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Determination and Evaluation of Metal Bioaccessibility in Some Nuts and Seeds by in Vitro Gastro-Intestinal Method

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Abstract

Metal bioaccessibility in some nuts and seeds has been determined by performing a physiologicallybased extraction test. Nine trace metallic analytes (B, Ca, Co, Cu, Fe, Mn, Mg, Ni and Zn) in gastric and intestinal phase extractions of nuts and seeds were determined using inductively coupled plasma-atomic emission spectrometry and inductively coupled plasma-mass spectrometry. Hazelnuts, almonds, sunflower seeds, peanuts, cashew nuts, Brazil nuts, walnuts, chickpeas, pumpkin seeds and pistachio nuts were used as the materials in this study. The bioaccessible portions of magnesium and calcium were found to be higher than for the other elements whereas the B bioaccessibility was the lowest for each of the different types of nuts and seeds. Based on an ingestion rate of 10 g day⁻¹, the amounts of B, Ca, Co, Cu, Fe, Mn, Mg, Ni and Zn from the nuts and seeds accessible to the body were found to be lower than the Tolerable Upper Intake Levels. The data were also subjected to chemometric evaluation using tools such as Principal component analysis (PCA) and linear

discriminant analyses (LDA) in an attempt to classify the nuts and seeds according to these elements bioaccessibility and to find out which elements are more bioaccessible in gastric and intestinal ingestions.

Keywords: Nuts and seeds, metals, bioaccessibility, principal component analysis, linear discriminant analyses (LDA)

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Introduction

Nuts and seeds are not only rich in fiber and protein, but they also contain numerous other nutrients. These include high levels of mono and polyunsaturated fats, omega-3 fatty acids as well as other bioactive compounds including several antioxidants, which are important for heart health and can lower cholesterol levels and improve cardiovascular outcomes through their lipid lowering, anti-inflammatory, antioxidant, vasoactive, and anti-arrhythmic effect. The American Heart Foundation recommends including some nuts and seeds in the diet daily because of these apparent benefits to heart health (Tupper 2012; Yang et al. 2009; Lee et al. 2011). Nuts and seeds are also rich in micronutrients such as folic acid and niacin, vitamins (E and B₆) and minerals (Ca, Cr, Mg, Mn, Cu, Fe, Zn, Se, P and K). Vitamin E, folate, manganese, and selenium are very important in the body as they help fight damage-causing free-radicals and are therefore thought to protect against cancers. Nuts are also a good source of minerals such as zinc and magnesium and the B vitamins which are essential for energy (Tupper 2012; Nascimento et al. 2010; Naozuka et al. 2011). In addition, nuts and seeds also contain micronutrients such as Copper, chromium, iron, zinc and selenium that are essential for human health. Some toxic elements such as Pb, Cd and Hg can also be transferred to nuts

and seeds through handling, including food processing and packaging and can cause potential health effects in human body (Rodushkin et al. 2008).

Thedetermination of elements that are either essential or that have toxic effects in the human body is therefore very important for nutritional and toxic analyses in food samples. Total elemental concentrations of Al, Ag, Ba, Ca, Co, Fe, Hg, Mg, Mn, Mo, Pb, Se, Sr, and Zn in nuts and seeds have been determined in several types of nut, including the Brazil nut in numerous papers previously (Kafaoglu et al. 2014; Naozuka et al. 2011, Rodushkin et al. 2008; Cabrera et al. 2003; Naozuka et al. 2010; Kannamkumarath et al. 2004; Wuilloud et al. 2004).

In terms of nutrition, it is not sufficient to measure only the total concentrations of metal ions. Instead, it is also important to know the bioavailability, i.e.the amount adsorbed and used by the organism or the bioaccessibility; that is the fraction of a metal which is solubilized from a sample under simulated gastrointestinal conditions (Nascimento et al. 2010; Intawongse & Dean 2006a).

Information obtained for the bioavailable fraction from in vivo studies can be difficult to interpret because of physiological discrepancies between humans and the experimental animals adopted. Such problems led to the development of several in vitro systems based on gastrointestinal extraction that give an indication of the levels of metals accessible to the body by either intentional or unintentional ingestion of foods or soils. These systems include the so called physiologically-based extraction test (PBET) (Intawongse & Dean 2008). Several in vitro methods have been developed and are reported in the literature (Miller et al. 1981; Crews et al. 1983; Ruby et al. 1993; Hack & Selenka 1996). *In-vitro* testing methods have been used most for assessing oral bioaccessibility of total trace metals in soil and food samples. These methods were reviewed by Intawongse and Dean (Intawongse & Dean

2006b). *In-vitro* extraction procedures to assess bioaccessibility seek to mimic processes that occur in typically two (or occasionally three) distinct, but linked, areas of the human digestive system; (i.e. stomach and small intestine (and sometimes the mouth) (Intawongse & Dean 2006a).

The present study will focus on the determination of the bioaccessible amounts of trace elements (B, Ca, Co, Cu, Fe, Mn, Mg, Ni and Zn) in some nuts and seeds using an in-vitro gastrointestinal method that employs enzymes and dilute hydrochloric acid; the modified PBET method.. This procedure was first described by Ruby et al. (1996). Two modifications have been adopted that make the test more reproducible and easier to undertake (Rodriguez et al. 1999; Ruby et al. 1999, Medlin 1997). The original method useddialysis tubing containing sodium carbonate or bicarbonate which raised the pH of the digest ready for the small intestine extraction step. This was replaced by simply titrating the stomach extract directly with saturated sodium carbonate or bicarbonate solution to bring the pH to 7 (Rodriguez et al. 1999). Other workers (Medlin 1997, Ruby et al. 1999) showed that it was not necessary to maintain anaerobic conditions in the extraction solutions and the extraction could be carried out in screw top polypropylene vessels. Agitation of the sample solution mixture could then be reproducibly carried out by end over end shaking in a water bath (Medlin 1997). A similar modified procedure has been used in several other research studies (Intawongse & Dean, 2008; Wragg et. al. 2007; Cave et al. 2002).

2. Experimental

2.1. Reagents and solutions

Doubly de-ionized water (18.2 M Ω cm), obtained from a combined Prima and Maxima water system (Elga, Buckinghamshire, UK) was used throughout the experiment. Nitric acid (Trace Analysis grade, Fisher, Loughborough, UK) was used to digest the nut samples prior to total

metal concentration determination. Stock standard solutions of individual metals (1000 or $10,000 \text{ mg L}^{-1}$) were supplied by Fisher.

2.2. Instrumentation

An ICP-MS instrument (X Series 2, Thermo Scientific, Hemel Hempstead, UK) was used for the determination of Co and Ni. Operating conditions for the ICP-MS instrument were: forward power 1.40 kW, coolant gas flow rate 13 L min⁻¹, auxiliary gas flow rate 0.75 L min⁻¹ and nebulizer gas flow rate 0.9 L min⁻¹. The dwell time per isotope was 10 ms and 50 sweeps were used. An ICP-OES instrument (Varian 725-ES, Melbourne, Australia) was used for the determination of B, Ca, Cu, Fe, Mn, Mg, and Zn in the samples. Operating conditions for the ICP-OES instrument were: forward power 1.4 kW, plasma (coolant) gas flow rate 15 1 min⁻¹, auxiliary gas flow rate 1.5 1 min⁻¹; nebuliser gas flow rate 0.68 1 min⁻¹; the viewing height was 8 mm above the load coil and the replicate read time was 4 s. For both instruments,the sample was introduced via a V-groove nebulizer and a Sturman-Masters spray chamber.

2.3. Procedure

2.3.1. Sample preparation

Hazelnuts, almonds, sunflower seeds, peanuts, cashew nuts, Brazil nuts, walnuts, chickpeas, pumpkin seeds and pistachio nuts were purchased from a shop in Bursa, Turkey. Some of the samples were imported: e.g. almonds from Spain, cashew nuts from India, walnuts from Chile and Brazil nuts from Brazil. Other samples were produced from different parts of Turkey. The samples were ground using a pestle and mortar. The pulverised and powdered or caked nut and seed samples were then transferred into plastic bags. All nuts and seeds were treated in an identical manner.

2.3.2. Modified PBET Method

A mass of 0.4 g of the pulverized nuts and seeds was weighed accurately into a widemouthed HDPE (high density polyethylene) bottle. A volume of 40 mL of simulated gastric solution (1.25 g pepsin (Aldrich), 0.50 g sodium malate (Aldrich), 0.50 g sodium citrate (Aldrich), 420 µL lactic acid (Aldrich) and 500 µL acetic acid (BDH) made up to 1 L with freshly prepared de-ionised water, adjusted to pH 2.5 with concentrated hydrochloric acid (BDH AristaR grade)) was added to each bottle. The bottles were placed in an end over end shaker within a temperature controlled water bath set at 37°C. After one hour at 37°C, a 5.0 mL aliquot was removed and centifuged for 15 min at 3000 rpm. The liquid phase was decanted into a 15 mL capacitypolyethylene tube. This extraction sample is known as the gastric phase sample. Then, 5.0 mL of the original gastric solution was back-flushed through the filter into the HDPE bottle (to retain the original solid:solution ratio). The conditions in the vessel were then altered from those that simulate the stomach to those in the small intestineby titration to pH 7.0 with saturated sodium bicarbonate and the addition of 175 mg bile salts (Aldrich) and 50 mg pancreatin (Aldrich). The samples were then incubated in the water bath for a further four hours when a second 5 mL aliquot was removed and filtered. This sample is known as *intestinal phase sample*.

2.3.3 Preparation of samples for total metal analysis

Etc. etc

2.3.4. Sample analysis

The concentrations of Co and Ni were determined using ICP-MS and B, Ca, Cu, Fe, Mn, Mg, Ni and Zn concentrations were determined using ICP-OES for all sample types. As an internal standard for ICP-MS determinations, a mixture of indium and iridium was added to each digest to give a final concentration of 100 μ g L⁻¹ after dilution to 25 mL. Similarly, the internal standard mixture was also added to all blanks and standards. All results are expressed

as the mean of the three replicates. All statistical calculations were made using the IBM SPSS Statistics version 21 software (1989–2012)) package.

For in vitro results, the bioaccessible metal concentrations for the stomach and intestinal digestions are calculated by dividing the metal ions' concentrations measured in the in vitro gastric phase or the in vitro intestinal phase solutions by total concentrations of metal ions as described by the following equation (Intawongse and Dean 2008):

In vitro bioaccessible metal ion,
$$\% = \frac{[In - vitro metal]}{[Toplam metal]} x100$$

3. Results and Discussion

An in vitro gastro-intestinal method was applied to determine the trace element concentrations that can dissolve in gastric and intestinal solutions inform samples of different nuts and seeds. The concentrations of elements (B, Ca, Co, Cu, Fe, Mn, Mg, Ni and Zn) obtained using total sample digestion (Kafaoğlu et al. 2014) and the in-vitro gastrointestinal experiments in nuts and seeds (mean and standard deviation) and the proportion of the trace elements that are bioaccessible are given in Table 1. The data given in this Table indicate that the vast proportion of the trace elements are not bioaccessible. The bioaccessible ratio is the highest for Zn in most of the nuts and seeds. The highest bioaccessible ratio in gastric phase was observed in hazelnut (20.6%) whereas 36.1 % of Zn is bioavailable in pistachio nut after the intestinal phase digestion. The lowest bioaccessible ratios were generally obtained for B and Cu in most of the nuts and seeds samples where the highest bioavailable ratio for B is 11.3 % in the gastric phase in walnut and for Cu it is 11.1% in the intestinal phase of peanuts. Magnesium and calcium concentration in gastric and intestinal solutions were higher than other element's concentrations for most of the nuts and seeds. It was shown that nuts and seeds are very healthy snacks because of their magnesium and calcium contents. These results also show that when the nuts and seeds are digested, even though the bioaccessible fractions are not large, they are also still supplying very beneficial levels of Fe, Mn and Zn which are essential elements for humans. Therefore the nuts and seeds are very good food supplements.

Daily B, Ca, Co, Cu, Fe, Mn, Mg, Ni and Zn ingestion amounts from nuts and seeds have been calculated based on an ingestion rate of 10 g of nut or seed day⁻¹ using the in vitro intestinal bioavailability. The results are given in Table 2 which also compares the levels found with the Dietary Reference Intakes (DRIs) given as Recommended Dietary Allowances and Adequate Intakes (USDA (a), 2010; WHO, 1996) and Tolerable Upper Intake Levels (USDA (b), 2010). Although no safe Recommended Dietary Allowance (RDA) for cobalt has been established yet, the average adult intake of cobalt is 5 to 8 µg per day (University of UTAH Health Care). If cobalt is present in nutritional supplements, it is usually given in micrograms (µg). Recommended intakes of cobalt have not been set as the only form of cobalt required by the body is vitamin B12, of which cobalt is an integral part (Food Standards Agency, 2003). In the UK, COMA has set a RNI value for vitamin B12 of 1.5 µg/day for adults, including pregnant women (COMA, 1991). The average daily intake of cobalt from food is estimated to be 5 to 40 µg/day (EPA, 2000). As can be seen form Table 2, the concentrations of these elements on an ingestion rate of 10 g day⁻¹ of nuts and seeds using the in-vitro intestinal bioavailability results much lower than the Tolerable Upper Intake Levels. For most of the elements, consumption of 1 kg of seeds and nuts in a day would still be insufficient to reach the Tolerable Upper Intake Levels.

Further investigations were performed using Principal Component Analyses and Linear Discriminant Analysis (LDA) to investigate whether there is a relationship between metal ions in gastric and intestinal phases and nuts and seeds

2.1 Principal Component Analyses

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Principal Component Analysis was applied to the average values for each element for each matrix (nut or seed) for gastric and intestinal ingestions given in Table 1. The principal components which have eigenvalues higher than 1 were extracted for each gastric and intestinal ingestions separately. The observations from PCA analyses were described as the score values for the matrix (nut or seed) and loading values for each trace element. The components were rotated using Varimax rotation. This led to the formation of three principal components for each in-vitro gastric ingestions. The first component accounted for 41.6 %, the second for 27.0 % and the third for 13.6 % of the total variation of the data. The first three components account for 82.2 % of variances for all of the data given for in-vitro gastric ingestion. Table 3 gives the loadings and the scores of the three rotated principal components for in-vitro gastric ingestion of these nuts and seeds. From Table 3, it can be seen that the concentrations of the first group of elements, B, Ca, Cu, Mn and Zn on the first principal component are higher for walnut and hazelnut than for the other nuts and seeds. This is because these samples have higher score values for the first principal component. The score and loading values of the second principal component were evaluated and demonstrated that the Brazil nuts have higher concentrations of Co, Mg and Ni than the other nuts and seeds. When the third principal component is interpreted, it is evident that the concentrations of Fe and Mn are higher in hazelnut than other nuts and seeds for in-vitro gastric ingestions of these nuts and seeds.

The classification of the different nuts and seeds in the gastric phase from the viewpoint of metal contents can be made using three ways PC loading and score graphs. Fig. 1a shows the two way PCA loadings graphs (PC 1-2) and Fig. 1b shows the two way PCA score graphs (PC 1-2). The PC 1-2 graph shows the highest percentage of total variance of about 68.6%. Using Figure (1a and b), the nuts and seeds can be classified into five groups in gastric phase. These groups include:

Group 1: Brazil nuts

Group 2: Walnuts, Hazelnuts,

Group 3: Almonds, pistachio nuts, chickpeas,

Group 4: Sunflower seeds, cashew nuts,

Group 5: Pumpkin seeds, peanuts

For in-vitro intestinal ingestions, the first component accounted for 44.1 %, the second for 19.6 %, the third for 13.0 % and the fourth for 12.0 % of the total variation of the data. The first four components account for 88.7 % of variances for all of the data given for in-vitro intestinal ingestion. The first component represents the maximum variation of the data set. Table 4 gives the loadings and the scores of the four rotated principal components for in-vitro intestinal ingestion of these nuts and seeds. From Table 4, when the score and loading values of the first principal components were evaluated, it was demonstrated that Cu, Fe, Mg, Ni and Zn are higher for white sunflower seeds (the highest), and cashew than for the other nuts and seeds. The concentrations of Co and Cu on the second principal component are higher in Brazil nuts than the other nuts and seeds. When the third principal component is interpreted, it is evident that the concentrations of Ca are higher in almond and pistachio than in other nuts and seeds for in-vitro intestinal ingestions. Interpretation of the fourth principal component demonstrates that the concentrations of B and Mn are higher in hazelnut (the highest) and sunflower seeds. Three ways PC loading and score graphs were made for the classification of the different nuts and seeds in intestinal phase from the viewpoint of metal contents. Fig. 2 a shows the three way PCA loadings graphs (PC 1-2-3) and Fig. 2 b shows the three way PCA score graphs (PC 1-2-3). The PC 1-2-3 graph shows the highest percentage of total variance of about 76.7. Using Figure (2a and 2b), the nuts and seeds can be classified into four groups in the intestinal phase. These groups include: Group 1: Chickpeas, Pumpkin seeds, cashew nuts, sunflower seeds

Group 2: Brazil nuts,

Group 3: Hazelnuts, Walnuts,

Group 4: Almonds, pistachio nuts, peanuts

The recognition of the groups made to classify the nuts and seeds for gastric and intestinal ingestions using PCA were done by introducing these groupings to Linear Discriminant Analysis (LDA), below.

Linear discriminant analysis

Linear discriminant analysis can be used to show how these group members made by PCA above for gastric and intestinal ingestions may correctly be classified as a percentage of the original group. The linear discriminant analysis was performed for the 9 elements on each of the groups for gastric and intestinal ingestions resulting from PCA as discussed above. The recognition of the groups was highly satisfactory for gastric digestions using LDA; withall the group members determined using PCA being predicted to be 100.0% correctly classified. Also, cross-validation segments for the LDA model validation were performed for all the data sets for nuts and seeds for the gastric digestions with 90.0% of the cross-validated grouped cases being correctly classified using PCA. From the results of the cross-validation, only the Brazil nuts were mis-classified using the PCA interpretations made above. This nut should have been included in Group 2 according to the cross-validated groups.

Four canonical discriminant functions with Eigenvalues greater than 1 were obtained from the data. The first canonical discriminant function explains 95.7 % of the variance. The discriminant equation of the first function is:

Z = -10.06 + (3.10 x Co) + (53.01 x Cu) + (-9.51 x Mn) + (-0.21 x Ni)

The concentrations of each of the metals were inserted to this equation and the figure obtained for each nut or seed type is given in the parentheses below enabling the first canonical discriminant function for each of the nuts and seeds from gastric indigestions to make clear groupings :

Group 1: Brazil nuts (129.3)

Group 2: Walnuts (6.43), Hazelnuts (6.55),

Group 3: Almonds (-5.40), pistachio nuts (-8.00), chickpeas (-6.95),

Group 4: Sunflower seeds (-38.9), cashew nuts (-39.9),

Group 5: Pumpkin seeds (-22.3), peanuts (-20.8)

The group members for intestinal digestions determined by PCA are in the predicted group that 100.0% of original grouped cases were correctly classified. From cross-validation segments for the LDA model validation, 90.0 % of cross validated grouped cases were correctly classified above by PCA. Only Brazil nuts were mis-classified by PCA interpretations made above from the results of the cross-validation. This nut should have been included in Group 1 according to the cross-validated groups.

Three canonical discriminant functions with Eigenvalues greater than 1 were obtained by linear discriminant analyses. The first canonical discriminant function explains nearly 100 % of the variance. The discriminant equation of the first function is:

Z = -79.0 + (0.025 x Ca) + (2.73 x Co) + (-22.7 x Mn) + (10.2 x Zn)

The concentrations of each of the metals in the intestinal digestions were inserted to the equation above and the figure obtained for each nut or seed type is given in the parentheses

below. The calculated values given in the parentheses from first canonical discriminant function for each of the nuts and seeds enables clear groupings to be made:

Group 1: Chickpeas (-14.2), Pumpkin seeds (-15.7), cashew nuts (-12.4), sunflower seeds (-4.2)

Group 2: Brazil nuts (269.0),

Group 3: Hazelnuts (-62.3), Walnuts (-62.9),

Group 4: Almonds (-28.7), pistachio nuts (-29.5), peanuts (-29.1)

The linear discriminant analyses therefore proved that the groupings for gastric and intestinal digestions made by PCA are highly accurate because the figures for each of the nuts in each of the groups are very close to each other.

Conclusions

The trace element concentrations that can dissolve in gastric and intestinal solutions obtained from different nuts and seeds were determined using an in vitro gastro-intestinal method. The bioaccessible ratio of elements (B, Ca, Co, Cu, Fe, Mn, Mg, Ni and Zn) in nuts and seeds was calculated using concentrations found using total digestion and the in-vitro gastrointestinal experiments in nuts and seeds. The bioaccessible ratio as expressed as a percentage was highest for Zn compared with other elements while the ratio was the lowest for B in gastric and intestinal solutions for each of the nuts and seeds. Magnesium and calcium concentrations in gastric and intestinal solutions were higher than other element's concentrations for most of the nuts and seeds. The concentrations of B were the lowest for the other elements in gastric and intestinal solutions for each of the nuts and seeds.

Relationships between nuts and seeds from the perspective of metal concentrations in gastric and intestinal solutions were demonstrated using PCA interpretations. The chemometric tool of LDA demonstrated that these groupings made using PCA in gastric and

intestinal digestions were 100% correctly classified. The interpretations between nuts and seeds using their heavy metal concentrations are based solely on the statistical analysis of the analytical data obtained.

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References

Cabrera C, Lloris F, Gimenez R, Olalla M, Lopez MC. 2003. Mineral content in legumes and nuts: contribution to the Spanish dietary intake. Sci Total Environ. 308: 1–14.

Cave MR, Wragg J, Palumbo B, Klinck BA. 2002. Measurement of the bioaccessibility of arsenic in UK soils, technical report. R&D Technical Report P5-062/TR02, British Geological Survey, Environmental Agency.

COMA 1991. Dietary Reference Values for Food Energy and Nutrients for the United Kingdom. Report of the Panel on Dietary Reference Values, Committee on Medical Aspects of Food and Nutrition Policy. HMSO, London.

Crews HM, Burrell JA, McWeeny DJ. 1983. Preliminary enzymolysis studies on trace element extractability from food. J Sci Food Agric. 34: 997–1004.

EPA, 2000. United Stated Environmental Agency, Cobalt Compounds. http://www.epa.gov/ttn/atw/hlthef/cobalt.html. (accessed 4.09.2014)

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Food Standards Agency, Safe Upper Levels for Vitamins and Minerals, Expert Group on Vitamins and Minerals, May, 2003 http://tna.europarchive.org/20110911090542/http://cot.food.gov.uk/pdfs/vitmin2003.pdf (accessed 4.09.2014)

Hack A, Selenka F. 1996. Mobilization of PAH and PCB from contaminated soil using a digestive tract Model. Toxicol Lett. 88:199–210.

Intawongse M, Dean, JR. 2006a. Uptake of heavy metals by vegetable plants grown on contaminated soil and their bioavailability in the human gastrointestinal tract. Food Addit. Contam. 23: 36-48.

Intawongse M, Dean JR. 2006b. *In-vitro* testing for assessing oral bioaccessibility of trace metals in soil and food samples. TrAC-Trend Anal Chem. 25: 876–886.

Intawongse M, Dean JR. 2008. Use of the physiologically-based extraction test to assess the oral bioaccessibility of metals in vegetable plants grown in contaminated soil. Environ Pollut. 152: 60–72.

Kafaoğlu B, Fisher A, Hill S, Kara D. 2014. Chemometric evaluation of trace metal concentrations in some nuts and seeds. Food Addit. Contam. A. 31: 1529–1538.

Kannamkumarath SS, Wuilloud RG, Caruso JA. 2004. Studies of various elements of nutritional and toxicological interest associated with different molecular weight fractions in Brazil nuts. J Agric Food Chem. 52: 5773–5780.

Lee JH, Lavie CJ, O'Keefe JH, Milani R. 2011. Chapter 8 – Nuts and Seeds in Cardiovascular Health. Nuts and Seeds in Health and Disease Prevention. 75–82.

Medlin EA. 1997. An in vitro method for estimating the relative bioavailability of lead in humans. Master's Thesis, Department of Geological Sciences, University of Colorado at Boulder.

Miller DD, Schricker BR, Rasmussen RR, Campen DV. 1981. An in vitro method for estimation of iron availability from meals. Am J Clin Nutr. 34:2248–2256.

Naozuka J, Marana SR, Oliveira PV. 2010. Water-soluble Cu, Fe, Mn and Zn species in nuts and seeds. J Food Comp Anal. 23:78–85.

Naozuka J, Vieira EC, Nascimento AN, Oliveira PV. 2011. Elemental analysis of nuts and seeds by axially viewed ICP OES. Food Chem. 124: 1667-1672.

Nascimento AN, Naozuka J, Oliveira PV. 2010. In vitro evaluation of Cu and Fe bioavailability in cashew nuts by off-line coupled SEC-UV and SIMAAS. Microchem. J. 96: 58-63.

Rodriguez RR, Basta NT, Casteel S, Pace L. 1999. An in vitro gastrointestinal method to estimate bioavailable arsenic in contaminated soils and solid media. Environ Sci Technol. 33: 642-649.

Rodushkin I, Engström E, Sörlin D, Baxter D. 2008. Levels of inorganic constituents in raw nuts and seeds on the Swedish market. Sci Total Environ. 392: 290–304.

Ruby MV, Davis A, Link TE, Schoof R, Chaney RL, Freeman GB, Bergstrom P. 1993. Development of an in vitro screening test to evaluate the in vivo bioaccessibility of ingested mine-waste lead. Environ Sci Technol. 27:2870–2877. Ruby MV, Davis A, Schoof R, Eberle S, Sellstone CM. 1996. Estimation of lead and arsenic bioavailability using a physiologically based extraction test. Environ Sci Technol. 30:422–430.

Ruby MV, Schoof R, Brattin W, Goldade M, Post G, Harnois M, Mosby DE, Casteel SW, Berti W, Carpenter M, Edwards D, Cragin D, Chappell W. 1999. Advances in evaluating the oral bioavailability of inorganics in soil for use in human health risk assessment. Environ Sci Technol. 33: 3697-3705.

Tupper N. 2012. Best Healthy Nuts and Seeds for Weight Loss. Retrieved August 28, 2013 from: http://www.caloriesecrets.net/best-healthy-nuts-and-seeds-for-weight-loss/

University of Utah Health Care, Health Library, cobalt, (http://healthcare.utah.edu/healthlibrary/related/doc.php?type=19&id=cobalt)

USDA (a). 2010. Food and Nutrition Board, Institute of Medicine, National Academies, Dietary Reference Intakes (DRIs): Recommended Dietary Allowances and Adequate Intakes http://www.nal.usda.gov/fnic/DRI/DRI_Tables/UL_vitamins_elements.pdf (accessed 01.05.2010)

USDA (b). 2010. Food and Nutrition Board, Institute of Medicine, National Academies, Dietary Reference Intakes (DRIs): Adequate Intakes and Tolerable Upper Intake Levels http://www.nal.usda.gov/fnic/DRI/DRI_Tables/recommended_intakes_individuals.pdf (accessed 01.05.2010)

WHO. 1996. Trace elements in human nutrition and health, World Health Organization Geneva ISBN 92 4 156173 4, Macmillan/Ceuterick. Wragg J, Cave M, Nathanail P. 2007. A Study of the relationship between arsenic bioaccessibility and its solid-phase distribution in soils from Wellingborough, UK. J Environ Sci Health A Tox Hazard Subst Environ Eng 42: 1303-1315.

Wuilloud RG, Kannamkumarath SS, Caruso JA. 2004. Speciation of nickel, copper, zinc, and manganese in different edible nuts: a comparative study of molecular size distribution by SEC-UV–ICP-MS. Anal Bioanal Chem. 379: 495–503.

Yang J, Liu RH, Halim L. 2009. Antioxidant and antiproliferative activities of common edible nut seeds. LWT - Food Sci Technol. 42:1–8.

Legends to Figures

Figure 1. Two way Principal Component Analysis loadings and score plot for gastric ingestions

Figure 2. Three way Principal Component Analysis loadings and score plot for intestinal ingestions

Table 1. Concentrations of elements obtained using total digestion and the in-vitro gastrointestinal experiments in nuts and seeds (mean and standard deviation)

		B (mg/kg)	Ca (mg/kg)	Co (µg/kg)	Cu (mg/kg)	Fe (mg/kg)	Mg (mg/kg)	Mn (mg/kg)	Ni (µg/kg)	Zn (mg/kg)
Hazelnut	Total*	15.98 ± 0.47	1436.52±71.17	269.44 ± 9.71	13.71±0.41	25.95 ± 1.02	1276.00 ± 18.65	53.47±1.73	1497.25±57.75	16.02 ± 0.15
	Gastric	1.29 ± 0.28	116.28±9.49	22.40±1.12	0.78 ± 0.08	1.76±0.31	172.50±12.44	5.98 ± 0.46	179.54 ± 8.17	3.31±0.15
	Intestinal	1.57 ± 0.15	121.15±5.99	35.01±0.78	1.27±0.09	3.05±0.23	187.76±6.25	6.02 ± 0.07	211.69±22.16	5.40±0.37
	Gastric phase (%)	8.1	8.1	8.3	5.7	6.8	13.5	11.2	12.0	20.6
	Intestinal phase (%)	9.8	8.4	13.0	9.3	11.8	14.7	11.3	14.1	33.7
Almond	Total*	27.76 ± 4.04	2708.78±99.25	69.03 ± 8.82	11.65±0.83	20.57 ± 1.92	2057.19±135.34	21.13±0.94	1040.03±118.14	27.31 ±0.98
	Gastric	0.95±0.26	119.40±39.12	2.76±0.45	0.45±0.11	1.15±0.06	136.95±23.12	1.10±0.27	83.29±7.65	3.35±0.61
	Intestinal	1.40 ± 0.09	187.30±16.73	10.62±0.71	1.20 ± 0.10	3.03±0.19	258.63±18.22	2.09±0.14	238.61±91.07	6.31±0.56
	Gastric phase (%)	3.4	4.4	4.0	3.9	5.6	6.7	5.2	8.0	12.3
	Intestinal phase (%)	5.0	6.9	15.4	10.3	14.7	12.6	9.9	22.9	23.1
Pistachio	Total*	10.82±2.12	1792.70±11.26	10.66 ± 1.00	10.24±0.30	24.92 ± 3.87	1109.24±22.99	9.11±0.23	1244.85±41.99	16.81 ± 0.65
	Gastric	0.50±0.30	159.48±8.57	1.94±0.11	0.51±0.11	0.80±0.01	117.81±7.41	0.78±0.01	114.73±11.49	3.05±0.20
	Intestinal	0.90±0.19	212.21±8.50	3.21±0.41	1.07 ± 0.18	1.14±0.06	174.39±6.63	1.16±0.02	181.32±5.76	6.07±0.19
	Gastric phase (%)	4.6	8.9	18.2	5.0	3.2	10.6	8.6	9.2	18.2
	Intestinal phase (%)	8.3	11.8	30.1	10.4	4.6	15.7	12.7	14.6	36.1
Peanut	Total*	19.65±1.43	676.48 ±2.62	52.00 ± 7.62	7.00 ± 0.26	13.72 ± 0.58	1549.15±29.89	16.38±0.93	1597.57±145.99	22.15 ± 1.40
	Gastric	0.53±0.12	31.26±3.18	2.54±0.56	0.26±0.10	0.74±0.04	79.22±6.56	0.54±0.10	130.49±38.86	2.18±0.15
	Intestinal	1.78 ± 0.70	66.32±5.54	9.96±1.35	0.78 ± 0.04	2.69±0.16	164.93±10.91	1.59±0.10	253.41±26.56	5.62±0.17
	Gastric phase (%)	2.7	4.6	4.9	3.7	5.4	5.1	3.3	8.2	9.8
	Intestinal phase (%)	9.0	9.8	19.2	11.1	19.6	10.6	9.7	15.9	25.4
Cashew	Total*	9.06±0.30	386.28 ±9.77	52.56 ± 6.38	15.96±0.23	53.04 ± 3.22	2025.25 ± 38.98	12.39±0.31	3812.37±102.44	39.12 ± 0.64
	Gastric	0.37±0.13	25.31±2.10	4.10±0.44	0.43±0.05	1.06±0.12	127.05±13.42	0.68 ± 0.07	283.61±21.77	3.48±0.31
	Intestinal	0.53±0.04	39.70±1.74	10.78±1.25	1.29±0.10	5.60±0.65	227.42±1.66	1.30±0.03	568.29±38.51	6.45±0.09
	Gastric phase (%)	4.0	6.6	7.8	2.7	2.0	6.3	5.5	7.4	8.9
	Intestinal phase (%)	5.9	10.3	20.5	8.1	10.6	11.2	10.5	14.9	16.5
Brazil	Total*	9.79±0.29	1315.27±17.41	926.96±14.37	18.69±0.73	21.29 ± 1.36	3323.20±24.59	11.98±0.16	5126.52±39.60	32.06 ± 1.15
nut	Gastric	0.45 ± 0.11	90.54±1.34	69.18±7.45	0.71±0.02	0.85±0.02	284.94 ± 6.45	0.89 ± 0.02	503.74±19.43	3.62 ± 0.08
	Intestinal	0.85 ± 0.06	62.89±1.31	113.79±13.58	1.86±0.33	3.61±0.33	345.23±19.31	1.16 ± 0.12	673.09±25.37	6.10±0.69
	Gastric phase (%)	4.6	6.9	7.5	3.8	4.0	8.6	7.4	9.8	11.3
	Intestinal phase %)	8.7	4.8	12.3	10.0	17.0	10.4	9.7	13.1	19.0
Walnut	Total*	15.5 ± 1.35	793.04±66.92	29.11 ± 3.44	11.73±0.56	22.08±1.01	1034.96 ± 61.66	24.23±1.62	718.66±80.32	23.25 ± 0.50
	Gastric	1.75 ± 0.52	92.01±4.05	4.33±0.19	1.05 ± 0.15	0.34±0.09	143.15±8.55	3.18±0.24	107.91±16.05	4.51±0.13
	Intestinal	1.37±0.07	93.10±5.40	7.57±0.58	1.08 ± 0.18	1.32±0.13	139.20±7.82	2.12±0.03	145.38±49.15	4.06±0.20
	Gastric phase (%)	11.3	11.6	14.9	9.0	1.5	13.8	13.1	15.0	19.4
	Intestinal phase %)	8.8	11.7	26.0	9.2	6.0	13.4	8.8	20.2	17.5

Table 1. Continued

		B (mg/kg)	Ca (mg/kg)	Co (µg/kg)	Cu (mg/kg)	Fe (mg/kg)	Mg (mg/kg)	Mn (mg/kg)	Ni (µg/kg)	Zn (mg/kg)
Chick pea	Total*	7.26±0.76	515.49±12.25	102.36 ± 5.63	7.42 ± 0.12	28.96 ± 3.22	805.50±14.91	14.35±0.36	1904.52±57.93	19.99 ±0.35
	Gastric	0.71±0.28	60.14±7.25	11.50 ± 1.70	0.35±0.03	1.02±0.10	103.47±13.97	1.53±0.22	177.16±30.20	3.64±0.34
	Intestinal	0.37 ± 0.08	73.05±1.70	17.65±1.21	0.71±0.11	4.23±0.27	116.08±3.65	1.72±0.07	283.78±4.74	5.29±0.23
	Gastric phase (%)	9.8	11.7	11.2	4.8	3.5	12.8	10.7	9.3	18.2
	Intestinal phase (%)	5.1	14.2	17.2	9.6	14.6	14.4	12.0	14.9	26.5
Pumpkin	Total*	12.41 ± 1.91	314.69±9.89	116.86±10.91	10.65±0.31	61.30 ± 2.34	4111.59±154.92	42.66±1.03	1929.99±125.82	59.16 ±2.31
seed	Gastric	0.25 ± 0.07	16.01±1.06	2.09±0.24	0.15±0.02	0.66±0.09	105.09±13.56	0.79 ± 0.10	91.49±8.77	2.15±0.17
	Intestinal	0.72 ± 0.12	36.21±2.30	16.13±1.33	0.80 ± 0.15	4.00±0.31	276.25±21.07	2.23±0.14	312.51±11.37	6.79±0.08
	Gastric phase (%)	2.0	5.1	1.8	1.4	1.1	2.6	1.8	4.7	3.6
	Intestinal phase (%)	5.8	11.5	13.8	7.5	6.5	6.7	5.2	16.2	11.5
Sunflower	Total*	20.28 ± 2.60	900.72 ±1.58	92.70 ± 8.19	18.20±0.75	45.85 ± 2.23	2826.43 ± 58.72	25.11±0.10	5535.38±85.34	45.82 ± 0.89
seed	Gastric	1.12 ± 0.77	50.41±8.34	6.19±1.76	0.45 ± 0.02	1.30±0.27	138.33±13.00	0.10 ± 0.03	342.49 ± 83.87	3.40±0.37
	Intestinal	1.61±0.39	98.07±2.10	12.50±0.55	1.89 ± 0.07	5.34±0.09	326.87±2.07	2.28±0.09	701.99±44.52	7.87±0.44
	Gastric phase (%)	5.5	5.6	6.7	2.5	2.8	4.9	0.4	6.2	7.4
	Intestinal phase (%)	7.9	10.9	13.5	10.4	11.6	11.6	9.1	12.7	17.2

*These data were obtained from Kafaoğlu et al. 2014.

	В	Ca	Со	Cu	Fe		Mg		Mn		Ni	Zn
Hazelnut	15.7	1211	0.35	12.7	30.5		1877		60.2		2.12	54.00
Almond	14.0	1873	0.11	12.0	30.3		2586		20.9		2.39	63.10
Pistachio	8.97	2122	0.03	10.7	11.4		1743		11.6		1.81	60.72
Peanut	17.8	663	0.10	7.76	26.9		1649		15.9		2.53	56.24
Cashew	5.33	396	0.11	12.8	55.9		2274		12.9		5.68	64.47
Brazil nuts	8.52	628	1.14	18.6	36.1		3452		11.6		6.73	60.97
Walnut	13.7	930	0.08	10.8	13.2		1392		21.2		1.45	40.61
Chickpea	3.72	730	0.18	7.08	42.3		1160		17.2		2.84	52.93
Pumpkin seed	7.21	362	0.16	7.96	40.0		2762		22.3		3.13	67.86
Sunflower seed	16.1	980	0.13	18.9	53.4		3268		22.8		7.02	78.73
Recommended Dietary	**1.52	*1000	*2.4 µg/day as	*900	*8	mg/	*420 mg/	day	*2.3	mg/	**150	*11 mg/ day for
Allowances and	mg/day	mg/day	Vitamin B12	μg/	day	for	for male 2	320	day	for	µg/day	male 8 mg/day for
Adequate Intakes				day	male	18	mg/day	for	male	1.8		female
					mg/day	7	female		mg/day	for		
					for fem	ale			female			
Tolerable Upper Intake	20	2500	ND	10	45		350		11		1	40
Levels* (mg/day)												

Table 2 Amounts (μ g/day) of metal ingested from the assumption of a nuts or seeds ingestion rate of 10 g day⁻¹ for different nuts and seed samples calculated from values taken by in-vitro intestinal bioavailability results.

*Dietary Reference Intakes (DRIs) were given as Recommended Dietary Allowances and Adequate Intakes by USDA, (a) and Tolerable Upper Intake Levels by USDA, (b).

** These values are given by World Health Organization Geneva 1996 as average daily intakes (WHO, 1996).

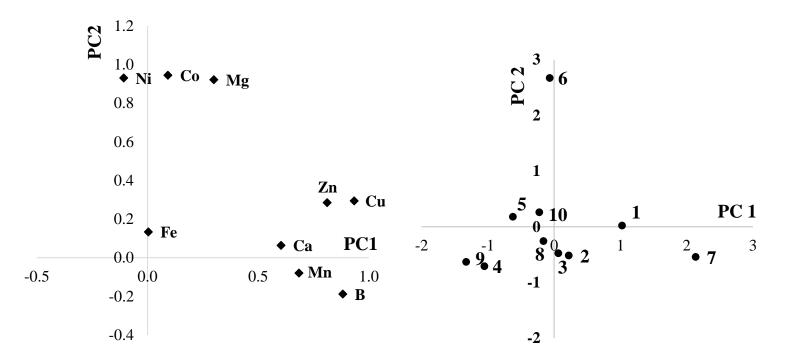
ND = Not determinable due to lack of data of adverse effects in this age group and concern with regard to lack of ability to handle excess amounts. Source of intake should be from food only to prevent high levels of intake (USDA (b)).

The Load	ings			The scores							
Element	PC1	PC2	PC3	Nut or Seed	PC1	PC2	PC3				
В	0.88	-0.19	0.08	Hazelnut	1.03	0.02	2.44				
Ca	0.60	0.06	0.25	Almond	0.22	-0.52	0.36				
Co	0.09	0.94	0.09	Pistachio	0.06	-0.48	-0.17				
Cu	0.93	0.29	-0.05	Peanut	-1.05	-0.71	-0.25				
Fe	0.00	0.13	0.93	Cashew	-0.62	0.18	-0.21				
Mg	0.30	0.92	0.08	Brazil Nut	-0.06	2.67	-0.45				
Mn	0.68	-0.08	0.56	Walnut	2.14	-0.54	-1.57				
Ni	-0.11	0.93	-0.01	Chickpeas	-0.16	-0.26	-0.02				
Zn	0.81	0.28	-0.22	Pumpkin Seed	-1.33	-0.63	-0.31				
				Sunflower Seed	-0.22	0.26	0.17				

Table 3. The loadings and the scores of the three rotated principal components for gastric phase

The Load	ings				The scores						
Element	PC1	PC2	PC3	PC4	Nut or Seed	PC1	PC2	PC3	PC4		
В	0.10	-0.01	0.41	0.75	Hazelnut	-0.62	0.32	-0.27	2.32		
Ca	-0.03	-0.15	0.91	0.09	Almond	0.46	-0.38	1.30	0.12		
Co	0.05	0.95	-0.16	-0.06	Pistachio	-0.23	-0.45	1.91	-1.00		
Cu	0.65	0.65	0.13	0.17	Peanut	-0.49	-0.54	-0.05	0.34		
Fe	0.67	-0.05	-0.70	-0.05	Cashew	0.62	-0.45	-1.23	-0.77		
Mg	0.80	0.49	-0.02	0.00	Brazil Nut	0.24	2.69	-0.14	-0.71		
Mn	-0.15	-0.01	-0.11	0.90	Walnut	-1.47	0.20	0.35	0.19		
Ni	0.77	0.45	-0.35	-0.17	Chickpeas	-0.90	-0.58	-1.05	-0.90		
Zn	0.96	-0.10	-0.03	-0.07	Pumpkin Seed	0.23	-0.62	-0.93	-0.29		
					Sunflower Seed	2.16	-0.20	0.11	0.70		

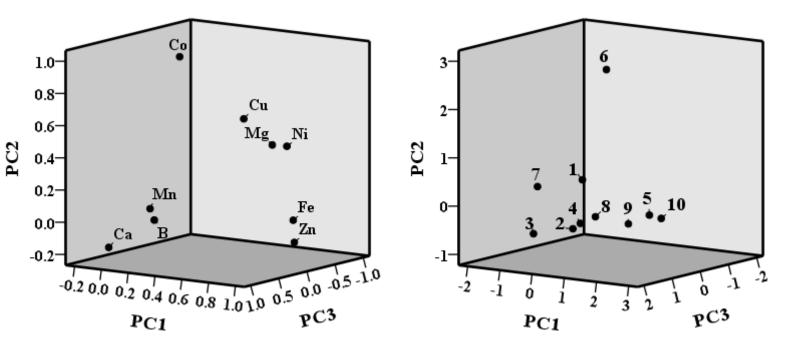
Table 4. The loadings and the scores of the three rotated principal components for intestinal phase



b) Score graph (1.hazelnuts, 2. almonds, 3. pistachio nuts, 4. peanuts, 5. cashew nuts, 6. Brazil nuts, 7. walnuts, 8. chickpeas, 9. pumpkin seeds, 10. sunflower seeds)

a) Loading graph

Figure 1. Two way Principal component analysis loadings and score plot for gastric ingestions



a) Loading graph

b) Score graph (1.hazelnuts, 2. almonds, 3. pistachio nuts,4. peanuts, 5. cashew nuts, 6. Brazil nuts, 7. Walnuts 8.chickpeas, 9. pumpkin seeds, 10. sunflower seeds)

Figure 2. Three way Principal component analysis loadings and score plot for intestinal ingestions