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Upregulated Kynurenine Pathway Enzymes in Aortic Atherosclerotic Aneurysm: Macrophage Kynureninase Downregulates Inflammation

Masanori Nishimura[1, 2], Atsushi Yamashita[2], Yunosuke Matsuura[3], Junichi Okutsu[4], Aiko Fukahori[4], Tsuyoshi Hirata[4], Tomohiro Nishizawa[5], Hirohito Ishii[1], Kazunari Maekawa[2], Eriko Nakamura[2], Kazuo Kitamura[3], Kunihide Nakamura[1] and Yujiro Asada[2]

[1]Division of Cardiovascular Surgery, Department of Surgery, Faculty of Medicine, University of Miyazaki, Miyazaki, Japan
[2]Department of Pathology, Faculty of Medicine, University of Miyazaki, Miyazaki, Japan
[3]Department of Internal Medicine, Faculty of Medicine, University of Miyazaki, Miyazaki, Japan
[4]Translational Research Department, Daiichi Sankyo RD Novare Co., Ltd., Tokyo, Japan
[5]Specialty Medicine Research Laboratories I, Daiichi Sankyo Co., Ltd., Tokyo, Japan

Aims: Inflammation and hypertension contribute to the progression of atherosclerotic aneurysm in the aorta. Vascular cell metabolism is regarded to modulate atherogenesis, but the metabolic alterations that occur in atherosclerotic aneurysm remain unknown. The present study aimed to identify metabolic pathways and metabolites in aneurysmal walls and examine their roles in atherogenesis.

Methods: Gene expression using microarray and metabolite levels in the early atherosclerotic lesions and aneurysmal walls obtained from 42 patients undergoing aortic surgery were investigated (early lesion $n=11$, aneurysm $n=35$) and capillary electrophoresis–time-of-flight mass spectrometry (early lesion $n=14$, aneurysm $n=38$). Using immunohistochemistry, the protein expression and localization of the identified factors were examined (early lesion $n=11$, non-aneurysmal advanced lesion $n=8$, aneurysm $n=11$). The roles of the factors in atherogenesis were analyzed in macrophages derived from human peripheral blood mononuclear cells.

Results: Enrichment analysis using 35 significantly upregulated genes (log2 ratio, >3) revealed the alteration of the kynurenine pathway. Metabolite levels of tryptophan, kynurenine, and quinolinic acid and the kynurenine-to-tryptophan ratio were increased in the aneurysmal walls. Gene and protein expression of kynureninase and kynurenine 3-monooxygenase were upregulated and localized in macrophages in the aneurysmal walls. The silencing of kynureninase in the cultured macrophages enhanced the expression of interleukin-6 and indoleamine 2,3-dioxygenase 1.

Conclusion: Our study suggests the upregulation of the kynurenine pathway in macrophages in aortic atherosclerotic aneurysm. Kynureninase may negatively regulate inflammation via the kynurenine pathway itself in macrophages.

Key words: Atherosclerosis, Aneurysm, Kynureninase, Kynurenine pathway, Macrophage

Introduction

Worldwide, cardiovascular disease is the leading cause of death. In 2015, there were an estimated 8.9 million deaths due to ischemic heart disease, 6.3 million deaths due to cerebrovascular disease, and 170,000 deaths due to aortic aneurysm globally[1)]. Atherosclerosis and its complications are the major pathology underlying these cardiovascular diseases. Atherosclerosis is a gradually progressive, inflammatory disease caused by chronic endogenous or exogenous endothelial injuries, or both, and the accumulation of

Address for correspondence: Yujiro Asada, Department of Pathology, Faculty of Medicine, University of Miyazaki, 5200 Kihara, Kiyotake, Miyazaki 889-1692, Japan E-mail: yasada@med.miyazaki-u.ac.jp

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blood contents.

Aortic atherosclerosis is clinically silent, and most atherosclerotic aneurysms are incidentally found during imaging study and suddenly become symptomatic upon rupture or embolism. An aortic angioscopic study reported widely distributed but silent aortic atherosclerosis with vulnerable morphology, plaque rupture, and mural thrombus formation in patients with stable angina pectoris[2]. The mortality rate of ruptured aneurysms is extremely high, and 30 day mortality was found to occur in 27% or 42% of patients treated with endovascular repair or open repair[3]. Optimal medical therapies and a regular follow-up imaging study to measure the diameter are standard medical management; however, no medical drugs have managed to stabilize aortic aneurysm[4]. Thus, a deeper understanding of the mechanisms underlying aortic atherosclerosis and aneurysm formation is required.

The alteration of vascular cell metabolism is considered to contribute to the development of atherosclerosis and reflect a high-risk condition. Clinical and basic studies have reported the accumulation of ^{18}F-fluorodeoxyglucose (FDG) in advanced atherosclerotic plaques, as shown by positron emission tomography, which indicates enhanced glucose uptake in the vascular cells[5, 6]. Recent metabolomic studies have identified metabolites that deepen our understanding of atherogenesis[7], and we have confirmed increased levels of glycolysis, pentose phosphate pathway, and citric acid cycle metabolites in rabbit macrophage-rich atherosclerotic lesions[8]. The increased levels of FDG uptake and expression of a glycolytic enzyme, hexokinase II, were shown to reflect vascular wall hypoxia and thrombogenicity[9, 10]. Amino acid metabolites, such as arginine and homoarginine, play significant roles in endothelial anti-thrombosis and vascular tone, and endothelial dysfunction via nitric oxide and reactive oxygen species synthesis, respectively[11]. Although the transcriptomic analysis of abdominal aortic aneurysm identified alterations of inflammatory-, tissue remodeling-, and hypoxia-related genes[12, 13], knowledge of the altered metabolic pathways and metabolites in aortic atherosclerotic aneurysm is limited.

The present study aimed to identify metabolic pathways and metabolites in aneurysmal walls and investigate their roles in atherogenesis.

Materials and Methods

Patient Population

The present study examined aortic tissue (n=52) removed from 42 patients who underwent aortic surgery for aneurysm between April 2013 and November 2015. The risk factors for atherosclerosis and aneurysm, medications, and data of blood cell counts and blood chemistry were collected from the clinical records. The ethics committees of the participating institutions approved the study protocol (2012–013).

Tissue Processing

Tissue samples were taken from aneurysmal lesions and non-aneurysmal anastomotic sites, and each sample was separated for microarray, metabolomic, and histological assays. The samples were immersed in RNA Later (Qiagen, Hilden, Germany) for microarray assay and stored at −80℃ until this assay, were frozen in liquid nitrogen and stored at −80℃ for metabolomic assay, and then were fixed in 4% paraformaldehyde and embedded in paraffin for histological evaluation. Four-micrometer-thick sections were stained with hematoxylin and eosin. The histological specimens were identified according to the AHA atherosclerosis classification[14]. The specimens from the aortic lesions were classified as early atherosclerotic lesions (AHA I to III), non-aneurysmal advanced lesion (AHA V and VI), and aneurysmal lesion (AHA V and VI). Tissues with dense calcified atherosclerotic lesions were excluded from the present study considering their low cellularity.

RNA Microarray Analysis and Quantitative Polymerase Chain Reaction (PCR)

Total RNA was extracted from 52 aortic tissues using TRIzol reagent (Sigma, St. Louis, MO, USA) and was purified using the PureLink RNA Mini kit (Thermo Fisher Scientific, Waltham, MA, USA), following the manufacturer's instructions. Six samples with an RNA integrity number of less than seven were excluded. The early atherosclerotic lesions (n=11) and aneurysmal lesions (n=35) were subjected to transcriptomic examination using Affymetrix GeneChip Human Genome U133 Plus 2.0 array (Affymetrix, Santa Clara, CA, USA). The fluorescent signals were quantified by the Robust Multi-array Average method. To identify altered pathways, enrichment analysis was conducted using lists of differentially expressed genes (log2 ratio, >3.0 and <-3.0; Q value, <0.05) (Enrichr_1GO biological process 2018)[15].

Single-stranded, complementary DNA was synthesized from the RNA using PrimeScript RT reagent kits (Perfect Real Time, Takara Bio, Kusatsu, Japan) and used for real-time PCR. The gene

expression of indoleamine 2,3-dioxygenase (IDO)1, tryptophan 2,3-dioxygenase (TDO2), kynurenine 3-monooxygenase (KMO), kynureninase, 3-hydroxyanthranilate 3,4-dioxygenase, interleukin (IL)-6, IL-10, matrix metalloproteinase (MMP)-2, MMP-9, tissue factor, and β-actin was measured using a LightCycler 96 (Roche Diagnostics GmbH, Mannheim, Germany), SYBR Premix EX Taq II (Perfect Real Time, Takara Bio), and specific primers **(Supplementary Table 1)**. The gene expression was normalized to that of β-actin in aortic tissue or was expressed as the fold change relative to the control level using the $\Delta\Delta$Ct method in a cell culture experiment.

Metabolomic Profiling of Aortic Tissue using Capillary Electrophoresis–Time-of-Flight Mass Spectrometry (CE-TOFMS)

Metabolomic analysis was conducted using the early atherosclerotic lesions (n=14) and aneurysmal lesions (n=38). Metabolite extraction and metabolome analysis were conducted at Human Metabolome Technologies, Inc. (HMT) (Tsuruoka, Japan). Aortic samples were weighed, and then, the tissues (30 mg) were added to 500 µL of methanol, containing 50 µM internal standard and homogenized, using a tissue disruptor. Homogenates were mixed with 200 µL of Milli-Q water and 500 µL of chloroform and then separated by centrifugation at 2,300×g for 5 min at 4℃. The upper aqueous layer (400 µL) was passed through a filter (5 kDa cut-off; Millipore) by centrifugation at 9,100×g for 120 min at 4℃ to remove proteins. The filtrates were lyophilized and suspended in 50 µL of Milli-Q water for metabolomic analysis.

Metabolomic analysis was conducted with the Basic Scan package of HMT using CE-TOFMS based on methods described previously[16]. Briefly, CE-TOFMS analysis was conducted using an Agilent CE capillary electrophoresis system equipped with an Agilent 6210 time-of-flight mass spectrometer, Agilent 1100 isocratic HPLC pump, Agilent G1603A CE-MS adapter kit, and Agilent G1607A CE-ESI-MS sprayer kit (Agilent Technologies, Waldbronn, Germany). The systems were controlled by Agilent G2201AA ChemStation software version B.03.01 for CE (Agilent Technologies) and connected by a fused silica capillary (50 µm i.d. ×80 cm total length) with commercial electrophoresis buffer (H3301-1001 and H3302-1021 for cation and anion analyses, respectively; HMT) as the electrolyte. The spectrometer scanned from m/z 50 to 1,000. Peaks were extracted using MasterHands, automatic integration software (Keio University Tsuruoka Town Campus, Tsuruoka, Japan) to obtain peak information including m/z, peak area, and migration time (MT). Signal peaks corresponding to isotopomers, adduct ions, and other product ions of known metabolites were excluded, and the remaining peaks were annotated according to the HMT metabolite database based on their m/z values with the MTs. The areas of the annotated peaks were then normalized based on internal standard levels and sample volumes to obtain relative levels of each metabolite. Hierarchical clustering analysis and principal component analysis were conducted using HMT's proprietary software, PeakStat and SampleStat, respectively. Detected metabolites were plotted on metabolic pathway maps using VANTED software.

Histology and Immunohistochemistry

Four-micrometer-thick sections of early atherosclerotic lesions (n=11), non-aneurysmal advanced lesions (n=8), and aneurysmal lesions (n=11) were immunohistochemically stained using antibodies against the smooth muscle cell marker smooth muscle actin (mouse monoclonal, clone 1A4; DAKO, Agilent Technologies, Santa Clara, CA, USA), the macrophage marker CD68 (mouse monoclonal, clone PGM-1; DAKO), the endothelial marker CD31 (mouse monoclonal, clone CJ70A; DAKO), the T-cell marker CD3 (mouse monoclonal, clone F7.2.38; DAKO), the B-cell marker CD20 (mouse monoclonal, clone L26; DAKO), the plasma cell marker CD138 (mouse monoclonal, clone MI15; DAKO), kynureninase (rabbit polyclonal; GenTex, Irvine, CA, USA), and KMO (rabbit polyclonal; Atras Antibodies AB, Stockholm, Sweden). The negative controls included non-immune mouse or rabbit IgG (Jackson ImmunoResearch, Baltimore, MA, USA). The sections were stained with EnVision anti-mouse or rabbit immunoglobulin (DAKO). Horseradish peroxidase activity was visualized using 3,3'-diaminobenzidine containing hydrogen peroxide, and the sections were counterstained with Meyer's hematoxylin. The microscopic images of immunopositive areas were captured using an image analysis software (cellSens Standard; Olympus, Tokyo, Japan). The areas of the antigens were semi-quantified using a color image analysis system (WinRoof; Mitani, Fukui, Japan). These areas are expressed as ratios of positively stained areas per vascular area, as described by Kuroiwa *et al.*[17].

Isolation of Human Peripheral Blood Mononuclear Cells (PBMCs) and Macrophage Differentiation

PBMCs were obtained from healthy volunteers using SepMate™-50 (StemCell Technologies, Vancouver, Canada), following the manufacturer's

instructions with some modifications. Briefly, Ficoll-Paque™ PLUS (GE Healthcare, Chicago, IL, USA) was filled through the insert in the SepMate™ tube and pipetted onto the tube insert with a 30 mL blood sample diluted 50% in Dulbecco's phosphate-buffered saline containing 2% fetal bovine serum (Sigma-Aldrich, St. Louis, MO, USA) and 1 mM ethylenediaminetetraacetic acid (Wako, Tokyo, Japan). After centrifugation at 1,200 × g for 15 min at room temperature (RT), the plasma layer containing PBMCs was transferred to another new Falcon tube. Then, platelets were removed through further centrifugation at 200 × g for 10 min at RT. Following supernatant removal, cell pellets were resuspended in RPMI1640 containing 10% heat-inactivated fetal bovine serum and antibiotics (Zellshield®; Minerva Biolabs, Berlin, Germany), and cells were counted and uniformly seeded into tissue culture plates. Then, a cocktail of recombinant human granulocyte colony-stimulating factor (2 ng/mL; PEPROTECH, Rocky Hill, NJ, USA) and recombinant human macrophage colony-stimulating factor (50 ng/mL; PEPROTECH) was added to the PBMCs and incubated for 7 days for their differentiation into macrophages. After the removal of non-adherent cells, macrophages were washed three times with Dulbecco's phosphate-buffered saline and used for the following experiments. The institutional review board at the University of Miyazaki approved the present study (Approval No. O-0272).

IFN-γ Stimulation and Kynureninase Inhibition by Small Interfering RNA (siRNA) in Macrophages Derived from PBMCs

Macrophages were stimulated with recombinant human interferon-γ (rhIFN-γ; R & D Systems, Minneapolis, MN, USA) at a final concentration of 10 ng/mL in serum- and antibiotic-free RPMI1640 for 24 h. Macrophages were transfected by incubating human kynureninase siRNA (Silencer® Select, siRNA ID; s17103 and/or s17104; Life Technologies, Carlsbad, CA, USA) or negative control siRNA (Silencer® Select, Negative Control #2 siRNA; Life Technologies) at a final concentration of 37.5 nM under the serum- and antibiotic-free RPMI1640 for 24 h, using Viromer® Blue transfection reagent (Lipocalyx). To examine siRNA-induced cell damage, lactate dehydrogenase (LDH) in the medium was measured using the LDH cytotoxicity assay kit (Nacalai Tesque, Kyoto, Japan). First, 100 µL of the supernatant from each well in which macrophages had been cultured under the indicated conditions was transferred to an optically transparent 96-well plate. A total of 100 µL of substrate solution was added and incubated for 20 min at RT under protection from light to each well of the above plate. Finally, 50 µL of stop solution was added to each well, and the absorbance was measured at 490 nm using a microplate reader. The results are presented as absorbance values and expressed as mean ± standard deviation (SD).

IL-6 ELISA

The level of IL-6 in culture medium was measured by Quantikine® ELISA (R & D Systems), following the manufacturer's instructions.

Statistical Analysis

Data are expressed as median and range or mean ± SD. Differentially expressed genes were calculated using unpaired *t*-test with Benjamini–Hochberg multiple testing correction for the microarray assay results. Differences between the groups on quantitative PCR, metabolite levels, and the results of cell culture experiment were analyzed using the Mann–Whitney *U* test. Differences among the groups were analyzed using the Kruskal–Wallis test with Dunn's multiple comparison test. Q value (adjusted *P* value using the Benjamini–Hochberg method) of <0.05 and *P* value of <0.05 were deemed to indicate statistical significance.

Results

Patient Characteristics

Supplementary Table 2 shows the clinical characteristics of the 42 patients with aortic aneurysm. Their mean age was 73 yr, and 76% were male. The aneurysms were located in thoracic aorta ($n=17$, 40%) and abdominal aorta ($n=25$, 60%). Generally, 95% of the patients had hypertension, 79% had dyslipidemia, and 45% had diabetes or impaired glucose tolerance. In histological sections, plaque disruption with mural thrombus formation was observed in 63% of atherosclerotic aneurysms and 25% of non-aneurysmal advanced atherosclerotic lesions but in no early atherosclerotic lesions.

Gene Expression in Early Atherosclerosis and Atherosclerotic Aneurysm

To conduct a comprehensive evaluation of gene expression in the early atherosclerotic and atherosclerotic aneurysmal lesions, we conducted transcriptome analysis with microarray in the aortic lesions. **Table 1** shows a list of 35 differentially expressed genes (log2 ratio, >3; Q value, <0.05). Enrichment analysis using the 35 upregulated genes revealed alterations of the kynurenine pathway,

Table 1. Differentially expressed upregulated genes (Log2 ratio >3, Q<0.05)in aneurysm

Probeset_Id	Gene_Symbol	Log2Ratio	Q value
204580_at	*MMP12*	6.57	1.86164E-08
236028_at	*IBSP*	5.85	1.07746E-10
209395_at	*CHI3L1*	5.56	4.52479E-07
203936_s_at	*MMP9*	4.81	2.79815E-11
37892_at	*COL11A1*	4.50	1.5779E-14
223484_at	*C15orf48*	4.34	1.75656E-08
212942_s_at	*CEMIP*	4.32	8.46489E-13
212657_s_at	*IL1RN*	4.31	2.41151E-10
204475_at	*MMP1*	4.20	3.33309E-11
204259_at	*MMP7*	4.14	8.82392E-08
219434_at	*TREM1*	3.97	2.0773E-15
206134_at	*ADAMDEC1*	3.71	1.10951E-06
204430_s_at	*SLC2A5*	3.66	5.77635E-12
222939_s_at	*SLC16A10*	3.58	1.52386E-11
213909_at	*LRRC15*	3.58	8.62686E-12
205943_at	*TDO2*	3.44	2.70123E-13
210004_at	*OLR1*	3.43	2.81577E-05
206858_s_at	*HOXC6*	3.42	0.00037992
202888_s_at	*ANPEP*	3.41	8.62166E-17
205242_at	*CXCL13*	3.40	9.46235E-07
203665_at	*HMOX1*	3.39	1.72862E-13
209351_at	*KRT14*	3.25	1.47257E-07
211643_x_at	*IGK ///IGKC*	3.23	3.33309E-11
210845_s_at	*PLAUR*	3.18	7.53482E-11
211645_x_at	*IGKV1-17////IGKV1-17*	3.18	3.63523E-11
205568_at	*AQP9*	3.18	3.35934E-10
209875_s_at	*SPP1*	3.17	2.38724E-05
227566_at	*LOC102725271///NTM*	3.14	6.51337E-08
211634_x_at	*IGHM*	3.14	1.32342E-08
32128_at	*CCL18*	3.09	0.003213613
204638_at	*ACP5*	3.08	4.84916E-06
202859_x_at	*CXCL8*	3.06	8.59447E-07
230748_at	*SLC16A6*	3.05	1.87046E-08
209555_s_at	*CD36*	3.02	1.91741E-08
205306_x_at	*KMO*	3.02	1.03391E-08

cysteine-type endopeptidase activity involved in the apoptotic signaling pathway, extracellular matrix disassembly, and B-cell chemotaxis, among others **(Supplementary Table 3)**. **Table 2** shows a list of the 21 differentially expressed genes (log2 ratio, <−3; Q value, <0.05). Enrichment analysis using the 21 downregulated genes revealed alterations of the regulation of bone resorption, blood pressure, idolealkylamine metabolic pathway, synaptic transmission, and smooth muscle cell differentiation, among others **(Supplementary Table 4)**. The results suggest enhancement of kynurenine pathway activity in atherosclerotic aneurysm among metabolic pathways. Real-time PCR assay confirmed the overexpression of kynurenine pathway enzymes, namely, IDO1, TDO2, kynureninase, and KMO, and the downregulation of 3-hydroxyanthranilate 3,4-dioxygenase in atherosclerotic aneurysm, compared with the levels in the early atherosclerotic lesions **(Fig. 1)**. The mRNA of IDO1, TDO2, kynureninase, and KMO also overexpressed in non-aneurysmal advanced atherosclerotic lesions, compared with the levels in the early atherosclerotic lesions **(Supplementary Fig. 1)**. There were no genes that were differentially expressed (Q<0.05) between non-diabetics and those with diabetes mellitus/impaired glucose tolerance.

Table 2. Differentially expressed downregulated genes (Log2 ratio <-3, Q<0.05) in aneurysm

Probeset_Id	Gene_Symbol	Log2Ratio	q.value
229339_at	*MYOCD*	-3.01	1.04239E-09
206163_at	*MAB21L1///MIR548F5*	-3.02	2.86195E-05
237094_at	*FAM19A5*	-3.08	4.73173E-08
206339_at	*CARTPT*	-3.21	0.001015519
210302_s_at	*MAB21L2*	-3.22	2.66037E-05
226281_at	*DNER*	-3.22	8.05926E-07
205358_at	*GRIA2*	-3.27	7.54217E-11
219791_s_at	*HAND2-AS1*	-3.34	9.79795E-07
213228_at	*PDE8B*	-3.39	5.33939E-09
228796_at	*CPNE4*	-3.44	5.30392E-06
209793_at	*GRIA1*	-3.47	1.57857E-09
227209_at	*CNTN1*	-3.56	1.49466E-09
210292_s_at	*PCDH11X///PCDH11Y*	-3.69	4.09956E-07
214961_at	*MTUS2*	-3.80	8.21645E-11
1563933_a_at	*PLD5*	-3.80	2.20896E-12
231626_at	*TPH1*	-3.83	2.18209E-15
219837_s_at	*CYTL1*	-4.04	2.69301E-08
239913_at	*SLC10A4*	-4.08	8.77746E-06
213745_at	*ATRNL1*	-4.12	4.89661E-13
229831_at	*CNTN3*	-4.45	9.43889E-14
223869_at	*SOST*	-4.65	1.55716E-08

Metabolic Differences in Early Atherosclerosis and Atherosclerotic Aneurysm

To conduct a comprehensive evaluation of metabolic status in early atherosclerotic and atherosclerotic aneurysmal lesions, we conducted metabolomic analysis with CE-TOFMS in the aortic lesions. This analysis identified 258 metabolites (149 cationic and 109 anionic metabolites) in the aortic tissues. **Fig. 2** and **Supplementary Table 5** show the hierarchical clustering analysis results of metabolites in the early atherosclerotic and aneurysmal lesions. The hierarchical tree indicates the relative distance among the metabolite peaks. The red or green densities indicate high or low concentrations of metabolites, respectively. The hierarchical clustering analysis revealed metabolic differences between early atherosclerotic and atherosclerotic aneurysmal lesions.

Supplementary Table 6 shows metabolites whose relative levels were more than 1.5-fold (19 metabolites) or less than one-half (22 metabolites) of the levels in the atherosclerotic aneurysmal lesions and whose levels were detectable as at least one-half of the samples in aneurysmal lesions (for "abundant" metabolites) or early atherosclerotic lesions (for "deficient" metabolites). The abundant metabolites were associated with the kynurenine pathway (kynurenine and quinolinic acid), glycolysis/gluconeogenesis (dihydroxyacetone phosphate and fructose 6-phosphate), and individual pathway. By contrast, the deficient metabolites were associated with purine metabolism (inosine, adenine, adenosine triphosphate, adenosine monophosphate, adenosine diphosphate, and adenosine), glycolysis/gluconeogenesis (3-phosphoglyceric acid, phosphoenolpyruvic acid, and 2,3-diphophoglyceric acid), pentose phosphate pathway (sedoheptulose 7-phosphate and 6-phosphogluconic acid), and individual pathway.

Relative metabolite levels of tryptophan, kynurenine, and quinolinic acid were higher in aneurysmal lesions than those in the early atherosclerotic lesions **(Fig. 1, Supplementary Table 7)**. Among early atherosclerotic lesions, quinolinic acid was detectable in only one lesion. The kynurenine-to-tryptophan ratio was also higher in aneurysmal lesions (0.025 ± 0.015, $n=38$) than in the early atherosclerotic lesions (0.016 ± 0.005, $n=14$; $p<0.001$).

Macrophages Express Kynureninase and KMO in Atherosclerotic Aneurysm

The transcriptomic and metabolomic analyses revealed a significant alteration of the kynurenine pathway in atherosclerotic aneurysmal lesions. We examined the protein expression and localization of kynureninase and KMO in the aortic lesions. **Fig. 3, Supplementary Fig. 2, and Table 3** show cellular

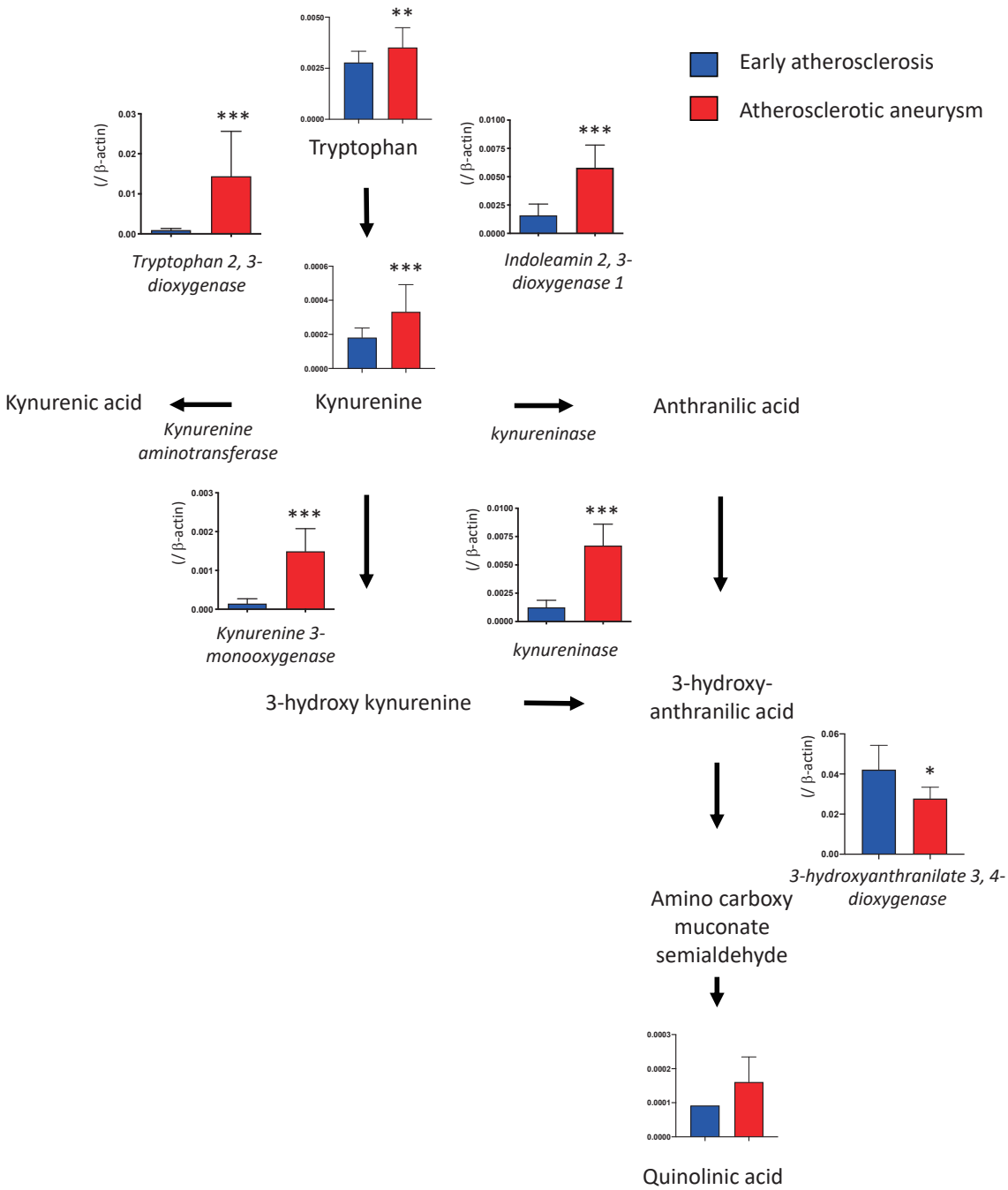

Fig. 1.

mRNA expression of kynurenine pathway enzymes and relative amounts of metabolites in the early atherosclerotic lesion and atherosclerotic aneurysm (enzymes, $n=11$ in each; metabolites, early atherosclerotic lesions $n=14$, atherosclerotic aneurysms $n=38$; in early lesions, quinolinic acid was detected in one lesion). $^{*}p<0.01$, $^{**}p<0.001$, $^{***}p<0.0001$, Mann–Whitney U test

components and the expression of kynurenine pathway enzymes in early atherosclerotic, non-aneurysmal advanced atherosclerotic, and atherosclerotic aneurysmal lesions. The areas immunopositive for macrophages, T cells, kynureninase, and KMO were larger in the non-aneurysmal advanced atherosclerotic and atherosclerotic aneurysmal lesions than those in the early atherosclerotic lesions. The areas immunopositive for B cells and plasma cells were larger in the atherosclerotic aneurysmal lesions than those in the early atherosclerotic lesions. The area immunopositive for SMCs progressively decreased

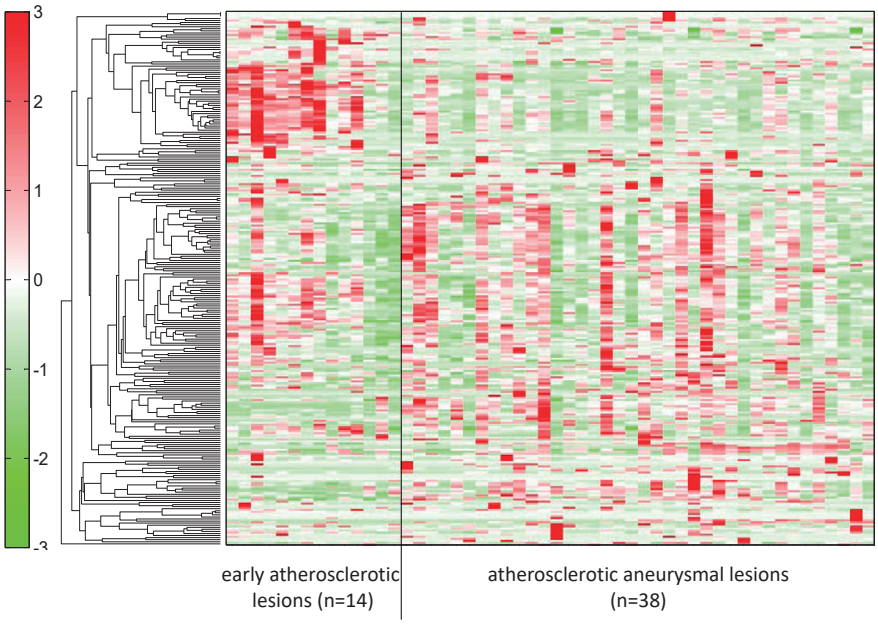

Fig. 2. Metabolomic analysis of early atherosclerotic lesions and atherosclerotic aneurysms.

Representative heatmap created using hierarchical clustering analysis shows metabolic differences between groups. Supplementary Table 5 shows the original data for each metabolite

and was significantly smaller in the atherosclerotic aneurysmal wall than that in the early atherosclerotic aorta **(Table 3)**. Kynureninase was predominantly expressed in macrophages in non-aneurysmal advanced atherosclerotic lesions and atherosclerotic aneurysm **(Fig. 3A, Supplementary Fig. 2)**. KMO was expressed in macrophages in early atherosclerotic intima, non-aneurysmal advanced atherosclerotic lesion, and the aneurysmal wall **(Fig. 3A, Supplementary Fig. 2)** and also expressed in adventitial peripheral nerve bundles with lymphoid aggregates in atherosclerotic aneurysm **(Fig. 3B)**. The lymphoid aggregates comprised B cells, plasma cells, T cells, macrophages, and capillary vessels **(Fig. 3B)**.

Upregulation of IDO1 and IL-6 by Kynureninase Inhibition in Cultured Macrophages

To examine the roles of kynureninase in atherogenesis, we conducted cell culture experiments with macrophages derived from PBMCs. Macrophages differentiated with M-CSF and GM-CSF expressed kynurenine pathway enzymes, namely, TDO2, kynureninase, and KMO. IFN-γ stimulation significantly upregulated the expression of IDO1, TDO2, kynureninase, IL-6, and tissue factor, whereas KMO expression did not alter such expression **(Fig. 4)**. Kynureninase siRNA suppressed the mRNA levels to approximately 25%. The kynureninase inhibition increased the expression of IDO1 and IL-6 but not of TDO2 and KMO, MMP-2, MMP-9, IL-10, and tissue factor **(Fig. 5A)**. Kynureninase inhibition also increased the protein expression of IL-6 **(Fig. 5B)**. The kynureninase inhibition and increases in the expression of IDO1 and IL-6 were confirmed with two different kynureninase siRNAs in macrophages. The siRNA did not alter the release of LDH from the macrophages **(Supplementary Fig. 3)**.

Discussion

The transcriptomic and metabolomic assays identified an enhanced activity of the kynurenine pathway in aortic atherosclerotic aneurysm. Kynureninase and KMO were upregulated in the advanced atherosclerotic lesions and localized in macrophages. Kynureninase inhibition in the cultured macrophages enhanced the expression of IDO1 and IL-6.

Kynurenine pathway enzymes and metabolites were shown to be increased in non-aneurysmal advanced atherosclerotic lesions and atherosclerotic aneurysmal wall compared with the levels in early atherosclerosis. Tryptophan, an essential amino acid, is used for protein synthesis or metabolized to bioactive metabolites via the kynurenine and serotonin pathways. Tryptophan is considered to be constitutively metabolized to kynurenine via TDO2 in the liver and brain or metabolized via Th1-cytokine-inducible

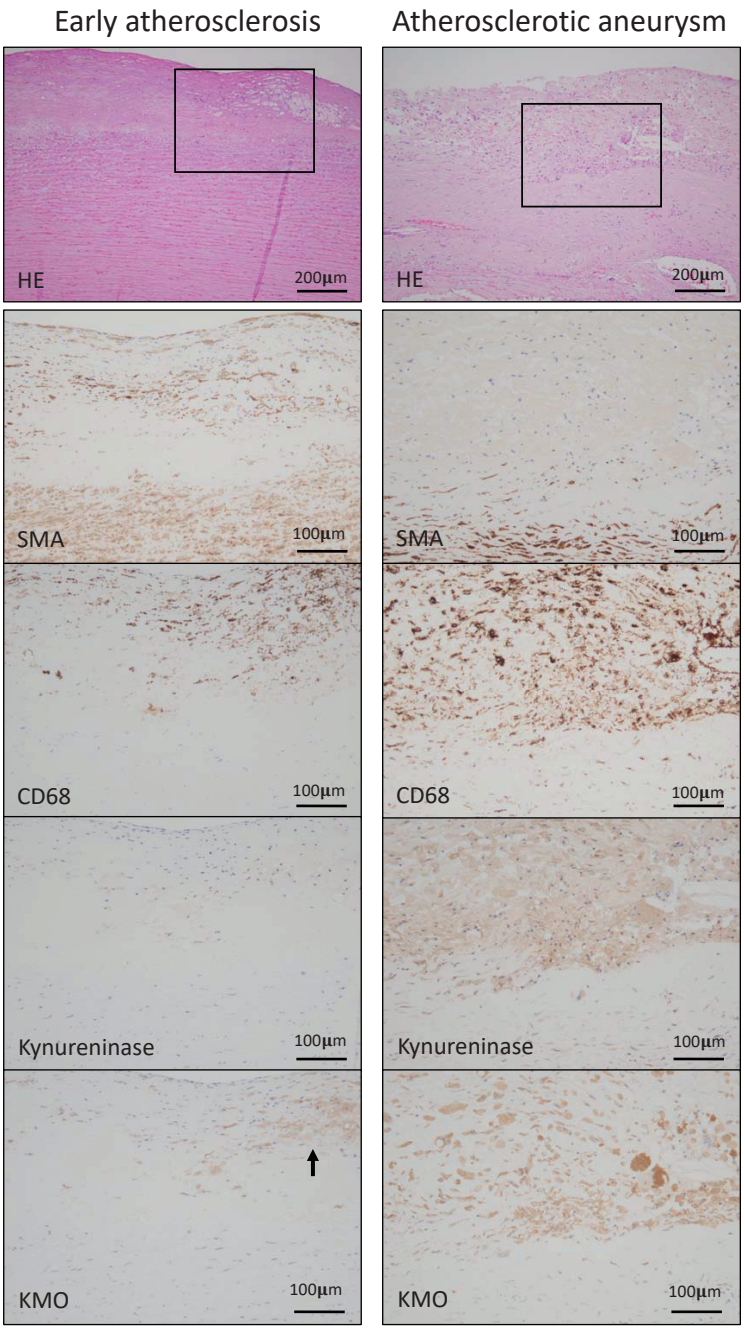

Figure 3A

Fig. 3. Representative histological and immunohistochemical images for cells and kynurenine pathway enzymes in the early atherosclerotic lesion and atherosclerotic aneurysm

A. Hematoxylin–eosin (HE) staining and immunohistochemistry for smooth muscle cells (SMA), macrophages (CD68), kynureninase, and kynurenine 3-monooxygenase (KMO) in the early atherosclerotic lesion (left column) and atherosclerotic aneurysm (right column). Squares in HE staining indicate areas of high magnification of immunohistochemical images. The left column shows fibrous intimal thickening with the accumulation of foamy macrophages. The intimal macrophages express KMO (arrow) but not kynureninase. The right column shows advanced atherosclerotic plaque formation and medial degeneration and angiogenesis. Macrophages express both kynureninase and KMO.

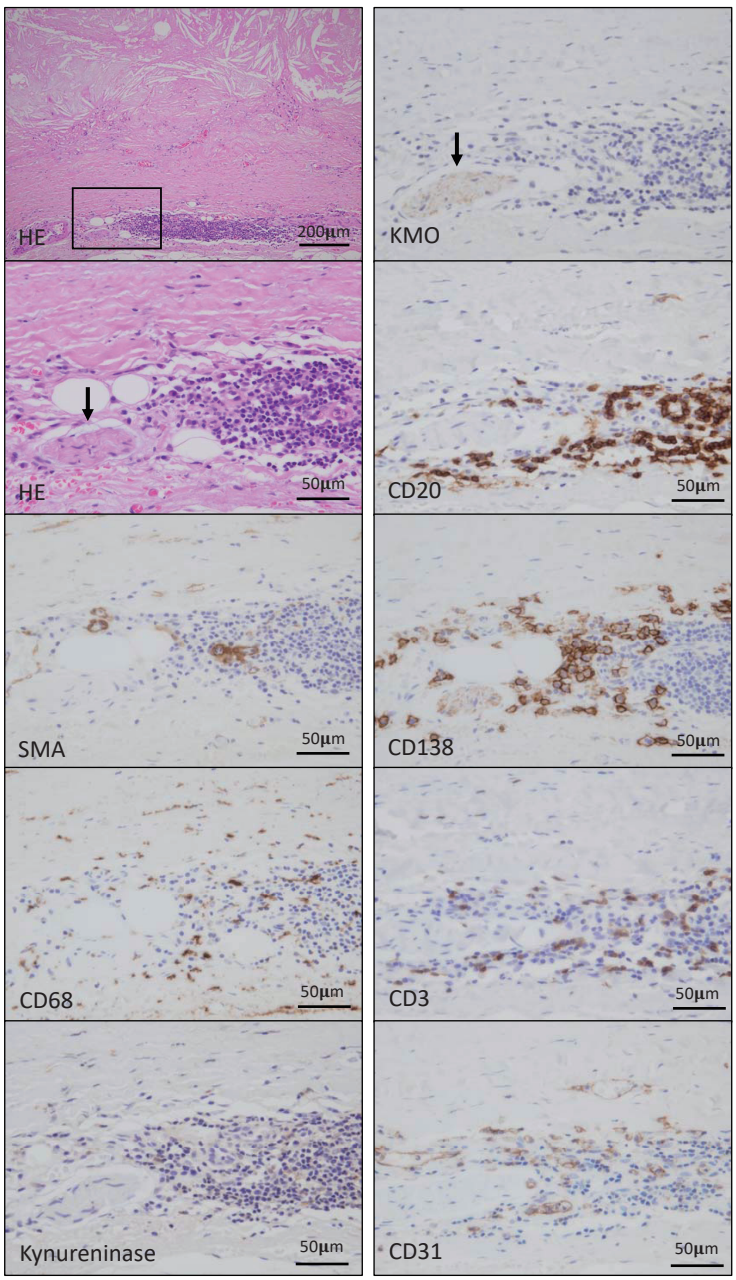

Figure 3B

Fig. 3. Representative histological and immunohistochemical images for cells and kynurenine pathway enzymes in the early atherosclerotic lesion and atherosclerotic aneurysm

B. HE staining and immunohistochemistry for smooth muscle cells (SMA), macrophages (CD68), kynureninase, KMO, B cells (CD20), plasma cells (CD138), T cells (CD3), and endothelial cells (CD31) in atherosclerotic aneurysm. Square in HE staining indicates an area of other high-magnification images. Advanced atherosclerotic change with medial degeneration and accumulation of lymphoid cells in adventitia. A peripheral nerve bundle expresses KMO (arrows) but not kynureninase and closely localizes with the lymphoid aggregate. Lymphoid aggregate comprises B cells, plasma cells, T cells, and macrophages with capillary vessels.

Table 3. Immunopositive area of cells and kynurenine enzymes in aorta

	A. early lesion ($n=11$)	B. non-aneurysmal advanced lesion ($n=8$)	C. aneurysmal lesion ($n=11$)	*p* value		
				A vs. B	A vs. C	B vs. C
SMA (smooth muscle cell)	30.8 (10.8 - 42.4)	16.2 (12.7 - 25.5)	8.1 (0.2 - 26.6)	0.12	<0.001	0.36
CD68 (macrophage)	0.01 (0 - 0.08)	1.6 (0.6 - 5.5)	4.9 (0.3 - 11.9)	<0.01	<0.0001	1
CD3 (T cell)	0.004 (0 - 0.03)	0.04 (0.01 - 0.2)	0.03 (0 - 0.7)	<0.01	<0.01	1
CD20 (B cell)	0.007 (0 - 0.02)	0.01 (0.003 - 0.07)	0.42 (0.001 - 5.1)	1	<0.05	0.13
CD138 (plasma cell)	0.004 (0.001 - 0.2)	0.019 (0.006 - 0.11)	0.16 (0.01 - 0.53)	0.12	<0.001	0.29
CD31 (endothelial cell)	1.2 (0.7 - 3.4)	0.8 (0.3 - 2.0)	3.2 (0.9 - 6.3)	0.87	<0.05	<0.01
Kynureninase	0.004 (0 - 0.19)	0.4 (0.2 - 3.5)	1.4 (0.004 - 15.1)	<0.05	<0.001	1
Kynurenine 3-monooxygenase	0.02 (0 - 0.15)	0.5 (0.2 - 2.7)	0.6 (0 - 4.8)	<0.01	<0.01	1

Data are expressed as median and range. SMA (smooth muscle actin) Kruskal-Wallis test with Dunn's multiple comparison test.

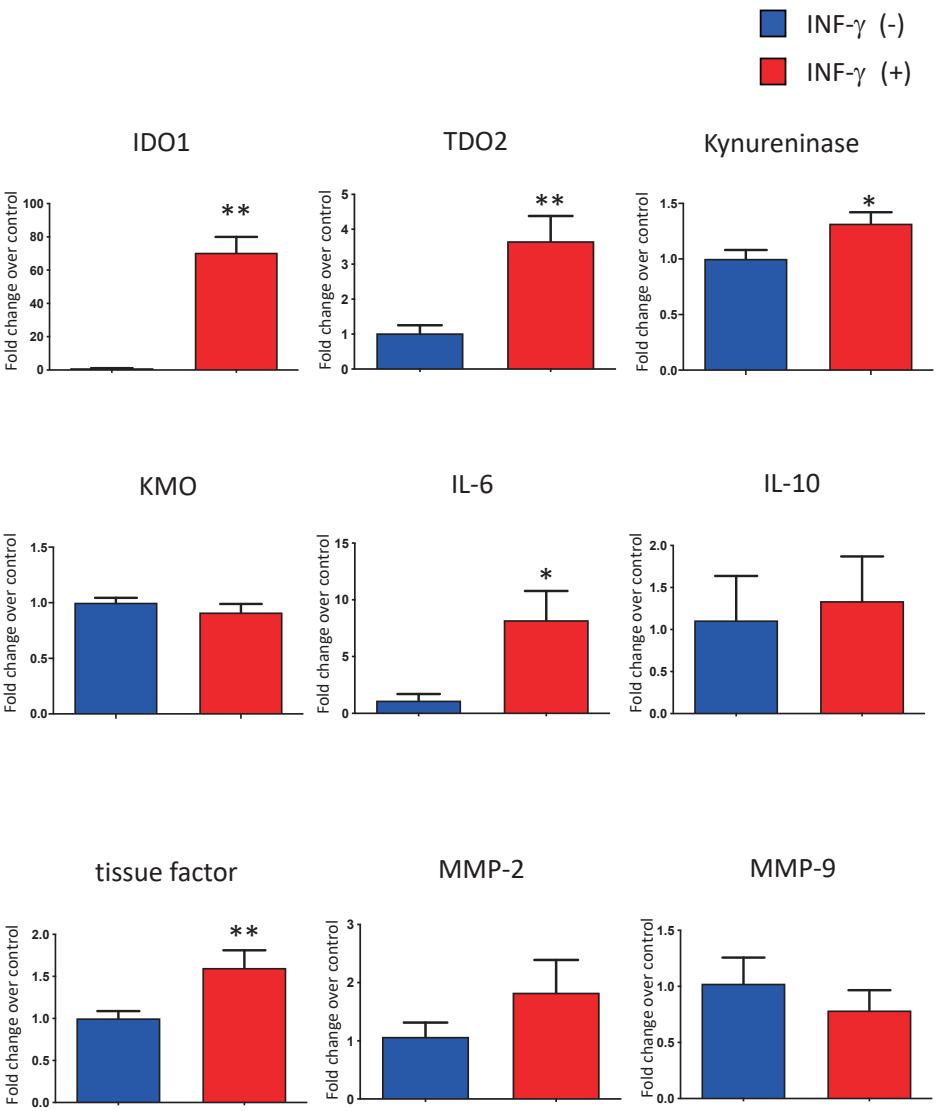

Fig. 4.

Effect of IFN-γ stimulation on the expression of genes encoding kynurenine pathway enzymes, IL-6, IL-10, tissue factor, MMP-2, and MMP-9, in human PBMCs-derived macrophages. $^{*}p<0.05$, $^{**}p<0.01$, $n=4$ or 5 in each, Mann–Whitney *U* test

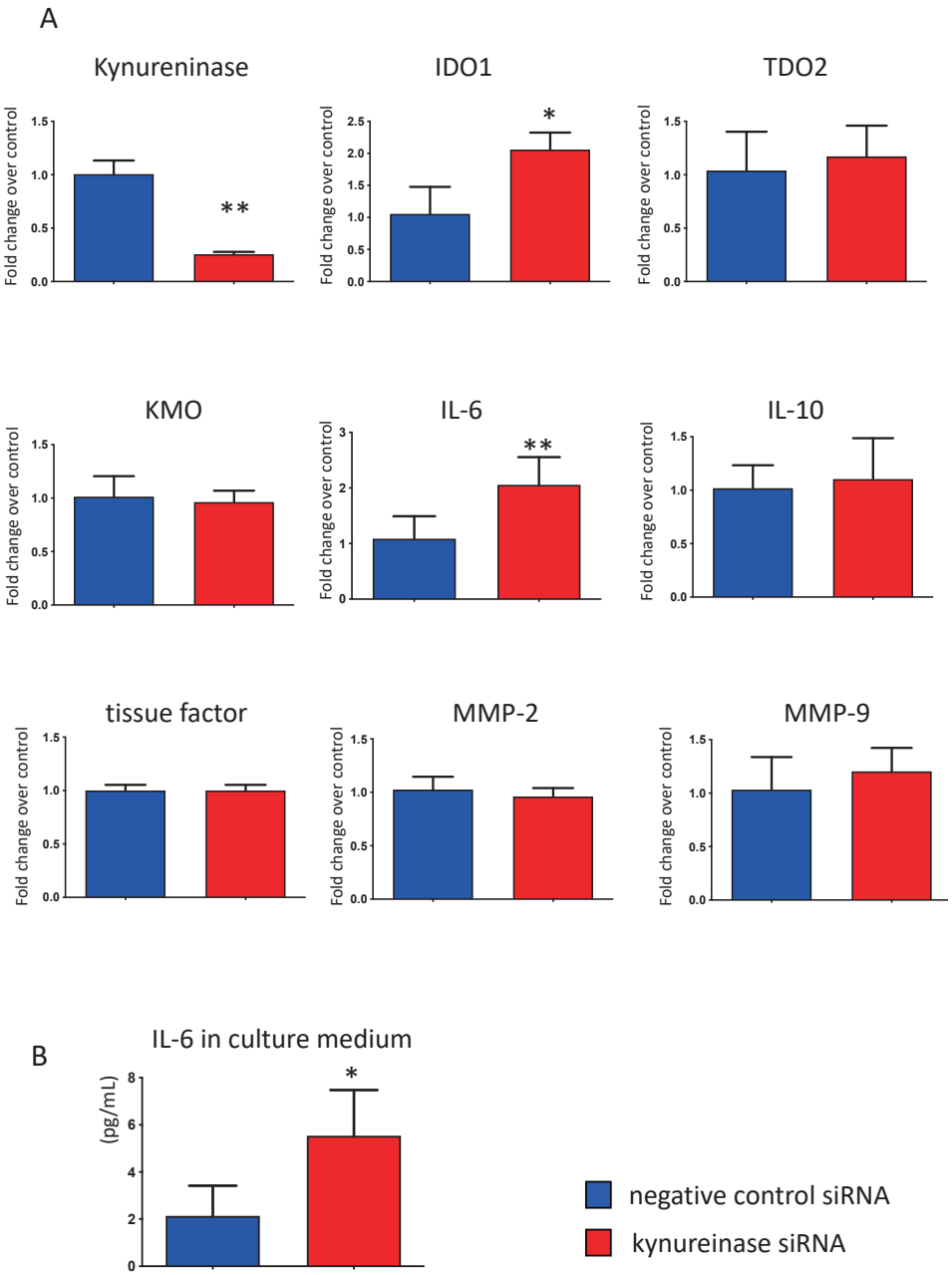

Fig. 5. Effect of kynureninase inhibition by siRNA

(A) The effect of kynureninase inhibition by siRNA for 24 h on the expression of genes encoding kynurenine pathway enzymes, IL-6, IL-10, tissue factor, MMP-2, and MMP-9, in human PBMCs-derived macrophages. (B) The protein levels of IL-6 in culture medium treated with kynureninase siRNA and negative control siRNA for 24 h. $^{*}p<0.05$, $^{**}p<0.01$, $n=4$ or 5 in each, Mann–Whitney U test

IDO1[18]. Clinical studies have shown increases in the serum kynurenine level and kynurenine-to-tryptophan ratio in ischemic heart disease[19] and indicated that downstream kynurenine metabolites, anthranilic acid, hydroxyanthranilic acid, hydroxykynurenine, and kynurenic acid **(Fig. 1)**, were associated with an increased risk of acute myocardial infarction[20]. Plasma levels of quinolinic acid were also found to be significantly associated with the intima to media ratio of the carotid artery in patients with end-stage renal disease[21]. These lines of evidence suggested that the kynurenine pathway is upregulated in cardiovascular disease, but the sources of the metabolites were unclear. Jung *et al.*[22] conducted metabolomic analysis

using aortic tissue with or without plaque from thoracic aortic aneurysm and identified increased levels of tryptophan, kynurenine, and kynurenine-to-tryptophan ratio in aorta with atherosclerotic plaques compared with the levels in control aorta. Our data are comparable with those of that study and support the notion that advanced atherosclerotic wall is a source of kynurenine metabolites. However, another metabolomic study using carotid and femoral artery plaques and intimal thickening did not reveal alterations in the levels of kynurenine metabolites[23]. Several studies examined the transcriptome in abdominal aortic aneurysm and showed the upregulation of genes related to the immune system/inflammation, tissue remodeling, and hypoxia[12, 13, 24]. However, these studies did not show kynurenine pathway enzymes as being differentially expressed, except for TDO2 in one study[24]. The different study designs and differences in control samples could have caused this discrepancy in results. TDO2 and IDO1 were upregulated in the non-aneurysmal advanced atherosclerotic lesion and aneurysmal wall and unstimulated cultured macrophages expressed TDO2. Therefore, tryptophan could be constitutively and inducibly metabolized to kynurenine via two enzymes in the advanced atherosclerotic lesions.

IDO1 may affect atherogenesis, aneurysmal formation, and atherothrombus formation. Presently, there is conflicting evidence on its effects on atherogenesis. The inhibition or gene deletion of IDO1 promoted inflammation and atherosclerosis in $Apoe^{-/-}$ mice fed a chow diet[25, 26]. Meanwhile, the gene deletion of IDO1 inhibited inflammation and atherosclerosis in $Ldlr^{-/-}$ mice fed with a chow diet[27]. The gene deletion of IDO1 reduced the formation of angiotensin II-induced abdominal aortic aneurysm in $Apoe^{-/-}$ and $Ldlr^{-/-}$ mice[28, 29]. Our results support the involvement of the kynurenine pathway in the formation of advanced atherosclerotic lesions in humans. IDO1 was localized in coronary macrophages and was a regulator of the thrombogenic potential of atherosclerotic plaque via the expression of tissue factor, a blood coagulation initiator[30]. As previously reported[30], IFN-γ stimulation markedly upregulated IDO1 and enhanced tissue factor expression. Meanwhile, the inhibition of a downstream enzyme, kynureninase, did not affect tissue factor expression in macrophages. Hence, IDO1 may contribute to the thrombogenic potential of advanced atherosclerotic lesions. This could affect mural thrombus formation on the aortic advanced atherosclerotic lesions.

Kynureninase was upregulated in the non-aneurysmal advanced atherosclerotic lesion and aneurysmal wall and localized in macrophages. Kynureninase was shown to be differentially expressed in a microarray analysis of carotid atherosclerotic plaque[31]. Kynureninase protein expression was increased in aneurysmal walls, as revealed by western blotting[28]. The distribution of kynureninase and its expression in IFN-γ-stimulated macrophages suggest that IFN-γ is a stimulant of kynureninase expression in macrophages in the advanced atherosclerotic aorta.

The distribution of KMO in early, non-aneurysmal advanced, and aneurysmal lesions suggests its constitutive expression in aortic macrophages, as is also found in murine macrophages[32]. Interestingly, we identified KMO expression in adventitial peripheral nerve bundles with lymphoid aggregates and capillary vessels. The neuronal KMO may play a role in inflammation. The activation of neuronal KMO and IL-1 signaling were required in the development of depression-like behavior in the spared nerve injury model[33]. Immunological and neuroscientific research has identified close interaction and communication between the immune system and the nervous system[34]. Postganglionic fibers innervate the blood vessels and other organs and release predominantly norepinephrine, which can regulate the adaptive immune response. Locally released norepinephrine affects lymphocyte traffic, circulation, and proliferation and modulates cytokine production and the functional activity of different lymphoid cells via their adrenergic receptors[35]. These lines of evidence and our data suggest that an enhanced kynurenine pathway modulates plaque, adventitial inflammation, and immune response via macrophages and peripheral nerves.

The inhibition of kynureninase in the cultured macrophages enhanced the expression of IDO1 and IL-6. Kynureninase metabolizes kynurenine to anthranilic acid and 3-hydroxykynurenine to 3-hydroxyanthranilic acid **(Fig. 1)**. Kynureninase inhibition with the transfection of kynureninase siRNA reduced angiotensin II-induced aneurysmal formation, MMP2 activity, and the concentration of a catabolite of kynureninase, 3-hydroxyanthranilic acid, in $Apoe^{-/-}$ mouse aorta. Additionally, exogenous 3-hydroxyanthranilic acid promoted MMP2 expression in human aortic SMCs and mouse aorta[28]. By contrast, 3-hydroxyanthranilic acid inhibited inflammasome activation and IL-1β secretion by mouse bone marrow-derived macrophages and reduced the atherosclerotic lesion burden in $Ldlr^{-/-}$ mice[36]. Our results appear to be comparable with these latter findings, and kynureninase in macrophages may play protective roles in atherogenesis and aneurysmal formation by inhibiting the kynurenine pathway itself and IL-6 and be not directly involved in

matrix remodeling. Kynureninase is expressed in various organs, such as the liver, lung, brain, kidney, skeletal muscle, heart, and pancreas[37]. Therefore, the differences in anti-inflammatory action from the study by Wang *et al.*[28] may be due to systemic or cellular inhibition of kynureninase and differences in cell types.

Although significant differences were observed in inflammatory cell contents between early and aneurysmal atherosclerotic aorta, the metabolomic analysis showed no significant difference in the level of lactic acid, a marker of glycolysis, in atherosclerotic lesions. The results are not comparable with our previous study that reported increased levels of metabolites of the glycolysis pathway in a rabbit model of atherosclerotic arteries, and their association with vascular and macrophage hypoxia[8]. The metabolites of glycolysis were also reported to decrease in alloxan-induced diabetic atherosclerosis in rabbit[38]; however, the transcriptomic analysis did not show any significant difference in diabetic status. This could have been influenced by the heterogeneity of patients, medications, and insulin dependence.

The present study had several limitations. First, microarray and metabolomic analyses were conducted only one time point in each patient; therefore, it is possible that the data do not always reflect the chronic, dynamic process of atherogenesis. Second, we did not conduct microarray and metabolomic analyses in the non-aneurysmal advanced atherosclerotic lesion. Instead, we examined mRNA and protein expression of the kynurenine enzymes and found that most of the kynurenine enzymes upregulate before aneurysmal formation. Third, we also did not conduct *in vivo* experiments on the contributions of kynureninase and KMO to atherogenesis and aneurysmal formation.

Conclusions

In conclusion, our study suggests that the kynurenine pathway is progressively upregulated in macrophages in aortic advanced atherosclerosis and aneurysm. Kynureninase may downregulate inflammation via the kynurenine pathway itself and IL-6 in macrophages.

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Disclosures

Junichi Okutsu, Aiko Fukahori, and Tsuyoshi Hirata are employees of Daiichi Sankyo RD Novare Co., Ltd. Tomohiro Nishizawa is an employee of Daiichi Sankyo Co., Ltd. The other authors declare no conflicts of interest.

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Supplementary Table 1. Sequences of primer used for qPCR analysis

Gene		
Homo sapiens, tryptophan 2, 3-dioxygenase 2	forward	5'-CAGTGATCCTGAAACTGCTGGTG-3'
	reverse	5'-TGCAAACTCTGGAAGCCTGATG-3'
Homo sapiens, indoleamine 2, 3-dioxygenase 1	forward	5'-TCCTGATTCCTGCAAGCCAG-3'
	reverse	5'-AGTGCCTCCAGTTCCTTTGG-3'
Homo sapiens, kynureninase	forward	5'-CAGCACTTTAATATTCCTGCCATCA-3'
	reverse	5'-GCATGTGCTAGATCAAAGCCAAC-3'
Homo sapiens, kynurenine 3-monooxygenase	forward	5'-AGATGATCACGCGATTTCAGACC-3'
	reverse	5'-CAATGCCAACGCTGCACA-3'
Homo sapiens, 3-hydroxyanthranilate 3,4-dioxygenase	forward	5'-CAGGAGCAGCTCAAAGTCATGTTC-3'
	reverse	5'-ATGACCACATCCCGGTGTTTC-3'
Homo sapiens, interleukin 6	forward	5'-AAGCCAGAGCTGTGCAGATGAGTA-3'
	reverse	5'-TGTCCTGCAGCCACTGGTTC-3'
Homo sapiens, interleukin 10	forward	5'-GAGATGCCTTCAGCAGAGTGAAGA-3'
	reverse	5'-AAGGCTTGGCAACCCAGGTA-3'
Homo sapiens, matrix metalloproteinase-2	forward	5'-CTTCCAAGTCTGGAGCGATGTG-3'
	reverse	5'-ATGAGCCAGGAGTCCGTCCTTA-3'
Homo sapiens, matrix metalloproteinase-9	forward	5'-TGGCACCACCACAACATCAC-3'
	reverse	5'-GCAAAGGCGTCGTCAATCA-3'
Homo sapiens, tissue factor	forward	5'-TGACCTCACCGACGAGATTGTGAA-3'
	reverse	5'-TCTGAATTGTTGGCTGTCCGAGGT-3'
Homo sapiens, human β-actin	forward	5'-TGGCACCCAGCACAATGAA-3'
	reverse	5'-TAAGTCATAGTCCGCCTAGAAGCA-3'

Supplementary Table2. Clinical characteristics of study patients ($n=42$)

	n (%) or mean ± SD
Age	72.9 ± 7.8
Male gender	32 (76%)
Body mass index (Kg/m^2)	23.3 ± 9.8
Systolic blood pressure (mmHg)	131 ± 13
Diastolic blood pressure (mmHg)	76 ± 12
Aneurysm maximum diameter (mm)	56.4 ± 2.3
Hypertension	40 (95)
Diabetes mellitus or impaired glucose tolerance	19 (45)
Dyslipidemia	33 (79)
Chronic obstructive pulmonary disease	4 (10)
Smoking	31 (74)
Coronary artery disease	11 (26)
Cerebrovascular disease	11 (26)
Sample location	
Thoracic aorta	17 (40)
Abdominal aorta	25 (60)
Complete blood count and blood chemistry	
White blood cell (/μL)	6376 ± 1466
Hemoglobin(g/dL)	13.1 ± 1.6
Hematocrit (%)	39.7 ± 4.8
Platelet ($\times 10^4$/μL)	18.1 ± 5.3
Total Cholesterol (mg/dL)	179.1 ± 33.4
High density lipoprotein cholesterol (mg/dL)	44.9 ± 10.2
Low density lipoprotein cholesterol (mg/dL)	111.5 ± 31.4
Triglyceride (mg/dL)	122.3 ± 64.3
Fasting blood glucose (mg/dL)	85.9 ± 0.6
Fasting insulin (μU/mL)	9.0 ± 10.0
HemoglobinA$_{1C}$(%)	5.8 ± 0.6
Homeostasis model assessment insulin resistance	2.07 ± 2.87
Creatinine (mg/dL)	1.03 ± 0.3
C-reactive protein (mg/dL)	0.20 ± 0.24
Medication	
Angiotensin-converting enzyme inhibitors	5 (12)
Angiotensin receptor blockers	23 (55)
β-blocker	12 (29)
Calcium channel blockers	31 (74)
Aspirin	11 (26)
Statins	25 (60)
Anti-diabetics	4 (10)

Supplementary Table 3. Biological processes associated with the upregulated genes

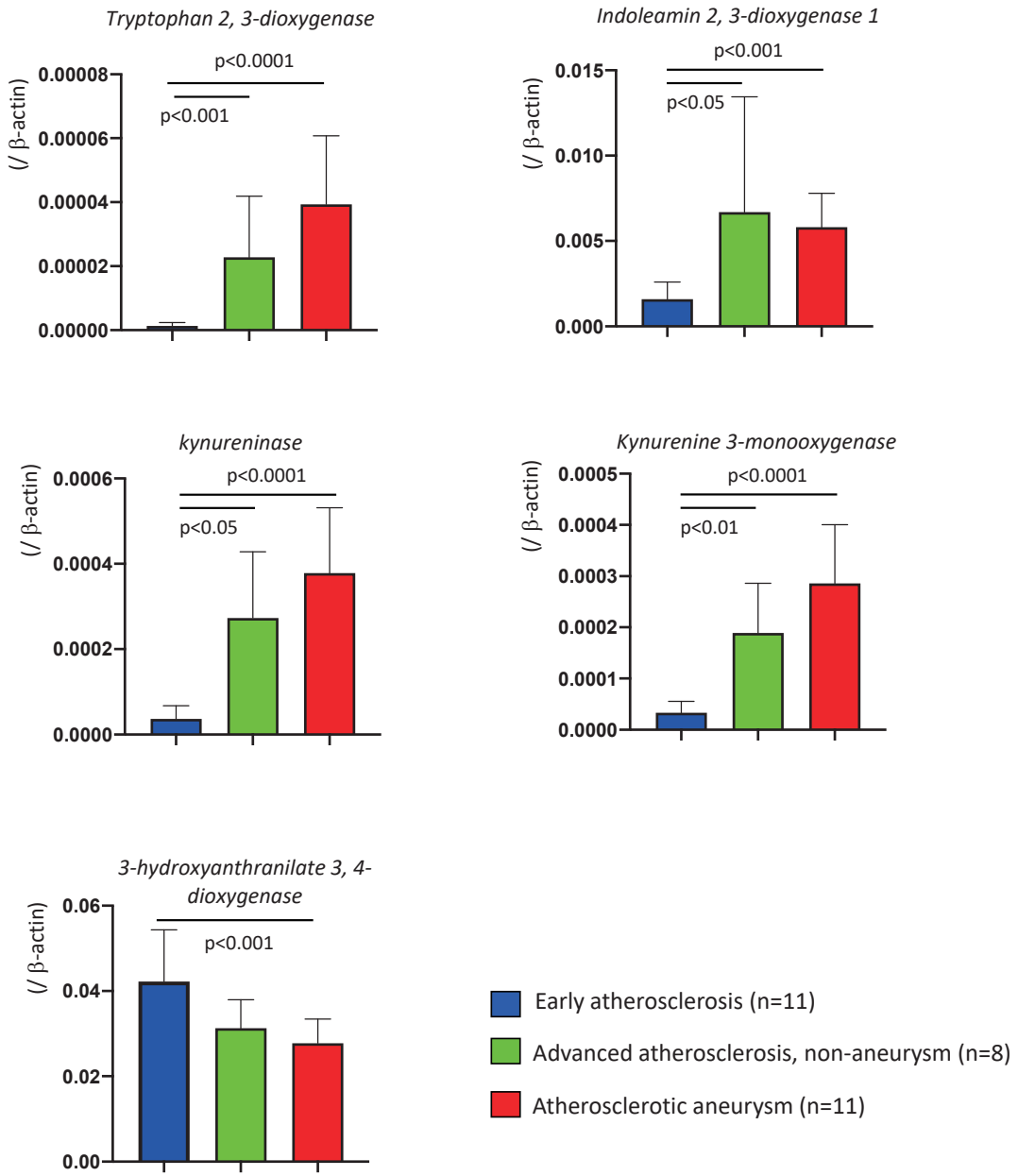
Term	Overlap	*p*-value	Q-value	Combined Score
kynurenine metabolic process (GO: 0070189)	2/8	8.28E-05	4.69E-02	1342.81
regulation of cysteine-type endopeptidase activity involved in apoptotic signaling pathway (GO: 2001267)	2/9	1.06E-04	4.52E-02	1161.83
indolalkylamine catabolic process (GO: 0046218)	2/9	1.06E-04	4.17E-02	1161.83
tryptophan catabolic process (GO: 0006569)	2/10	1.33E-04	4.84E-02	1020.27
tryptophan metabolic process (GO: 0006568)	2/10	1.33E-04	4.51E-02	1020.27
intestinal absorption (GO: 0050892)	2/13	2.29E-04	7.31E-02	736.76
extracellular matrix disassembly (GO: 0022617)	5/78	2.35E-07	3.99E-04	559.16
B cell chemotaxis (GO: 0035754)	1/6	1.05E-02	7.02E-01	434.35
intermediate filament bundle assembly (GO: 0045110)	1/6	1.05E-02	6.93E-01	434.35
sterol import (GO: 0035376)	1/6	1.05E-02	6.84E-01	434.35
positive regulation of receptor binding (GO: 1900122)	1/6	1.05E-02	6.75E-01	434.35
cellular response to bacterial lipopeptide (GO: 0071221)	1/6	1.05E-02	6.67E-01	434.35
cholesterol import (GO: 0070508)	1/6	1.05E-02	6.59E-01	434.35
positive regulation of keratinocyte migration (GO: 0051549)	1/6	1.05E-02	6.51E-01	434.35
aromatic amino acid family catabolic process (GO: 0009074)	2/20	5.54E-04	1.35E-01	428.46

Enrichment analysis using upregulated genes (Log2 ratio > 3, Q < 0.05)

Supplementary Table 4. Biological processes associated with downregulated genes

Term	Overlap	*p*-value	Q-value	Combined Score
negative regulation of bone resorption (GO: 0045779)	1/6	6.28E-03	1	804.72
positive regulation of blood pressure (GO: 0045777)	1/6	6.28E-03	1	804.72
positive regulation of catecholamine secretion (GO: 0033605)	1/6	6.28E-03	1	804.72
positive regulation of cardiac muscle tissue development (GO: 0055025)	1/6	6.28E-03	1	804.72
indolalkylamine metabolic process (GO: 0006586)	1/6	6.28E-03	1	804.72
synaptic transmission, glutamatergic (GO: 0035249)	2/21	2.18E-04	1	764.77
regulation of transcription from RNA polymerase II promoter involved in myocardial precursor cell differentiation (GO: 0003256)	1/7	7.33E-03	1	668.85
regulation of smooth muscle cell differentiation (GO: 0051150)	1/7	7.33E-03	1	668.85
positive regulation of neurological system process (GO: 0031646)	1/7	7.33E-03	1	668.85
negative regulation of bone remodeling (GO: 0046851)	1/8	8.37E-03	1	569.41
regulation of positive chemotaxis (GO: 0050926)	1/8	8.37E-03	1	569.41
negative regulation of appetite (GO: 0032099)	1/8	8.37E-03	1	569.41
regulation of cell communication (GO: 0010646)	2/71	2.50E-03	1	160.76
modulation of chemical synaptic transmission (GO: 0050804)	2/82	3.32E-03	1	132.62
negative regulation of multicellular organismal process (GO: 0051241)	2/125	7.53E-03	1	74.50

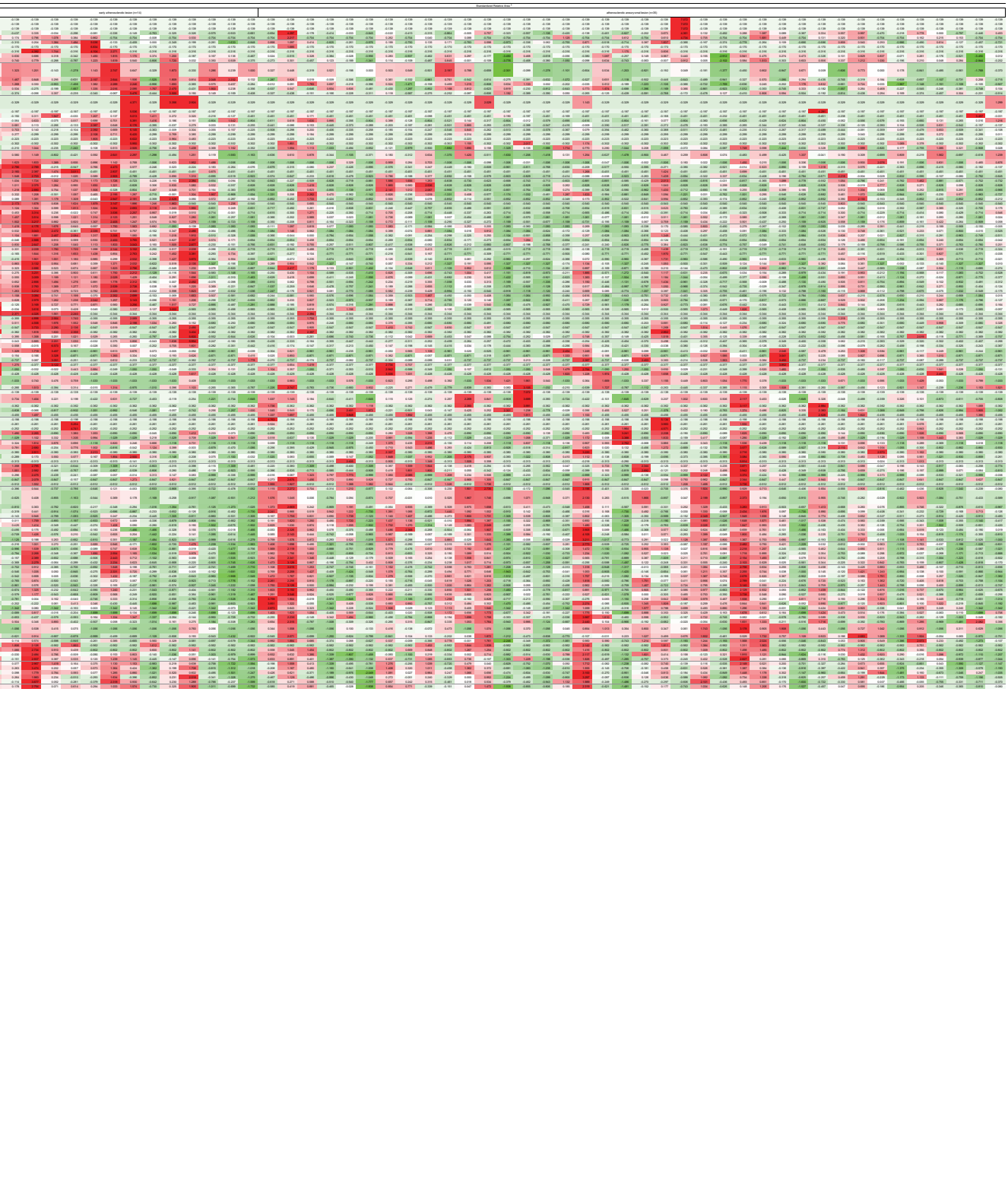
Enrichment analysis using downregulated genes (Log2 ratio > -3, Q < 0.05)



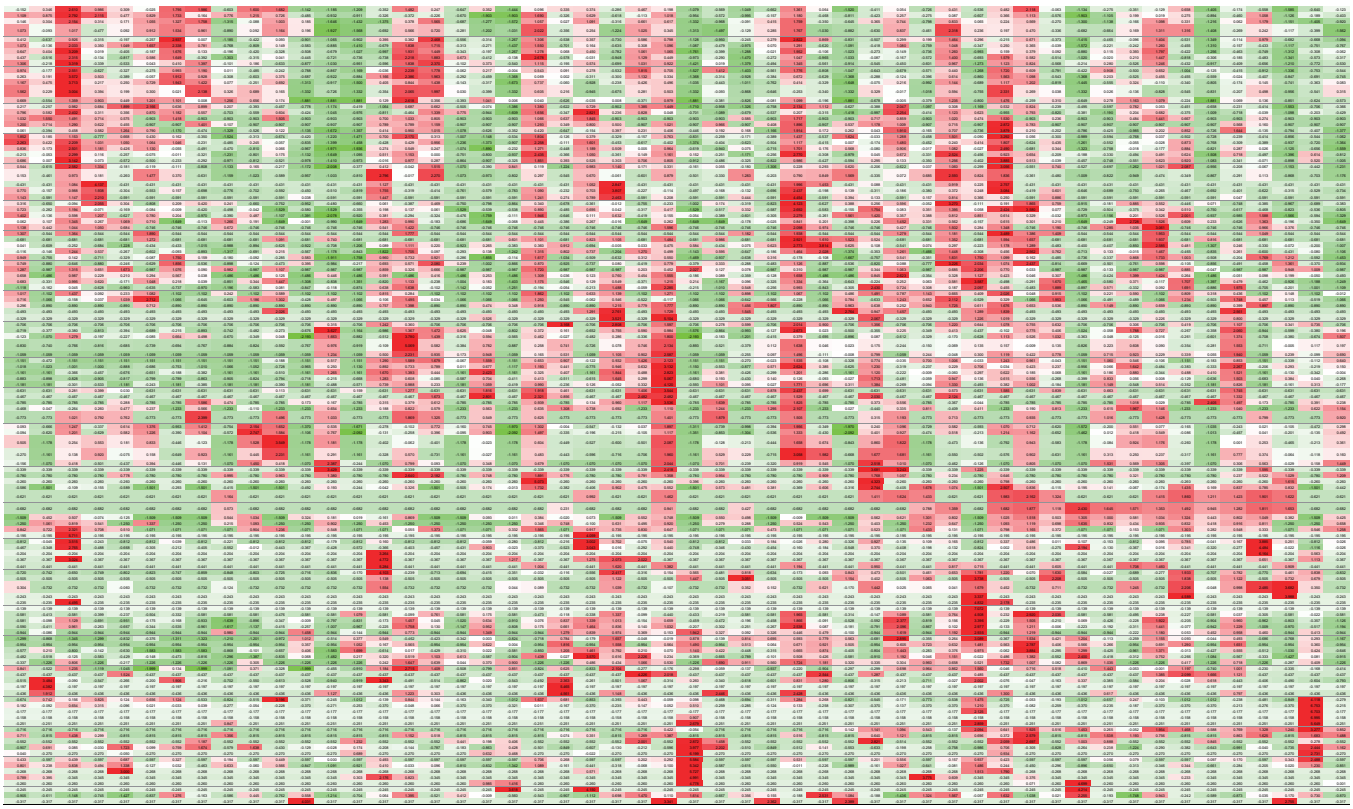
Supplementary Fig. 1.

The mRNA expression of kynurenine pathway enzymes in early atherosclerosis, advanced atherosclerosis non-aneurysm, and atherosclerotic aneurysm. Kruskal-Wallis test with Dunn's multiple comparison tests. Data are expressed as mean and standard deviation.

Supplementary Table 5. Hierarchical clustering analysis of early atherosclerotic lesion and atherosclerotic aneurysmal lesion



(Cont. Supplementary Table 5)



Supplementary Table 6. Altered metabolites in atherosclerotic aneurysmal lesions compared with early atherosclerotic lesions

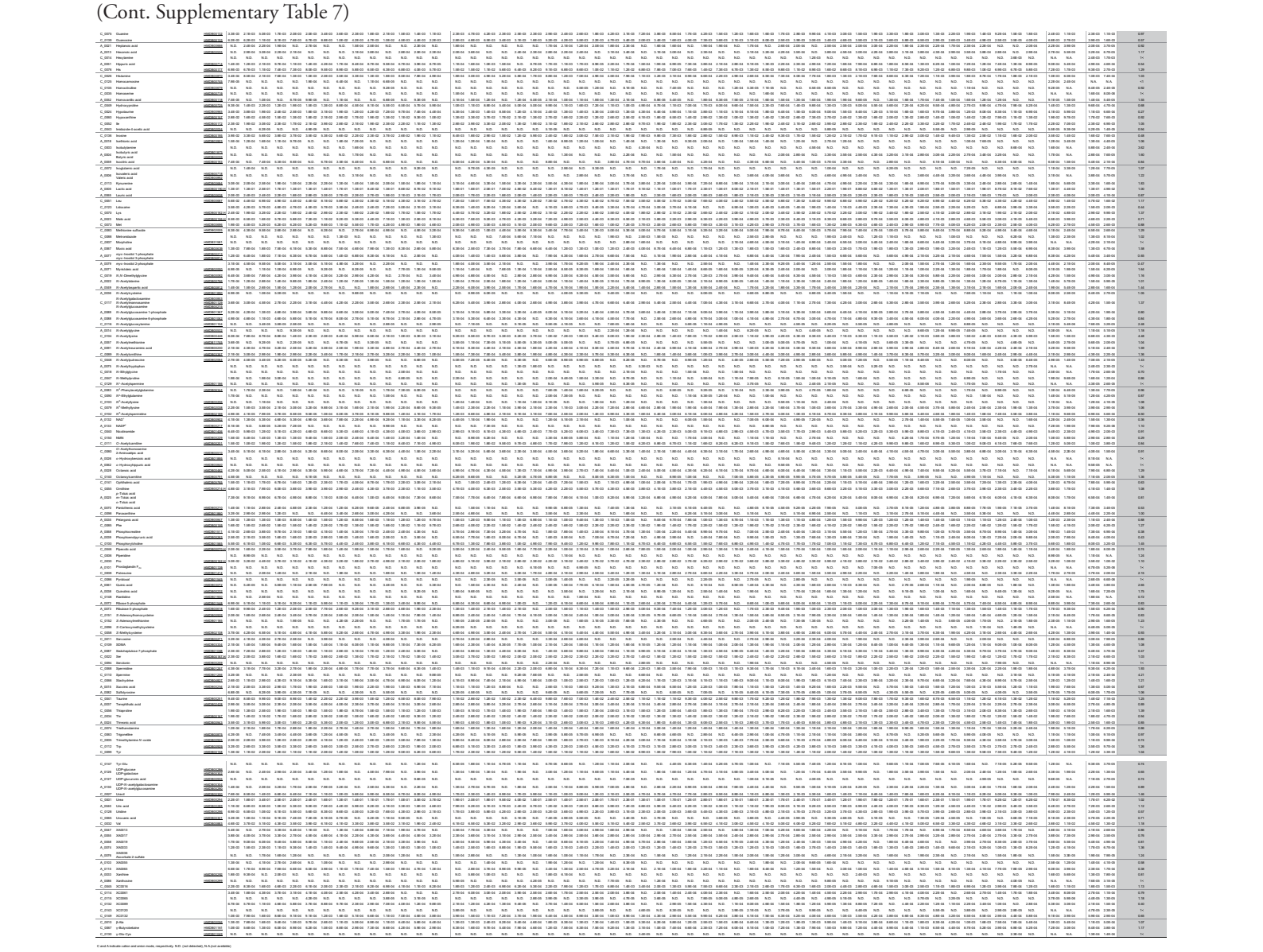
Metabolite	KEGG ID	ratio	pathway
aneurysm/early atherosclerotic lesion >1.5 (relative mean value)			
Cystine	C00491, C01420	6.61	cysteine and methionine metabolism
N^8-Acetylspermidine	C01029	6.50	
N-Acetyllysine	C12989	4.48	
Daminozide	C10996	2.85	
Ala-Ala	C00993		
Glycerophosphocholine	C00670	2.83	glycerophospholipid metabolism
Quinic acid	C00296	2.65	phenylalanine, tyrosine and tryptophan biosynthesis
Putrescine	C00134	2.15	arginine and proline metabolism
4-Methyl-2-oxovaleric acid	C00233	2.01	
3-Methyl-2-oxovaleric acid	C00671, C03465		
Spermidine	C00315	1.93	polyamine metabolism
Glu-Glu	C01425	1.87	
Kynurenine	C00328, C01718	1.83	kynurenine metabolism
Dihydroxyacetone phosphate	C00111	1.80	glycolysis/gluconeogenesis
Fructose 6-phosphate	C05345, C00085	1.76	glycolysis/gluconeogenesis
Quinolinic acid	C03722	1.75	kynurenine metabolism
Asymmetric dimethylarginine	C03626	1.71	
Myristoleic acid	C08322	1.64	
Gly-Gly	C02037	1.60	
N-Acetylneuraminic acid	C00270	1.59	amino sugar and nucleotide sugar metabolism
Mucic acid	C00879, C01807	1.58	ascorbate and aldarate metabolism
aneurysm/early athrosclerotic lesion <0.5 (relative mean value)			
Inosine	C00294	0.48	purine metabolism
myo-Inositol 2-phosphate	NoID	0.47	
Sedoheptulose 7-phosphate	C05382	0.47	pentose phosphate pathway
3-Phosphoglyceric acid	C00197	0.45	glycolysis/gluconeogenesis
Lidocaine	C07073	0.44	
Phosphoenolpyruvic acid	C00074	0.43	glycolysis/gluconeogenesis
2, 3-Diphosphoglyceric acid	C01159	0.41	glycolysis/gluconeogenesis
Adenine	C00147	0.40	purine metabolism
Adenosine triphosphate	C00002	0.39	purine metabolism
Adenosine monophosphate	C00020	0.38	purine metabolism
Cytidine	C00475	0.38	pyrimidine metabolism
Phosphocreatine	C02305	0.38	arginine and proline metabolism
Nicotinamide adenine dinucleotide	C00003	0.36	coenzyme
gamma-aminobutyric acid	C00334	0.32	
3-Aminoisobutyric acid	C03284, C05145	0.29	valine, leucine and isoleucine degradation
Nicotinamide mononucleotide	C00455	0.29	nicotinate and nicotinamide metabolism
Hypotaurine	C00519	0.27	taurineandhypotaurinemetabolism
6-Phosphogluconic acid	C00345	0.26	pentose phosphate pathway
Anserine_divalent	C01262	0.24	histidine metabolism
Glutathione (GSSG)_divalent	C00127	0.20	glutathione metabolism
Adenosine diphosphate	C00008	0.18	purine metabolism
Adenosine	C00212	0.14	purine metabolism

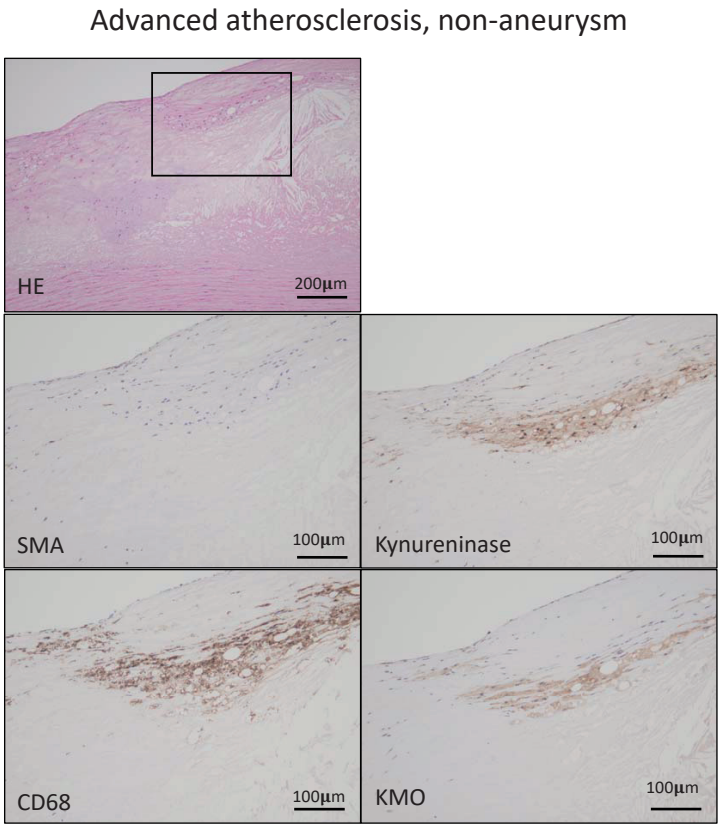
Supplementary Table 7. Relative levels of metabolites in early atherosclerotic lesions and atherosclerotic aneurysm

ID	Compound name	HMDB ID	Relative Area: Early atherosclerotic lesion (n=14) 1	2	3	4	5	6	7	8	9	10	11	12	13	14	Atherosclerotic aneurysmal lesion (n=38) 1	2	3	4	5	6	7	8	9	10	11	12	13
C_0013	1-Methyl-2-pyrrolidone		9.9E-04	9.2E-04	1.7E-03	9.2E-04	8.1E-04	N.D.	N.D.	N.D.	N.D.	1.0E-03	1.2E-03	N.D.	8.3E-04	N.D.	N.D.	5.8E-04	1.3E-03	N.D.	N.D.	7.2E-04	1.4E-03	N.D.	1.0E-03	9.3E-04	9.8E-04	8.8E-04	N.D.
C_0138	1-Methyladenosine	HMDB03331	9.5E-05	6.3E-05	1.4E-04	7.5E-05	7.9E-05	6.9E-05	9.6E-05	9.3E-05	5.6E-05	6.3E-05	8.3E-05	6.3E-05	4.9E-05	2.9E-05	8.6E-05	8.7E-05	7.6E-05	3.3E-05	3.3E-05	3.9E-05	8.1E-05	4.9E-05	9.2E-05	5.0E-05	8.4E-05	6.4E-05	3.5E-05
C_0042	1-Methylhistamine	HMDB00898	6.7E-05	1.7E-04	1.6E-04	1.9E-04	1.1E-04	1.0E-04	1.4E-04	1.1E-04	7.1E-05	1.5E-04	1.5E-04	6.0E-05	1.2E-04	5.4E-05	1.8E-04	2.8E-04	8.5E-05	7.1E-05	1.2E-04	1.1E-04	1.2E-04	1.6E-04	8.2E-05	1.6E-04	2.0E-04	2.5E-04	7.6E-05
C_0087	1-Methylhistidine 3-Methylhistidine	HMDB00001 HMDB00479	1.0E-03	6.7E-04	1.5E-03	1.1E-03	7.9E-04	1.9E-03	1.2E-03	5.6E-04	2.4E-04	3.2E-04	5.9E-04	3.6E-04	3.3E-04	4.5E-04	2.7E-03	9.4E-04	2.3E-03	2.9E-04	3.5E-04	4.6E-04	1.1E-03	6.2E-04	1.4E-03	9.1E-04	5.8E-04	1.5E-03	6.4E-04
C_0061	1-Methylnicotinamide	HMDB00699	7.6E-04	9.6E-04	1.4E-03	9.5E-04	5.7E-04	1.1E-03	1.2E-03	2.0E-03	7.5E-04	1.1E-03	1.6E-03	1.2E-03	1.5E-03	9.0E-04	8.3E-04	1.6E-03	1.5E-03	6.5E-04	8.5E-04	7.9E-04	9.9E-04	6.7E-04	8.5E-04	3.3E-04	1.7E-03	1.1E-03	5.4E-04
C_0028	1-Pyrroline 5-carboxylic acid	HMDB01301	N.D.	N.D.	7.8E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.2E-04	N.D.	N.D.	N.D.	N.D.	3.6E-05	N.D.	N.D.	3.8E-05	5.6E-05	5.5E-05	N.D.	N.D.
C_0120	2'-Deoxycytidine	HMDB00014	4.7E-05	3.7E-05	4.1E-05	2.8E-05	5.0E-05	N.D.	4.7E-05	N.D.	N.D.	N.D.	N.D.	1.9E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
A_0084	2,3-Diphosphoglyceric acid	HMDB01294	8.5E-03	1.1E-03	8.7E-03	1.3E-03	2.1E-03	1.3E-03	3.8E-04	3.5E-03	1.4E-03	7.4E-03	N.D.	N.D.	2.3E-04	N.D.	N.D.	N.D.	2.4E-03	4.8E-04	3.6E-03	N.D.	8.4E-04	5.2E-04	N.D.	1.4E-03	N.D.	N.D.	N.D.
C_0101	2-(Creatinine-3-yl)propionic acid		N.D.	N.D.	4.5E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
C_0039	2-Amino-2-(hydroxymethyl)-1,3-propanediol		3.9E-03	3.2E-03	5.5E-03	3.6E-03	3.1E-03	4.3E-03	3.7E-03	N.D.	2.0E-03	3.0E-03	3.3E-03	2.3E-03	8.1E-04	N.D.	N.D.	N.D.	3.9E-03	N.D.	N.D.	N.D.	4.6E-03	2.4E-03	5.8E-03	N.D.	N.D.	N.D.	N.D.
C_0016	2-Aminoisobutyric acid 2-Aminobutyric acid	HMDB01906 HMDB00452	5.8E-03	3.9E-03	5.7E-03	3.3E-03	4.2E-03	5.5E-03	6.5E-03	5.6E-03	2.7E-03	4.2E-03	5.9E-03	4.4E-03	1.0E-03	1.6E-03	3.0E-03	5.0E-03	5.2E-03	3.6E-03	2.2E-03	2.4E-03	7.3E-03	3.5E-03	4.5E-03	2.1E-03	5.7E-03	4.6E-03	2.0E-03
A_0008	2-Hydroxybutyric acid	HMDB00008	2.5E-03	1.2E-03	1.9E-03	1.2E-03	1.5E-03	1.7E-03	3.1E-03	1.9E-03	1.7E-03	1.7E-03	1.6E-03	2.2E-03	3.3E-04	4.5E-04	2.5E-03	2.4E-03	2.5E-03	1.6E-03	3.7E-04	1.7E-03	3.0E-03	1.0E-03	1.4E-03	1.3E-03	2.0E-03	2.2E-03	6.7E-04
A_0031	2-Hydroxyglutaric acid	HMDB00606,H	2.4E-04	N.D.	2.8E-04	1.9E-04	1.9E-04	2.0E-04	2.3E-04	1.7E-04	N.D.	N.D.	1.8E-04	N.D.	2.2E-04	N.D.	2.4E-04	N.D.	2.2E-04	N.D.	2.0E-04	N.D.	3.2E-04	1.7E-04	1.8E-04	2.5E-04	2.2E-04	3.4E-04	1.5E-04
A_0009	2-Hydroxyisobutyric acid	HMDB00729	N.D.	N.D.	N.D.	N.D.	N.D.	1.9E-03	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	4.9E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
A_0016	2-Hydroxyvaleric acid	HMDB01863	2.1E-04	2.2E-04	3.7E-04	2.4E-04	3.3E-04	2.9E-04	N.D.	1.9E-04	3.6E-04	2.3E-04	N.D.	2.0E-04	8.1E-05	N.D.	3.6E-04	3.3E-04	1.8E-04	1.2E-04	N.D.	1.9E-04	2.6E-04	1.6E-04	2.4E-04	2.0E-04	N.D.	2.4E-04	N.D.
A_0012	2-Oxoisovaleric acid	HMDB00019	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
A_0053	2-Phosphoglyceric acid	HMDB03391	1.5E-03	1.1E-03	1.8E-03	7.8E-04	1.1E-03	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.1E-03	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
A_0094	3'-CMP 2'-CMP	HMDB11692	2.6E-04	2.9E-04	3.8E-04	2.3E-04	2.3E-04	1.5E-04	2.8E-04	1.1E-04	1.5E-04	1.9E-04	1.7E-04	5.2E-05	1.1E-04	N.D.	3.3E-04	3.4E-04	2.3E-04	1.2E-04	1.5E-04	N.D.	2.2E-04	2.2E-04	1.7E-04	2.4E-04	4.8E-04	3.1E-04	9.1E-05
C_0017	3-Aminobutyric acid		9.7E-04	N.D.	2.6E-04	2.6E-04	3.8E-04	4.9E-04	2.0E-04	N.D.	N.D.	2.6E-04	1.8E-04	3.6E-04	3.0E-04	1.6E-04	5.5E-04	3.6E-04	N.D.	1.7E-04	1.8E-04	1.4E-04	1.0E-03	1.2E-04	1.8E-04	N.D.	3.0E-04	1.1E-04	N.D.
C_0020	3-Aminoisobutyric acid	HMDB03911	1.3E-03	1.6E-03	9.4E-04	5.3E-04	6.6E-04	3.6E-03	1.6E-03	8.0E-03	3.6E-04	7.7E-04	1.1E-03	8.4E-04	9.6E-04	5.2E-04	1.9E-04	4.3E-04	1.0E-03	3.1E-04	3.9E-04	4.8E-04	5.6E-04	6.0E-04	4.9E-04	1.4E-04	1.8E-03	4.7E-04	1.8E-04
A_0007	3-Hydroxybutyric acid	HMDB00011,H	7.0E-03	8.2E-04	1.7E-03	8.2E-04	3.2E-03	1.8E-03	1.1E-02	2.1E-03	7.5E-04	3.7E-03	2.4E-03	5.7E-03	4.2E-03	3.3E-03	6.8E-03	6.3E-03	1.2E-02	5.1E-03	3.5E-04	8.7E-04	7.5E-03	6.5E-04	1.4E-03	4.5E-03	2.6E-03	7.1E-03	2.7E-03
A_0069	3-Indoxylsulfuric acid	HMDB00682	1.3E-04	N.D.	2.1E-04	1.3E-04	2.3E-04	2.0E-04	N.D.	N.D.	1.2E-04	N.D.	1.7E-04	N.D.	1.9E-04	N.D.	1.5E-04	3.8E-05	1.1E-04	N.D.	N.D.	8.1E-05	2.0E-04	4.3E-05	2.2E-04	9.0E-05	N.D.	7.7E-05	N.D.
A_0032	3-Phenylpropionic acid	HMDB00764	6.9E-04	9.4E-04	9.4E-04	2.8E-04	2.1E-04	5.0E-04	1.3E-03	4.6E-04	2.8E-04	3.3E-04	2.8E-04	3.7E-04	2.7E-04	3.1E-04	1.8E-04	2.6E-04	3.1E-04	3.8E-04	2.4E-04	1.8E-04	2.0E-04	2.5E-04	3.8E-04	2.3E-04	2.9E-04	2.0E-04	1.2E-04
A_0054	3-Phosphoglyceric acid	HMDB00807	8.0E-03	6.7E-03	1.2E-02	4.4E-03	5.5E-03	6.3E-03	8.5E-03	6.1E-03	4.3E-03	5.0E-03	6.2E-03	N.D.	2.1E-03	N.D.	1.8E-03	2.3E-03	5.4E-03	2.9E-03	2.9E-03	N.D.	4.8E-03	2.6E-03	N.D.	2.8E-03	1.6E-03	1.6E-03	N.D.
A_0020	4-Methyl-2-oxovaleric acid 3-Methyl-2-oxovaleric acid	HMDB00695 HMDB00491	5.5E-04	N.D.	N.D.	N.D.	3.6E-04	N.D.	N.D.	N.D.	3.2E-04	N.D.	N.D.	N.D.	N.D.	N.D.	1.2E-03	N.D.	N.D.	N.D.	N.D.	4.2E-04	4.4E-04	N.D.	N.D.	9.5E-04	N.D.	3.3E-04	N.D.
C_0082	5-Hydroxylysine	HMDB00450	3.0E-05	N.D.	6.9E-05	N.D.	3.4E-05	N.D.	N.D.	N.D.	N.D.	5.2E-05	N.D.	N.D.	N.D.	N.D.	5.0E-05	9.8E-05	6.3E-05	4.3E-05	5.4E-05	7.2E-05	N.D.	N.D.	5.8E-05	5.6E-05	7.8E-05	6.0E-05	N.D.
A_0066	5-Methoxyindoleacetic acid	HMDB04096	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.0E-04	N.D.	N.D.	N.D.	N.D.	N.D.	5.0E-05	N.D.	N.D.	7.1E-05	N.D.	6.3E-05	N.D.
A_0019	5-Oxoproline	HMDB00267	2.1E-03	1.6E-03	4.2E-03	1.6E-03	1.1E-03	1.1E-03	1.4E-03	1.3E-03	7.2E-04	8.6E-04	1.1E-03	7.2E-04	3.5E-04	7.2E-04	1.5E-03	2.4E-03	1.8E-03	8.2E-04	7.8E-04	1.3E-03	3.1E-03	1.9E-03	2.0E-03	1.8E-03	2.1E-03	2.2E-03	7.8E-04
A_0085	6-Phosphogluconic acid	HMDB01316	4.6E-03	1.5E-03	5.3E-03	1.3E-03	9.2E-04	1.9E-03	5.6E-03	3.9E-03	2.2E-03	9.7E-04	1.6E-03	N.D.	9.4E-04	N.D.	1.3E-04	5.5E-04	2.6E-03	6.5E-04	4.8E-04	N.D.	1.9E-03	6.7E-04	N.D.	3.5E-04	5.0E-04	6.0E-04	N.D.
C_0084	7-Methylguanine	HMDB00897	4.0E-05	N.D.	4.1E-05	N.D.	N.D.	5.3E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	5.0E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
C_0059	Adenine	HMDB00034	2.0E-04	9.2E-05	5.4E-04	2.4E-04	1.9E-04	2.4E-04	4.6E-04	6.7E-04	1.8E-04	1.9E-04	4.6E-04	1.9E-04	1.3E-04	N.D.	N.D.	N.D.	1.7E-04	N.D.	N.D.	N.D.	5.8E-05	N.D.	9.0E-05	N.D.	7.3E-05	5.8E-05	N.D.
C_0135	Adenosine	HMDB00050	4.5E-03	3.7E-03	9.3E-03	1.8E-02	1.0E-02	3.2E-03	5.9E-03	9.1E-03	3.1E-03	1.4E-02	4.3E-02	1.8E-03	2.2E-03	7.7E-04	5.0E-04	1.2E-03	2.3E-03	1.0E-03	4.2E-04	4.5E-04	1.6E-03	1.3E-03	1.7E-03	3.4E-04	1.6E-03	1.7E-03	2.6E-04
A_0030	Adipic acid	HMDB00448	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
C_0108	ADMA	HMDB01539	1.1E-04	9.8E-05	1.8E-04	1.0E-04	9.7E-05	1.4E-04	1.3E-04	1.4E-04	8.2E-05	5.8E-05	1.0E-04	7.4E-05	6.4E-05	8.2E-05	2.6E-04	3.5E-04	1.8E-04	9.3E-05	2.5E-04	1.2E-04	1.5E-04	2.3E-04	1.3E-04	2.0E-04	2.3E-04	2.8E-04	1.1E-04
A_0113	ADP	HMDB01341	1.9E-03	1.9E-03	2.0E-03	1.4E-03	2.5E-03	2.2E-03	3.7E-03	3.4E-03	2.8E-03	2.0E-03	2.7E-03	N.D.	1.0E-03	N.D.	1.1E-04	3.2E-04	1.0E-03	7.8E-04	2.1E-04	N.D.	5.6E-04	3.7E-04	7.1E-05	1.7E-04	3.3E-04	2.3E-04	N.D.
A_0128	ADP-glucose GDP-fucose	HMDB06557 HMDB01095	N.D.	N.D.	6.2E-05	5.4E-05	5.3E-05	N.D.	N.D.	1.1E-04	N.D.	N.D.	7.8E-05	N.D.	6.2E-05	N.D.	N.D.	9.1E-05	7.2E-05	N.D.	4.5E-05	N.D.	4.8E-05	N.D.	N.D.	N.D.	N.D.	7.5E-05	N.D.
A_0125	ADP-ribose	HMDB01178	1.5E-04	N.D.	2.5E-04	9.4E-05	1.0E-04	2.4E-04	1.0E-04	2.0E-04	N.D.	4.2E-04	2.5E-04	N.D.	5.7E-04	1.4E-04	N.D.	3.1E-04	1.9E-04	N.D.	2.4E-04	N.D.	2.9E-04	N.D.	N.D.	N.D.	N.D.	5.2E-04	N.D.
C_0009	Ala	HMDB00161,H	5.6E-02	4.9E-02	8.7E-02	6.7E-02	5.0E-02	5.3E-02	6.3E-02	6.3E-02	3.5E-02	5.1E-02	5.6E-02	3.8E-02	2.5E-02	2.7E-02	4.5E-02	6.9E-02	5.3E-02	3.0E-02	4.1E-02	3.8E-02	6.5E-02	4.5E-02	6.2E-02	3.8E-02	5.4E-02	4.8E-02	3.7E-02
A_0099	AMP	HMDB00045	3.0E-04	2.7E-04	5.2E-04	2.6E-04	4.8E-04	5.4E-04	5.4E-04	2.2E-03	8.0E-04	3.6E-04	3.4E-04	N.D.	8.5E-04	N.D.	7.0E-05	5.1E-04	8.8E-04	6.1E-04	1.6E-04	N.D.	4.3E-04	2.4E-04	N.D.	1.0E-04	3.4E-04	1.8E-04	N.D.
C_0037	Anserine_divalent		1.1E-04	8.1E-05	N.D.	N.D.	N.D.	3.4E-04	1.4E-04	9.4E-04	N.D.	1.5E-04	9.6E-05	N.D.	N.D.	N.D.	N.D.	N.D.	6.8E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	8.6E-05	N.D.	N.D.
C_0091	Arg	HMDB00517,H	1.9E-02	1.7E-02	3.0E-02	2.0E-02	1.8E-02	1.8E-02	3.0E-02	2.0E-02	1.5E-02	1.6E-02	1.9E-02	1.8E-02	1.4E-02	1.3E-02	3.2E-02	4.2E-02	2.8E-02	1.9E-02	1.9E-02	1.9E-02	2.3E-02	2.0E-02	2.0E-02	2.7E-02	3.1E-02	3.0E-02	1.2E-02
C_0144	Arg-Glu		8.4E-05	4.6E-05	1.4E-04	8.1E-05	8.0E-05	8.6E-05	1.2E-04	1.1E-04	6.2E-05	7.5E-05	9.1E-05	N.D.	1.6E-04	5.4E-05	6.9E-05	2.4E-04	1.5E-04	7.7E-05	1.1E-04	6.6E-05	1.1E-04	8.8E-05	7.0E-05	1.0E-04	7.6E-05	1.6E-04	N.D.
C_0142	Argininosuccinic acid	HMDB00052	1.3E-04	1.3E-04	4.0E-04	1.5E-04	7.9E-05	1.1E-04	2.7E-04	1.9E-04	8.6E-05	6.2E-05	1.4E-04	5.5E-05	3.6E-05	3.9E-05	2.1E-04	3.0E-04	2.7E-04	8.7E-05	7.4E-05	N.D.	1.2E-04	1.1E-04	8.6E-05	8.6E-05	2.0E-04	1.4E-04	N.D.
A_0050	Ascorbic acid	HMDB00044	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
C_0053	Asn	HMDB00168	6.2E-03	5.2E-03	1.0E-02	5.4E-03	5.9E-03	7.0E-03	8.6E-03	8.4E-03	3.8E-03	5.4E-03	7.1E-03	6.3E-03	4.2E-03	5.7E-03	1.0E-02	1.1E-02	9.1E-03	5.4E-03	5.7E-03	5.3E-03	8.7E-03	6.2E-03	8.4E-03	8.3E-03	9.5E-03	9.9E-03	4.7E-03
C_0057	Asp	HMDB00191,H	1.7E-02	2.1E-02	3.9E-02	2.6E-02	2.1E-02	1.8E-02	3.2E-02	3.4E-02	1.3E-02	3.1E-02	3.1E-02	9.0E-03	8.6E-03	8.4E-03	1.5E-02	3.0E-02	2.0E-02	1.3E-02	2.1E-02	6.6E-03	1.9E-02	2.1E-02	2.1E-02	1.6E-02	2.2E-02	2.0E-02	9.5E-03
A_0123	ATP	HMDB00538	1.4E-03	9.1E-04	4.0E-03	2.0E-03	1.7E-03	9.4E-04	1.2E-03	2.2E-03	1.3E-03	7.2E-04	6.4E-04	N.D.	3.3E-04	N.D.	N.D.	N.D.	1.3E-03	3.3E-04	N.D.	N.D.	8.1E-04	2.2E-04	N.D.	N.D.	N.D.	1.5E-04	N.D.
A_0056	Azelaic acid	HMDB00784	2.0E-04	1.3E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.1E-04	4.5E-04	2.1E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	9.4E-04
A_0017	Benzoic acid	HMDB01870	8.1E-04	4.8E-04	8.2E-04	9.8E-04	6.0E-04	8.6E-04	N.D.	N.D.	3.8E-04	N.D.	6.8E-04	N.D.	N.D.	N.D.	N.D.	1.6E-03	N.D.	N.D.	N.D.	N.D.	5.3E-04	N.D.	4.1E-04	N.D.	4.3E-04	N.D.	N.D.
C_0033	Betaine	HMDB00043	1.0E-02	4.8E-03	9.4E-03	9.7E-03	6.4E-03	5.9E-03	1.0E-02	4.6E-03	3.5E-03	5.9E-03	6.8E-03	5.2E-03	6.6E-03	4.3E-03	1.1E-02	1.4E-02	1.2E-02	7.7E-03	1.1E-02	5.2E-03	1.4E-02	7.7E-03	8.7E-03	1.2E-02	8.6E-03	1.4E-02	3.9E-03
C_0036	Betaine aldehyde_+H_2O		N.D.	N.D.	6.5E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	5.9E-05	7.1E-05	N.D.	N.D.	1.4E-04	N.D.	N.D.	7.8E-05	N.D.	N.D.	N.D.	N.D.	N.D.
C_0077	Betonicine		N.D.	N.D.	1.2E-04	3.5E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.2E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.1E-04	2.5E-04	N.D.	N.D.	N.D.
C_0122	Butyrylcarnitine	HMDB02013	1.1E-04	7.2E-05	1.1E-04	1.1E-04	8.7E-05	N.D.	N.D.	N.D.	8.7E-05	N.D.	N.D.	N.D.	N.D.	N.D.	7.9E-05	1.3E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.4E-04	N.D.
C_0107	Caffeine	HMDB01847	3.2E-04	8.4E-04	9.8E-04	1.6E-03	N.D.	N.D.	5.7E-04	6.0E-04	3.8E-04	3.5E-04	3.2E-04	N.D.	N.D.	3.4E-04	2.4E-04	5.8E-04	N.D.	N.D.	2.6E-04	N.D.	3.3E-04	5.6E-04	7.5E-04	N.D.	3.6E-04	1.6E-04	N.D.
C_0081	Carnitine	HMDB00062	1.7E-02	1.2E-02	1.9E-02	1.7E-02	1.3E-02	1.9E-02	1.5E-02	1.4E-02	8.9E-03	1.5E-02	2.1E-02	6.5E-03	9.0E-03	6.3E-03	9.1E-03	1.9E-02	1.2E-02	8.1E-03	1.4E-02	6.0E-03	2.0E-02	1.1E-02	1.4E-02	1.6E-02	1.1E-02	2.0E-02	1.5E-02
C_0119	Carnosine	HMDB00033	4.7E-04	1.9E-04	4.5E-04	4.1E-04	4.4E-04	1.6E-04	7.7E-04	6.2E-04	1.0E-04	7.1E-04	1.9E-04	2.9E-04	1.1E-04	1.3E-04	N.D.	8.1E-05	5.2E-04	8.7E-05	N.D.	N.D.	2.7E-04	9.9E-05	1.9E-04	N.D.	1.5E-04	6.1E-05	N.D.
A_0090	cCMP 2',3'-cCMP	HMDB11691	7.7E-05	N.D.	6.9E-05	5.9E-05	7.0E-05	N.D.	N.D.	N.D.	N.D.	N.D.	5.6E-05	N.D.	N.D.	N.D.	N.D.	7.9E-05	N.D.	N.D.	6.3E-05	N.D.	7.2E-05	5.2E-05	6.0E-05	5.6E-05	N.D.	1.0E-04	9.6E-05
A_0121	CDP-choline	HMDB01413	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.5E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	8.6E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	5.8E-05	N.D.	N.D.
A_0111	Cholic acid	HMDB00619	N.D.	N.D.	N.D.	N.D.	2.2E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
C_0021	Choline	HMDB00097	5.9E-02	5.8E-02	7.1E-02	5.6E-02	6.1E-02	7.0E-02	5.9E-02	4.8E-02	3.1E-02	2.8E-02	4.7E-02	7.8E-02	2.5E-02	8.3E-02	8.6E-02	9.3E-02	1.1E-01	9.6E-02	5.5E-02	4.5E-02	1.2E-01	7.6E-02	2.0E-01	9.6E-02	1.5E-01	8.3E-02	6.3E-02
C_0131	Cimetidine		N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.0E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
A_0048	*cis*-Aconitic acid	HMDB00072	4.9E-04	N.D.	4.9E-04	N.D.	6.1E-04	N.D.	4.4E-04	N.D.	3.7E-04	N.D.	4.9E-04	N.D.	2.8E-04	N.D.	5.1E-04	N.D.	N.D.	N.D.	N.D.	N.D.	6.1E-04	4.1E-04	N.D.	N.D.	3.8E-04	4.2E-04	2.9E-03
A_0060	Citric acid	HMDB00094	1.7E-02	1.3E-02	1.7E-02	1.5E-02	2.1E-02	1.1E-02	1.2E-02	8.5E-03	1.6E-02	1.1E-02	1.5E-02	5.9E-03	4.3E-03	5.4E-03	1.4E-02	1.1E-02	1.2E-02	6.2E-03	6.0E-03	2.6E-03	1.9E-02	1.1E-02	8.7E-03	9.5E-03	1.1E-02	1.2E-02	4.8E-02
C_0093	Citrulline	HMDB00904	2.2E-03	1.8E-03	2.5E-03	2.7E-03	3.3E-03	2.8E-03	1.0E-03	1.7E-03	8.9E-04	1.6E-03	2.2E-03	1.1E-03	5.7E-04	8.6E-04	2.5E-03	3.0E-03	3.1E-03	1.1E-03	1.5E-03	1.8E-03	2.5E-03	1.5E-03	2.0E-03	2.5E-03	2.3E-03	2.5E-03	1.7E-03
A_0131	CMP-*N*-acetylneuraminate	HMDB01176	9.4E-05	1.0E-04	2.1E-04	1.1E-04	9.5E-05	8.0E-05	1.3E-04	1.6E-04	3.3E-05	8.7E-05	1.6E-04	N.D.	5.5E-05	N.D.	7.7E-05	1.1E-04	8.9E-05	N.D.	5.6E-05	2.8E-05	6.6E-05	7.9E-05	7.3E-05	N.D.	7.2E-05	9.1E-05	N.D.
C_0050	Creatine	HMDB00064	4.1E-02	4.0E-02	8.0E-02	5.1E-02	5.1E-02	6.4E-02	5.1E-02	8.3E-02	2.8E-02	4.9E-02	5.0E-02	2.5E-02	1.7E-02	1.9E-02	1.9E-02	3.4E-02	4.4E-02	2.1E-02	1.3E-02	1.4E-02	3.3E-02	2.4E-02	4.2E-02	1.7E-02	3.2E-02	3.2E-02	1.1E-02
C_0029	Creatinine	HMDB00562	2.1E-02	1.2E-02	2.0E-02	1.5E-02	1.5E-02	1.5E-02	1.5E-02	8.7E-03	7.5E-03	1.1E-02	9.6E-03	5.9E-03	3.2E-03	3.0E-03	7.8E-03	1.2E-02	1.4E-02	4.0E-03	3.4E-03	5.6E-03	2.0E-02	9.3E-03	1.7E-02	7.7E-03	6.9E-03	7.9E-03	3.4E-03
C_0038	Cys	HMDB00574,H	2.1E-04	1.0E-04	3.8E-04	2.6E-04	1.0E-04	1.5E-04	1.3E-04	1.5E-04	3.5E-05	1.2E-04	5.3E-05	N.D.	3.7E-05	4.4E-05	N.D.	1.2E-04	1.6E-04	N.D.	N.D.	1.8E-05	1.4E-04	7.6E-05	N.D.	5.1E-05	1.9E-04	1.6E-04	3.3E-04
C_0118	Cystathionine	HMDB00099	N.D.	N.D.	1.3E-04	5.7E-05	N.D.	1.2E-04	7.1E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
A_0041	Cysteic acid	HMDB02757	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	8.1E-05	N.D.	1.2E-04	6.9E-05	9.4E-05	N.D.	N.D.	9.7E-05	5.9E-05	N.D.	1.7E-04	N.D.	1.5E-04	N.D.	9.4E-05	1.5E-04
C_0153	Cysteine glutathione disulfide	HMDB00656	1.5E-03	1.5E-03	2.8E-03	2.1E-03	1.4E-03	1.1E-03	1.6E-03	8.4E-04	4.6E-04	1.0E-03	1.8E-03	1.1E-03	5.8E-04	8.9E-04	N.D.	3.2E-03	1.7E-03	1.4E-03	1.8E-03	5.3E-04	2.8E-03	2.4E-03	2.3E-03	5.6E-04	2.8E-03	2.1E-03	6.1E-04
A_0034	Cysteinesulfinic acid	HMDB00996	9.8E-05	1.7E-04	7.6E-05	1.4E-04	9.3E-05	N.D.	N.D.	N.D.	7.0E-05	1.4E-04	9.0E-05	1.9E-04	N.D.	3.2E-04	9.4E-05	1.6E-04	1.1E-04	1.2E-04	1.6E-04	9.8E-05	7.1E-05	2.7E-04	3.0E-04	2.3E-04	1.7E-04	1.6E-04	2.7E-04
C_0124	Cystine	HMDB00192	5.7E-04	5.1E-04	7.9E-04	4.8E-04	3.2E-04	2.5E-04	4.9E-04	1.4E-04	1.8E-04	3.2E-04	5.6E-04	5.9E-04	2.1E-04	2.3E-03	1.5E-02	2.1E-03	5.9E-04	6.5E-04	1.5E-03	1.7E-03	2.7E-03	2.4E-03	1.1E-03	1.0E-02	1.8E-03	3.3E-03	4.5E-03
C_0127	Cytidine	HMDB00089	2.5E-03	2.6E-03	3.4E-03	4.1E-03	3.6E-03	2.8E-03	5.0E-03	2.5E-03	1.2E-03	2.5E-03	3.5E-03	8.3E-04	1.0E-03	2.4E-04	9.9E-04	1.3E-03	1.4E-03	5.3E-04	6.8E-04	2.4E-04	9.9E-04	1.3E-03	1.1E-03	1.1E-03	1.0E-03	1.7E-03	1.9E-04
C_0025	Cytosine	HMDB00630	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
C_0078	Daminozide Ala-Ala	HMDB03459	2.3E-04	3.1E-04	2.9E-04	2.5E-04	3.5E-04	3.1E-04	3.4E-04	2.8E-04	1.9E-04	2.4E-04	4.2E-04	2.8E-04	1.7E-03	1.6E-04	8.1E-04	4.9E-03	1.4E-03	5.9E-04	1.5E-03	1.9E-04	1.1E-03	1.5E-03	3.2E-04	9.6E-04	1.5E-03	2.6E-03	1.8E-04
A_0046	Decanoic acid	HMDB00511	N.D.	3.3E-04	1.8E-03	2.4E-04	N.D.	N.D.	3.6E-04	N.D.	2.5E-04	N.D.	N.D.	N.D.	2.7E-04	N.D.	4.1E-04	N.D.	N.D.	N.D.	3.2E-04	2.3E-04	N.D.	2.5E-04	1.6E-03	6.4E-04	3.1E-04	5.7E-04	N.D.
C_0023	Diethanolamine	HMDB04437	4.3E-04	3.8E-04	5.1E-04	6.7E-04	4.3E-04	3.9E-04	5.0E-04	2.7E-03	3.5E-04	1.9E-03	6.0E-04	5.9E-04	4.6E-04	3.6E-04	4.0E-04	3.5E-04	4.7E-04	4.6E-04	4.4E-04	4.0E-04	5.8E-04	2.9E-04	4.2E-04	4.3E-04	4.9E-04	2.9E-04	1.0E-03
A_0042	Dihydroxyacetone phosphate	HMDB01473	5.6E-04	8.0E-04	1.2E-03	4.6E-04	3.7E-04	6.9E-04	5.9E-04	4.8E-04	8.5E-04	6.1E-04	9.1E-04	4.6E-04	7.2E-04	N.D.	2.3E-03	1.5E-03	1.3E-03	1.5E-03	1.7E-03	N.D.	1.8E-03	1.6E-03	4.9E-04	1.8E-03	1.6E-03	1.6E-03	4.7E-04
C_0133	Dyphylline		1.3E-03	1.5E-03	1.1E-03	1.4E-03	8.7E-04	1.3E-03	2.9E-03	1.8E-03	1.5E-03	2.2E-03	1.4E-03	2.3E-03	1.0E-03	1.4E-03	2.0E-03	2.9E-03	1.5E-03	2.2E-03	1.9E-03	1.5E-03	1.5E-03	1.3E-03	1.1E-03	2.4E-03	3.0E-03	2.0E-03	3.1E-03
C_0121	Ergothioneine	HMDB03045	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.4E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.3E-03	N.D.	1.0E-03	N.D.	1.4E-03	N.D.	1.5E-03	5.6E-04	N.D.
C_0002	Ethanolamine	HMDB00149	3.4E-03	3.7E-03	7.6E-03	4.0E-03	3.3E-03	3.6E-03	3.8E-03	3.2E-03	1.7E-03	2.4E-03	4.1E-03	1.9E-03	1.0E-03	2.0E-03	4.6E-03	5.7E-03	4.0E-03	2.5E-03	2.4E-03	2.2E-03	7.8E-03	3.2E-03	5.5E-03	3.2E-03	4.0E-03	5.7E-03	4.0E-03
A_0027	Ethanolamine phosphate	HMDB00224	1.8E-02	1.3E-02	2.7E-02	1.6E-02	1.6E-02	1.6E-02	2.1E-02	1.8E-02	7.2E-03	1.1E-02	1.9E-02	3.7E-03	7.7E-03	1.1E-03	5.9E-03	1.2E-02	1.4E-02	7.2E-03	8.1E-03	5.7E-04	1.3E-02	9.0E-03	7.0E-03	5.6E-03	1.1E-02	1.2E-02	1.3E-03
A_0106	FAD_divalent		N.D.	5.1E-05	5.7E-05	N.D.	5.2E-05	N.D.	N.D.	N.D.	4.6E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	4.3E-05	N.D.	4.7E-05	5.3E-05	N.D.
A_0097	Fructose 1,6-diphosphate	HMDB01058	5.1E-04	6.3E-04	1.7E-03	2.7E-04	3.7E-04	4.4E-04	5.1E-04	4.8E-04	6.3E-04	5.5E-04	5.2E-04	N.D.	3.2E-04	N.D.	4.7E-04	5.2E-04	1.1E-03	8.7E-04	1.3E-03	N.D.	9.9E-04	1.5E-03	N.D.	1.1E-03	6.0E-04	7.3E-04	N.D.
A_0081	Fructose 6-phosphate	HMDB00124	3.3E-04	3.5E-04	6.8E-04	1.6E-04	6.4E-05	1.1E-04	4.7E-04	3.2E-04	2.6E-04	N.D.	1.5E-04	1.3E-04	2.3E-04	N.D.	4.4E-04	9.2E-04	6.4E-04	6.0E-04	9.2E-04	5.5E-05	7.4E-04	8.2E-04	2.5E-04	4.7E-04	9.3E-04	9.1E-04	N.D.
A_0011	Fumaric acid	HMDB00134	6.2E-04	5.2E-04	9.8E-04	4.8E-04	3.5E-04	N.D.	N.D.	N.D.	3.9E-04	N.D.	5.4E-04	N.D.	N.D.	N.D.	3.8E-04	5.1E-04	6.3E-04	N.D.	N.D.	N.D.	N.D.	4.1E-04	4.6E-04	N.D.	5.8E-04	5.7E-04	N.D.
C_0015	GABA	HMDB00112	1.2E-03	1.6E-03	9.4E-04	5.0E-04	6.6E-04	3.6E-03	1.6E-03	8.0E-03	3.6E-04	7.7E-04	1.1E-03	8.5E-04	9.6E-04	5.2E-04	3.0E-04	7.1E-04	9.9E-04	3.0E-04	3.8E-04	4.8E-04	5.8E-04	6.0E-04	4.9E-04	2.3E-04	1.8E-03	4.7E-04	1.5E-04
A_0114	GDP	HMDB01201	1.4E-04	1.5E-04	1.2E-04	7.6E-05	1.2E-04	1.4E-04	2.0E-04	1.3E-04	9.8E-05	1.3E-04	N.D.	N.D.	9.8E-05	N.D.	N.D.	N.D.	6.6E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
A_0129	GDP-glucose GDP-mannose GDP-galactose	HMDB03351 HMDB01163	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.1E-04	N.D.	8.5E-05	7.5E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	5.4E-05	N.D.
C_0069	Gln	HMDB00641,H	4.7E-02	5.0E-02	8.2E-02	5.0E-02	4.8E-02	6.3E-02	6.8E-02	7.5E-02	2.2E-02	4.3E-02	6.2E-02	4.8E-02	1.6E-02	2.0E-02	2.1E-02	5.1E-02	7.2E-02	3.3E-02	2.3E-02	1.7E-02	6.3E-02	4.5E-02	6.4E-02	3.9E-02	5.6E-02	5.5E-02	2.2E-02
C_0071	Glu	HMDB00148,H	1.4E-01	1.2E-01	2.6E-01	1.2E-01	1.5E-01	1.3E-01	1.9E-01	2.1E-01	5.1E-02	9.9E-02	1.8E-01	7.7E-02	5.1E-02	5.0E-02	9.3E-02	1.2E-01	1.6E-01	7.7E-02	7.9E-02	3.8E-02	1.4E-01	9.0E-02	1.4E-01	7.0E-02	1.5E-01	1.2E-01	5.0E-02
C_0137	Glu-Glu		N.D.	N.D.	3.7E-05	3.3E-05	2.3E-05	2.4E-05	4.6E-05	3.6E-05	N.D.	N.D.	3.0E-05	N.D.	1.1E-04	N.D.	1.3E-04	1.2E-04	7.5E-05	N.D.	1.7E-04	N.D.	7.0E-05	7.3E-05	N.D.	1.4E-04	7.7E-05	1.9E-04	N.D.
A_0063	Gluconic acid	HMDB00625	1.4E-03	3.5E-03	4.8E-03	1.0E-03	6.6E-04	1.1E-03	4.9E-04	9.4E-04	6.2E-04	1.3E-03	1.0E-03	1.4E-03	4.7E-04	8.8E-04	1.1E-03	9.5E-04	7.3E-04	8.9E-04	1.1E-03	4.9E-04	2.1E-03	3.9E-03	3.8E-03	1.3E-03	1.7E-03	1.2E-03	1.5E-03
C_0095	Gluconolactone	HMDB00150	6.2E-04	1.4E-03	2.2E-03	5.2E-04	3.5E-04	5.3E-04	N.D.	5.3E-04	2.7E-04	5.2E-04	4.8E-04	6.3E-04	N.D.	4.4E-04	5.7E-04	4.9E-04	N.D.	4.3E-04	4.8E-04	N.D.	8.8E-04	1.5E-03	1.7E-03	6.2E-04	9.1E-04	5.4E-04	6.1E-04
C_0097	Glucosamine	HMDB01514	N.D.	N.D.	N.D.	N.D.	6.9E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.8E-05	N.D.	N.D.	N.D.	1.3E-04
A_0080	Glucose 1-phosphate	HMDB01586	2.4E-04	3.1E-04	4.7E-04	1.8E-04	1.7E-04	2.0E-04	4.7E-04	2.6E-04	2.0E-04	1.4E-04	2.7E-04	N.D.	1.3E-04	N.D.	3.7E-04	4.1E-04	3.8E-04	2.6E-04	4.3E-04	N.D.	5.2E-04	4.4E-04	3.0E-05	2.6E-04	4.1E-04	5.7E-04	1.3E-04
A_0078	Glucose 6-phosphate	HMDB01401	1.8E-03	1.6E-03	3.5E-03	5.6E-04	2.2E-04	5.1E-04	2.3E-03	1.6E-03	1.2E-03	5.7E-04	6.5E-04	5.7E-04	8.1E-04	N.D.	1.8E-03	3.4E-03	2.9E-03	2.1E-03	3.5E-03	1.9E-04	3.1E-03	2.8E-03	9.7E-04	1.5E-03	3.8E-03	3.1E-03	2.6E-04
C_0146	Glutathione (GSH)	HMDB00125	1.7E-04	6.5E-05	1.6E-04	5.2E-04	9.0E-05	3.6E-04	1.3E-04	1.2E-03	4.0E-04	1.6E-04	1.7E-04	5.1E-05	7.0E-05	N.D.	N.D.	N.D.	1.3E-04	N.D.	N.D.	N.D.	N.D.	N.D.	7.6E-05	N.D.	N.D.	N.D.	1.4E-04
C_0145	Glutathione (GSSG)_divalent		2.8E-02	2.0E-02	4.2E-02	2.5E-02	2.3E-02	2.3E-02	3.2E-02	4.7E-02	1.2E-02	2.8E-02	4.5E-02	7.3E-03	8.2E-03	7.8E-04	N.D.	7.7E-03	1.9E-02	6.9E-03	3.9E-03	2.9E-04	5.0E-03	3.5E-03	9.4E-03	5.8E-05	7.5E-03	2.1E-03	2.8E-04
C_0004	Gly	HMDB00123	5.7E-02	4.9E-02	1.0E-01	4.8E-02	4.6E-02	5.4E-02	6.7E-02	5.2E-02	2.3E-02	4.8E-02	7.3E-02	3.2E-02	2.1E-02	2.9E-02	4.2E-02	6.4E-02	5.3E-02	2.6E-02	3.5E-02	2.9E-02	6.8E-02	5.0E-02	6.1E-02	3.8E-02	6.7E-02	5.1E-02	3.4E-02
C_0106	Gly-Asp		6.5E-05	3.8E-05	9.5E-05	6.3E-05	5.6E-05	N.D.	N.D.	6.9E-05	N.D.	N.D.	5.5E-05	N.D.	5.9E-05	N.D.	1.2E-04	1.2E-04	1.0E-04	4.9E-05	8.5E-05	4.0E-05	4.9E-05	6.5E-05	4.5E-05	1.1E-04	8.9E-05	1.3E-04	3.8E-05
C_0054	Gly-Gly	HMDB11733	N.D.	1.1E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.5E-04	N.D.	2.6E-04	N.D.	3.8E-04	4.3E-04	4.4E-04	1.7E-04	4.2E-04	N.D.	2.8E-04	3.2E-04	1.6E-04	3.2E-04	4.0E-04	5.1E-04	N.D.
A_0010	Glyceric acid	HMDB00139,H	6.4E-04	6.5E-04	7.7E-04	4.5E-04	3.8E-04	5.5E-04	7.2E-04	4.3E-04	7.7E-04	7.1E-04	4.6E-04	4.3E-04	4.6E-04	2.5E-04	8.9E-04	1.1E-03	8.1E-04	5.1E-04	6.4E-04	4.4E-04	7.6E-04	6.0E-04	5.2E-04	8.3E-04	6.2E-04	1.1E-03	N.D.
C_0012	Glycerol	HMDB00131	1.7E-01	2.4E-01	2.0E-01	2.0E-01	1.3E-01	2.8E-01	3.1E-01	3.2E-01	3.0E-01	3.3E-01	1.9E-01	3.1E-01	2.2E-01	1.9E-01	1.8E-01	2.4E-01	1.8E-01	2.6E-01	2.4E-01	1.9E-01	1.9E-01	1.4E-01	1.7E-01	2.7E-01	2.5E-01	1.8E-01	2.5E-01
A_0045	Glycerol 2-phosphate		N.D.	1.1E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.4E-04	N.D.	N.D.	3.0E-04	N.D.	1.0E-04	N.D.	2.1E-04	N.D.	N.D.	2.9E-04	N.D.	N.D.	2.8E-04	2.2E-04	2.4E-04	1.6E-04	1.4E-04	3.6E-04
A_0044	Glycerol 3-phosphate	HMDB00126	1.5E-02	1.4E-02	2.5E-02	1.2E-02	1.3E-02	1.6E-02	1.8E-02	1.4E-02	6.1E-03	7.0E-03	1.4E-02	7.8E-03	4.0E-03	4.4E-03	5.4E-03	7.6E-03	1.9E-02	1.5E-02	4.6E-03	4.7E-04	2.1E-02	8.3E-03	2.8E-02	1.2E-02	1.6E-02	1.1E-02	6.1E-03
C_0134	Glycerophosphocholine	HMDB00086	4.0E-03	2.2E-02	2.0E-03	4.0E-03	2.3E-02	4.2E-02	3.4E-03	3.0E-03	7.7E-03	8.0E-03	1.8E-02	1.0E-01	3.7E-02	7.4E-02	8.1E-02	3.9E-02	4.0E-02	4.4E-02	5.8E-02	2.4E-02	2.8E-02	9.1E-02	5.1E-02	1.3E-01	5.6E-02	5.7E-02	9.4E-02
A_0116	Glycocholic acid	HMDB00138	N.D.	N.D.	N.D.	N.D.	3.1E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	4.5E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
A_0102	GMP	HMDB01397	N.D.	N.D.	N.D.	N.D.	5.5E-05	N.D.	N.D.	N.D.	6.2E-05	N.D.	6.1E-05	N.D.	7.6E-05	N.D.	N.D.	1.2E-04	1.0E-04	N.D.	5.4E-05	N.D.	9.4E-05	5.5E-05	N.D.	N.D.	5.5E-05	7.5E-05	N.D.
A_0124	GTP	HMDB01273	6.8E-05	N.D.	1.8E-04	9.8E-05	7.1E-05	N.D.	N.D.	8.2E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	6.9E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.
C_0092	Guanidinosuccinic acid	HMDB03157	6.5E-05	4.3E-05	8.8E-05	6.0E-05	5.4E-05	N.D.	5.8E-05	4.1E-05	5.3E-05	N.D.	5.5E-05	N.D.	N.D.	N.D.	3.5E-05	N.D.	N.D.	N.D.	N.D.	2.6E-05	7.4E-05	4.6E-05	1.0E-04	3.0E-05	3.7E-05	4.0E-05	N.D.
C_0031	Guanidoacetic acid	HMDB00128	2.4E-04	3.0E-04	3.0E-04	3.3E-04	3.7E-04	3.6E-04	2.7E-04	2.7E-04	1.2E-04	2.1E-04	2.7E-04	2.1E-04	2.2E-04	2.1E-04	7.8E-04	4.8E-04	3.2E-04	1.6E-04	3.7E-04	2.6E-04	4.0E-04	4.4E-04	2.3E-04	3.6E-04	3.8E-04	3.6E-04	1.5E-04

ID	Atherosclerotic aneurysmal lesion (n=38) 14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	Early Mean	Early S.D.	Aneurysm Mean	Aneurysm S.D.	Comparative Analysis: Aneurysm/ Early Ratio
C_0013	N.D.	N.D.	8.0E-04	N.D.	N.D.	N.D.	8.2E-04	N.D.	1.3E-03	6.2E-04	N.D.	9.6E-04	1.2E-03	7.4E-04	N.D.	N.D.	6.3E-04	N.D.	1.2E-03	7.0E-04	8.3E-04	7.2E-04	N.D.	8.3E-04	1.2E-03	1.1E-03	3.0E-04	9.2E-04	2.4E-04	0.87
C_0138	9.5E-05	3.8E-05	5.3E-05	1.1E-04	5.4E-05	3.6E-05	4.8E-05	7.5E-05	6.0E-05	7.6E-05	5.4E-05	1.3E-04	7.3E-05	6.8E-05	4.5E-05	7.4E-05	4.8E-05	6.9E-05	5.8E-05	5.8E-05	6.1E-05	6.7E-05	4.4E-05	4.5E-05	4.1E-05	7.5E-05	2.6E-05	6.3E-05	2.3E-05	0.84
C_0042	2.0E-04	6.4E-05	9.3E-05	2.9E-04	1.4E-04	7.4E-05	1.2E-04	9.2E-05	1.3E-04	1.3E-04	9.5E-05	2.5E-04	2.2E-04	1.2E-04	1.0E-04	2.4E-04	9.1E-05	1.1E-04	8.6E-05	8.0E-05	2.2E-04	7.4E-05	4.1E-05	2.7E-04	1.6E-04	1.2E-04	4.3E-05	1.4E-04	6.9E-05	1.20
C_0087	7.5E-04	1.7E-03	8.2E-04	1.4E-03	1.5E-03	1.9E-03	7.4E-04	9.9E-04	1.4E-03	2.5E-03	1.5E-03	2.1E-03	7.3E-04	6.6E-04	3.3E-04	4.4E-04	3.7E-04	6.6E-04	7.4E-04	4.2E-04	7.7E-04	1.0E-03	4.2E-04	5.2E-04	2.3E-04	7.9E-04	4.9E-04	1.0E-03	6.5E-04	1.29
C_0061	1.1E-03	4.4E-04	1.7E-03	1.3E-03	1.1E-03	6.5E-04	1.1E-03	6.2E-04	1.0E-03	1.1E-03	8.2E-04	1.7E-03	1.3E-03	5.0E-04	8.0E-04	1.2E-03	6.7E-04	1.6E-03	7.3E-04	1.1E-03	7.5E-04	1.4E-03	1.2E-03	7.0E-04	1.0E-03	1.1E-03	3.7E-04	1.0E-03	3.7E-04	0.88
C_0028	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.3E-05	N.D.	3.4E-05	N.D.	7.8E-05	N.A.	5.2E-05	3.2E-05	0.66
C_0120	N.D.	N.D.	N.D.	2.5E-05	N.D.	N.D.	N.D.	2.7E-05	N.D.	N.D.	N.D.	1.6E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	3.8E-05	1.1E-05	2.3E-05	5.5E-06	0.59
A_0084	N.D.	N.D.	N.D.	N.D.	4.9E-04	N.D.	1.2E-03	1.4E-04	N.D.	3.4E-03	2.4E-04	5.0E-03	N.D.	3.3E-04	N.D.	N.D.	4.5E-04	N.D.	1.3E-03	N.D.	9.9E-05	1.4E-03	8.1E-04	N.D.	1.1E-03	3.3E-03	3.3E-03	1.3E-03	1.3E-03	0.41
C_0101	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	4.8E-05	2.3E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	4.5E-05	N.A.	3.6E-05	1.8E-05	0.79
C_0039	N.D.	N.D.	1.9E-03	N.D.	N.D.	2.3E-04	N.D.	3.2E-03	N.D.	N.D.	1.9E-03	N.D.	N.D.	1.6E-03	N.D.	N.D.	2.3E-03	1.4E-03	2.8E-03	N.D.	1.9E-03	N.D.	1.6E-03	N.D.	N.D.	3.2E-03	1.2E-03	2.5E-03	1.4E-03	0.79
C_0016	1.7E-03	3.9E-03	4.5E-03	6.9E-03	2.4E-03	2.7E-03	3.1E-03	5.4E-03	3.3E-03	7.7E-03	4.4E-03	4.4E-03	4.8E-03	3.5E-03	3.0E-03	3.0E-03	4.3E-03	6.2E-03	6.2E-03	1.8E-03	3.6E-03	4.5E-03	3.9E-03	3.4E-03	1.6E-03	4.3E-03	1.7E-03	4.0E-03	1.6E-03	0.92
A_0008	1.1E-03	1.3E-03	2.0E-03	1.7E-03	1.7E-03	5.2E-04	1.2E-03	2.2E-03	1.5E-03	3.2E-03	3.1E-03	2.6E-03	1.9E-03	1.4E-03	1.2E-03	1.5E-03	2.5E-03	3.1E-03	2.6E-03	1.4E-03	1.6E-03	3.1E-03	2.7E-03	1.3E-03	6.8E-04	1.7E-03	7.3E-04	1.8E-03	7.8E-04	1.11
A_0031	1.8E-04	1.3E-04	1.5E-04	3.6E-04	N.D.	N.D.	2.6E-04	4.7E-04	1.3E-04	2.1E-04	3.0E-04	2.2E-04	1.7E-04	2.0E-04	N.D.	1.2E-04	3.3E-04	3.3E-04	2.0E-04	N.D.	1.6E-04	1.8E-04	1.9E-04	1.6E-04	1.1E-04	2.1E-04	3.4E-05	2.2E-04	8.2E-05	1.04
A_0009	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.9E-03	N.A.	4.9E-04	N.A.	0.26
A_0016	N.D.	1.8E-04	2.0E-04	3.1E-04	2.3E-04	1.5E-04	2.3E-04	3.8E-04	2.4E-04	2.7E-04	1.8E-04	2.8E-04	2.4E-04	2.3E-04	N.D.	1.7E-04	5.4E-04	3.9E-04	2.8E-04	2.3E-04	1.3E-04	3.7E-04	1.8E-04	1.6E-04	N.D.	2.5E-04	8.5E-05	2.5E-04	9.0E-05	0.99
A_0012	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.5E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	3.3E-04	N.D.	N.D.	N.D.	2.9E-04	N.D.	N.D.	N.A.	N.A.	2.9E-04	4.3E-05	1<
A_0053	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.2E-03	3.9E-04	1.1E-03	N.A.	0.90
A_0094	6.6E-04	1.5E-04	1.1E-04	1.5E-04	1.9E-04	N.D.	1.3E-04	2.3E-04	3.3E-04	2.8E-04	2.7E-04	4.6E-04	2.6E-04	2.0E-04	N.D.	2.4E-04	2.0E-04	1.4E-04	1.6E-04	2.4E-04	2.2E-04	9.5E-05	1.3E-04	1.2E-04	1.0E-04	2.0E-04	8.9E-05	2.3E-04	1.2E-04	1.13
C_0017	N.D.	4.1E-04	6.5E-04	9.8E-04	N.D.	5.9E-04	N.D.	4.3E-04	2.8E-04	2.1E-04	1.8E-04	1.7E-04	4.0E-04	6.4E-04	2.1E-04	N.D.	1.3E-04	1.4E-04	3.7E-04	N.D.	1.1E-04	5.3E-04	3.3E-04	N.D.	2.3E-04	3.5E-04	2.3E-04	3.5E-04	2.5E-04	1.00
C_0020	3.6E-04	1.1E-04	3.5E-04	8.7E-04	3.2E-04	2.4E-04	3.6E-04	3.3E-04	1.8E-04	3.5E-04	2.2E-04	5.1E-04	4.2E-04	4.7E-04	4.1E-04	2.1E-04	2.6E-04	6.7E-04	3.6E-04	8.9E-04	2.2E-04	1.5E-03	1.9E-04	3.8E-04	6.5E-04	1.6E-03	2.0E-03	4.7E-04	3.5E-04	0.29
A_0007	5.4E-03	3.1E-03	2.3E-03	8.4E-03	7.9E-04	2.1E-03	1.0E-03	4.5E-03	1.2E-03	1.6E-02	1.2E-02	8.3E-03	1.3E-02	1.1E-03	1.6E-03	2.3E-03	1.3E-03	6.4E-03	3.8E-03	9.1E-04	2.3E-03	2.4E-03	9.3E-03	2.8E-03	6.1E-04	3.2E-03	3.0E-03	4.5E-03	4.0E-03	1.41
A_0069	2.0E-04	6.4E-05	N.D.	2.2E-04	1.0E-04	4.3E-04	4.2E-05	2.8E-04	8.5E-05	1.4E-04	9.2E-05	2.3E-04	N.D.	8.2E-05	N.D.	5.2E-05	1.5E-04	N.D.	8.2E-05	N.D.	2.1E-04	2.4E-04	1.4E-04	N.D.	N.D.	1.7E-04	4.1E-05	1.4E-04	9.1E-05	0.83
A_0032	1.5E-04	1.6E-04	1.9E-04	2.8E-04	2.0E-04	2.1E-04	2.0E-04	2.7E-04	2.4E-04	2.5E-04	2.2E-04	3.3E-04	4.7E-04	1.3E-04	3.6E-04	6.2E-04	2.6E-04	4.6E-04	2.8E-04	2.5E-04	1.8E-04	2.4E-04	2.0E-04	2.9E-04	3.0E-04	5.1E-04	3.3E-04	2.6E-04	1.0E-04	0.51
A_0054	1.3E-03	N.D.	N.D.	N.D.	2.6E-03	N.D.	2.8E-03	4.2E-03	N.D.	4.3E-03	2.4E-03	4.4E-03	1.8E-03	1.8E-03	N.D.	N.D.	3.7E-03	N.D.	2.8E-03	N.D.	2.3E-03	3.1E-03	2.1E-03	N.D.	2.9E-03	6.3E-03	2.5E-03	2.8E-03	1.1E-03	0.45
A_0020	N.D.	5.8E-04	4.7E-04	N.D.	7.2E-04	3.0E-04	1.2E-03	4.0E-04	4.3E-04	4.1E-04	1.3E-03	6.3E-04	N.D.	7.7E-04	N.D.	N.D.	1.1E-03	N.D.	1.6E-03	3.7E-04	9.2E-04	1.7E-03	2.4E-03	5.8E-04	N.D.	4.1E-04	1.2E-04	8.3E-04	5.3E-04	2.01
C_0082	9.9E-05	7.3E-05	6.1E-05	1.0E-04	8.2E-05	5.3E-05	5.3E-05	5.2E-05	7.4E-05	6.4E-05	5.9E-05	1.0E-04	7.6E-05	4.5E-05	7.1E-05	7.7E-05	N.D.	N.D.	5.6E-05	N.D.	6.6E-05	N.D.	3.2E-05	5.6E-05	N.D.	4.6E-05	1.8E-05	6.6E-05	1.7E-05	1.42
A_0066	N.D.	N.D.	N.D.	5.6E-05	N.D.	N.D.	4.8E-05	N.D.	N.D.	4.3E-05	4.0E-05	N.D.	N.D.	3.6E-05	N.D.	N.D.	7.4E-05	6.5E-05	N.D.	N.D.	N.D.	N.D.	4.6E-05	N.D.	N.D.	N.A.	N.A.	6.6E-05	4.3E-05	1<
A_0019	1.1E-03	1.3E-03	8.5E-04	2.4E-03	1.2E-03	1.0E-03	1.8E-03	1.7E-03	1.3E-03	2.6E-03	1.2E-03	4.8E-03	2.0E-03	1.5E-03	1.2E-03	9.1E-04	1.1E-03	2.1E-03	2.3E-03	6.3E-04	N.D.	1.5E-03	7.9E-04	2.0E-03	7.0E-04	1.3E-03	9.4E-04	1.6E-03	8.1E-04	1.21
A_0085	1.7E-04	2.1E-04	N.D.	6.5E-05	3.0E-04	N.D.	6.1E-04	4.3E-04	N.D.	9.9E-04	2.9E-04	5.2E-04	2.7E-04	5.0E-04	N.D.	N.D.	1.3E-03	N.D.	6.4E-04	N.D.	4.0E-04	1.1E-03	3.8E-04	N.D.	1.7E-03	2.6E-03	1.8E-03	6.8E-04	5.9E-04	0.26
C_0084	N.D.	3.7E-05	N.D.	5.4E-05	N.D.	N.D.	N.D.	3.9E-05	N.D.	3.7E-05	N.D.	8.7E-05	4.2E-05	4.2E-05	N.D.	N.D.	N.D.	5.4E-05	N.D.	N.D.	N.D.	3.4E-05	N.D.	N.D.	N.D.	4.5E-05	6.9E-06	4.8E-05	1.5E-05	1.07
C_0059	2.7E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	8.0E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.2E-04	1.7E-04	N.D.	7.2E-05	N.D.	1.4E-04	N.D.	N.D.	9.3E-05	2.9E-04	1.8E-04	1.2E-04	6.1E-05	0.40
C_0135	7.1E-04	2.7E-04	1.2E-03	1.7E-03	5.7E-04	3.7E-04	9.4E-04	6.7E-03	5.8E-04	1.3E-03	6.9E-04	9.7E-04	8.8E-04	1.1E-03	7.3E-04	1.0E-03	3.8E-03	2.0E-03	7.8E-04	1.2E-03	1.3E-03	1.4E-03	7.2E-04	6.9E-04	1.6E-03	9.2E-03	1.1E-02	1.3E-03	1.1E-03	0.14
A_0030	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.5E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.A.	N.A.	2.5E-04	N.A.	1<
C_0108	1.9E-04	1.2E-04	1.2E-04	2.7E-04	1.7E-04	1.2E-04	2.0E-04	1.4E-04	1.8E-04	2.6E-04	1.3E-04	4.0E-04	2.2E-04	1.2E-04	1.1E-04	2.6E-04	1.1E-04	1.8E-04	1.6E-04	9.1E-05	2.1E-04	1.3E-04	9.3E-05	1.1E-04	8.6E-05	1.0E-04	3.4E-05	1.8E-04	7.5E-05	1.71
A_0113	1.9E-04	1.5E-04	1.2E-04	1.7E-04	2.0E-04	N.D.	3.0E-04	1.4E-03	6.8E-05	3.6E-04	4.2E-04	1.9E-04	2.8E-04	4.1E-04	N.D.	8.0E-05	8.4E-04	1.3E-04	8.1E-04	4.6E-04	2.3E-04	1.2E-03	4.8E-04	1.2E-04	9.6E-04	2.3E-03	7.7E-04	4.0E-04	3.4E-04	0.18
A_0128	9.1E-05	N.D.	N.D.	7.8E-05	N.D.	N.D.	3.7E-05	6.8E-05	N.D.	5.1E-05	N.D.	N.D.	5.9E-05	N.D.	N.D.	6.2E-05	N.D.	7.6E-05	N.D.	N.D.	N.D.	5.4E-05	N.D.	N.D.	5.8E-05	7.0E-05	2.1E-05	6.4E-05	1.6E-05	0.92
A_0125	2.9E-04	1.4E-04	1.2E-04	3.3E-04	2.7E-04	N.D.	6.0E-04	3.5E-04	N.D.	1.0E-04	1.6E-04	N.D.	3.1E-04	N.D.	N.D.	4.3E-04	2.7E-04	3.9E-04	1.1E-04	N.D.	2.3E-04	2.7E-04	1.7E-04	2.0E-04	4.2E-04	2.3E-04	1.5E-04	2.8E-04	1.3E-04	1.22
C_0009	3.8E-02	3.4E-02	4.9E-02	7.1E-02	4.7E-02	3.2E-02	4.7E-02	5.8E-02	4.6E-02	4.9E-02	4.1E-02	7.6E-02	5.5E-02	5.1E-02	3.9E-02	4.4E-02	4.4E-02	5.7E-02	5.7E-02	3.7E-02	3.7E-02	4.9E-02	2.7E-02	3.3E-02	3.3E-02	5.1E-02	1.7E-02	4.7E-02	1.2E-02	0.91
A_0099	2.9E-04	8.1E-05	3.9E-05	6.8E-05	1.6E-04	N.D.	2.3E-04	3.2E-04	N.D.	1.6E-04	5.4E-05	6.5E-05	1.4E-04	6.5E-05	N.D.	1.3E-04	2.7E-04	9.0E-05	8.1E-05	2.2E-04	5.2E-05	3.4E-04	1.9E-04	3.0E-04	7.3E-04	6.2E-04	5.4E-04	2.4E-04	2.0E-04	0.38
C_0037	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	4.8E-05	N.D.	N.D.	8.0E-05	N.D.	N.D.	4.4E-05	2.7E-04	3.1E-04	6.5E-05	1.9E-05	0.24
C_0091	2.1E-02	1.7E-02	1.6E-02	2.7E-02	1.5E-02	1.7E-02	2.7E-02	1.9E-02	2.5E-02	2.8E-02	1.7E-02	3.5E-02	2.8E-02	2.2E-02	1.6E-02	3.1E-02	1.6E-02	2.1E-02	2.7E-02	1.5E-02	2.6E-02	1.7E-02	1.5E-02	1.8E-02	1.7E-02	1.9E-02	5.0E-03	2.3E-02	6.6E-03	1.18
C_0144	8.2E-05	4.0E-05	7.3E-05	1.0E-04	6.1E-05	5.3E-05	9.3E-05	6.5E-05	7.6E-05	8.3E-05	6.4E-05	1.3E-04	1.1E-04	1.3E-04	7.5E-05	8.8E-05	8.4E-05	8.9E-05	7.9E-05	6.5E-05	1.5E-04	6.1E-05	7.4E-05	6.2E-05	1.5E-04	9.1E-05	3.3E-05	9.4E-05	3.9E-05	1.03
C_0142	8.5E-05	3.6E-05	8.0E-05	1.7E-04	6.0E-05	N.D.	8.7E-05	1.0E-04	7.9E-05	1.6E-04	4.1E-05	2.4E-04	2.0E-04	6.7E-05	N.D.	1.3E-04	6.8E-05	1.5E-04	7.4E-05	6.6E-05	1.1E-04	7.8E-05	4.2E-05	5.5E-05	5.0E-05	1.3E-04	9.9E-05	1.1E-04	6.8E-05	0.86
A_0050	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	3.9E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.A.	N.A.	3.9E-04	N.A.	1<
C_0053	5.7E-03	5.2E-03	6.6E-03	1.0E-02	4.9E-03	5.3E-03	9.1E-03	9.4E-03	7.2E-03	8.8E-03	8.3E-03	1.2E-02	9.7E-03	6.7E-03	6.1E-03	8.1E-03	5.9E-03	8.5E-03	8.2E-03	5.0E-03	7.9E-03	6.2E-03	4.7E-03	6.0E-03	5.1E-03	6.4E-03	1.8E-03	7.5E-03	2.0E-03	1.17
C_0057	1.4E-02	9.7E-03	1.3E-02	2.9E-02	1.9E-02	6.0E-03	1.5E-02	1.9E-02	1.2E-02	2.2E-02	1.4E-02	2.2E-02	3.5E-02	1.8E-02	9.0E-03	1.6E-02	1.5E-02	1.7E-02	2.3E-02	6.9E-03	1.7E-02	1.4E-02	5.4E-03	1.3E-02	1.7E-02	2.2E-02	1.0E-02	1.7E-02	6.5E-03	0.75
A_0123	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	3.6E-04	1.2E-03	N.D.	4.3E-04	4.2E-04	3.4E-04	N.D.	2.7E-04	N.D.	N.D.	1.1E-03	N.D.	2.9E-04	N.D.	1.8E-04	1.3E-03	2.8E-04	N.D.	7.7E-04	1.4E-03	9.8E-04	5.7E-04	4.1E-04	0.39
A_0056	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	7.3E-04	1.7E-04	N.D.	N.D.	1.3E-04	N.D.	N.D.	N.D.	N.D.	N.D.	1.0E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.5E-04	4.6E-05	3.9E-04	3.3E-04	2.62
A_0017	N.D.	N.D.	1.0E-03	N.D.	N.D.	7.2E-04	N.D.	7.2E-04	3.0E-03	7.7E-04	N.D.	N.D.	1.3E-03	6.3E-04	N.D.	4.6E-04	5.6E-04	9.0E-04	N.D.	N.D.	4.7E-04	5.0E-04	4.7E-04	N.D.	N.D.	7.0E-04	2.0E-04	8.5E-04	6.4E-04	1.22
C_0033	8.7E-03	6.7E-03	8.6E-03	2.0E-02	5.2E-03	6.0E-03	9.8E-03	8.8E-03	1.3E-02	1.4E-02	1.2E-02	1.4E-02	1.1E-02	1.1E-02	5.2E-03	8.1E-03	5.6E-03	1.3E-02	8.3E-03	4.9E-03	1.3E-02	6.9E-03	5.9E-03	6.9E-03	5.2E-03	6.7E-03	2.3E-03	9.5E-03	3.6E-03	1.43
C_0036	7.8E-05	N.D.	N.D.	5.2E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.0E-04	N.D.	7.5E-05	N.D.	N.D.	N.D.	N.D.	N.D.	6.1E-05	N.D.	6.5E-05	N.A.	8.0E-05	2.8E-05	1.24
C_0077	N.D.	N.D.	N.D.	1.9E-04	1.4E-04	N.D.	4.0E-05	N.D.	N.D.	1.0E-04	5.0E-05	2.4E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.3E-04	1.7E-04	1.4E-04	7.6E-05	0.60
C_0122	N.D.	N.D.	N.D.	1.7E-04	1.1E-04	N.D.	N.D.	7.2E-05	N.D.	1.4E-04	6.3E-05	1.3E-04	N.D.	1.2E-04	N.D.	1.2E-04	1.1E-04	2.3E-04	N.D.	N.D.	N.D.	1.0E-04	6.9E-05	N.D.	N.D.	9.7E-05	1.7E-05	1.2E-04	4.4E-05	1.23
C_0107	N.D.	N.D.	1.2E-03	8.1E-04	3.2E-04	N.D.	3.9E-04	N.D.	2.0E-04	3.4E-04	5.9E-04	2.0E-04	N.D.	1.1E-03	1.4E-04	5.3E-04	4.4E-04	8.3E-04	5.6E-04	N.D.	3.1E-04	4.4E-04	N.D.	2.6E-04	N.D.	6.3E-04	4.2E-04	4.8E-04	2.9E-04	0.76
C_0081	7.3E-03	1.4E-02	1.2E-02	2.0E-02	1.1E-02	1.0E-02	1.2E-02	1.5E-02	1.4E-02	1.6E-02	3.2E-02	1.1E-02	1.2E-02	2.2E-02	5.7E-03	1.0E-02	1.5E-02	1.3E-02	2.2E-02	6.1E-03	1.6E-02	1.1E-02	8.5E-03	7.2E-03	6.8E-03	1.4E-02	4.7E-03	1.3E-02	5.5E-03	0.98
C_0119	N.D.	N.D.	1.0E-04	2.1E-04	N.D.	6.7E-05	4.3E-05	3.7E-04	N.D.	9.3E-05	1.4E-04	N.D.	1.2E-04	N.D.	N.D.	N.D.	2.5E-04	6.0E-04	1.4E-04	1.0E-04	N.D.	9.1E-05	N.D.	N.D.	1.3E-04	3.6E-04	2.3E-04	1.8E-04	1.5E-04	0.50
A_0090	1.2E-04	6.1E-05	N.D.	5.5E-05	1.1E-04	N.D.	5.3E-05	8.5E-05	5.8E-05	7.2E-05	8.1E-05	1.1E-04	5.5E-05	N.D.	N.D.	N.D.	6.6E-05	N.D.	6.4E-05	N.D.	9.6E-05	3.8E-05	N.D.	7.1E-05	N.D.	6.6E-05	8.4E-06	7.4E-05	2.1E-05	1.11
A_0121	9.3E-05	N.D.	N.D.	5.9E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	5.5E-05	2.7E-05	N.D.	N.D.	N.D.	N.D.	2.5E-04	N.A.	6.3E-05	2.4E-05	0.25
A_0111	N.D.	N.D.	N.D.	N.D.	N.D.	9.6E-05	N.D.	2.1E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	4.8E-05	N.D.	N.D.	N.D.	N.D.	2.2E-04	N.A.	1.2E-04	8.3E-05	0.54
C_0021	5.3E-02	4.9E-02	7.2E-02	9.3E-02	9.0E-02	5.6E-02	4.4E-02	1.2E-01	7.7E-02	1.1E-01	7.5E-02	9.2E-02	1.2E-01	5.7E-02	1.6E-01	6.1E-02	6.5E-02	1.0E-01	6.6E-02	3.9E-02	5.6E-02	6.4E-02	2.7E-02	7.8E-02	4.4E-02	5.5E-02	1.8E-02	8.3E-02	3.6E-02	1.50
C_0131	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	7.9E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.0E-04	N.A.	7.9E-05	N.A.	0.78
A_0048	N.D.	N.D.	N.D.	5.3E-04	N.D.	N.D.	3.8E-04	5.4E-04	N.D.	3.6E-04	7.2E-04	4.8E-04	N.D.	N.D.	N.D.	3.1E-04	3.2E-04	N.D.	N.D.	3.3E-04	3.6E-04	N.D.	4.4E-04	1.5E-03	N.D.	4.5E-04	1.0E-04	6.4E-04	6.3E-04	1.41
A_0060	7.7E-03	7.5E-03	7.9E-03	1.2E-02	1.0E-02	5.0E-03	1.1E-02	1.9E-02	7.3E-03	1.1E-02	2.2E-02	1.3E-02	8.6E-03	9.6E-03	5.6E-03	7.3E-03	9.5E-03	9.3E-03	1.4E-02	7.2E-03	9.7E-03	1.0E-02	1.2E-02	2.0E-02	5.9E-03	1.2E-02	4.9E-03	1.1E-02	7.3E-03	0.92
C_0093	2.3E-03	2.3E-03	1.0E-03	3.9E-03	2.3E-03	2.4E-03	2.2E-03	3.9E-03	2.0E-03	2.8E-03	1.4E-03	5.7E-03	2.3E-03	1.9E-03	1.4E-03	1.7E-03	1.2E-03	2.3E-03	2.9E-03	1.4E-03	3.6E-03	2.0E-03	1.4E-03	1.7E-03	8.4E-04	1.8E-03	8.5E-04	2.2E-03	9.4E-04	1.24
A_0131	N.D.	N.D.	5.5E-05	1.2E-04	5.8E-05	N.D.	5.1E-05	1.1E-04	3.9E-05	4.9E-05	3.9E-05	6.9E-05	7.0E-05	9.0E-05	N.D.	8.1E-05	6.6E-05	8.0E-05	N.D.	6.3E-05	5.7E-05	5.6E-05	N.D.	N.D.	6.3E-05	1.1E-04	4.9E-05	7.0E-05	2.1E-05	0.63
C_0050	1.4E-02	1.4E-02	2.3E-02	3.4E-02	1.9E-02	1.2E-02	2.4E-02	4.4E-02	1.6E-02	2.4E-02	2.0E-02	2.7E-02	1.6E-02	2.0E-02	1.5E-02	1.5E-02	3.9E-02	3.8E-02	2.6E-02	9.0E-03	2.0E-02	4.1E-02	1.0E-02	1.6E-02	2.8E-02	4.6E-02	2.0E-02	2.4E-02	1.0E-02	0.51
C_0029	7.9E-03	6.9E-03	7.5E-03	1.5E-02	7.9E-03	9.9E-03	9.0E-03	1.7E-02	7.7E-03	1.1E-02	1.2E-02	1.8E-02	6.9E-03	1.2E-02	3.9E-03	7.2E-03	9.6E-03	9.7E-03	1.4E-02	6.2E-03	8.4E-03	8.0E-03	5.4E-03	6.4E-03	3.4E-03	1.1E-02	5.6E-03	9.2E-03	4.3E-03	0.80
C_0038	4.2E-05	3.2E-05	9.2E-05	1.9E-04	4.2E-05	2.4E-05	N.D.	8.8E-05	N.D.	N.D.	5.9E-04	N.D.	1.1E-04	4.2E-05	N.D.	5.1E-05	6.9E-05	N.D.	N.D.	N.D.	5.9E-05	3.5E-05	N.D.	2.7E-03	5.8E-05	1.4E-04	1.0E-04	2.2E-04	5.3E-04	1.60
C_0118	N.D.	N.D.	N.D.	N.D.	N.D.	4.1E-05	N.D.	3.2E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	9.5E-05	3.6E-05	3.6E-05	6.6E-06	0.38
A_0041	N.D.	9.0E-05	6.3E-05	1.0E-04	1.2E-04	9.0E-05	N.D.	9.6E-05	1.0E-04	9.8E-05	1.5E-04	9.3E-05	2.3E-04	8.4E-05	9.4E-05	7.5E-05	N.D.	1.0E-04	1.4E-04	N.D.	1.6E-04	N.D.	N.D.	8.1E-05	N.D.	1.0E-04	3.0E-05	1.1E-04	3.9E-05	1.07
C_0153	2.2E-03	1.1E-03	1.5E-03	3.0E-03	3.5E-03	4.1E-04	1.3E-03	2.8E-03	3.2E-04	3.0E-03	1.4E-03	2.1E-03	4.0E-03	2.3E-03	1.0E-03	1.8E-03	1.6E-03	3.0E-03	1.2E-03	8.9E-04	1.9E-03	1.1E-03	1.3E-03	1.6E-03	6.5E-04	1.3E-03	6.2E-04	1.8E-03	9.4E-04	1.35
A_0034	2.0E-04	1.1E-04	1.2E-04	2.2E-04	2.4E-04	1.1E-04	7.6E-05	2.9E-04	1.3E-04	1.9E-04	2.5E-04	1.5E-04	5.3E-04	1.3E-04	2.8E-04	1.1E-04	6.1E-05	2.9E-04	1.5E-04	1.1E-04	2.1E-04	1.0E-04	5.4E-05	2.0E-04	9.1E-05	1.4E-04	7.6E-05	1.7E-04	9.3E-05	1.24
C_0124	1.3E-03	5.3E-03	8.4E-04	2.3E-03	3.1E-03	5.4E-03	4.3E-03	1.2E-03	4.2E-03	4.8E-03	8.3E-03	6.6E-03	3.0E-03	7.8E-03	9.9E-04	2.7E-03	6.7E-04	1.9E-03	8.8E-03	6.1E-04	5.2E-03	4.2E-04	4.2E-03	5.4E-03	2.3E-04	5.5E-04	5.3E-04	3.6E-03	3.2E-03	6.61
C_0127	7.9E-04	4.6E-04	1.0E-03	1.7E-03	7.1E-04	4.1E-04	7.2E-04	2.9E-03	9.1E-04	9.5E-04	8.8E-04	7.7E-04	5.8E-04	7.7E-04	3.2E-04	7.0E-04	2.0E-03	1.6E-03	1.4E-03	4.9E-04	1.0E-03	1.1E-03	3.4E-04	5.7E-04	8.3E-04	2.6E-03	1.3E-03	9.6E-04	5.3E-04	0.38
C_0025	1.7E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.A.	N.A.	1.7E-04	N.A.	1<
C_0078	4.8E-04	5.9E-04	9.9E-04	2.2E-03	9.3E-04	9.2E-04	1.0E-03	7.0E-04	7.8E-04	8.4E-04	1.0E-03	1.0E-03	8.9E-04	1.0E-03	2.4E-04	1.1E-03	1.4E-03	9.7E-04	6.1E-04	6.7E-04	2.1E-03	3.3E-04	8.9E-04	9.9E-04	1.1E-03	3.8E-04	3.8E-04	1.1E-03	8.3E-04	2.85
A_0046	N.D.	3.6E-04	4.2E-04	3.3E-04	4.6E-04	2.1E-04	7.4E-04	2.9E-04	3.9E-04	4.1E-04	N.D.	4.9E-04	5.5E-04	3.5E-04	3.9E-04	2.8E-04	N.D.	3.8E-04	6.8E-04	3.3E-04	4.2E-04	2.0E-03	4.5E-04	N.D.	3.6E-04	5.5E-04	6.3E-04	5.0E-04	3.9E-04	0.92
C_0023	3.7E-04	3.7E-04	5.5E-04	4.2E-04	4.8E-04	3.5E-04	2.9E-04	2.1E-04	4.6E-04	3.4E-04	5.8E-04	3.6E-04	8.5E-04	7.7E-04	3.3E-04	4.5E-04	2.0E-04	3.5E-04	6.5E-04	6.0E-04	3.8E-04	3.4E-04	6.5E-04	4.4E-04	3.2E-04	7.4E-04	7.0E-04	4.5E-04	1.7E-04	0.61
A_0042	1.1E-03	6.0E-04	7.2E-04	1.2E-03	1.1E-03	N.D.	1.3E-03	5.5E-04	1.0E-03	1.7E-03	9.5E-04	2.4E-03	2.1E-03	1.0E-03	N.D.	1.5E-03	5.9E-04	1.0E-03	7.5E-04	7.9E-04	8.5E-04	5.7E-04	8.9E-04	1.4E-03	9.1E-04	6.7E-04	2.4E-04	1.2E-03	5.1E-04	1.80
C_0133	1.4E-03	2.2E-03	2.0E-03	1.8E-03	3.8E-03	2.7E-03	1.8E-03	8.6E-04	2.9E-03	3.1E-03	1.9E-03	3.1E-03	1.2E-03	1.9E-03	3.6E-03	1.8E-03	6.3E-04	2.6E-03	1.7E-03	1.5E-03	1.3E-03	1.9E-03	2.0E-03	2.5E-03	1.3E-03	1.6E-03	5.6E-04	2.1E-03	7.6E-04	1.32
C_0121	N.D.	N.D.	N.D.	7.8E-05	N.D.	1.9E-05	N.D.	N.D.	N.D.	N.D.	N.D.	5.2E-03	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.8E-04	N.D.	1.1E-03	N.D.	N.D.	N.D.	N.D.	1.4E-04	N.A.	1.3E-03	1.5E-03	9.35
C_0002	3.4E-03	1.8E-03	3.3E-03	8.3E-03	3.3E-03	2.1E-03	3.8E-03	4.9E-03	3.2E-03	7.9E-03	3.0E-03	5.3E-03	4.8E-03	3.4E-03	3.5E-03	3.2E-03	3.0E-03	4.6E-03	4.8E-03	1.3E-03	5.0E-03	2.9E-03	1.5E-03	4.8E-03	2.4E-03	3.3E-03	1.6E-03	4.0E-03	1.7E-03	1.21
A_0027	6.3E-03	4.2E-03	5.9E-03	1.7E-02	8.9E-03	8.7E-04	7.4E-03	1.6E-02	3.5E-03	9.2E-03	6.6E-03	7.3E-03	1.3E-02	8.9E-03	1.1E-03	6.3E-03	1.4E-02	1.3E-02	7.1E-03	3.1E-03	8.0E-03	9.4E-03	3.9E-03	5.1E-03	7.4E-03	1.4E-02	7.2E-03	7.9E-03	4.2E-03	0.56
A_0106	N.D.	N.D.	N.D.	5.8E-05	N.D.	N.D.	N.D.	5.3E-05	N.D.	4.2E-05	N.D.	4.0E-05	5.4E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.8E-05	N.D.	N.D.	N.D.	N.D.	5.2E-05	4.5E-06	4.6E-05	9.3E-06	0.90
A_0097	3.7E-04	1.9E-04	N.D.	N.D.	3.3E-04	N.D.	4.2E-04	2.1E-04	N.D.	1.5E-03	5.3E-04	2.1E-03	2.8E-04	5.6E-04	N.D.	N.D.	3.7E-04	N.D.	6.6E-04	N.D.	4.3E-04	2.3E-04	3.3E-04	N.D.	4.7E-04	5.8E-04	3.6E-04	7.0E-04	4.8E-04	1.21
A_0081	5.5E-04	1.5E-04	2.8E-04	6.3E-04	3.7E-04	N.D.	2.9E-04	9.5E-05	3.5E-04	1.1E-03	1.2E-04	9.8E-04	8.9E-04	4.4E-04	1.4E-04	6.3E-04	2.7E-04	5.9E-04	2.9E-04	2.4E-04	3.1E-04	1.2E-04	2.7E-04	1.2E-04	2.5E-04	2.7E-04	1.7E-04	4.8E-04	3.0E-04	1.76
A_0011	N.D.	N.D.	N.D.	6.2E-04	N.D.	N.D.	N.D.	4.5E-04	N.D.	5.0E-04	N.D.	6.4E-04	6.4E-04	N.D.	N.D.	N.D.	4.4E-04	5.9E-04	N.D.	N.D.	3.2E-04	N.D.	N.D.	N.D.	N.D.	5.5E-04	2.1E-04	5.1E-04	1.0E-04	0.93
C_0015	3.6E-04	1.8E-04	3.3E-04	8.0E-04	3.6E-04	2.1E-04	3.6E-04	5.0E-04	2.6E-04	6.0E-04	3.7E-04	5.1E-04	4.1E-04	4.2E-04	4.2E-04	4.3E-04	4.2E-04	6.7E-04	3.6E-04	9.3E-04	2.0E-04	1.5E-03	1.9E-04	6.3E-04	6.5E-04	1.6E-03	2.0E-03	5.2E-04	3.4E-04	0.32
A_0114	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	7.4E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	1.3E-04	3.3E-05	7.0E-05	5.4E-06	0.55
A_0129	N.D.	N.D.	N.D.	3.4E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	3.8E-05	8.9E-05	1.7E-05	4.2E-05	1.1E-05	0.47
C_0069	2.8E-02	4.3E-02	5.2E-02	7.5E-02	3.9E-02	3.4E-02	5.1E-02	5.8E-02	3.1E-02	5.9E-02	4.6E-02	4.9E-02	5.5E-02	4.0E-02	2.2E-02	2.5E-02	4.2E-02	6.4E-02	5.0E-02	2.4E-02	4.6E-02	6.5E-02	2.5E-02	2.0E-02	2.9E-02	5.0E-02	2.0E-02	4.3E-02	1.6E-02	0.87
C_0071	6.6E-02	5.9E-02	9.5E-02	1.9E-01	7.8E-02	4.5E-02	7.9E-02	1.6E-01	7.5E-02	1.2E-01	7.3E-02	1.1E-01	1.1E-01	9.1E-02	5.8E-02	8.4E-02	1.1E-01	1.5E-01	9.5E-02	4.8E-02	7.8E-02	1.1E-01	5.4E-02	7.2E-02	8.0E-02	1.3E-01	6.3E-02	9.5E-02	3.7E-02	0.73
C_0137	7.2E-05	3.4E-05	4.0E-05	1.1E-04	4.1E-05	N.D.	1.1E-04	4.4E-05	5.4E-05	5.2E-05	6.8E-05	8.4E-05	6.3E-05	5.8E-05	N.D.	6.3E-05	9.6E-05	3.8E-05	7.7E-05	7.4E-05	1.3E-04	N.D.	4.8E-05	3.8E-05	5.4E-05	4.3E-05	3.0E-05	8.0E-05	3.9E-05	1.87
A_0063	1.1E-03	8.6E-04	1.5E-03	1.1E-03	1.8E-03	6.0E-04	4.9E-03	1.5E-03	1.9E-03	1.9E-03	2.2E-03	6.4E-03	2.0E-03	1.1E-03	1.2E-03	8.6E-04	6.3E-04	2.4E-03	1.3E-03	1.8E-03	1.8E-03	3.8E-04	1.7E-03	1.5E-03	6.1E-04	1.4E-03	1.2E-03	1.7E-03	1.2E-03	1.20
C_0095	5.5E-04	3.9E-04	7.2E-04	5.2E-04	1.1E-03	2.7E-04	1.9E-03	6.3E-04	8.7E-04	7.9E-04	8.1E-04	2.7E-03	8.3E-04	5.7E-04	4.0E-04	4.6E-04	2.5E-04	1.1E-03	6.7E-04	7.9E-04	7.7E-04	2.1E-04	6.5E-04	5.5E-04	2.8E-04	7.1E-04	5.4E-04	7.6E-04	5.0E-04	1.08
C_0097	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	3.8E-05	4.4E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	6.9E-05	N.A.	5.6E-05	4.8E-05	0.82
A_0080	2.7E-04	1.2E-04	1.5E-04	4.8E-04	1.9E-04	N.D.	2.1E-04	1.2E-04	1.1E-04	5.7E-04	1.1E-04	5.4E-04	4.0E-04	2.8E-04	N.D.	1.8E-04	1.7E-04	3.0E-04	2.2E-04	1.2E-04	2.7E-04	N.D.	1.6E-04	1.4E-04	1.7E-04	2.5E-04	1.1E-04	2.8E-04	1.5E-04	1.10
A_0078	1.9E-03	5.8E-04	1.2E-03	2.5E-03	1.4E-03	1.8E-04	1.2E-03	2.3E-04	1.5E-03	3.7E-03	4.1E-03	3.4E-03	3.3E-03	2.1E-03	4.2E-04	2.1E-03	1.0E-03	2.2E-03	1.2E-03	9.2E-04	1.2E-03	4.3E-04	1.2E-03	2.8E-04	1.1E-03	1.2E-03	9.3E-04	1.7E-03	1.1E-03	1.40
C_0146	5.5E-05	N.D.	6.1E-05	N.D.	N.D.	N.D.	N.D.	5.7E-05	N.D.	N.D.	4.7E-05	N.D.	N.D.	N.D.	N.D.	N.D.	4.6E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	8.4E-04	9.1E-05	3.1E-04	3.3E-04	1.5E-04	2.4E-04	0.57
C_0145	7.6E-03	7.6E-04	5.5E-03	7.1E-03	7.1E-03	5.6E-05	6.7E-04	1.5E-02	1.0E-04	2.3E-03	6.8E-04	8.9E-04	9.4E-03	1.1E-03	2.2E-03	2.0E-03	1.2E-02	8.1E-03	3.9E-04	3.0E-03	9.1E-04	1.5E-02	8.1E-04	6.1E-04	1.2E-02	2.4E-02	1.4E-02	4.9E-03	5.0E-03	0.20
C_0004	2.7E-02	2.5E-02	3.5E-02	7.0E-02	4.3E-02	3.4E-02	4.2E-02	5.2E-02	3.5E-02	5.3E-02	3.9E-02	6.7E-02	5.1E-02	4.3E-02	3.0E-02	3.9E-02	3.8E-02	6.0E-02	5.1E-02	2.3E-02	4.1E-02	3.9E-02	2.4E-02	3.6E-02	2.6E-02	5.0E-02	2.1E-02	4.3E-02	1.4E-02	0.86
C_0106	7.7E-05	5.7E-05	5.5E-05	1.3E-04	7.0E-05	N.D.	9.7E-05	7.7E-05	7.6E-05	8.0E-05	7.0E-05	1.3E-04	8.7E-05	7.0E-05	N.D.	1.2E-04	8.2E-05	8.3E-05	7.1E-05	4.9E-05	8.4E-05	6.2E-05	6.8E-05	N.D.	9.0E-05	6.3E-05	1.6E-05	8.1E-05	2.6E-05	1.29
C_0054	N.D.	1.4E-04	1.8E-04	4.2E-04	1.6E-04	1.3E-04	3.0E-04	1.7E-04	2.9E-04	3.3E-04	1.7E-04	3.7E-04	3.3E-04	1.7E-04	N.D.	3.4E-04	2.6E-04	1.6E-04	N.D.	1.5E-04	3.1E-04	N.D.	1.3E-04	1.6E-04	2.8E-04	1.7E-04	7.9E-05	2.7E-04	1.1E-04	1.60
A_0010	5.2E-04	2.4E-04	3.3E-04	7.7E-04	7.8E-04	4.1E-04	9.5E-04	7.8E-04	2.4E-04	9.6E-04	2.9E-04	1.0E-03	1.2E-03	4.6E-04	3.3E-04	3.9E-04	6.3E-04	8.7E-04	7.7E-04	2.4E-04	1.3E-03	6.8E-04	4.8E-04	2.7E-04	5.9E-04	5.5E-04	1.6E-04	6.6E-04	2.9E-04	1.21
C_0012	2.0E-01	1.7E-01	1.8E-01	2.5E-01	2.9E-01	1.7E-01	1.4E-01	1.5E-01	2.3E-01	1.7E-01	1.6E-01	1.6E-01	1.9E-01	1.7E-01	2.3E-01	2.0E-01	1.3E-01	2.2E-01	2.3E-01	2.0E-01	1.8E-01	2.0E-01	1.9E-01	1.7E-01	2.1E-01	2.4E-01	6.5E-02	2.0E-01	3.9E-02	0.82
A_0045	1.6E-04	1.3E-04	1.6E-04	1.7E-04	1.8E-04	1.1E-04	N.D.	3.2E-04	N.D.	2.7E-04	4.8E-04	N.D.	2.7E-04	N.D.	N.D.	N.D.	9.0E-05	2.6E-04	1.2E-04	1.7E-04	2.4E-04	1.8E-04	N.D.	1.7E-04	N.D.	1.9E-04	9.9E-05	2.2E-04	8.9E-05	1.15
A_0044	3.7E-03	6.2E-03	7.1E-03	1.2E-02	8.5E-03	1.7E-03	4.3E-03	2.4E-02	7.9E-03	1.7E-02	1.4E-02	4.5E-03	1.5E-02	5.4E-03	8.1E-03	3.1E-03	1.2E-02	1.5E-02	8.1E-03	4.2E-03	6.0E-03	1.0E-02	3.6E-03	9.2E-03	7.8E-03	1.2E-02	5.9E-03	9.8E-03	6.3E-03	0.80
C_0134	5.2E-02	9.2E-02	8.7E-02	8.3E-02	9.0E-02	8.1E-02	3.0E-02	1.8E-01	4.3E-02	6.9E-02	1.9E-01	4.3E-02	1.2E-01	7.5E-02	1.1E-01	5.3E-02	4.7E-02	1.1E-01	6.2E-02	5.6E-02	7.8E-02	2.9E-02	2.6E-02	7.0E-02	9.5E-03	2.5E-02	3.0E-02	7.0E-02	3.9E-02	2.83
A_0116	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	2.4E-04	N.D.	N.D.	N.D.	1.2E-04	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	4.8E-05	N.D.	N.D.	N.D.	N.D.	3.1E-04	N.A.	1.1E-04	9.2E-05	0.37
A_0102	6.1E-05	N.D.	N.D.	N.D.	5.9E-05	N.D.	8.9E-05	6.4E-05	N.D.	3.6E-05	N.D.	5.0E-05	N.D.	N.D.	N.D.	N.D.	3.9E-05	N.D.	N.D.	N.D.	3.6E-05	4.6E-05	6.3E-05	N.D.	7.0E-05	6.3E-05	8.9E-06	6.5E-05	2.4E-05	1.02
A_0124	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	5.6E-05	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	9.9E-05	4.5E-05	6.3E-05	9.2E-06	0.63
C_0092	5.8E-05	N.D.	3.7E-05	6.0E-05	5.1E-05	1.2E-04	3.1E-05	6.0E-05	2.1E-05	4.4E-05	N.D.	7.9E-05	4.7E-05	N.D.	N.D.	3.7E-05	5.4E-05	3.7E-05	N.D.	4.2E-05	2.2E-05	N.D.	3.4E-05	N.D.	N.D.	5.7E-05	1.4E-05	4.9E-05	2.3E-05	0.85
C_0031	2.7E-04	2.7E-04	2.3E-04	5.9E-04	3.1E-04	2.1E-04	4.8E-04	5.0E-04	3.0E-04	3.5E-04	3.8E-04	4.2E-04	4.2E-04	2.5E-04	3.2E-04	4.4E-04	2.2E-04	3.1E-04	4.3E-04	2.3E-04	3.3E-04	4.4E-04	3.0E-04	2.2E-04	1.8E-04	2.6E-04	6.8E-05	3.5E-04	1.3E-04	1.31

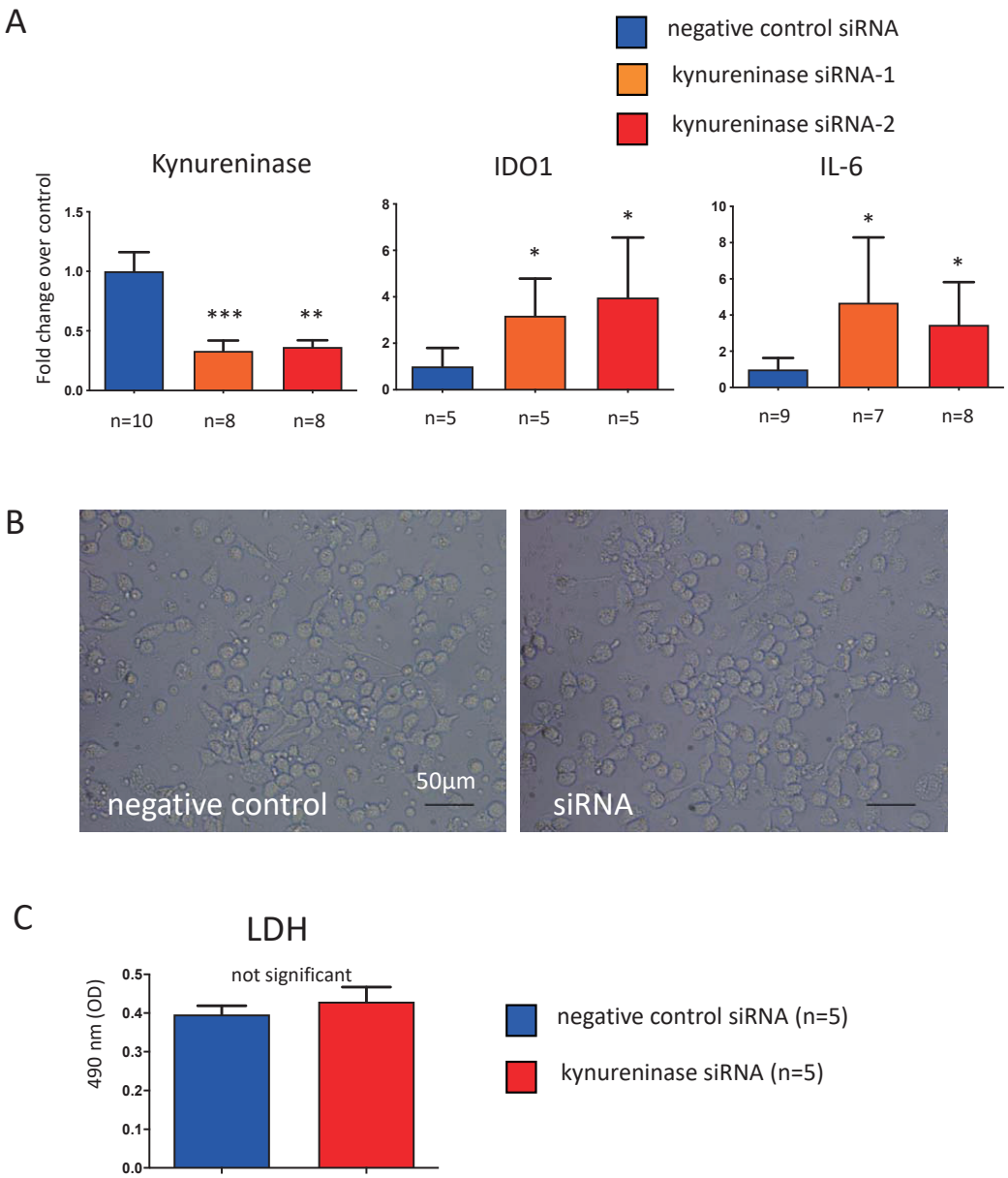
(Cont. Supplementary Table 7)





Supplementary Fig. 2.

Hematoxylin eosin and immunohistochemistry for smooth muscle actin (SMA), macrophage (CD68), kynureninase, and kynurenine 3-monooxygenase (KMO) in non-aneurysmal advanced atherosclerotic lesion.
Square in HE staining indicates an area of other high-magnification images. Macrophages express kynureninase and KMO.



Supplementary Fig. 3.

The effect of 24h-siRNA treatment against kynureninase on the expression of mRNA of kynureninase, Indoleamine 2, 3-dioxygenase (IDO)-1 and interleukin (IL)-6, cell morphology and lactate dehydrogenase (LDH) release in human peripheral blood mononuclear cells (PBMCs)-derived macrophages

(A) Significant downregulation of kynureninase mRNA or increase of IDO-1, and IL-6 mRNA in human PBMCs-derived macrophages treated with two different siRNAs against kynureninase. *$p<0.05$ vs. control, **$p<0.01$ vs. control, ***$p<0.0001$ vs. control, Kruskal-Wallis test with Dunn's multiple comparison tests.

(B) Microscopic images of human PBMCs-derived macrophages treated with negative control siRNA and kynureninase siRNA (siRNA) for 24 hours. No apparent morphological changes between them.

(C) LDH levels in culture medium of siRNA-treated macrophages. Mann-Whitney *U* test