

Supporting Information for:

Recipient Plasma Metabolomics as a Predictor of Heart Transplant Severe Primary Graft Dysfunction

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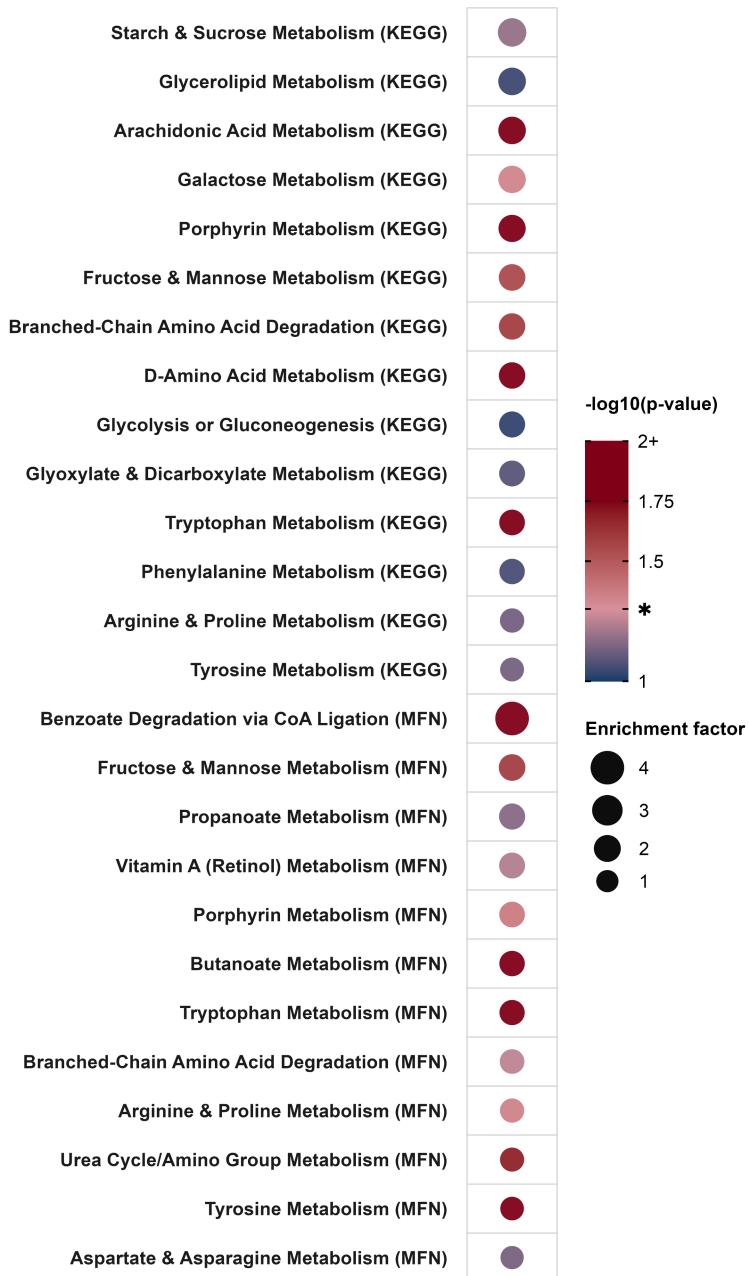
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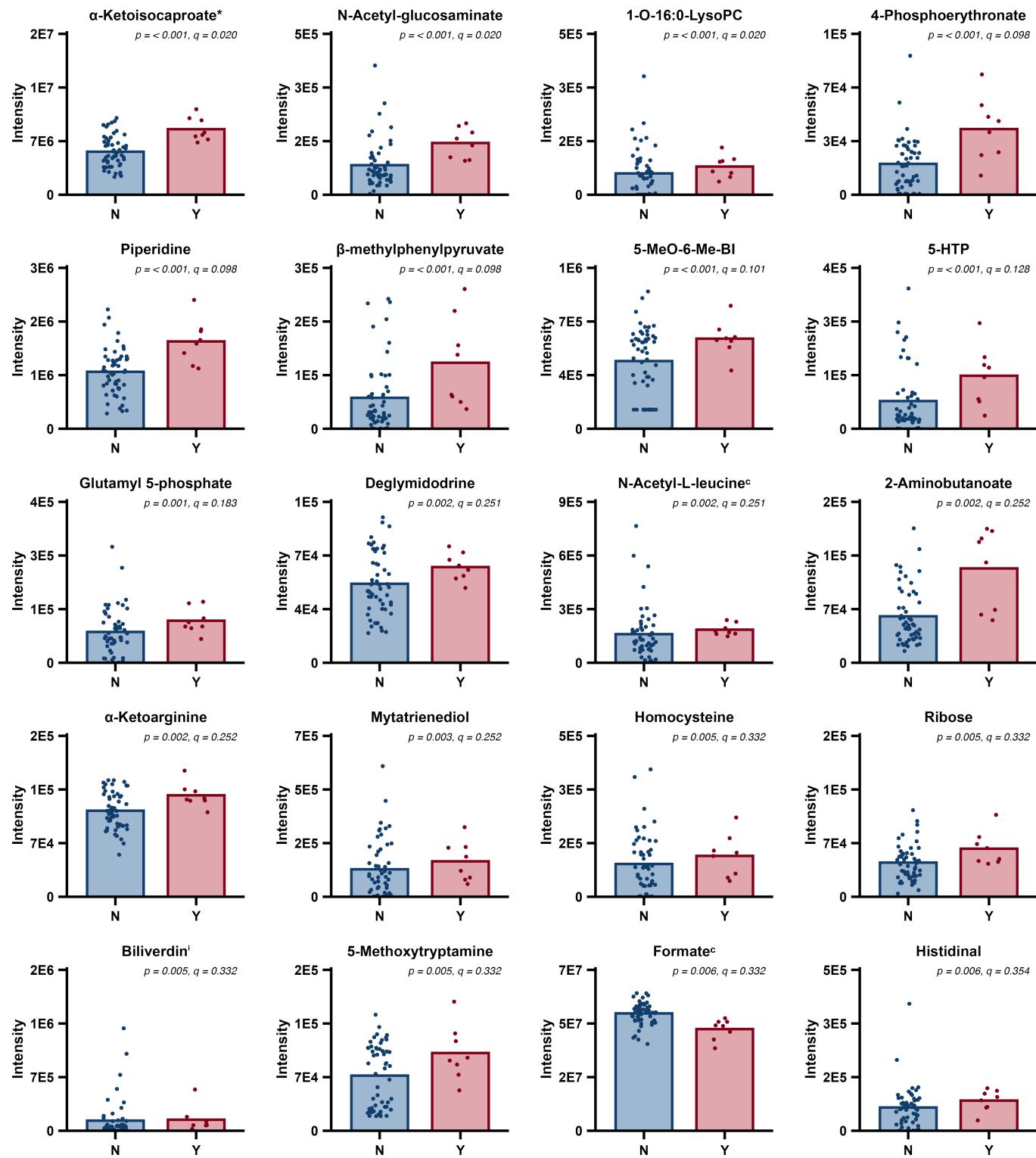
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Supplemental Figure 1



Supplemental Figure 2.1

Markers on X-axis indicate Y/N severe PGD status.



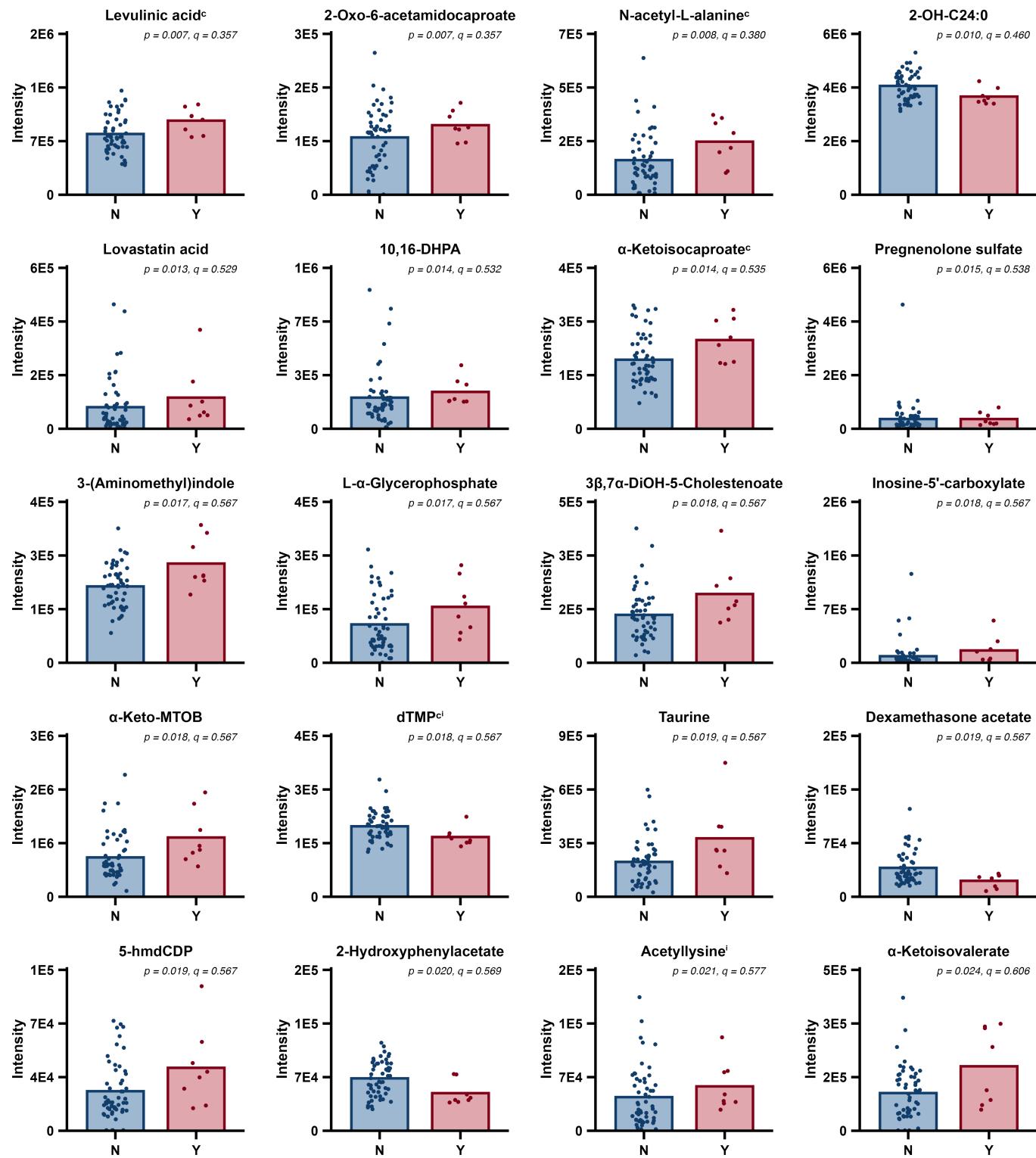
*Identified with authentic standard as M-H+HCOONa adduct; see Figure.Row.Column 2.2.2.3.

^c Identity confirmed with authentic chemical standard.

ⁱ Isomer annotated or isomer present in library of authentic chemical standards; see Figure 2.5 for details.

Supplemental Figure 2.2

Markers on X-axis indicate Y/N severe PGD status.

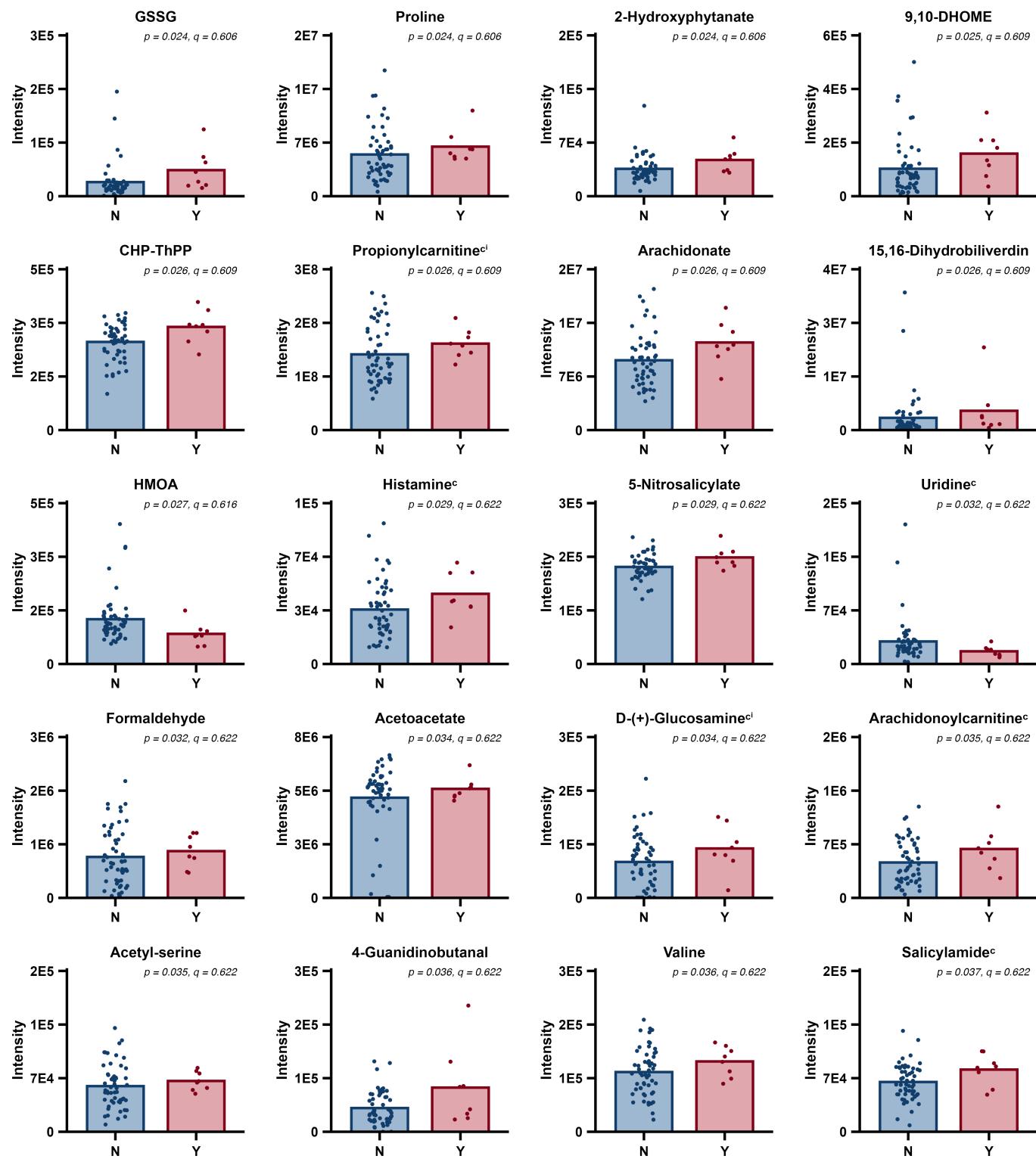


^c Identity confirmed with authentic chemical standard.

ⁱ Isomer annotated or isomer present in library of authentic chemical standards; see Figure 2.5 for details.

Supplemental Figure 2.3

Markers on X-axis indicate Y/N severe PGD status.

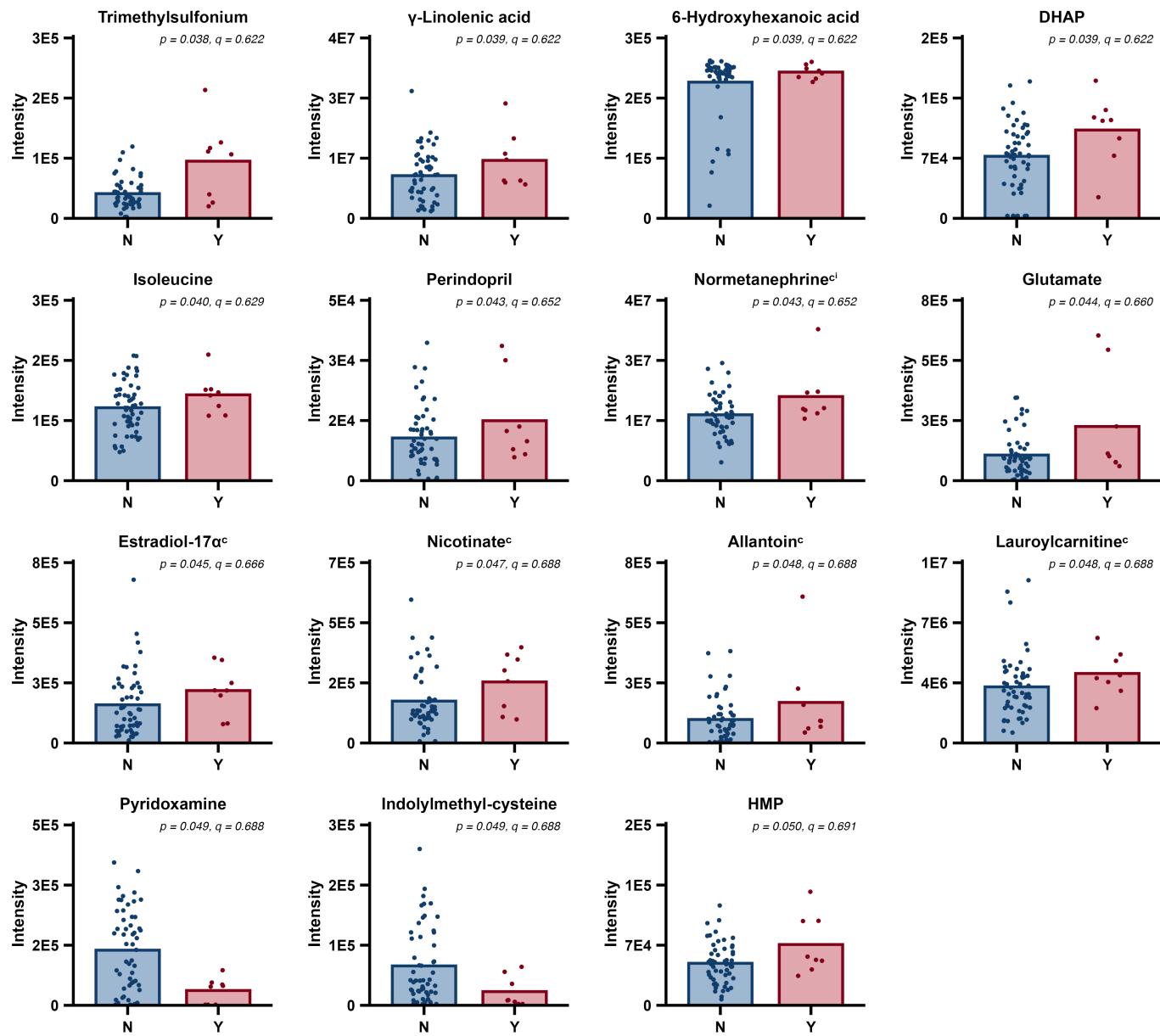


^c Identity confirmed with authentic chemical standard.

^l Isomer annotated or isomer present in library of authentic chemical standards; see Figure 2.5 for details.

Supplemental Figure 2.4

Markers on X-axis indicate Y/N severe PGD status.



^c Identity confirmed with authentic chemical standard.

^l Isomer annotated or isomer present in library of authentic chemical standards; see Figure 2.5 for details.

Supplemental Figure 2.5

Abbreviations and Adducts Table

Figure.Row.Column	Abbreviated/Displayed Name	Longform, Other, or Isomer Name(s)	Adduct
2.1.1.1	α -Ketoisocaproate*	-	M-H
2.1.1.2	N-Acetyl-glucosamine	-	M+H+HCOONa
2.1.1.3	1-O-16:0-LysoPC	1-O-Hexadecyl-lyso-sn-glycero-3-phosphocholine	M+FA-H
2.1.1.4	4-Phosphoerythronate	-	M-H
2.1.2.1	Piperidine	-	M+H
2.1.2.2	β -methyl/phenylpyruvate	(3S)-2-Oxo-3-phenylbutanoate	M+H+HCOONa
2.1.2.3	5-MeO-6-Me-BI	5-Methoxy-6-methylbenzimidazole	M+H
2.1.2.4	5-HTP	5-Hydroxy-L-tryptophan	M-H
2.1.3.1	Glutamyl 5-phosphate	-	M-H
2.1.3.2	Deglycidodine	-	M+H
2.1.3.3	N-Acetyl-L-leucine ^c	-	M+H
2.1.3.4	2-Aminobutanoate	-	M+H-NH3
2.1.4.1	α -Ketoarginine	5-Guanidino-2-oxopentanoate	M-H
2.1.4.2	Myoatrienediol	-	M+H
2.1.4.3	Homocysteine	-	M+2Na-H
2.1.4.4	Ribose	-	M+H-2H2O
2.1.5.1	Biliverdin ^c	-	M-H
2.1.5.2	5-Methoxytryptamine	-	M+H
2.1.5.3	Formate ^c	-	M+H+HCOONa
2.1.5.4	Histidinal	-	M+H
2.2.1.1	Levulinic acid ^c	-	M-H
2.2.1.2	2-Oxo-6-acetamidocaproate	6-Acetamido-2-oxohexanoate	M+H
2.2.1.3	N-Acetyl-L-alanine ^c	-	M+H-H2O
2.2.1.4	2-OH-C24:0	2-Hydroxytetracosanoic acid	M+Cl
2.2.2.1	Lovastatin acid	-	M+Hac-H
2.2.2.2	10,16-DHPA	10,16-Dihydroxyhexadecanoic acid	M+H
2.2.2.3	α -Ketoisocaproate ^c	-	M+H+HCOONa
2.2.2.4	Pregnenolone sulfate	3beta-Hydroxypregn-5-en-20-one sulfate	M-H
2.2.3.1	3-(Aminomethyl)indole	-	M+H-NH3
2.2.3.2	L- α -Glycerophosphate	sn-Glycerol 1-phosphate	M+H
2.2.3.3	3 β ,7 α -DIOH-5-Cholestenoate	3beta,7alpha-Dihydroxy-5-cholestenoate	M-H
2.2.3.4	Inosine-5'-carboxylate	-	M+H-H2O
2.2.4.1	α -Keto-MTOB	4-Methylthio-2-oxobutanoic acid	M+Cl
2.2.4.2	dTMP ^b	Thymidine 5'-monophosphate	M+FA-H
2.2.4.3	Taurine	-	M+H
2.2.4.4	Dexamethasone acetate	Dexamethasone acetate anhydrous	M-H
2.2.5.1	5-hmdCDP	2'-Deoxy-5-hydroxymethylcytidine-5'-diphosphate	M+H+HCOONa
2.2.5.2	2-Hydroxyphenylacetate	-	M+H
2.2.5.3	Acetylysine ^c	-	M-H
2.2.5.4	α -Ketoisovalerate	-	M+2Na-H
2.3.1.1	GSSG	Glutathione disulfide	M+FA-H
2.3.1.2	Proline	-	M+H
2.3.1.3	2-Hydroxyphytanate	-	M-H
2.3.1.4	9,10-DHOME	-	M+Hac-H
2.3.2.1	CHP-ThPP	3-Carboxy-1-hydroxypropyl-ThPP	M+Cl
2.3.2.2	Propionylcarnitine ^c	-	M+H-NH4
2.3.2.3	Arachidonate	-	M-H
2.3.2.4	15,16-Dihydroobilverdin	-	M+H
2.3.3.1	HMOA	4-Hydroxy-4-methyl-2-oxoadipate	M+Cl
2.3.3.2	Histamine ^c	-	M+H
2.3.3.3	5-Nitrosalicylate	-	M-H
2.3.3.4	Uridine ^c	-	M-H+HCOONa
2.3.4.1	Formaldehyde	-	M+Hac-H
2.3.4.2	Acetacetate	-	M-H
2.3.4.3	D-(+)-Glucosamine ^c	-	M+H
2.3.4.4	Arachidonoylcarnitine ^c	-	M+H
2.3.5.1	Acetyl-serine	-	M-H
2.3.5.2	4-Guanidinobutanal	-	M+H
2.3.5.3	Valine	-	M-H+HCOONa
2.3.5.4	Salicylamide ^c	-	M-H2O-H
2.4.1.1	Trimethylsulfonium	-	M+Hac-H
2.4.1.2	y-Linolenic acid	(6Z,9Z,12Z)-Octadecatrienoic acid	M-H
2.4.1.3	6-Hydroxyhexanoic acid	-	M-H
2.4.1.4	DHAP	Glycerone phosphate	M+H
2.4.2.1	Isoleucine	-	M-H+HCOONa
2.4.2.2	Perindopril	-	M-H
2.4.2.3	Normetanephrine ^c	-	M+H-H2O
2.4.2.4	Glutamate	-	M-H
2.4.3.1	Estradiol-17 α ^c	-	2M-3H
2.4.3.2	Nicotinate ^c	-	M+H
2.4.3.3	Allantoin ^c	-	M+2Na-H
2.4.3.4	Lauroylcarnitine ^c	-	M+H
2.4.4.1	Pyridoxamine	-	M+Hac-H
2.4.4.2	Indolylmethyl-cysteine	S-(Indolylmethylthiohydroximoyl)-L-cysteine	M+H
2.4.4.3	HMP	4-Amino-5-hydroxymethyl-2-methylpyrimidine	M+H

Footnote Table

Figure.Row.Column	Abbreviated/Displayed Name	Footnote
2.1.5.1	Biliverdin ^c	Annotated isomers = Biliverdin-IX- δ and Biliverdin-IX- β (M-H adducts)
2.2.4.2	dTMP ^b	Standard library isomers = 2'-Deoxyuridine 5'-monophosphate (M+Hac-H adduct)
2.2.5.3	Acetylysine ^c	Annotated isomer = 6-Acetamido-3-aminohexanoate (M-H adduct)
2.3.2.2	Propionylcarnitine ^c	Standard library isomers = 5-Aminopentanoic acid (M+H adduct)
2.3.4.3	D-(+)-Glucosamine ^c	Standard library isomers = D-Mannosamine (M+H adduct)
2.4.2.3	Normetanephrine ^c	Standard library isomers = 1,3-Dihydro-(2H-indol-2-one) (M+CH4O+H adduct)

*Identified with authentic standard as M-H+HCOONa adduct; see Figure.Row.Column 2.2.2.3.

^c Identity confirmed with authentic chemical standard.

^b Isomer annotated or isomer present in library of authentic chemical standards. See Footnote Table for details.

Expanded Metabolomics Methods and Other Technical Details

In keeping with consensus reporting standards for metabolomics experiments,¹ we provide in-depth metabolomics details as follows. In addition, we provide expanded details on statistical analyses and other technical details:

Sample Collection and Processing

Blood samples were collected immediately before transplant into K2EDTA plasma preparation tubes. Once samples were collected from patients, they were immediately placed on ice. The samples remained on ice and were centrifuged at 2500 x g for 15 minutes at 4°C within 5 hours of collection (the mean time from collection to processing for all samples = 1 hour). Following centrifugation, three aliquots were made from the supernatant, which were stored at -80°C shortly after.

Metabolomics Profiling with Liquid Chromatography-Mass Spectrometry

High-resolution metabolomics was performed in the Emory Clinical Biomarkers Laboratory as previously described.² Plasma samples were thawed, and 10 µL of plasma was added to 190 µL of acetonitrile containing 2% internal standards. Samples were then vortexed and allowed to incubate on ice for 30 minutes. Following incubation, samples were centrifuged 20,817 x g for 10 minutes. Supernatants were then transferred into autosampler vials. These were loaded onto autosampler plates and were kept at -20°C until analysis. Pooled control plasma (Equitech-Bio, SHP45) and National Institute of Standards and Technology (NIST) reference plasma were prepared and analyzed in tandem.

Metabolomics analysis was conducted with a liquid chromatography-mass spectrometry system (LC-MS) consisting of a Vanquish Duo UHPLC coupled to an Orbitrap ID-X Tribrid Mass Spectrometer (Thermo Scientific). Autosampler plates containing samples were maintained at 4°C in the autosampler of the LC-MS throughout the analysis. Samples were analyzed in triplicate on a 5-minute method using C18 chromatography coupled to negative electrospray ionization (ESI) (C18-) and hydrophilic interaction liquid chromatography coupled to positive ESI (HILIC+). Analyte separation for HILIC was performed with a Waters Acuity BEH Amide HILIC column (2.1 mm x 100 mm, 1.7 µm particle size) and gradient elution with LCMS-grade solvents and additives. The HILIC mobile phases included (Buffer A) water with 1 mM ammonium acetate and 0.1% formic acid and (Buffer B) 95% acetonitrile with 1 mM ammonium acetate and 0.1% formic acid. For the HILIC gradient, an initial 0.5 min hold at 90% B was followed by a linear decrease to 20% B from 0.6 to 2.55 min, a 2 min hold, and a 5-minute re-equilibration period. C18 chromatography was performed on a Thermo Hypersil Gold C18 column (2.1 mm x 100 mm, 1.9 µm particle size). The C18 mobile phases included (Buffer A) water with 1 mM ammonium acetate and (Buffer B) 99% acetonitrile with 1 mM ammonium acetate. For the C18 gradient, an initial 0.5 min hold at 1% B was followed by a 0.75 min linear increase to 99% B, held for 3.75 min, and a 5-minute re-equilibration period. The flow rate for both methods was 0.3 mL/min, and the column compartment was heated to 45 °C. The mass spectrometer was operated at 120k resolution and MS1 scans were collected for m/z 85-1,275. Tune parameters consisted of sheath gas at 50, auxiliary gas at 10, and sweep gas at 1. The spray voltage was set to 3.50 kV for ESI+ and -2.75 kV for ESI-.

Feature Extraction, Annotation, and Identification

ProteoWizard v3³ was used to convert raw spectral files to .mzXML files. Untargeted feature tables were then generated from extracted data using established laboratory workflows, softwares⁴⁻⁶ (namely, apLCMS and xMSAnalyzer), and an in-house R package, *MetaboJanitoR*, which is available upon request. All feature tables were corrected for batch effects using ComBat⁷ from the sva package⁸.

Untargeted feature tables comprised all HILIC+ and C18- features observed with a unique mass to charge ratio (m/z) and corresponding retention time (RT). Targeted feature tables were made up Level 1⁹ and Level 3⁹ identifications. Level 1 identifications were obtained by ion dissociation mass spectrometry relative to authentic standards (mass error = \pm 5 ppm, retention time threshold = \pm 30 seconds). Level 3 identifications were made using an in-house annotation algorithm (see <https://github.com/jamesjiadazhan> for the future release of this R package upon its publication), which generates annotations based on precursor-product correlations (note: this version of the algorithm chose a single best adduct for the feature table based on the strength of the precursor-product correlation alone). All raw data files (.mzXML format) and feature tables analyzed for this study will be available on October 31, 2026, through the NIH Common Fund's National Metabolomics Data Repository (NMDR) website, Metabolomics Workbench (Project ID PR002742; Study ID ST004328; Project DOI <http://dx.doi.org/10.21228/M87549>).

Quality Control and Data Transformation

Features that were not detected in at least 80% of samples were excluded from analysis, but were included as background for pathway enrichment analysis. All '0' values (indicating non-detection by the instrument) were replaced with $\frac{1}{2}$ the minimum value for that feature detected across all samples.¹⁰ Following this, all intensity (peak area) values were \log_2 transformed prior to further analysis. Annotated/identified features in targeted feature tables that significantly differed between study groups (Severe PGD versus No Severe PGD) were manually inspected and evaluated for biological plausibility and annotation quality control. Only annotated features encompassing endogenous, microbiome-derived, or relevant pharmacological metabolites were included.

Statistical Analysis and Data Visualization

The complete analytical pipeline and source code used for data processing and statistical analyses are publicly available at <https://github.com/jdpreston30/PGD-preop-metabolomics>. Data were analyzed and visualized using Microsoft Excel (Mac v16.101.294, Microsoft Corporation, 2025), R (v4.5.1),¹¹ and MetaboAnalyst (v6.0)¹² via the *MetaboAnalystR* package.¹³ All code was written, compiled, and run in Visual Studio Code (v1.104.2, Microsoft Corporation, 2025). All data visualizations were created using *ggplot2*¹⁴ and compiled using *cowplot*.¹⁵ Epidemiologic statistics were performed using the *TernTablesR* package.¹⁶ For all analyses, statistical significance was set at $\alpha = 0.05$.

Partial Least Squares Discriminant Analysis and Volcano Plots

Partial Least Squares Discriminant Analysis (PLS-DA) was performed using the R package mixOmics¹⁷. Volcano plots were created by calculating p-values (based on Welch's t-tests on transformed data) to compare the means of each (untargeted) feature between groups, while fold changes were determined using untransformed data. All p-value calculations were performed using R's stats package. For purposes of visualization, p-values were then $-\log_{10}$ transformed, and fold changes were \log_2 transformed.

Pathway Enrichment Analysis

All pathway enrichment analysis was performed using Mummichog¹⁸. A full list of p-values comparing means of all features between groups was first generated in R using the stats package via Welch's t-tests. These features, including the m/z, retention time, and ESI mode, along with their corresponding p-values, were then analyzed using MetaboAnalyst's¹² 'Functional Analysis' tool for LC-MS, which utilizes Mummichog¹⁸ as its core algorithm. Notably, this tool allows for the usage of Mummichog v2.0, which takes into account

retention time and ESI mode, whereas the original algorithm simply used m/z. The parameters for the pathway enrichment included the following: mixed mode, 5 ppm mass tolerance, primary ions enforced, p-value cutoff = default for top 10% of peaks, KEGG and MFN *homo sapiens* databases. A bubble plot was then constructed based on the p-values (Fisher's) and enrichment factors generated by the algorithm output.

Targeted Metabolomics Analysis

For targeted data, Welch's t-tests were employed to determine the p-values comparing means between each group. FDR adjustments were performed using the Benjamini-Hochberg (BH) procedure. A representative set of various families of metabolites was selected and displayed in the main text figures (**Figure 3A-D**), but all significantly differing annotated or identified features are displayed in **Supplemental Figure 2**. Both raw p-values and FDR q-values are displayed on all plots displaying annotated/identified metabolites.

Sensitivity Analysis

Given the exploratory and hypothesis-generating nature of the untargeted metabolomics study performed herein, which measured over 20,000 individual metabolic features, a conventional power analysis is not applicable. However, we conducted a quantitative post-hoc sensitivity analysis (further details of which can be found in the GitHub repository) to determine the minimum detectable effect size based on the group sizes in our study. With 62 patients (8 with severe PGD and 54 without), we had ~80% power ($\alpha=0.05$) to detect a minimum standardized mean difference of Cohen's $d \approx 1.08$. Based on observed within-group variance (calculated using the filtered, i.e. post-QC, untargeted feature table), the median minimum detectable fold-change was 1.87 (IQR 1.46–2.99 \times) across all features. Considering this, this study was powered to detect very large metabolic differences between groups, which is consistent with the exploratory and hypothesis-generating design. More subtle differences would require larger cohorts moving forward.

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