Metabolomic Profiling of Arginine Metabolome Links Altered Methylation to Chronic Kidney Disease Accelerated Atherosclerosis

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Abstract

Atherosclerotic cardiovascular disease is the leading cause of death in patients with chronic kidney disease (CKD), but the mechanisms underlying vascular disease has not been fully understood. As the nitrogen donor in nitric oxide (NO) synthesis, arginine and its metabolic products are integrally linked to vascular health and inflammation. We hypothesized that derangements in this pathway could explain, in part, increased atherosclerotic risk in CKD. We developed a targeted metabolomic platform to profile quantitatively arginine metabolites in plasma by liquid chromatography tandem mass spectrometry (LC/MS). Male low-density lipoprotein receptor deficient (LDLr-) mice at age 6 weeks were subjected to sham or 5/6 nephrectomy surgery to induce CKD. Subsequently, the animals were maintained on high fat diet for 24 weeks. Targeted metabolomic analysis of arginine metabolites in plasma was performed by isotope dilution LC/MS including asymmetric dimethyl arginine (ADMA), symmetric dimethyl arginine (SDMA), N-mono-methylarginine (NMMA), arginine and citrulline. Although elevated plasma levels of ADMA and SDMA were found in the CKD mice, only higher ADMA level correlated with degree of atherosclerosis. No significant differences were noted in levels of NMMA between the groups. CKD mice had high levels of citrulline and arginine, but ADMA levels had no correlation with either of these metabolites. These findings strongly implicate altered arginine methylation and accumulation of ADMA, may in part contribute to CKD accelerated atherosclerosis. It raises the possibility that interrupting pathways that generate ADMA or enhance its metabolism may have therapeutic potential in mitigating atherosclerosis.

Keywords: Chronic kidney disease; Animal model; Arginine methylation; Atherosclerosis; Asymmetric dimethyl arginine

Abbreviations: ADMA: Asymmetric Dimethyl Arginine; BUN: Blood Urea Nitrogen; CKD: Chronic Kidney Disease; CAD: Coronary Artery Disease; CV: Cardiovascular; DDAH: Dimethyl Arginine Dimethyl Amino Hydrolase; eNOS: endothelial Nitric Oxide Synthetase; EIC: Extracted Ion Chromatogram; FIA: Flow Injection Analysis; HILIC: Hydrophilic Interaction Liquid Chromatography; LC/MS: Liquid Chromatography/Mass Spectrometry; LDLr: Low Density Lipoprotein Receptor Deficient; MRM: Multiple Reaction Monitoring; NMMA: N Mono Methylarginine; PRMT: Protein Arginine Methyltransferases; SDMA: Symmetric Dimethyl Arginine

Introduction

Coronary artery disease (CAD) and cardiovascular (CV) events associates with chronic kidney disease (CKD) and is the leading cause of death in patients with CKD (>10-fold mortality). Indeed, CV events and mortality are more likely outcome than progression to end-stage renal disease (ESRD) in CKD. Sectional studies have demonstrated that the traditional risk factors are only partially predictive of CAD in CKD subjects, implying the presence of additional CKD-specific risk factors. In response to physiologic stimuli, endothelial cells dynamically regulate arterial vascular tone by producing vasodilators and vasoconstrictors. Risk factors for atherosclerosis, such as CKD, interfere with this response, promoting endothelial dysfunction and atherosclerosis. One key regulator is nitric oxide (NO), which is generated from L-arginine by endothelial nitric oxide synthase (eNOS) in the presence of cofactors such as tetrahydrobiopterin. Gaseous NO diffuses to vascular smooth muscle cells and activates guanylate cyclase, which in turn elevates cyclic guanosine monophosphate to promote vasodilation. NO’s antithrombotic nature prevents platelet aggregation, promotes fibrinolysis and decreases smooth muscle 2015

Asymmetric dimethylarginine (ADMA:20’5) is the naturally occurring dimethylated modification of arginine and is a known inhibitor of NOS. One pathway for producing ADMA is proteolysis of methylated proteins which are formed by protein arginine methyl transferases (PRMT) once released into plasma, ADMA can inhibit eNOS and decrease NO bioavailability, causing endothelial dysfunction. The enzyme dimethyl arginine dimethyl amino hydrolase (DDAH) metabolizes ADMA to generate dimethylamine and has two isoforms of which DDAH 1 is thought to be the primary enzyme responsible for ADMA degradation. The proximal tubules of kidneys can reabsorb almost all of the filtered L-arginine, very little ADMA is reabsorbed or excreted into urine. The majority of filtered ADMA is degraded into citrulline and dimethylamine by the renal DDAH as the kidneys has abundant amount of DDAH. In CKD, loss of DDAH 1 activity may limit ADMA breakdown and sterol isomer of ADMA is also produced by proteolysis following PRMT methylation of protein-bound arginine, but has no NO inhibitory activity and is renally excreted. 20’5-L-arginine (NMMA) is the precursor of both ADMA and

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Reagents and materials

Male C57BL/6 2015 mice were purchased from Jackson Labs, Bar Harbor, ME. Authentic and isotopically labelled standards were purchased from the following vendors: NMMA, SDMA and 2,15SDMA (Santa Cruz, CA2015ADMA (Novachem, Australia); creatinine, D, creatinine, arginine and citrulline (Sigma-Aldrich); 31Citra, Citrulline and 31Arginine (Cambridge Isotope Laboratories, MA). All LC reagents were purchased from Sigma Aldrich, St. Louis, MO. Rodent diets were purchased from 2015and high fat diet from Harlan Teklad.

Mouse models

All animal procedures were approved by the University of Michigan Committee on Use and Care of Animals. Six week old male C57BL/6 2015 mice were fed 2015 standard rodent diet that has 200 ppm cholesterol, 28.5% protein, 13.5% fat and 58.0% carbohydrates by calories. Mice were housed in a climate-controlled, light-regulated facility with a 12:12 hour light-dark cycle and water ad libitum. At age 7 weeks, mice were subjected to either sham surgery (Control, n=11) or to 5/6 nephrectomy to induce CKD (CKD, n =11). This was accomplished by removing entire right kidney in a first procedure and then subsequent removal of two thirds left kidney by dissection after one week interval. At 9 weeks of age, mice in each group were fed on high fat diet containing 19.5% protein, 40.5% fat, 0.5% cholesterol and 40.0% carbohydrates from Harlan Teklad (TD00243). Murine systolic blood pressure was measured by the IITC Life Science blood pressure system (Woodland Hills, CA) with a highly sensitive photoelectric sensor. The accuracy of measurements with five or more successful readings was obtained and secured by regular calibration of the pressure transducer. Blood was collected from saphenous veins in living mice with tubes containing dry ethylenediaminetetraacetate (EDTA). Hematocrit was measured by 2015 Micro-Hematocrit centrifuge with Digital Hematocrit Reader (2015 Company).

Analysis of kidney function

 Plasma creatinine levels was measured by liquid chromatography electron spray ionization and tandem mass spectrometry (LC/ESI/MS) using an Agilent 6410 MS coupled with 1200 LC system (Agilent, New Castle, DE), as described previously 2015of plasma (n=11, each group) and 2015of deuterated creatinine (10 2015were added into 2015of 2015m ammonium acetate solution, subjected to protein precipitation by 85% 2015of the supernatant was then injected for LC/MS analysis. A hydrophilic interaction chromatography (HILIC) was performed utilizing Luna Phenomenex column (2015, 2015 Torrance, CA) with an isocratic gradient of 85% acetonitrile with 20 mM ammonium acetate for 5 min at flow rate of 0.3 mL/min. The ion transition of 2015114 to 201544 for creatinine and 2015117 to 201547 creatinine was monitored in the multiple reaction monitoring (MRM) mode. The creatinine concentration in each plasma sample was determined by comparing the peak areas of the creatinine 2015-creatinine for the above transitions. Blood urea nitrogen (BUN) was measured directly on IDEXX VetTest 8008 chemistry analyzer (Westbrook, Maine) using dry slide technology.

Analysis of atherosclerosis

At 24 weeks mice were anesthetized, and the thoracic cavity was exposed and a small incision was made in the right cardiac auricle, and a cannula was inserted into the left ventricle. Through the left ventricle, the animal was perfused with phosphate-buffered saline until the eluent from the right auricle became clear, and then the left ventricle was injected with 3 mL of 10% buffered formalin. Finally, the entire mice were immersed in the fixative 2015. Each aortic tree was microdissected to remove adventitial fat and stained with Oil Red O (Sigma) to visualize neutral lipids, pinned on wax plates. The images of the aorta were captured on a digital camera. 2015 Quantification was performed with Image Pro software (Media Cybernetics, Bethesda, MD). The lesional areas are represented as ratios between surface area of atherosclerotic lesion stained with Oil Red O to the surface area of the entire aortic tree (n=11, each group).

Arginine metabolome profiling by LC/MS

The detailed method development and chromatography

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optimization strategy for arginine metabolomic profiling are discussed in Results. Targeted metabolomic analysis of arginine metabolome in plasma was performed by LC/MS in the positive mode (n=11, each group). Briefly, 2015of ethylenediaminetetraacetic acid anticoagulated plasma was spiked with 2,15ADMA (10pM/sample), 2,15(10pM/sample), 3133Arginine (600pM/sample) 2015citrulline (600pM/sample). Protein was precipitated with 2015acetoniitrile. 2015of the supernatant was subjected to HILIC with a Phenomenex Luna 3 2015column using an Agilent 1200 LC, at a flow rate of 2015. Solvent A was 10mM ammonium formate and solvent B was acetoniitrile with 0.1% formic acid. The column was equilibrated with 95% solvent B and 5% solvent A initially. The gradient was: 95-15% solvent B over 8 min, 15% solvent B for 6 min, 15-95% solvent B for 1 min and then finally 95% solvent B for 10 min.

The eluent from the LC was subjected MS analysis using an Agilent 6410 Triple Quadrupole MS system) connected in series to the LC, equipped with an electrospray source. Positive LC/ESI/MS was performed using following parameters: spray voltage 4000 V, drying gas flow 15 L/min, drying gas temperature 2015and nebulizer pressure 40 psi. Flow injection analysis (FIA) using 2015scan was used to optimize the fragmentor voltage and collision energy for arginine, citrulline, NMMA, ADMA, and SDMA. To obtain the best signal-to-noise ratio for quantitation, the most abundant ions from each compound were chosen for use in MRM mode 2015Limits of detection (LOD) were calculated using peak areas corresponding to greater than five times signal to noise ratio. Data analysis was performed with Agilent Mass Hunter Analyst software (Version B6, Agilent, Santa Clara, CA). In preliminary studies, we found that the correlation between the peak areas of labelled standards and authentic compounds remained linear and greater than 0.95 irrespective of internal standard concentrations being a log fold higher or lower than the authentic compound concentration for all the arginine metabolites. The amount of isotope labelled standard spiked was subsequently individualized to each metabolite based on roughly 1:1 ratio of what is expected in the physiological range. Ratios of the peak area of metabolites to the spiked isotope labelled compounds were used for quantification since the amount of spike was known for each analyte that was measured. The most abundant fragment ion was chosen as the MRM transitions for each 2015

Statistical analysis

Results are given as 2015. Differences between the groups were considered significant at p<0.05 using the program GraphPad Prism version 6.00 for Windows (La Jolla, California, USA) for the independent t test when comparing characteristics of the different groups. Pearson’s correlation was used to measure the strength of a linear association between two variables using SPSS software for version 22 (SPSS Inc., Chicago, Illinois). A two-sided p<0.05 was considered significant.

Results

Development of a targeted arginine metabolome profiling platform by LC/MS

Selection of column and LC conditions for optimal separation of the analytes: To select the most appropriate column and chromatography technique, we attempted several different columns with both HILIC and reverse phase to optimize the separation. The columns and LC conditions were based on previous literature. We first examined the Phenomenex phenyl hexyl column (2.0 2015, 2015, Torrance, CA) with 10mM ammonium formate as solvent A and 201510.1% formic acid (1:3) as solvent B: at a flow rate of 0.25 mL/min. The column was equilibrated with 100% solvent A initially and then decreased to 0% from 1-3 min, and then finally 100% solvent B from 3-5 min. This resulted in very broad peak shapes. We then used the Agilent C18 reverse phase column (2.0 2015+ 0.1% formic acid as solvent A and acetoniitrile with 0.1% formic acid as solvent B at flow rate of 0.3 mL/min. The gradient was 100% solvent A to begin with which was decreased to 0% at 6 min and 100% solvent B was maintained between 4 min and 6 min. The peak areas were sharp but all compounds eluted in the first column volume consistent with non-retention of the analytes to the column. We then tested the Waters Symmetry C18 column (2.1 2015mm, 3.5μm, Milford, MA) with 10mM ammonium acetate as solvent A and 100% acetoniitrile as solvent B at 0.3 mL/min flow rate. The gradient began with 100% solvent A for 0.5 min, increased linearly between 0.5 min and 5 min to 100% solvent B and stayed at 100% solvent B between 5 and 6 min. This method produced very poor peak shapes. While for the most part peak shapes were acceptable, reverse phase separation resulted non-retention of the analytes as they were eluted within either void or two-column volumes and therefore were sub-optimal. Finally, we attempted Phenomenex HILIC 2015column (Torrance, CA)) at a flow rate of 2015with solvent A 10mM ammonium formate and solvent B acetoniitrile with 0.1% formic acid. The column was equilibrated with 95% solvent B and 5% solvent A initially. The gradient was: 95-15% solvent B over 8 min, 15% solvent B for 6 min, 15-95% solvent B for 1 min and then finally 95% solvent B for 10 min. This method produced optimal peaks, retention times and shapes for ADMA, SDMA, NMMA, arginine and citrulline. Therefore, this column and method was chosen for optimizing MS conditions.

Optimization of MS parameters: Using FIA of authentic standards, the fragmentor voltage, cell acceleration voltage and collision energy was optimized for each individual compound on an Agilent 6410 triple quadrupole MS. The molecular ion [M + H]2015arginine and 2015Arginine are m2015175 and 181 respectively and the loss of 2015105 yields an intense ion at 20154and 74 respectively. The molecular ion [M 2015ADMA, 2,15ADMA, SDMA and 2015SDMA are 2015, 210, 203 and 209 respectively. The loss of 2015133 is the major fragment ion yielding product ions at 201570, 77, 70 and 76 respectively. The molecular ion [M 202015citrulline and 2015Citrulline are 2015182 and 181 respectively and the loss 2015175 yields fragments at 2015and 164 respectively. These transitions form ideal candidates for MRM 2015The limit of detection (LOD) using this methodology was 64 fmol for arginine, 21 fmol for ADMA, 2 fmol for SDMA, 14 fmol for NMMA and 20 fmol for citrulline. The interassay and intra assay variability were 2 to 10% and less than 3% respectively for all compound to internal standard ratios.

<table>
<thead>
<tr>
<th>Analyte</th>
<th>Molecular ion (m/z)</th>
<th>Daughter ions (m/z)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ADMA</td>
<td>203</td>
<td>46*, 70, 88, 116, 158</td>
</tr>
<tr>
<td>d12ADMA</td>
<td>210</td>
<td>46*, 77, 71, 86, 165</td>
</tr>
<tr>
<td>SDMA</td>
<td>203</td>
<td>70*, 116, 88, 165</td>
</tr>
<tr>
<td>d12SDMA</td>
<td>209</td>
<td>70*, 77, 94, 60, 165</td>
</tr>
<tr>
<td>NMMA</td>
<td>189</td>
<td>70*, 57</td>
</tr>
<tr>
<td>Arginine</td>
<td>175</td>
<td>70*, 60, 43</td>
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<tr>
<td>13C1Arginine</td>
<td>181</td>
<td>74*, 61, 44, 74</td>
</tr>
<tr>
<td>Citrulline</td>
<td>176</td>
<td>159*, 70,113</td>
</tr>
<tr>
<td>13C1Citrulline</td>
<td>181</td>
<td>164*, 75, 118</td>
</tr>
</tbody>
</table>

Table 1: MS spectral characteristics following collision induced dissociation (CID) of arginine metabolites. Following ESI/MS in positive mode, MS spectrum was obtained for each arginine metabolite. The molecular ion and major daughter ions following CID are represented. *represents the most intense daughter ion that was chosen for MRM transition monitoring for quantification.
2015 depicts the extracted ion chromatogram (EIC) for the MRM transitions for ADMA (Panel A), SDMA (Panel C), D5-SDMA (Panel D), NMMMA (Panel E), arginine (Panel F), 3H-arginine (Panel G), citrulline (Panel H) and D15-SDMA (Panel I) with the optimized HILIC separation. The MRM transitions noted in 2015 utilized for quantitative measurements of arginine metabolite in plasma.

**CKD mouse model has biochemical evidence of CKD and increased atherosclerosis**

The CKD mice at 24 weeks had significantly higher plasma creatinine (1.75 ± 0.34 mg/dL; n = 11; p<0.001) and BUN (44.2015 ± 17 mg/dL; n = 11; p<0.001). The CKD mice had significantly lower body weight and hematocrit. The CKD mice did not show significant differences in cholesterol levels, mineral metabolism (Calcium, Phosphorus and intact parathyroid hormone) or blood 2015. We performed 2015 analysis of the entire aorta and stained with Oil Red O to determine lesion area. Figure 4 Panel A and Panel B depict a representative control and CKD mouse aortic following Oil Red O stain. The CKD mice had increased atherosclerotic lesion area compared with control mice (0.201504 vs 0.201501; (n=9); p<0.05) (Figure 4: panel C; n=11 per group). The data strongly support induction of CKD as a major factor that accelerates atherosclerosis in this model.

**Table 2:** Physiological and Biochemical characteristics of study animals (n=11; each group)

<table>
<thead>
<tr>
<th></th>
<th>CONTROL (n=11) Mean ± SD</th>
<th>CKD (n=11) Mean ± SD</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Body weight (g)</td>
<td>33.52 ± 1.5</td>
<td>27.39 ± 0.6</td>
<td>&lt;0.01</td>
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<td>Blood Pressure (mm Hg)</td>
<td>102.70 ± 4.1</td>
<td>116.30 ± 9.2</td>
<td>NS</td>
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<tr>
<td>Hematocrit (%)</td>
<td>61.0 ± 0.9</td>
<td>52.91 ± 1.2</td>
<td>&lt;0.0001</td>
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<tr>
<td>Serum Creatinine (mg/dL)</td>
<td>0.97 ± 0.34</td>
<td>1.75 ± 0.54</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Blood urea nitrogen (mg/dL)</td>
<td>28.25 ± 1.17</td>
<td>44.17 ± 1.79</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

**Altered arginine methylation in CKD mice**

Plasma ADMA, SDMA and NMMMA levels were measured in plasma collected from the Control and CKD mice (n = 11 for each group). The plasma ADMA level was elevated in CKD mice compared to control mice (0.2015 ± 0.2015 vs. 0.2015 ± 0.2015 p<0.01). Similarly plasma SDMA was elevated in CKD mice compared to control mice (0.2015 ± 0.2015 vs. 0.2015 ± 0.2015 p<0.05). The values for NMMMA were not different between the two 2015 arginine methylation index, defined as the ratio of dimethylated arginines to the monomethylated precursor (ADMA+SDMA)/NMMA 2015 was significantly higher in the CKD mice, when compared to the control mice (29.2015 ± 20.2015 vs. 20.2015 ± 20.2015 p<0.05; 2015 Figure 3 Panels E and F depict elevations of plasma arginine (80.66 ± 2015 vs. 62.2015 ± 2015 p<0.05) and Citrulline (5.79 ± 2015 vs. 5.79 ± 2015 p<0.05). The data strongly support that CKD alters arginine metabolism, raising the possibility that such alterations may diminish NO bioavailability, contributing to vascular dysfunction.

**Levels of altered arginine methylation products and their substrates correlate with each other**

The ADMA levels of both control and CKD mice together correlate with SDMA levels (r = 0.549, p < 0.01) while both arginine and citrulline levels correlate with each other (r = 0.449, p < 0.05; 2015). Using Pearson correlation, the substrate for ADMA production—arginine and the by-product of ADMA degradation—citrulline levels do not correlate with ADMA levels (arginine r=0.36 p=0.09; citrulline r=0.24; p=0.582015). This implies that plasma citrulline is predominantly derived from arginine and not from ADMA. We also performed Spearman correlation analysis between ADMA, SDMA, NMMMA and arginine and between citrulline and arginine which showed similar results (Data not shown).

**ADMA levels, but not other arginine metabolites, are strongly associated with atherosclerotic burden**

We tested whether arginine metabolites correlated with atherosclerotic burden by comparing levels of the metabolites with lesion area, a measure of degree of atheroma in control and CKD mice by performing Pearson Correlation 2015). Only ADMA levels correlated with the lesion area (r = 0.64, p<0.01) while SDMA levels did not (r = 0.02, p>0.05). Similarly, NMMMA and arginine methylation index did not correlate with degree of atherosclerosis (Data not shown). The association of ADMA to atherosclerosis was stronger in the CKD mice. This data strongly supports the notion that ADMA alters arginine metabolism, raising the possibility that such alterations may diminish NO bioavailability, contributing to vascular dysfunction.

**Discussion**

In this study, we utilized a mouse model of atherosclerosis to...
address whether alterations in arginine metabolism, in part accounts for increased atherosclerosis observed in CKD. We utilized LC/MS to profile and quantitatively measure arginine and its methylated derivatives in plasma. We established that ADMA, SDMA, NMMA levels and the arginine methylation index are elevated in CKD mice. ADMA levels are directly related to the extent of atherosclerosis in CKD mice, suggesting a central role for altered NO bioavailability in atheroma formation in this model. Finally, we provide evidence that the ADMA elevation is not entirely related to availability of its substrate arginine or correlated to levels of its byproduct citrulline but might be a result of potential enzymatic pathways that lead to formation of methylated proteins or its degradation.

LC/MS methodology for measurement of these methylated arginines remains the gold standard for accurately measuring these methylated arginines. Arginine and its methylated metabolites are extremely polar and are not well separated with standard reverse phase chromatography. Using HILIC, we are able to achieve optimal separation. Importantly, with the use of tandem spectrometry we are able to measure levels of methylated arginines in the low femtomolar range, which makes it highly sensitive especially for rodent studies where sample availability is limiting. We used four isotopically labeled standards; D7 ADMA, D5 arginine and D5 citrulline to accurately determine the instrument response and to account for ion suppression and matrix effects for each specific compound in contrast to previously published methods that use one labeled standard for many 2015. This method is the first to use labeled SDMA to accurately determine its concentration independent of ADMA. Using the appropriate isotope labeled standard, this methodology enables a very accurate measurement of the arginine metabolites with minimal interassay and intraassay variability. This method also eliminates derivatization, solid phase extraction and other complicated procedures like ultrafiltration and uses a straightforward protein precipitation process, minimizing sample loss and expense.

The LDL receptor deficient mouse is a well-characterized model for atherosclerosis that develops extensive lesions in aortic root and branches and perivascular system as described in 2015. The effect of CKD in this mouse model has also been described previously and mainly results in acceleration of the lesions similar to non-CKD 2015. The mouse however does not develop coronary atherosclerosis and hence we measured aortic root lesion area as this is the most reproducible measurement. Following 5/6 nephrectomy, the mice demonstrate features of CKD like increased BUN and creatinine. The mice do not have increased blood pressure or changes in the calcium phosphorus metabolism, making CKD the sole variable that could potentially

<table>
<thead>
<tr>
<th></th>
<th>ADMA</th>
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<th>NMMA</th>
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<td>.124</td>
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<td>SDMA</td>
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<td>.123</td>
<td>.449*</td>
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**. Correlation is significant at the 0.01 level (2-tailed).
*. Correlation is significant at the 0.05 level (2-tailed).
contribute to atherosclerosis. The atherosclerosis in this model is accelerated with high fat diet and similar models in the 2015 are previously 2015

Interestingly, a study by Jacobi et al reported no change in ADMA levels in subtotal nephrectomized mice in 2015 background fed on chow diet for 12-2015These findings are in contrast to earlier publications that show increased ADMA levels in CKD models and our current 2015 Also surprisingly in that work, 2015 CKD mice did not have higher atherosclerotic burden when compared with DDAH I overexpressing mice on an 2015 background. Over expression of DDAH in this model however predictably decreased ADMA levels but did not change atherosclerotic lesions. It is likely that these differences are attributable to strain differences, more modest reduction in renal function and diet compared to our study. In our work, CKD mice have accelerated atherosclerosis, manifested as increased luminal lipid accumulation, elevated aortic plaque, necrotic core, fibrosis as well as greater luminal narrowing. Selective alterations of arginine methylation were identified in our study and may in part account for propensity towards increased atherosclerosis in this model. Indeed, the atherosclerotic burden correlated only with ADMA levels, but not SDMA, NMMA or arginine methylation index.

CKD is associated with decreased NO bioavailability either due to decrease in production due to substrate (arginine) limitation, increased levels of ADMA or increased utilization or presence of increased 2015 We demonstrate in our work that changes in ADMA in our model is not related to free arginine and citrulline in plasma suggesting that ADMA levels are not entirely just a consequence of increased arginine levels due to decreased renal function. Elevated ADMA levels are probably a result of increased flux in this pathway, partly due to increased methylation of proteins by PRMT, decreased renal excretion and decreased DDAH1 activity. In a previous study, it was demonstrated that ADMA levels are elevated in puromycin-induced CKD rats due to reduced DDAH 2015 Studies in mongrel dogs have also revealed microvascular endothelial changes in CKD dogs that was associated with increased ADMA levels and down regulation of DDAH- II 2015 changes in DDAH might be associated with loss-of-function polymorphisms of a2015 gene, functional inhibition of the enzyme by oxidative stress in CKD and end-stage renal disease, or 2015 DDAH1 overexpression in the 2015 in a previous study demonstrated decreased plaque area in a non CKD model associated with lower ADMA 2015 but this has not been tested in CKD mice. Future studies will need to focus on DDAH isoforms and the effect of DDAH overexpression in relevant CKD models.

In clinical studies higher ADMA levels predicted cardiovascular events when compared to control patients in pre-dialysis subjects 2015 In a large clinical study, Wang et al demonstrated that arginine methylation index correlated with CVD 2015 Therefore, we tested the utility of this measurement in our CKD mouse model, but this measure did not correlate with atherosclerotic burden, perhaps due to small numbers in our study. ADMA together with related markers of oxidative stress like myeloperoxidase could potentiate development of 2015 Renal cyclooxygenase 2 (COX-2) inhibition raises ADMA levels and could explain the increased cardiovascular morbidity of COX-2 inhibitors adding to ADMA’s role in cardiovascular 2015 Thus, ADMA together with inflammatory and oxidative markers could play a central role in the accelerated atherosclerotic burden in CKD.

Our model has several strengths. We demonstrate for the first time that in a pathophysiologically relevant model of CKD atherosclerosis altered arginine methylation and a high ADMA levels correlate with atherosclerotic burden. These changes occur even with modest CKD suggesting that these pathways could be relevant in early CKD. The mouse model does not have common associated features of CKD such as hypertension, and abnormal mineral metabolism which make it an ideal model to study effect of mild CKD alone. The limitations of this work include small numbers in this study and lack of manipulations to alter ADMA levels to show causality with atherosclerotic burden. Manipulation of PRMT, DDAH or the alternate enzyme alanine-glyoxalate aminotransferase-2 (AGXT2201) could provide such direct evidence and future studies need to focus on this issue. Finally, our findings raise the possibility that interrupting arginine methylation pathways could provide a therapeutic avenue for combating CKD-atherosclerosis.

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**References**


