107.5; 131.5)| 125.5 (115.0; 137.3)| <0.001 |
| Diastolic blood pressure (mmHg)       | 74 (68.5; 81.5)    | 73.5 (68.0; 80.5)| 76.0 (70.5; 84.0)| <0.001 |
| eGFR (ml/min/1.73 m²)                 | 88.4 (77.0; 98.9)  | 90.2 (78.4; 100.3)| 82.7 (72.0; 95.0)| <0.001 |
| Physically inactive, n (%)            | 657 (42)           | 452 (39)  | 205 (52)| <0.001 |
| Current smoking, n (%)                | 273 (18)           | 225 (19)  | 48 (12) | 0.002 |
| Former smoking, n (%)                 | 639 (41)           | 452 (39)  | 187 (48)| 0.003 |
| Previous myocardial infarction n (%)  | 52 (3)             | 35 (3)    | 17 (4)  | 0.274 |
| Previous stroke n (%)                 | 34 (2)             | 11 (3)    | 23 (2)  | 0.445 |
| hsCRP (mg/l)                          | 1.11 (0.55–2.43)   | 0.91 (0.47–1.88)| 2.01 (1.06–4.22)| <0.001 |
| MR-proADM (nmol/l)                    | 0.51 (0.42; 0.61)  | 0.48 (0.41; 0.57)| 0.59 (0.51; 0.71)| <0.001 |
| Leptin (ng/ml)                        | 11.8 (5.4; 23.9)   | 8.7 (4.4; 16.7)| 28.5 (15.7; 48.9)| <0.001 |
| RBP-4 (g/l)                           | 0.042 (0.035; 0.049)| 0.042 (0.035; 0.049)| 0.043 (0.037; 0.051)| <0.001 |
| Chemerin (ng/ml)                      | 172.2 (141.2; 205.7)| 166.5 (135.8; 197.7)| 194.9 (159.5; 230.2)| <0.001 |
| Omentin (ng/ml)                       | 480.7 (398.3; 580.1)| 491.0 (413.3; 589.5)| 458.4 (373.1; 556.5)| <0.001 |
| SFRP-5 (ng/ml)                        | 55.9 (42.5; 70.4)  | 57.9 (44.1; 73.0)| 50.2 (37.1; 66.4)| <0.001 |
| Wnt-5a (ng/ml)                        | 0.038 (0.021; 0.077)| 0.038 (0.021; 0.080)| 0.037 (0.019; 0.066)| 0.433 |
| Participants aged ≥ 62 years          | N = 606            | N = 422   | N = 184|         |
| Age (years)                           | 70.1 ± 5.5         | 70.0 ± 5.6| 70.3 ± 5.2| 0.391  |
| BMI (kg/m²)                           | 28.4 ± 4.2         | 26.2 ± 2.4| 33.4 ± 3.1| <0.001 |
| Waist circumference (cm)              | 97.7 ± 12.0        | 92.9 ± 9.6| 108.4 ± 9.9| <0.001 |
| MR-proADM (nmol/l)                    | 0.60 (0.52; 0.71)  | 0.58 (0.50; 0.67)| 0.67 (0.57; 0.79)| <0.001 |
| Adiponectin (μg/ml)                   | 10.0 (6.6; 15.3)   | 10.5 (7.1; 15.7)| 9.1 (6.1; 14.0)| 0.012 |
| Omentin (ng/ml)                       | 480.7 (398.3; 580.1)| 491.0 (413.3; 589.5)| 458.4 (373.1; 556.5)| <0.001 |
| SFRP-5 (ng/ml)                        | 55.9 (42.5; 70.4)  | 57.9 (44.1; 73.0)| 50.2 (37.1; 66.4)| <0.001 |
| Wnt-5a (ng/ml)                        | 0.038 (0.021; 0.077)| 0.038 (0.021; 0.080)| 0.037 (0.019; 0.066)| 0.433 |

(Continued)
participants with an increased waist circumference ($\beta: 0.39 \pm 0.05; p < 0.001$), but not in participants with a normal waist circumference ($\beta: 0.06 \pm 0.124; p = 0.644$) (S4 Table).

Association of MR-proADM with all-cause mortality in consideration of adipokines and biomarkers of subclinical inflammation

Finally, we examined a possible involvement of the selected adipokines and inflammation markers in the association between MR-proADM and all-cause mortality. Of the biomarkers that significantly associated with MR-proADM, leptin, chemerin, and IL-22 neither associated ($p > 0.05$) with all-cause mortality in the simple model (model 2; adjusted for sex, age, and BMI) nor in the fully adjusted model (model 4; adjusted for sex, age, BMI, arterial hypertension, diabetes, eGFR, HDL, smoking, and physical activity).

In contrast, RBP-4, hsCRP, adiponectin, IL-6, MPO, and IL-1RA associated with all-cause mortality in both model 2 and 4 ($p < 0.05$). Hence, we further examined their potential to affect the association of MR-proADM with all-cause mortality. Inclusion of the parameters into the regression models did not substantially attenuate the association of MR-proADM with all-cause mortality (S5 Table). Only the adjustment for IL-6 resulted in an attenuation of the association of MR-proADM with all-cause mortality (HR 1.89 (95% CI: 1.51–2.36)), which

### Table 1. (Continued)

| Characteristic       | Total study cohort | Non-obese | Obese  | p value * |
|----------------------|--------------------|-----------|--------|-----------|
| IL-1RA (pg/ml)       | 303.9 (235.2; 409.0)| 283.1 (222.3; 365.8)| 370.65 (300.9; 455.4)| < 0.001 d |

Non-obese: BMI $< 30$ kg/m$^2$, obese: BMI $\geq 30$ kg/m$^2$.

a data are given as mean ± SD, median (first quartile; third quartile), or number of participants (proportion in %);

b defined as systolic blood pressure $\geq 140$ mmHg and/or diastolic blood pressure $\geq 90$ mmHg and/or use of antihypertensive medication, given that the participants were aware of being hypertensive;

c t-test;

d Mann-Whitney U-test;

e Chi-square test.

https://doi.org/10.1371/journal.pone.0262330.t001

Table 2. Association of MR-proADM (as dependent variable) with cardiovascular risk factors (as independent variables).

|                          | $\beta$-coefficient ± SE | p-value | $\beta$-coefficient ± SE | p-value |
|--------------------------|--------------------------|---------|--------------------------|---------|
|                          | crude                    |         | Model 2a                  |         |
| Male sex                 | 0.01 ± 0.05              | 0.874   | -                        | -       |
| Age (years)              | 0.63 ± 0.02              | < 0.001 | -                        | -       |
| BMI (kg/m$^2$)           | 0.46 ± 0.03              | < 0.001 | -                        | -       |
| Waist circumference (cm) | 0.46 ± 0.03              | < 0.001 | -                        | -       |
| Type 2 diabetes (yes/no) | 0.97 ± 0.08              | < 0.001 | 0.25 ± 0.08              | 0.003   |
| Arterial hypertension (yes/no) | 0.86 ± 0.05 | < 0.001 | 0.19 ± 0.05              | < 0.001 |
| LDL cholesterol (mmol/l) | 0.07 ± 0.03              | 0.008   | -0.02 ± 0.02             | 0.203   |
| HDL cholesterol (mmol/l) | -0.14 ± 0.03             | < 0.001 | -0.09 ± 0.02             | 0.003   |
| eGFR (ml/min/1.73 m$^2$) | -0.63 ± 0.02             | < 0.001 | -0.39 ± 0.03             | < 0.001 |
| Current smoking (yes/no) | 0.12 ± 0.07              | 0.222   | 0.28 ± 0.05              | < 0.001 |
| Former smoking (yes/no)  | 0.17 ± 0.06              | 0.003   | 0.14 ± 0.04              | < 0.001 |
| Physical activity (yes/no) | -0.34 ± 0.05          | < 0.001 | -0.15 ± 0.04             | < 0.001 |

a Model 2: adjusted for sex, age, and BMI.

https://doi.org/10.1371/journal.pone.0262330.t002
Persisted in the fully adjusted model (model 4 + IL-6: HR 1.93 (95% CI: 1.41–2.70)). MR-proADM remained significantly associated with all-cause mortality in all models (p < 0.001).

**Discussion**

In this study, we observed associations of MR-proADM with all-cause and cardiovascular mortality in the population-based KORA cohort, which were independent of (abdominal) obesity and other "traditional" cardiovascular risk factors, adipokines, and biomarkers of subclinical inflammation. To our knowledge, the association of MR-proADM with a large number of adipokines and biomarkers of subclinical inflammation was analyzed for the first time.

ADM and also MR-proADM have been suggested to be involved in many physiological processes with a possible impact on cardiovascular and metabolic regulation [4]. ADM has direct effects on blood vessels but may also act via central and renal mechanisms. Renal effects may explain the inverse association between MR-proADM and the eGFR, which was also found in the present study. We further confirmed an association between MR-proADM concentrations and cardiovascular risk factors. We observed that MR-proADM levels were dependent on age, which in turn is the major risk factor for mortality. Nevertheless, after adjustment for cardiovascular and metabolic risk factors, MR-proADM was still related to mortality, leaving some of the associations unexplained.

Regarding metabolic regulation, hyperglycemia, obesity, and dyslipidemia are associated with ADM levels [4]. ADM and its receptor have been found to be expressed in adipose tissue and ADM has therefore been claimed to be an adipokine [5, 6]. Additionally, ADM acts on adipose tissue to decrease lipolysis via the inhibition of beta-adrenergic pathways and thereby...
Fig 1. MR-proADM levels stratified by survival status (survived vs. died) in different BMI (A) and waist circumference (B) groups. Non-obese: BMI < 30 kg/m², obese: BMI ≥ 30 kg/m². Increased waist circumference: ≥ 94 cm in men and ≥ 80 cm in women. Group comparison by Mann-Whitney U-test. *p < 0.001 for comparison survival vs. death.

https://doi.org/10.1371/journal.pone.0262330.g001
Fig 2. Unadjusted (A) and adjusted (B; adjusted for sex, age, BMI, arterial hypertension, diabetes, eGFR, HDL, smoking, and physical activity) estimates ($\beta$ coefficient per 1-standard deviation) of the association of MR-proADM with biomarkers of subclinical inflammation (circles) and adipokines (black dots). Exploratory level of significance: $p < 0.05$ (light grey area); corrected level of significance after Bonferroni correction for multiple testing $p < 0.0028 \times \frac{0.05}{18}$ (white area).

https://doi.org/10.1371/journal.pone.0262330.g002
possibly affects plasma lipid concentrations [5]. In line, we previously demonstrated that increased MR-proADM levels were associated with increased triglyceride and decreased HDL cholesterol concentrations [17]. These metabolic connections may explain the link between increased MR-proADM levels and the metabolic syndrome [17]. However, the association of MR-proADM levels with mortality was independent of metabolic risk factors.

We further examined whether abdominal rather than overall body fat affects the association between MR-proADM and mortality. Our results show a strong positive association between MR-proADM and waist circumference, a surrogate marker for abdominal fat deposition, yet adjustment for waist circumference did not substantially attenuate the association of MR-proADM with all-cause and cardiovascular mortality in our study. However, we found that the association between MR-proADM and mortality was stronger in obese than in non-obese participants. The literature provides some evidence for a link between metabolic characteristics of adipose tissue and (MR-pro)ADM levels. TNF-α was found to increase the amount of ADM mRNA in omental fat, linking (MR-pro)ADM to subclinical inflammation [30]. On the other hand, ADM was suggested to reduce proinflammatory cytokines and counteract the inflammatory effect of cytokines and reactive oxygen species [4, 28, 29]. Hence, it is conceivable that MR-proADM correlates with adverse metabolic derangements, such as an altered adipokine and cytokine profile along with low-grade inflammation of the adipose tissue, which might be more likely to be present in obese compared to non-obese individuals and may partly mediate the association between MR-proADM and increased mortality. In the current study, MR-proADM was associated with several adipokines (leptin, RBP-4, chemerin, and adiponectin) and biomarkers of subclinical inflammation (hsCRP, IL-6, MPO, IL-22, and IL-1RA). Some of these associations tended to be stronger in obese than in non-obese participants. We found a strong association between MR-proADM and IL-6 in participants with an increased waist circumference, but not in participants with a normal waist circumference. Preclinical [42–44] and clinical data [45] point towards a modulating effect of ADM on IL-6 under different inflammatory conditions. Furthermore, plasma ADM levels have been found to be related to a single nucleotide polymorphism of the IL6 gene [46]. In the present study, we observed that addition of IL-6 to the multivariable model attenuated the association between MR-proADM and all-cause mortality, although the association remained significant.

Besides, MR-proADM was associated with leptin and this association was significantly stronger in participants with an increased BMI or waist circumference compared to normal-weight participants, possibly indicating a different activity profile of fat tissue in obese individuals. Both ADM and leptin display vasoprotective properties on the one hand, but are also related to cardiovascular risk factors beyond BMI and waist circumference, including arterial hypertension and vascular alterations on the other hand [47, 48]. However, since leptin was not associated with mortality in our study, we did not examine a possible influence of leptin on the association of MR-proADM with mortality.

**Study limitations**

The KORA study was conducted in Southern Germany and thus included mainly Western European participants. Consequently, our results might not be applicable to individuals of other ethnicities. Several investigated variables were available only in a subgroup of participants aged ≥ 62 years, limiting the power and generalizability of these results due to the lower number and selected age group. Although we examined a total of 18 adipokines/biomarkers of subclinical inflammation, low-grade adipose tissue inflammation might still have not been represented comprehensively. Similarly, residual confounding by possible unmeasured confounders cannot be excluded. In case of medication with antihypertensive or glucose lowering
agents, the diagnoses arterial hypertension and diabetes mellitus were made independently of the measured blood pressure and oral glucose tolerance test, given that the patients were aware of the diagnoses. The observational nature of the study precludes statements about causality. Due to the relatively low number of cardiovascular deaths, subgroup analyses were limited. A major strength of the present study is its population-based design with a high variety of phenotypes and standardized sampling methods that enable high inter-individual comparability of the obtained data.

**Conclusions**

In sum, MR-proADM was strongly associated with all-cause and cardiovascular mortality in the population-based KORA F4 study. This association was independent of classical cardiovascular and metabolic risk factors, including BMI. Further, there was no evidence that abdominal obesity (surrogate waist circumference) was a stronger confounding factor than whole body fat (surrogate BMI). From the selected adipokines and biomarkers of subclinical inflammation, only IL-6 attenuated the association of MR-proADM with all-cause mortality. Future studies should examine whether further cytokines or metabolic alterations of subclinical inflammation might be involved in the association between (MR-pro)ADM and mortality, thereby illuminating its underlying pathophysiology.

**Supporting information**

S1 Table. Hazard ratios (95% confidence interval) of the association between MR-proADM and all-cause mortality (per standard deviation), stratified by BMI and waist circumference. Non-obese: BMI < 30 kg/m², obese: BMI ≥ 30 kg/m². Increased waist circumference: ≥ 94 cm in men and ≥ 80 cm in women. a Model 4: adjusted for sex, age, BMI, arterial hypertension, diabetes, eGFR, HDL cholesterol, smoking, and physical activity; b Model 5: adjusted for sex, age, waist circumference, arterial hypertension, diabetes, eGFR, HDL cholesterol, smoking, and physical activity.

S2 Table. Hazard ratios (95% confidence interval) of the association between MR-proADM and cardiovascular mortality (per standard deviation), stratified by BMI. Non-obese: BMI < 30 kg/m², obese: BMI ≥ 30 kg/m². No analysis of subgroups of waist circumference due to small group sizes. a Model 4: adjusted for sex, age, BMI, arterial hypertension, diabetes, eGFR, HDL cholesterol, smoking and physical activity; b Model 5: adjusted for sex, age, waist circumference, arterial hypertension, diabetes, eGFR, HDL cholesterol, smoking and physical activity.

S3 Table. Beta estimates (β ± standard error) of the association of MR-proADM with adipokines and biomarkers of subclinical inflammation. Bold indicates significance after multivariable adjustment and Bonferroni correction for multiple testing (p < 0.0028 (0.05 ÷ 18)). a Model 4: adjusted for sex, age, BMI, arterial hypertension, diabetes, eGFR, HDL cholesterol, smoking, and physical activity.

S4 Table. Beta estimates (β ± standard error) of the association of MR-proADM with adipokines and biomarkers of subclinical inflammation stratified by BMI and waist circumference. Non-obese: BMI < 30 kg/m², obese: BMI ≥ 30 kg/m². Increased waist circumference: ≥ 94 cm in men and ≥ 80 cm in women. Bold indicates significance after multivariable adjustment and Bonferroni correction for multiple testing for the interaction terms (p < 0.0028.
Adjusted for sex, age, BMI, arterial hypertension, diabetes, eGFR, HDL, smoking, and physical activity. In the model adjusted for BMI. In the model adjusted for waist circumference.

S5 Table. Hazard ratios (95% confidence interval) of the association between MR-proADM and all-cause mortality (per 1-standard deviation). Model 4: Adjustment for sex, age, BMI, arterial hypertension, diabetes, eGFR, HDL, smoking and physical activity.

Acknowledgments

We gratefully acknowledge the contribution of all field staff members conducting the KORA F4 study and thank all study participants.

Author Contributions

Conceptualization: Christina Gar, Barbara Thorand, Christian Herder, Margit Heier, Christa Meisinger, Annette Peters, Jochen Seissler, Cornelia Then.

Data curation: Barbara Thorand, Christian Herder, Margit Heier, Christa Meisinger, Annette Peters, Wolfgang Koenig, Wolfgang Rathmann, Michael Roden, Michael Stumvoll, Haifa Maalmi, Thomas Meitinger, Jochen Seissler, Cornelia Then.

Formal analysis: Holger Then, Cornelia Then.

Funding acquisition: Barbara Thorand, Annette Peters, Jochen Seissler.

Investigation: Barbara Thorand, Christa Meisinger, Annette Peters, Jochen Seissler.

Methodology: Chaterina Sujana, Cornelia Then.

Project administration: Annette Peters, Jochen Seissler.

Resources: Annette Peters.

Supervision: Barbara Thorand, Jochen Seissler, Cornelia Then.

Validation: Barbara Thorand, Chaterina Sujana, Cornelia Then.

Visualization: Christina Gar.

Writing – original draft: Christina Gar.

Writing – review & editing: Barbara Thorand, Christian Herder, Chaterina Sujana, Margit Heier, Christa Meisinger, Wolfgang Koenig, Wolfgang Rathmann, Michael Roden, Michael Stumvoll, Haifa Maalmi, Thomas Meitinger, Holger Then, Jochen Seissler, Cornelia Then.

References

1. Brouwers FP, de Boer RA, van der Harst P, Struck J, de Jong PE, de Zeeuw D, et al. Influence of age on the prognostic value of mid-regional pro-adrenomedullin in the general population. Heart. 2012; 98 (18):1348–53. https://doi.org/10.1136/heartjnl-2012-302390 PMID: 22821276

2. Eggers KM, Venge P, Lindahl B, Lind L. Associations of mid-regional pro-adrenomedullin levels to cardiovascular and metabolic abnormalities, and mortality in an elderly population from the community. Int J Cardiol. 2013; 168(4):3537–42. https://doi.org/10.1016/j.ijcard.2013.05.005 PMID: 23722054

3. Odermatt J, Meili M, Hersberger L, Bolliger R, Christ-Crain M, Briell M, et al. Pro-Adrenomedullin predicts 10-year all-cause mortality in community-dwelling patients: a prospective cohort study. BMC Cardiovasc Disord. 2017; 17(1):178. https://doi.org/10.1186/s12872-017-0605-3 PMID: 28676115
4. Bełtowski J, Jamroz A. Adrenomedul lin—what do we know 10 years since its discovery? Pol J Pharma col. 2004; 56(1):5–27. PMID: 15047974

5. Herder C, Bongaerts BW, Rathmann W, Heier M, Kowall B, Koenig W, et al. Association of subclinical inflammation with polyneuropathy in the older population: KORA F4 study. Diabetes Care. 2013; 36(11):3663–70. https://doi.org/10.2337/dc13-0382 PMID: 24009302

6. Herder C, Kannenberg JM, Huth C, Carstensen-Kirberg M, Rathmann W, Koenig W, et al. Myeloperoxidase, superoxide dismutase-3, cardiometabolic risk factors, and distal sensorimotor polyneuropathy: The KORA F4/FF4 study. Diabetes Metab Res Rev. 2018; 34(5):e3000. https://doi.org/10.1002/dmrr.3000 PMID: 29577557

7. Behnes M, Papassoti riou J, Walter T, Fiedler E, Sauer T, Lang S, et al. Long-term prognostic value of mid-regional pro-adrenomedullin and C-terminal pro-endothelin-1 in patients with acute myocardial infarction. Clin Chem Lab Med. 2008; 46(2):204–11. https://doi.org/10.1515/CCLM.2008.040 PMID: 18076360

8. Kato J, Tsuruda T, Kita T, Kitamura K, Eto T. Adrenomedul lin: a protective factor for blood vessels. Arterioscler Thromb Vasc Biol. 2005; 25(12):2480–7. https://doi.org/10.1161/01.ATV.0000184759.91369.f8 PMID: 16141406

9. Zudaire E, Cuttitta F, Martı́nez A. Regulation of pancreatic physiology by adrenomedullin and its binding protein. Regul Pept. 2003; 112(1–3):121–30. https://doi.org/10.1016/S0167-0115(03)00030-2 PMID: 12667633

10. Li Y, Jiang C, Wang X, Zhang Y, Shibahara S, Takahashi K. Adrenomedullin is a novel adipokine: adrenomedullin in adipocytes and adipose tissues. Peptides. 2007; 28(5):1129–43. https://doi.org/10.1016/j.peptides.2007.03.005 PMID: 17433499

11. Masson S, Latini R, Carbonieri E, Moretti L, Rossi MG, Ciricugno S, et al. The predictive value of stable precursor fragments of vasoactive peptides in patients with chronic heart failure: data from the GISSI-heart failure (GISSI-HF) trial. Eur J Heart Fail. 2010; 12(4):338–47. https://doi.org/10.1093/eurjhf/hfp206 PMID: 20097683

12. Lorubbio M, Conti AA, Ognibene A. Midregional pro-adrenomedullin (MR-ProADM) reference values in serum. Clin Biochem. 2018; 53:173–4. https://doi.org/10.1016/j.clinbiochem.2018.01.003 PMID: 29305834

13. Vila G, Riedl M, Maier C, Struck J, Morgenthaler NG, Handisurya A, et al. Plasma MR-ProADM correlates to BMI and decreases in relation to leptin after gastric bypass surgery. Obesity (Silver Spring). 2009; 17(6):1184–8.

14. Del Ry S, Cabiati M, Bianchi V, Caponi L, Di Cecco P, Marchi B, et al. Mid-regional pro-adrenomedullin plasma levels are increased in obese adolescents. Eur J Nutr. 2016; 55(3):1255–60. https://doi.org/10.1007/s00394-015-0938-6 PMID: 26018656

15. Kistorp C, Bliddal H, Goetze JP, Christensen R, Faber J. Cardiac natriuretic peptides in plasma increase after dietary induced weight loss in obesity. BMC Obes. 2014; 1:24. https://doi.org/10.1186/s40608-014-0024-2 PMID: 26217511

16. Koyama T, Kuriyama N, Uehara R. Midregional Proadrenomedullin Can Reflect the Accumulation of Visceral Adipose Tissue—A Key to Explaining the Obesity Paradox. Int J Environ Res Public Health. 2020; 17(11). https://doi.org/10.3390/ijerph17113968 PMID: 32503285

17. Seissler J, Feghelm N, Then C, Meisinger C, Herder C, Koenig W, et al. Vasoregulatory peptides pro-endothelin-1 and pro-adrenomedullin are associated with metabolic syndrome in the population-based KORA F4 study. Eur J Endocrinol. 2012; 167(6):847–53. https://doi.org/10.1530/EJE-12-0472 PMID: 23002189

18. Flegal KM, Kit BK, Orpana H, Graubard BI. Association of all-cause mortality with overweight and obesity using standard body mass index categories: a systematic review and meta-analysis. Jama. 2013; 309(1):71–82. https://doi.org/10.1001/jama.2012.113905 PMID: 23280227

19. Sinning C, Ojeda F, Wild PS, Schnabel RB, Schwarz M, Ohdah S, et al. Midregional proadrenomedullin and growth differentiation factor-15 are not influenced by obesity in heart failure patients. Clin Res Cardiol. 2017; 106(6):401–10. https://doi.org/10.1007/s00392-016-1066-x PMID: 28004184

20. Neumann JT, Tzikas S, Funke-Kaiser A, Wilde S, Appelbaum S, Keller T, et al. Association of MR-proadrenomedullin with cardiovascular risk factors and subclinical cardiovascular disease. Atherosclerosis. 2013; 228(2):451–9. https://doi.org/10.1016/j.atherosclerosis.2013.03.006 PMID: 23562132

21. Ohlsson T, Nilsson PM, Persson M, Melander O. Midregional proadrenomedullin predicts reduced blood pressure and glucose elevation over time despite enhanced progression of obesity markers. J Hypertens. 2019; 37(3):590–5. https://doi.org/10.1097/HJH.0000000000002183 PMID: 30540625

22. Wong HK, Tang F, Cheung TT, Cheung BM. Adrenomedullin and diabetes. World J Diabetes. 2014; 5(3):364–71. https://doi.org/10.4239/wjd.v5.i3.364 PMID: 24936257
23. Calabro P, Yeh ET. Obesity, inflammation, and vascular disease: the role of the adipose tissue as an endocrine organ. Subcell Biochem. 2007; 42:63–91. PMID: 17612046

24. Unamuno X, Gómez-Ambrosi J, Rodríguez A, Becerrí S, Fruhbeck G, Catalán V. Adipokine dysregulation and adipose tissue inflammation in human obesity. Eur J Clin Invest. 2018; 48(9):e12997. https://doi.org/10.1111/eci.12997 PMID: 29995306

25. Fantuzzi G. Adipose tissue, adipokines, and inflammation. J Allergy Clin Immunol. 2005; 115(5):911–9; quiz 20. https://doi.org/10.1016/j.jaci.2005.02.023 PMID: 15867843

26. Li Y, Zhong X, Cheng G, Zhao C, Zhang L, Hong Y, et al. Hs-CRP and all-cause, cardiovascular, and cancer mortality risk: A meta-analysis. Atherosclerosis. 2017; 259:75–82. https://doi.org/10.1016/j.atherosclerosis.2017.02.003 PMID: 28327451

27. Sook Lee E, Park SS, Kim E, Sook Yoon Y, Ahn HY, Park CY, et al. Association between adiponectin levels and coronary heart disease and mortality: a systematic review and meta-analysis. Int J Epidemiol. 2013; 42(4):1029–39. https://doi.org/10.1093/ije/dyt087 PMID: 23739486

28. Yang S, Zhou M, Fowler DE, Wang P. Mechanisms of the beneficial effect of adrenomedullin and adrenomedullin-binding protein-1 in sepsis: down-regulation of proinflammatory cytokines. Crit Care Med. 2002; 30(12):2729–35. https://doi.org/10.1097/00003246-200212000-00018 PMID: 12483065

29. Isumi Y, Kubo A, Katafuchi T, Kangawa K, Minamino N. Adrenomedullin suppresses interleukin-1beta-induced tumor necrosis factor-alpha production in Swiss 3T3 cells. FEBS Lett. 1999; 463(1–2):110–4. https://doi.org/10.1016/s0014-5793(99)01615-4 PMID: 10601648

30. Dong Y, Chauhan M, Betancourt K, Belfort M, Yallapalli C. Adipose Tissue Inflammation and Adrenomedullin Overexpression Contribute to Lipid Dysregulation in Diabetic Pregnancies. J Clin Endocrinol Metab. 2018; 103(10):3810–8. https://doi.org/10.1210/jc.2018-00905 PMID: 30020508

31. Rathmann W, Strassburger K, Heier M, Holle R, Thorand B, Giani G, et al. Incidence of Type 2 diabetes in the elderly German population and the effect of clinical and lifestyle risk factors: KORA S4/F4 cohort study. Diabet Med. 2009; 26(12):1212–9. https://doi.org/10.1111/j.1464-5491.2009.02863.x PMID: 20002472

32. Huth C, von Toerne C, Schederrecker F, de Las Heras Gala T, Herder C, Kronenberg F, et al. Protein markers and risk of type 2 diabetes and prediabetes: a targeted proteomics approach in the KORA F4/FF4 study. Eur J Epidemiol. 2019; 34(4):409–22. https://doi.org/10.1007/s10654-018-0475-8 PMID: 30599058

33. Alberti KG, Zimmet P, Shaw J. Metabolic syndrome—a new world-wide definition. A Consensus Statement from the International Diabetes Federation. Diabet Med. 2006; 23(5):469–80. https://doi.org/10.1111/j.1464-5491.2006.01858.x PMID: 16681555

34. Alberti KG, Zimmet PZ. Definition, diagnosis and classification of diabetes mellitus and its complications. Part 1: diagnosis and classification of diabetes mellitus provisional report of a WHO consultation. Diabet Med. 1998; 15(7):539–53. PMID: 9686693

35. Caruvel P, Mazier C, Kunde J, Morgenthaler NG, Darbouret B. Homogeneous time-resolved fluoromunocassay for the measurement of midregional proadrenomedullin in plasma on the fully automated system B.R.A.H.M.S KRYPTOR. Clin Biochem. 2009; 42(7–8):725–8. https://doi.org/10.1016/j.clinbiochem.2009.01.002 PMID: 19318039

36. Herder C, Kannenberg JM, Carstensen-Kirberg M, Huth C, Meisinger C, Koenig W, et al. Serum levels of interleukin-22, cardiometabolic risk factors and incident type 2 diabetes: KORA F4/FF4 study. Cardiovasc Diabetol. 2017; 16(1):17. https://doi.org/10.1186/s12933-017-0498-6 PMID: 28143481

37. Ebert T, Gebhardt C, Scholz M, Wohland T, Schleinitz D, Fasshauer M, et al. Relationship Between 12 Adipocytokines and Distinct Components of the Metabolic Syndrome. J Clin Endocrinol Metab. 2018; 103(3):1015–23. https://doi.org/10.1210/jc.2017-02085 PMID: 29325128

38. Wolf K, Popp A, Schneider A, Breitner S, Hampel R, Rathmann W, et al. Association Between Long-Term Exposure to Air Pollution and Biomarkers Related to Insulin Resistance, Subclinical Inflammation, and Adipokines. Diabetes. 2016; 65(11):3314–26. https://doi.org/10.2337/db15-1567 PMID: 27605624

39. Herder C, Kannenberg JM, Niersmann C, Huth C, Carstensen-Kirberg M, Wittenbecher C, et al. Independent and opposite associations of serum levels of omentin-1 and adiponectin with increases of glycemia and incident type 2 diabetes in an older population: KORA F4/FF4 study. Eur J Endocrinol. 2017; 177(4):277–86. https://doi.org/10.1530/EJE-17-0100 PMID: 28679518

40. Carstensen-Kirberg M, Kannenberg JM, Huth C, Meisinger C, Koenig W, Heier M, et al. Inverse associations between serum levels of secreted frizzled-related protein-5 (SFRP5) and multiple cardiometabolic risk factors: KORA F4 study. Cardiovasc Diabetol. 2017; 16(1):109. https://doi.org/10.1186/s12933-017-0591-x PMID: 28831362

41. Lefevre AS, Stevens LA, Schmid CH, Zhang YL, Castro AF 3rd, Feldman HI, et al. A new equation to estimate glomerular filtration rate. Ann Intern Med. 2009; 150(9):604–12. https://doi.org/10.7326/0003-4819-150-9-200905050-00006 PMID: 19414839
42. Yildirim NC, Yurekli M. The effect of adrenomedullin and cold stress on interleukin-6 levels in some rat tissues. Clin Exp Immunol. 2010; 161(1):171–5. https://doi.org/10.1111/j.1365-2249.2010.04156.x PMID: 20456410

43. Zhou PH, Hu W, Zhang XB, Wang W, Zhang LJ. Protective Effect of Adrenomedullin on Rat Leydig Cells from Lipopolysaccharide-Induced Inflammation and Apoptosis via the PI3K/Akt Signaling Pathway ADM on Rat Leydig Cells from Inflammation and Apoptosis. Mediators Inflamm. 2016; 2016:7201549. https://doi.org/10.1155/2016/7201549 PMID: 27212810

44. Castellani G, Paliuri G, Orso G, Paccagnella N, D'Amore C, Facci L, et al. An intracellular adrenomedullin system reduces IL-6 release via a NF-kB-mediated, cAMP-independent transcriptional mechanism in rat thymic epithelial cells. Cytokine. 2016; 88:136–43. https://doi.org/10.1016/j.cyto.2016.09.003 PMID: 27619517

45. Mandal J, Roth M, Papakonstantinou E, Fang L, Savic S, Tamm M, et al. Adrenomedullin mediates pro-angiogenic and pro-inflammatory cytokines in asthma and COPD. Pulm Pharmacol Ther. 2019; 56:8–14. https://doi.org/10.1016/j.pupt.2019.01.006 PMID: 30690080

46. Wong HK, Ong KL, Leung RY, Lam TH, Thomas GN, Lam KS, et al. A single nucleotide polymorphism of interleukin-6 gene is related to plasma adrenomedullin levels. Clin Endocrinol (Oxf). 2013; 79(4):504–9. https://doi.org/10.1111/ce n.12078 PMID: 23088295

47. Kang KW, Ok M, Lee SK. Leptin as a Key between Obesity and Cardiovascular Disease. J Obes Metab Syndr. 2020; 29(4):248–59. https://doi.org/10.7570/jomes20120 PMID: 33342767

48. D’Elia L, Giaquinto A, Iacone R, Russo O, Strazzullo P, Galletti F. Serum leptin is associated with increased pulse pressure and the development of arterial stiffening in adult men: results of an eight-year follow-up study. Hypertens Res. 2021. https://doi.org/10.1038/s41440-021-00718-x PMID: 34385686