Cerebral Metabolic Profiling of Hypothermic Circulatory Arrest with and Without Antegrade Selective Cerebral Perfusion: Evidence from Nontargeted Tissue Metabolomics in a Rabbit Model

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Abstract

Background: Antegrade selective cerebral perfusion (ASCP) is regarded to perform cerebral protection during the thoracic aorta surgery as an adjunctive technique to deep hypothermic circulatory arrest (DHCA). However, brain metabolism profile after ASCP has not been systematically investigated by metabolomics technology.

Methods: To clarify the metabolomics profiling of ASCP, 12 New Zealand white rabbits were randomly assigned into 60 min DHCA with (DHCA+ASCP [DA] group, n = 6) and without (DHCA [D] group, n = 6) ASCP according to the random number table. ASCP was conducted by cannulation on the right subclavian artery and cross-clamping of the innominate artery. Rabbits were sacrificed 60 min after weaning off cardiopulmonary bypass. The metabolic features of the cerebral cortex were analyzed by a nontargeted metabolic profiling strategy based on gas chromatography-mass spectrometry. Variable importance projection values exceeding 1.0 were selected as potentially changed metabolites, and then Student's t-test was applied to test for statistical significance between the two groups.

Results: Metabolic profiling of brain was distinctive significantly between the two groups (Q²Y = 0.88 for partial least squares-DA model). In comparing to group D, 62 definable metabolites were varied significantly after ASCP, which were mainly related to amino acid metabolism, carbohydrate metabolism, and lipid metabolism. Kyoto Encyclopedia of Genes and Genomes analysis revealed that metabolic pathways after DHCA with ASCP were mainly involved in the activated glycolytic pathway, subdued anaerobic metabolism, and oxidative stress. In addition, L-kynurenine (P = 0.0019), 5-methoxyindole-3-acetic acid (P = 0.0499), and 5-hydroxyindole-3-acetic acid (P = 0.0495) in tryptophan metabolism pathways were decreased, and citrulline (P = 0.0158) in urea cycle was increased in group DA comparing to group D.

Conclusions: The present study applied metabolomics analysis to identify the cerebral metabolic profiling in rabbits with ASCP, and the results may shed new lights that cerebral metabolism is better preserved by ASCP compared with DHCA alone.

Key words: Antegrade Selective Cerebral Perfusion; Cardiopulmonary Bypass; Metabolic Profiling; Metabolomics

Introduction

Cerebral protection during aortic arch repair is of great concern for clinic outcome. Antegrade selective cerebral perfusion (ASCP), an adjunctive technique to deep hypothermic circulatory arrest (DHCA), has been applied for cerebral protection during aortic arch repair surgery. Evidences from several studies have indicated that ASCP...
performs cerebral protection during the aortic arch repair surgery.\[1-5\] However, brain metabolism profile after ASCP has not been systematically investigated by metabolomics technology.

Metabolites in tissues could accurately reflect the status of the biological system as the end results of metabolism. The metabolic profile of brain after ASCP may reveal new insight into cerebral protective mechanisms. Metabolomics is regarded as a useful tool for revealing the cerebral metabolic profiling as the advantage of comprehensive and quantitative analysis of global metabolites. In the present study, we compared the metabolic characteristics of the brain tissues between DHCA rabbits with and without ASCP by the virtue of gas chromatography-mass spectrometry (GC/MS). We hypothesized that cerebral metabolism reflected by carbohydrates, amino acids (AAs), and lipids could be better preserved by ASCP in a rabbit model.

**Methods**

**Chemical**

All chemicals were reagent grades and obtained from commercial sources. Bis-trimethylsilyl-trifluoroacetamide (BSTFA), a derivatization reagent, was purchased from REGIS Technologies Inc., (USA), and 2-chloro-L-phenylalanine (number 103616-89-3) was purchased from Hengbo Corporation (Shanghai, China).

**Animals and grouping schedules**

The experimental protocols were reviewed and approved by the Animal Ethics Committee of the Animal Experimental Center in Fuwai Hospital (Ethical Code: 2013-4-150-GZR). Twelve adult male New Zealand rabbits (15-20 weeks, 2.1-3.7 kg) were randomly divided into DHCA group \( n = 6 \), D group) or DHCA + ASCP group \( n = 6 \), DA group) and another 12 rabbits were used as blood donors during cardiopulmonary bypass (CPB).

The procedure was performed according to the previous studies.\[6-8\] Animals were anesthetized with sodium pentobarbital (20 mg/kg, intravenous injection), then incubated with a 3.5 F trachea cannula, and ventilated with a respirator (ELISA, Kronberg, German). The left femoral artery was catheterized for arterial pressure monitoring and blood sample collection for blood gas analysis. The CPB included a twin-roller pump (Stöckert, Munich, German), a Capiox Baby FX05 membrane oxygenator, a heat-cooler system (Maquet, Jostra HCU30), and a reservoir. The priming solution was composed of succinylated gelatin, plasmalyte-A, sodium bicarbonate, and 45 ml fresh whole blood from donor rabbits, and the total priming volume was about 70 ml. After activated clotting time reached above 480 s, CPB was initiated by cannulation on right atrium and right subclavian for venous drainage and artery inflow, respectively. Cardiac arrest was conducted by perfusion of histidine-tryptophan-ketoglutarate solution (40 ml/kg) after aortic cross-clamping when the rectal temperature was decreased to 30°C. DHCA was initiated as well as the rectal temperature was decreased to 20°C. Rabbits in group D were performed DHCA alone, and rabbit in group DA were performed DHCA with ASCP by occluding the innominate artery with flow rate at 10 ml·kg\(^{-1}\)·min\(^{-1}\).\[6\] The duration of ASCP and DHCA were maintained for 60 min. The ascending aorta was reopened when the rectal temperature was recovered to 30°C. Rabbits were weaned off CPB when the rectal temperature was 37°C and hemodynamics were stable. The rabbit cerebral cortex was harvested for metabolomics analysis at 60 min after weaning off CPB [Figure 1].

**Tissue sample pretreatment for metabolomics**

Metabolomics analyses were performed by GC/MS, the methods of which have been described previously.\[6-11\] Rabbit brain tissue (100 mg) was collected in 2 ml sealed vials and stored at −80°C until it was removed into a 2 ml tube for analysis. Both 50 μl of L-2-chlorophenylalanine (0.1 mg/ml stock in dH₂O), as an internal standard, and 0.5 ml of the extraction liquid \( v_{methan} : v_{chloriform} = 3 : 1 \) was added into the tube. To homogenize the tissue, steel balls were placed into the sample tubes. The sample was homogenized for 7 min at 55 Hz, then centrifuged for 15 min at 12,000 r/min. The 0.4 ml supernatant was transferred into a fresh 2 ml GC/MS glass vial and lyophilized in a vacuum concentrator. The dried supernatant was mixed with 80 μl of methoxyamination reagent (20 mg/ml in pyridine) and incubated for 2 h at 37°C. Total 0.1 ml BSTFA regent (1% TMCS, v/v) was added to the sample aliquots, then incubated for 1 h at 70°C. GC/MS analysis was performed when the temperatures were decreased to the room temperature.

**Metabolomic analysis by gas chromatography-mass spectrometry**

GC/time-of-flight (TOF) MS analysis was performed applying an Agilent 7890A (Agilent Technologies, USA) gas chromatograph system coupled with a Pegasus HT time-of-flight mass spectrometer (LECO Chroma TOF Pegasus HT, LECO, USA). The system utilized a Agilent DB-5MS capillary column coated with 5% diphenyl cross-linked with 95% dimethylpolysiloxane (30 m × 250 μm inner diameter, 0.25 μm film thickness; J and W Scientific, Folsom, CA, USA). 1 μl aliquot of the analyte was injected in splitless mode. Helium was used as the carrier gas, and the front inlet purge flow was 3 ml/min, and the gas flow rate through the column was 20 ml/min. Before being raised to 330°C at a rate of 10°C/min, the initial temperature was kept at 50°C for 1 min, then the temperature was maintained for 5 min at 330°C. The injection, transfer line, and ion source temperatures were 280, 280, and 220°C, respectively. The energy was −70 eV in electron impact mode. The MS data were acquired in a full-scan mode with the m/z range of 30–600 at a rate of 20 spectra/s after a solvent delay of 366 s.

**Metabolite identification**

The LECO/Fiehn Metabolomics Library was used to identify the compounds,\[12\] which would give a similarity value for
the compound identification accuracy. If the similarity is more than 700, the metabolite identification is reliable. If the similarity is less than 200, the library would only use “analyte” for the compound name. If the similarity is between 200 and 700, the compound name is a putative annotation.

Data processing and statistical analysis
The data were analyzed according to the previous studies.\(^9,13\) Internal standard normalization method was employed in this data analysis. The three-dimensional data involving the peak null number, sample name, and normalized peak area were fed to SIMCA-P 13.0 software package (Umetrics, Umea, Sweden) for principal component analysis (PCA), and partial least squares-DA (PLS-DA). Several parameters, including \(R^2_X\), \(R^2_Y\), \(Q^2_Y\)-intercepts, were used to evaluate the validity and over-fitting of the model. Based on the orthogonal projections to latent structures-DA, a loading plot was constructed, which showed the contribution of variables to difference between two groups. It also showed the important variables which were situated far from the origin, but the loading plot is complex because of many variables. To refine this analysis, the first principal component (t1) of variable importance projection (VIP) was obtained. The VIP values exceeding 1.0 were selected as potentially changed metabolites.

The variables were then assessed by Student’s \(t\)-test using SPSS 16.0 (SPSS Inc., Chicago, IL, USA), \(P < 0.05\) was considered statistically significant, whereas variables with \(P > 0.05\) were discarded between two comparison groups.

Pathway analysis
To evaluate the metabolic change associated with ASCP, metabolic pathway analysis was performed using the significantly changed metabolites determined by Student’s \(t\)-test. In our study, Kyoto Encyclopedia of Genes and Genomes (KEGG) (http://www.genome.jp/kegg/) was utilized to search for the pathways of metabolite which is a commercial database and also used for pathway analysis previously.\(^9\)

RESULTS
Multivariate analysis
In the present study, metabolomics characteristic was analyzed by a nontargeted metabolomics strategy (GC-MS). The MS data indicating the metabolic characteristic were analyzed by SIMCA-P. PCA showed the distribution of origin data. In order to obtain a higher level of group separation and get a better understanding of variables responsible for classification, a supervised PLS-DA was applied. The results showed that group DA was separated from group D by first principal component (t1) [Figure 2]. The parameters of PLS-DA for the classification from the software were \(R^2_X = 0.441\), \(R^2_Y = 0.996\), and \(Q^2_Y = 0.88\). A leave-one-out validation was further used to estimate the robustness and the predictive ability of the model, which was proceeded in order to further validate the model. The \(R^2\) and \(Q^2\) intercepts were 0.922 and 0.229 after 200 permutations, respectively [Figure 3]. The values of \(Q^2\) intercept indicated the robustness of the models, thus which showed a low risk of overfitting and reliable. The results above strongly verified the model validity.

Differential metabolites between deep hypothermic circulatory arrest and deep hypothermic circulatory arrest + antegrade selective cerebral perfusion groups
The metabolic profiles in the DA group substantially differed from the D group. Missing values of raw data were filled up by half of the minimum value. Metabolites with both VIP more than one and \(P < 0.05\) would be selected for further identification. We found that 62 endogenous biochemical markers were identified to be significantly different between D group and DA group [Supplementary Table 1]. Major changes were observed in AAs, followed by carbohydrates and then lipids. Nucleotides, cofactors and vitamins, energy substrate, and peptides were also changed significantly, and the names of which were presented in Supplementary Table 1.
Metabolic pathway

All 62 different metabolites between the two groups were searched in an online database (KEGG, http://www.genome.jp/kegg), and 43 metabolites were found in the KEGG metabolic pathways [Supplementary Table 1]. A map of ASCP-related metabolic pathways was plotted according to the most relevant distinguishing metabolites [Figure 4].

Compared to D group, metabolic pathways in DA group were characterized by activation of the glycolytic pathway with the increase in glucose-6-phosphate and glucose-1-phosphate, whereas anaerobic metabolism reflecting by decreased lactic acid production was inhibited in the early stage after ASCP. The level of glucose and pyruvate were not changed between the two groups, whereas the glucose catabolites (such as sorbitol, D-myo-inositol, and fructose) were increased significantly in DA group. Metabolites on tricarboxylic acid cycle (TAC) were not significantly different after reperfusion between the two groups, whereas several intermediates including oxalic acid and 2-hydroxybutanoic acid were higher in DA group comparing to D group \([P\) values were displayed in Supplementary Table 1].

The levels of several AAs (serine, methionine, cysteine, glycine, glutamine, glutamic acid, and aspartate) were decreased in the DA group, some of which involved in oxidative stress, such as glutamine, glutamic acid, glycine, and aspartate \([P\) values were displayed in Supplementary Table 1]. Tryptophan known as a kind of aromatic amino acids showed no difference between the two groups, but its metabolites including L-kynurenine (KYN, \(P=0.0019\), 5-methoxyindole-3-acetic acid \((P=0.0499)\), and 5-hydroxyindole-3-acetic acid \((P=0.0495)\) were decreased significantly in DA group, which had been demonstrated to be related to cerebral injury.\[10\] Furthermore, citrulline in urea cycle was found to be increased \((P=0.0158)\).

There were also significant different changes in the metabolites involved in lipid metabolism between group DA and group D. Metabolomics analysis identified increased levels of 3,7,12-trihydroxycoprostanone, cis-gondoic acid, glucronic lactone, methyl jasmonate, and heptadecanoic acid. The level of O-phosphorylethanolamine, palmitoleic acid, and palmitic acid were decreased in DA group \([P\) values were displayed in Supplementary Table 1].

**Discussion**

In this study, we focused metabolism profile alteration in the early stage after DHCA with and without ASCP in a rabbit model and yielded fundamental insights into metabolic pathways related to ASCP. In the previous study, several methods have been applied for cerebral metabolism research during ASCP including positron emission tomography, diffusion-weighted imaging, proton magnetic resonance spectroscopy, and microdialysis.\[5,15\] However, few study explored cerebral metabolism after ASCP with metabolomics. Currently, metabolomics is applied for describing the metabolic profile in neurology researches such as stroke and Alzheimer’s disease,\[9,16\] as its comprehensive and quantitative advantage. In addition, GC-MS is of excellent sensitivity and resolution for chemical analysis in metabolomics\[17-19\] and applied in our study to explore the effects of ASCP on the metabolism profile in the brain. In the present study, 62 definable metabolites were varied significantly after ASCP, which were mainly related to amino acid metabolism, carbohydrate metabolism, and lipid metabolism.

Brain energy supply has been thought to depend on glucose oxidative metabolism from glycolytic processes. In Cavus et al. study, they found that the level of cerebral glucose and pyruvate were higher during ASCP compared to that during circulatory arrest, but there were no differences of glucose.

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**Figure 2:** Score plots of PLS-DA. Score plot of PLS-DA model was obtained from DHCA and DHCA + ASCP group. \(R^2_X = 0.441\), \(R^2_Y = 0.996\) and \(Q^2_Y = 0.88\). DHCA: Deep hypothermic circulatory arrest; PLS-DA: Partial least squares-discriminant analysis; ASCP: Antegrade selective cerebral perfusion.

**Figure 3:** Permutation test of a leave-one-out validation model. Validation was obtained from 200 permutations test for the same comparisons, and the resulting \(R^2\) (green triangle) and \(Q^2\) (blue square) values were plotted. The green line represents the regression line for \(R^2\) and the blue line for \(Q^2\). The \(R^2\) (goodness of the fit) and \(Q^2\) (goodness of the prediction) intercept values were 0.922 and 0.229, respectively.
and pyruvate between the reperfusion phase after DHCA and ASCP. Accordingly, we demonstrated that glucose and pyruvate showed no difference between the early stage after ASCP and DHCA, indicating that the energy supply in D group is recovered rapidly after the reperfusion is restored, thus there is no difference in the glucose and pyruvate between group AD and group D. In addition, we found that glucose-6-phosphate and glucose-1-phosphate in glycolytic pathway were increased significantly in DA group, as well as the TAC products such as oxalate and 2-hydroxybutanoic acid.

In our study the lactic acid from glycolytic pathway was observed to be decreased in group DA, and previous studies have demonstrated that lactic acid produced from anaerobic glycolysis in astrocytes provides the primary metabolic fuel for neurons[20] and is an essential element of neuron-glial metabolic interactions[21,22] as an alternative energy substrate for brain metabolism in the absence of oxygen.[20] In the piglet model with DHCA, the level of lactate was observed to be persistently increased during the early stage after DHCA, whereas ASCP could attenuate this trend.[23] Consistently, jugular bulb pH was found to be significantly higher after ASCP comparing with DHCA[24] that is accordance with Hagl’s et al. study, in which a lower pH could be observed even 4 h after DHCA, and ASCP could attenuate the acidosis.[25] Hence, the decrease of lactic acid in group DA indicated that ASCP may improve the acid-base balance and alleviated acidosis induced by DHCA.

Figure 4: Changed metabolites and altered metabolic pathways for the most relevant distinguishing metabolites in the brain during antegrade selective cerebral perfusion. Red metabolites with an underline are up regulated, and green metabolites with italic are downregulated in DA group comparing to D group. Dotted arrows represent multiple steps. TAC: Tricarboxylic acid cycle; DA: Discriminant analysis.

Oxidative stress has been demonstrated to be increased during CPB.[26] The brain is especially susceptible to oxidative stress due to abundant lipid content in myelin sheaths, and antioxidant level has been demonstrated to be related to postoperative delirium in patients undergoing CPB surgery.[27] Cerebral damage induced by oxidative stress during DHCA attracts lots of attention, and excitatory amino acids were observed to increase significantly after DHCA.[28] Glutamate excitotoxicity was observed to be related to neuronal apoptosis and necrosis after hypothermic circulatory arrest.[29] In our study, several excitatory amino acids relating to oxidative stress were decreased in the DA group, including glutamine, glutamic acid, and aspartate, in which glutamine is the main precursor for excitatory neuromediator glutamate and inhibitory neurotransmitter γ-aminobutyric acid.[30] The decrease of excitatory amino acid (EAA) after ASCP may provide further evidence for the protective effect of ASCP.

In addition, several studies have showed that tryptophan loading could induce oxidative stress in brain cortex of rats, and tryptophan metabolism may contribute to brain damage following stroke.[31,32] In the present study, we determined that several tryptophan metabolites were decreased during the early stage after ASCP that may alleviate brain oxidative damage. Besides, KYN has been known to play an important role in the process of neuronal damage and neurodegenerative disorders, and it could promote oxidative stress following a stroke.[31] In addition,
we found that citrulline in urea cycle was also increased at the early stage after ASCP, but the exact mechanism of this effect need further study.

In our study, the change of cerebral glycerol was not determined in the early stage after ASCP, but the alteration of several long-chain fatty acid and metabolites were observed. Palmitic acid was observed to be decreased after ASCP, which has been demonstrated to induce cell death associated with trauma in the central nervous system.[33]

There are also several limitations in the present study. First of all, although metabolomics was applied to observe brain tissues metabolism in the rabbit ASCP model, metabolites alteration in plasma and cerebrospinal fluid were not determined simultaneously by metabolomics. In the further study, metabolic profile between the two groups should be observed for a longer time. In addition, targeted metabolomics may be required for further verify of the metabolites with a significant difference in the present study, what is more underlying mechanism for the cerebral metabolism after ASCP required further studied.

In conclusion, the present study clarify the cerebral metabolic profiling in DHCA rabbits with and without ASCP using metabolomics analysis, and the results may shed new lights that cerebral metabolism is better preserved by DHCA with ASCP comparing to DHCA alone, but further studies are necessary to reveal the underlying mechanism.

Supplementary information is linked to the online version of the paper on the Chinese Medical Journal website.

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Conflicts of interest

There are no conflicts of interest.

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| Metabolite                              | tR (min) | Mass  | SD    | VIP   | P     | Log₂ fold change* | Pathway                                           |
|----------------------------------------|----------|-------|-------|-------|-------|-------------------|---------------------------------------------------|
| Glutamine                              | 15.67    | 257   | 3.23  | 2.59  | 0.0018 | −20.59            | Purine metabolism                                 |
| Aspartic acid                          | 12.37    | 116   | 2.64  | 1.22  | 0.0153 | −1.61             | Alanine, aspartate and glutamate metabolism       |
| Serine                                 | 10.14    | 143   | 2.87  | 1.50  | 0.0123 | −3.00             | Glycine, serine and threonine metabolism          |
| L-cysteine                             | 13.98    | 220   | 3.76  | 2.99  | 0.0013 | −3.17             | Glycine, serine and threonine metabolism          |
| Sarcosine                              | 8.39     | 267   | 3.09  | 2.88  | 0.0001 | 19.55             | Glycine, serine and threonine metabolism          |
| Creatine                               | 13.91    | 102   | 3.10  | 2.05  | 0.0371 | 20.34             | Glycine, serine and threonine metabolism          |
| Methionine                             | 13.54    | 176   | 3.63  | 2.92  | 0.0008 | −4.08             | Cysteine and methionine metabolism                |
| Citraconic acid                        | 11.58    | 144   | 3.10  | 2.46  | 0.0013 | 3.10              | Valine, leucine and isoleucine biosynthesis       |
| Saccharopine                           | 23.18    | 199   | 3.26  | 2.62  | 0.0008 | 20.77             | Lysine biosynthesis                               |
| Citrulline                             | 16.98    | 157   | 2.82  | 1.55  | 0.0158 | 1.31              | Map00330                                         |
| 5-hydroxyindole-3-acetic acid          | 20.72    | 290   | 2.72  | 1.56  | 0.0495 | −18.99            | Arginine and proline metabolism                   |
| L-kynurenine                           | 20.30    | 174   | 3.01  | 1.97  | 0.0019 | −3.70             | Map00380                                         |
| 5-methoxyindole-3-acetic acid          | 20.34    | 156   | 2.72  | 1.44  | 0.0499 | −18.72            | Tryptophan metabolism                             |
| N-methyl-L-glutamic acid               | 12.86    | 98    | 3.21  | 2.58  | 0.0013 | −20.48            | Tryptophan metabolism                             |
| 8-aminocaprylic acid                   | 18.14    | 223   | 3.13  | 2.92  | 0.0025 | 19.72             | Null number                                      |
| N(alpha),N(alpha)-dimethyl-L-histidine | 16.95    | 214   | 3.09  | 2.42  | 0.0025 | 19.32             | Null number                                      |
| Glutamic acid                          | 14.57    | 156   | 3.52  | 2.38  | 0.0042 | −2.57             | Null number                                      |
| Glucuronic acid                        | 17.74    | 182   | 3.37  | 2.72  | 0.0013 | 4.19              | Null number                                      |
| Fructose                               | 17.36    | 273   | 3.18  | 2.97  | 0.0146 | 20.18             | Pentose and glucuronate interconversions          |
| Sorbitol                               | 18.03    | 157   | 3.31  | 2.21  | 0.0055 | 4.65              | Fructose and mannose metabolism                   |
| Myo-inositol                           | 19.62    | 150   | 3.75  | 2.55  | 0.0022 | 2.63              | Fructose and mannose metabolism                   |
| Glycolic acid                          | 7.28     | 100   | 3.78  | 3.53  | 0.0025 | 24.00             | Galactose metabolism                              |
| Ethanolamine                           | 10.23    | 319   | 3.15  | 2.14  | 0.0205 | 20.63             | Chlorocyclohexane and chlorobenzene degradation   |
| Cellobiose                             | 24.60    | 214   | 2.64  | 1.83  | 0.0170 | 17.26             | Phosphonate and phosphinate metabolism             |
| Alpha-D-glucosamine 1-phosphate        | 16.75    | 232   | 2.65  | 1.76  | 0.0108 | 17.17             | Starch and sucrose metabolism                     |
| Dihydroxyacetone                       | 10.00    | 231   | 2.64  | 1.76  | 0.0363 | 17.26             | Amino sugar and nucleotide sugar metabolism       |
| Oxalic acid                            | 8.10     | 114   | 3.04  | 1.79  | 0.0097 | 4.03              | Glycerolipid metabolism                           |
| 2-hydroxybutanoic acid                 | 8.21     | 131   | 2.51  | 1.19  | 0.0082 | 1.96              | Chloroalkane and chloroalkene degradation         |

Contd...
### Supplementary Table 1: Contd...

| Metabolite                        | tR  (min) | Mass | SD      | VIP  | P     | Log2 fold change* | Pathway                                             |
|-----------------------------------|-----------|------|---------|------|-------|-------------------|-----------------------------------------------------|
| Lactic acid                       | 7.14      | 102  | 3.15    | 2.49 | 0.0014| −2.82            | Null number                                         |
| Lactitol                          | 24.98     | 191  | 3.00    | 1.94 | 0.0222| 2.56             | Null number                                         |
| Glucose-6-phosphate               | 21.59     | 123  | 3.21    | 2.11 | 0.0228| 4.53             | Null number                                         |
| DL-dihydrophosphinosine           | 22.82     | 204  | 2.93    | 1.49 | 0.0126| 2.43             | Null number                                         |
| Glucose-1-phosphate               | 16.37     | 217  | 1.29    | 1.29 | 0.0303| 1.67             | Null number                                         |
| Gluconic lactone                  | 17.76     | 204  | 2.56    | 1.14 | 0.0059| 1.22             | map00030                                            |
| Palmitoleic acid                  | 19.02     | 211  | 2.88    | 1.41 | 0.0376| −3.05            | map00061                                            |
| Palmitic acid                     | 19.20     | 225  | 3.24    | 2.50 | 0.0119| −20.51           | map00061                                            |
| O-phosphorylethanolamine          | 16.55     | 216  | 2.71    | 1.49 | 0.0211| −1.48            | map00564                                            |
| Methyl jasmonate                  | 15.53     | 180  | 2.87    | 1.84 | 0.0100| 2.66             | map00592                                            |
| 3,7,12-trihydroxyeycoseprostane   | 27.84     | 102  | 4.01    | 3.74 | 0.0019| 25.43            | Null number                                         |
| cis-gondoic acid                  | 22.42     | 221  | 3.00    | 1.96 | 0.0499| 1.74             | Null number                                         |
| Heptadecanoic acid                | 19.97     | 156  | 3.00    | 1.50 | 0.0307| 4.45             | Null number                                         |
| Adenine                           | 17.48     | 264  | 3.39    | 2.25 | 0.0075| 1.72             | map00230                                            |
| Inosine                           | 23.49     | 221  | 2.28    | 1.07 | 0.0074| 1.70             | map00230                                            |
| Cytidine-monophosphate            | 18.93     | 243  | 3.21    | 3.00 | 0.0000| 20.15            | map00240                                            |
| Methylmalonic acid                | 9.44      | 216  | 2.93    | 1.50 | 0.0161| 4.23             | map00240                                            |
| Isoxanthopterin                   | 19.95     | 166  | 3.02    | 2.39 | 0.0158| −3.10            | map00240                                            |
| 5,6-dihydrouracil                 | 13.17     | 243  | 2.89    | 1.46 | 0.0203| −2.05            | map00480                                            |
| Oxoproline                        | 13.60     | 160  | 3.50    | 3.26 | 0.0105| 22.21            | map00480                                            |
| Acetophenone                      | 9.52      | 162  | 2.72    | 1.04 | 0.0200| −2.16            | map00642                                            |
| Acenaphthenequinone               | 20.07     | 238  | 3.32    | 3.10 | 0.0041| −21.21           | map00642                                            |
| Ascorbate                         | 18.11     | 244  | 3.24    | 1.78 | 0.0213| 1.39             | map00190                                            |
| Phosphate                         | 10.38     | 98   | 3.80    | 3.55 | 0.0240| 24.23            | map000190                                           |
| 3-aminopropionitrile              | 13.33     | 199  | 3.33    | 2.12 | 0.0478| 2.71             | map00410                                            |
| 3-cyanoalanine                    | 11.66     | 141  | 3.02    | 1.37 | 0.0364| −2.76            | map00460                                            |
| 4-aminobenzoic acid               | 17.23     | 221  | 3.09    | 2.09 | 0.0034| 2.17             | map00627                                            |
| Maleamate                         | 12.54     | 216  | 2.94    | 1.87 | 0.0331| −2.32            | map00760                                            |
| Neothesperidin                    | 23.06     | 117  | 3.05    | 2.45 | 0.0006| −19.40           | map00941                                            |
| Piceatannol                       | 23.15     | 98   | 2.93    | 1.90 | 0.0039| −3.40            | map00945                                            |
| Aminomalonic acid                 | 12.90     | 218  | 3.28    | 2.19 | 0.0027| −1.90            | map00945                                            |
| Picolinic acid                    | 11.13     | 180  | 2.84    | 1.88 | 0.0120| −18.45           | map00945                                            |
| 2-amino-3-methyl-1-butanol        | 8.75      | 145  | 2.95    | 1.99 | 0.0091| −2.31            | map00945                                            |

*Fold changes when DHCA + ASCP group was compared to DHCA group; *P* value was calculated using Student’s t-test. VIP: Variable importance projection; tR: Retention time; DHCA: Deep hypothermic circulatory arrest; ASCP: Antegrade selective cerebral perfusion; SD: Standard deviation.