EDITORIAL

UNIVERSITY OF SARAJEVO MEDICAL FACULTY - 60 YEARS OF CONTINUOUS WORK

Six decades of continuous work is a jubilee rarely found among the institutions of high education in the former Yugoslavia or even in Europe. From the standpoint of history, 60 years may not constitute long period, but if we consider time in which University of Sarajevo Medical Faculty operated, and achieved results in the areas of education and research, then we can freely state that it is an institution with its own tradition. Medical Faculty in Sarajevo, for a long time, had a generation of professors who, following the academic tradition, transferred their knowledge and experience not only to many generations of students but also to their younger colleagues, who later became teachers themselves – professors. Furthermore, Sarajevo Medical Faculty, as one of the oldest institutions of high education in Bosnia and Herzegovina, cradled numerous health-oriented faculties, not only in Sarajevo, but also at Tuzla and Banja Luka Universities. If we add that the graduates of Sarajevo Medical Faculty founded the faculties in Foča and Mostar than this School may rightfully bare the title of the cradle of medical education in Bosnia and Herzegovina.

In 1960 Department of Dental Medicine was established within the Medical Faculty. By 1974 it grew into Faculty of Dental Medicine. During 1974 the Faculty joined forces with the Faculty of Science in Sarajevo and established Faculty of Pharmacy. In the same year, Medical Faculty started with the curriculum for the students of College of Health Studies. In 1976, Medical Faculty was established in Tuzla.

As early as October 1963, in the 16th year of Faculty existence, postgraduate studies started. That was the first form of postgraduate studies at Sarajevo University and in B&H in general. The Center for postgraduate studies and continuous education was formed in 1986. In 1993 there were 26 programs at postgraduate level. Until 31 December 2006 Postgraduate studies at the Medical Faculty University of Sarajevo resulted in 657 Master's Thesis, or 657 promoted Masters of Science.

Medical Faculty from its early beginning paid special attention to the scientific work as the foundation of medical education and high quality of the healthcare profession. During the last six decades more than 15 thousand scientific and professional medical publications, monographs, textbooks, laboratory handbooks and other educational texts intended for undergraduate students were published at the Faculty. More than 200 notable scientific research projects were conducted at its departments and clinics, many of which have met international promotion and approval.

The results of the scientific work at the Faculty are also 468 doctoral thesis prepared and presented until 31 December 2006. It is important to emphasize that notable number of graduate medical doctors and health-educated staff continued their professional and scientific carriers at the universities of Europe and all over the world.

Medical faculty as Alma mater has given a crucial contribution to the development of health services and medical science in our country. It has remained the basis for all other health oriented faculties, which educated not only professionals from Bosnia and Herzegovina but also Serbia, Croatia, Montenegro as well as many other foreign students.

Professor Bakir Mehić, M.D. Ph.D.

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URINARY HIPPURIC ACID AFTER INGESTION OF EDIBLE FRUITS

JASMIN TOROMANOVIC¹, ELVIRA KOVAČ-BEŠOVIĆ₂, AIDA ŠAPČANIN², ISMET TAHIROVIĆ³, ZLATAN RIMPAPA³, GERHARD KROYER ⁴, EMIN SOFIĆ¹,²

¹ Department of Chemistry, Faculty of Science, University of Sarajevo, Zmaja od Bosne 33-35, 71 000 Sarajevo, Bosnia and Herzegovina
² Faculty of Pharmacy, University of Sarajevo, Čekaluša 90, 71000 Sarajevo, Bosnia and Herzegovina
³ Faculty of Medicine, University of Sarajevo, Čekaluša 90, 71000 Sarajevo, Bosnia and Herzegovina
⁴ Department of Food Chemistry and Technology, Technical University, Getreidemarkt 9, A-1060, Vienna, Austria

* Corresponding author

ABSTRACT

Aim of this study was to evaluate the biotransformation of simple phenols after ingestion of edible fruits and mixed food. It was analyzed hippuric acid in urine as biomarker of conjugation in the liver cells of glycine with aromatic phenolic acids such benzoic and salicylic acid from ingested food. Measurement of hippuric acid in urine samples of 10 healthy individuals: 5 female and 5 male with a mean age 51.5 years were recruited to participate in this study. Urine samples were collected for 24 hours. The additional meals 300 g of fruits: blueberry, cherry, raspberry, melon, blackberry and mixed food were given immediately before the 24 hr urine sampling. Otherwise, the meals given during 24 hr was a usually food. Biotransformation of phenols in edible fruits, that are together with liver glycins precursors of hippuric acid biosynthesis, was evaluated by direct spectrophotometric measurement of excreted hippuric acid in urine at 410 nm. It was established that the highest quantity of hippuric acid was after ingestion of 300g of bilberry fruits (p<0.003), and same quantity of cherries (p<0.003). Concentration of excreted hippuric acid was twice higher after ingestion of these fruits in comparison with hippuric acid concentrations in urine after ingestion of common – mixed food. Quantity of biosynthesised hippuric acid was in direct correlation with the concentrations of its precursors, primarily phenol acids and other simple aromatic acids ingested with food.

KEY WORDS: hippuric acid, fruits, phenolic acid, biotransformation.
INTRODUCTION

Fruits and vegetables, black and green tea contain phytochemicals, which are thought to protect consumers against cancer (1, 2), diabetes, cardiovascular diseases (3) and diseases of aging (4). However, the conventional bioanalytical techniques have significantly underestimated the phytochemicals content of these foods. Vitamin C, vitamin E and β-carotene make up only a small portion of the total chemicals in fruits and vegetables. There are a number of other phytochemicals in the foods in the chemical form of phenolics, flavonoids, anthocyanins and etc. Little is known about biotransformation of these compounds in food after food ingestion. Hippuric acid is normally present in human urine as a metabolite of dietary components. Therefore, the analysis of hippuric acid in urine provides an diet and liver diagnostic test (5). In humans the synthesis of hippuric acid takes place almost entirely in the liver. Both ATP and CoA are required for this biosynthesis (Figure 1.).

Tracing these phytochemicals at physiologic concentrations has been hindered by a lack of quantitative sensitivity for chemical equivalent tracers that could be used safely in healthy people. Accelerator mass spectrometry is a relatively new technique that provides the necessary sensitivity in attomoles and measurement precision (< 3 %) towards 14C labelled phytochemicals for detailed kinetic studies in humans at dietary levels (6). Oxigen Radical Absorbance Capacity assay (7). HPLC-multichannel ECD, mass spectrometry and also classical methods as spectrophotometry can be use for biotransformation studies (8). The biotransformation of phytochemicals from food is very complicated, for example the biotransformation of quercetin included degycosylation of the flavanol glycoside, biotransformation (intestinal and hepatic) like methylation, glucuronidation, sulphation, glutathionylation and mixtures of conjugates. It is possible 18 types of metabolites possible and more than 30 when considering isomers. Quercetin metabolites indentified in urine are methyl-quercetin with 4 isomers, quercetin-mono-glucuronide with 2 isomers, quercetin-di-glucuronide with 1 isomer, methyl- quercetin-mono-glucuronide with 2 isomers, sulfate- quercetin-glucuronide with 1 isomers and sulfate- quercetin-glucoside with 2 isomers. Quercetin is the major flavonol present in vegetables and fruits (e.g. apples, onions, grapes, kale, tea, red wine, etc.), occurs as the glycosides in plants, extensively metabolised.

MATERIALS AND METHODS

1. Subjects: 10 healthy persons, (5 f, 5 m)
2. Biological samples: urine (24 hr)
3. Edible portions of fruits (300 g)

Ten healthy subjects (5 f, 5 m) with a mean age 51.5 years were recruited to participate in this study. All study participants were in good health as determined by a medical history questionnaire, physical
TABLE 1. The content of hippuric acid in urine (24 hr)

<table>
<thead>
<tr>
<th></th>
<th>Hippuric acid (mg/24 h urine)</th>
<th>± S.D.</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blueberry</td>
<td>2247.4</td>
<td>852.13</td>
<td>&lt;0.003</td>
</tr>
<tr>
<td>Cherry</td>
<td>2118.8</td>
<td>698.65</td>
<td>&lt;0.003</td>
</tr>
<tr>
<td>Raspberry</td>
<td>1235.2</td>
<td>508.64</td>
<td></td>
</tr>
<tr>
<td>Melon</td>
<td>1085.4</td>
<td>337.66</td>
<td></td>
</tr>
<tr>
<td>Blackberry</td>
<td>1040.4</td>
<td>494.69</td>
<td></td>
</tr>
<tr>
<td>Mixed food</td>
<td>1042.8</td>
<td>712.65</td>
<td></td>
</tr>
</tbody>
</table>

examination and normal results for clinical laboratory tests. Urine samples were collected for 24 hours. An additional meal (300 g of fruits) was given immediately before the 24 hr urine sampling. Otherwise, the meals given during 24 hr was a usually food.

a) Direct Spectrophotometric determination of hippuric acid in urine

Direct spectrophotometric method was used for analysis of hippuric acid in urine (9). Hippuric acid dissolved in pyridine – water (1:1) produces a colour when benzene sulphonyl chloride was added. With the method based on this colour reaction urinary hippuric acid follows the Beer-Lambert law up to 1 mg/ml at 410 nm and path length 1 cm.

b) Apparatus and chromatographic conditions (HPLC-UV/VIS)

A Waters Assoc. Model ALC/GPC 202 liquid chromatograph equipped with a µBondapak C18 prepacked column (30 cm x 4 mm I.D., Waters) at ambient temperature and a UV detector set at 254 nm, was employed for chromatographic analysis (8). The detector was operated at 0.05 absorbance units full scale for most samples. The eluting solvent was 20 % (v/v) methanol in 0.01 M potassium phosphate containing 0.5 % (v/v) acetic acid. The flow rate was 1.0 ml/min at a pressure of 13789520 Pa. Sample injections were made on-column through a Waters U6K septumless injector with a 10-µl syringe (Hamilton 701 N).

RESULTS

On the basis of quantitative analysis of excreted hippuric acid in urine samples of ten healthy individuals, five male and five female, it was established that the highest quantity of hippuric acid was after ingestion of 300g of bilberry fruits and cherries. Statistically significantly highest excretion of hippuric acid after ingestion of bilberry fruits (p<0.003) and cherries (p<0.003) is expressed in Table 1. and Graph 1. Increased excretion of hippuric acid after ingestion of raspberry fruits is also obvious, even it was not statistically significantly different in comparison with content of hippuric acid after ingestion of common mixed food.
However, ingestion of melon fruits and blackberries didn’t cause statistically significant differences in hippuric acid content in urine in comparison with same content in urine after ingestion of common mixed food. Excretion of hippuric acid after ingestion of 300 g of different fruits, measured every 2, 6, 10, 15 and 24 hours is showed in Graph 2. High concentration of hippuric acid have been observed in subjects when they ingested prunes and followed blueberries, black grape, mixed foods, strawberries, beets, cranberries, red grape, green tea (Graph 3). The concentration of excreted hippuric acid was twice higher after ingestion of these fruits in comparison with hippuric acid concentrations in urine after ingestion of common – mixed food.

**DISCUSSION**

Two common aromatic acids are benzoic acid and cinnamic acid (unsaturated side-chain), which are widely distributed in nature and often occur free and combined in considerable amounts in drugs such as balsams. Truxillic acid, a polymer of cinnamic acid, occurs in coca leaves. Other related acids of fairly common occurrence are those having phenolic or other groupings in addition to a carboxyl group; such are: salicylic acid (o-hydroxybenzoic acid), protocatechuic acid (3,4-dihydroxybenzoic acid), veratric acid (3,4-dimethoxybenzoic acid), gallic acid (3,4,5-trihydroxybenzoic acid) and 3,4,5-trimethoxybenzoic acid. Similarly, derived from cinnamic acid, one finds p-coumaric acid (p-hydroxycinnamic acid), ferulic acid (hydroxymethoxycinnamic acid), cafffeic acid (hydroxycinnamic acid) and 3,4,5-trimethoxy-cinnamic acid. Acids having an alcohol group are quinic acid (tetrahydroxyhexahydrobenzoic acid), which occurs in cinchona bark and in some gymnosperms; shikimic acid, which is found in Japanese star-anise and is an important intermediate metabolite; and mandelic acid, C₆H₅CHOHCOOH, which occurs in combination in cyanogenetic glycosides such as those of bitter almonds and other species of Prunus (10). Tropic acid and phenyllactic acid are two aromatic hydroxy acids which occur as esters in tropane alkaloids (q.v.). Aromatic acids and phenols in the fruits and vegetables are for examples: catechol, floroglucinol, salicylic alcohol, salicylic acid, p-anisaldehyde, E-anethol, myristicin, sinapic alcohol, benzoic acid, protocatechic acid, vanillic acid, gallic acid, veratric acid, shikimic acid, homogentisic acid, p-coumaric acid, caffeeic acid, ferulic acid, cinnamic acid, quinic acid, rozmamic acid, litospermic acid and chlorogenic acid (10). Chlorogenic or caffeotannic acid is a condensation product of caffeic acid and quinic acid. It occurs in mate, coffee and nux vomica and is convert-}

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**GRAPH 4.** Liquid chromatograms of extracts from (a) normal urine, (b) normal urine containing added hippuric acid, and (c) urine from a person who eat 300 g of edible portion of cranberries.
tion, when an aqueous extract is treated with ammonia and exposed to air. Hippuric acid is normally present in human urine as a metabolite of dietary components and therapeutic components such as acetylsalicylic acid. Acetylsalicylic acid is changed to salicylic acid. This change is a typical reaction of hydrolysis of an active drug to an active metabolite. In the phase II reactions involve conjugation of benzoic and salicyl acid, during which polar groups utilized for linkage to normal body constituents such as carbohydrates, amino acids, glucuronate, glutathione, acetate, alkyl, methyl or, sulfate groups. Conjugation usually but not always results in xenobiotics and drug inactivation (11). Benzyl alcohol as preservative in food and saline solutions is metabolized to benzoic acid and then to benzoilglycine. In neonates, this metabolism is not rapid enough to prevent a sometimes fatal acidosis (11). After exposure to toluene and also administration of acetylsalicylic acid or quercetin, large quantities of hippuric acid are extracted in the urine and quantities of it are correlated to toluene exposure and to quantities of administrated drug or to ingested quercetin in food samples (12). Therefore, the analysis of hippuric acid in urine provides an exposure test and also an diet test. Hippuric acid is so called because it was first found in the urine of horses. It is benzoilglycine, a compound of benzoic acid and glycine. This is a physiological metabolic reaction that results in the detoxication of benzoic acid and benzoates. The latter occur in vegetables and fruits food or are derived from the oxidation of aromatic substances. About 0.7 g of hippuric acid per day is eliminated on the average diet, but deviations from this are to be expected, depending on the amount of precursors in the diet. Glycine is usually present in sufficient amounts to combine with any quantity of benzoates that are likely to be ingested (13). Cranberries contain 0.05 to 0.09% benzoic acid, and other fruits and berries contain smaller amounts. Some foods, e.g., catsups, are permitted by law to have 0.1% sodium benzoate added as a preservative. This is, of course, also converted to hippuric acid (14). The synthetic action of these compounds has been used as a test for liver function but evidently is a test for only this particular liver function, not for all. Nevertheless, subnormal values have been claimed in patients with various types of hepatitis and other liver conditions but not in those with uncomplicated obstruction of the common bile duct; i.e., it may aid in differentiating between hepatic and obstructive jaundice (5, 15). In conclusion biotransformation of phenols in edible fruits, that are together with liver glycins precursors of hippuric acid biosynthesis was evaluated by spectrophotometric measurement of excreted hippuric acid in urine. On the basis of measurement of hippuric acid concentration in urine samples of 10 healthy individuals it was established that the highest quantity of hippuric acid was after ingestion of 300 g of bilberry fruits (p< 0.003) and same quantity of cherries (p< 0.003) (First experiment, spectrophotometric method). Furthermore, as high concentration of hippuric acid have been observed in subjects when they ingested prunes and followed blueberries, black grape, mixed foods, strawberries, beets, cranberries, red grape, green tea (HPLC-UV / VIS method). The concentration of excreted hippuric acid was twice higher after ingestion of these fruits in comparison with hippuric acid concentrations in urine after ingestion of common – mixed food. Quantity of biosynthesised hippuric acid was in direct correlation with the concentrations of its precursors, primarily phenol acids and other simple aromatic acids ingested with food. Increased consumption of fruits and vegetables can increase the plasma, serum and urine antioxidant capacity (16, 17). Increased consumption of fruits and vegetables has been associated with protection against various diseases, including cancer (18, 19, 20) and cerebrovascular diseases (21, 22). It is not know what dietary constituents are responsible for this association, but it is often assumed that antioxidants contribute to the protection (23). Finally, the associations between phenolics intakes, these markers and health outcomes must be studied.

CONCLUSION

- It was established that the highest quantity of hippuric acid was after ingestion of 300g of bilberry fruits (p< 0.003), and same quantity of cherries (p< 0.003).
- Concentration of excreted hippuric acid was twice higher after ingestion of these fruits in comparison with hippuric acid concentrations in urine after ingestion of common – mixed food.
- Quantity of biosynthesised hippuric acid was in direct correlation with the concentrations of its precursors, primarily phenol acids and other simple aromatic acids ingested with food.
REFERENCES


