The medicinal value and the numerous sources of vitamin c - a review

Abstract

Vitamin C is one of the nine water soluble vitamins which approximate a carbohydrate in structure. The transport takes place via glucose transporters and the absorption eventuates in the gastrointestinal tract. It has a myriad of biological roles in the body and due to which it has abundant medicinal roles. These medicinal roles of Vitamin C can be subdivided as prevented and curative. The systems which are benefitted by Vitamin C include cardiovascular system, nervous system, musculoskeletal system, gastrointestinal system to name some. The review embraces various medicinal uses and sources along with the effect of food processing on the concentration of Vitamin C.

Keywords: ascorbic acid, vitamin, cancer, cardiovascular disease, diabetes, fruits, vegetables

Abbreviations: VC, vitamin c; L-AA, l-ascorbic acid; DHA, l-dehydroascorbic acid; GLO, l-gulonolactone oxidase gene; LDL, low density lipoprotein; HDL, high density Lipoprotein; DV, dentate gyrus; FG, fasting; PBG, post meal blood glucose; GHb, glycosylated hemoglobin; AD, alzheimer’s disease

Introduction

Polish biochemist Kazimierz Funk while working on experimental beriberi, gave the name “Vitaimine” from the words vital and amine meaning amine of life but then later the word “Vitamin” was acceptable in 1920 under the standard scheme of nomenclature adopted by the Chemical Society. McCollum classified vitamins as either water-soluble or fat-soluble, the total numbers of vitamins in humans are 13, enlisting 4 fat soluble vitamins i.e. A, D, E, K and 9 water soluble vitamins i.e. B complex and C. Vitamin C (VC), a water soluble vitamin has an oblong crystalline structure which is white in colour. Its general formula is C6H6O5 and its structural formula resembles a carbohydrate. L-ascorbic acid (L-AA) is the main biologically active form of VC, reversible oxidation of L-AA results in formation of L-dehydroascorbic acid (DHA) which also has biological activity.

Human beings are unable to synthesize VC on their own due to mutation of the l-gulonolactone oxidase gene (GLO). The last step in the synthesis of VC is catalyzed by GLO and inability of humans to synthesize VC is due to GLO deficiency in them. Humans and rats (the species that are able to synthesize) have homologous GLO genes but in man the gene has been mutated several times making the gene inactive. On intake of VC, its level in a healthy individual varies between 0.7-1.5mg/100ml of plasma. Concentration of VC in blood cells is much higher than in plasma, i.e. 3-4times higher in red blood cells and 20-30 times higher in white blood cells. Adrenals contain the highest concentration of VC amongst organs and the concentration of Vitamin C decreases with age, alcoholism and with intake of contraceptive pills.

Metabolism

The two major forms of VC available in diet are L-AA, which is, as mentioned physiologically reducing agent and DHA, which is the oxidized form. DHA is absorbed by facilitated transport, which takes place with the help of glucose transporters particularly GLUT1, GLUT3 and GLUT9. The transport of L-AA is mediated by sodium dependent SVCT1 and SVCT2 and the absorption takes place in the gastrointestinal tract. A large number of factors have been observed to affect the absorption, these enlist route of ingestion, quantity of the vitamin ingested, and prior nutritional status of the individual. The metabolites of vitamin C in humans are oxalate, dehydro-ascorbic acid, 2,3-diketogulonic acid, ascorbic acid 2-sulphate and saccharo-ascorbic acid. Maximum concentration of these compounds have been isolated and identified in urine, whereas the average faecal excretion of VC or its metabolites was observed to be about 3% when administered orally in physiological amounts.

Medicinal role

Intake of VC has been associated with the prevention and treatment of an ample number of diseases due to its various biological properties. Its medicinal uses can be assorted as disease prevention and disease treatment.

Disease prevention

Cancer: VC intake has been associated with the prevention of a number of cancers, many clinical trials have reported in which consumption of VC helped in reducing the occurrence of tumours in oral cavity, larynx, oesophagus, lungs, pancreas, stomach, colorectal, bladder and brain. McLaughlin et al., studied VC consumption and incidence of oral cancer and the results presented that the reduced intake of vitamin C was associated with increased risk for cancer. Graham et al., carried out a study to observe effect of VC on cancer of larynx by carrying out a survey and reported that lower ingestion of the vitamin was associated with higher incidence of the cancer of larynx. Cancer of the oesophagus was also studied by a number of investigators and the studies have presented that VC or fruit intake has a strong inverse relationship with incidence of cancer of oesophagus independent of all factors.

Fontham et al., examined approximately 2500 subjects including lung cancer patients and controls. On adjusting with other factors it was observed that higher intake of VC was related to decrease in risk
for the disease. Kromhout et al. analysed the results of a prospective study and confirmed that incidence of lung cancer and intake of vitamin C has a strong inverse association. In a study Falk et al., investigated cases of pancreatic cancer and matched hospital controls. They carried out a survey with regards to the diet of the individuals and it was observed that those who consumed less than 70mg vitamin C/day had a relative risk of 2.6 for males, or 1.8 for females compared with those who consumed 59mg/d. Another study by Mills et al. stated that frequent consumption was significantly protective against pancreatic cancer. Along with these literature also suggests preventive role of VC against carcinomas of stomach, cervix, colon and rectum.

N-Nitroso compounds including dialkyl-N-nitroso-mines, N-alkyl-N-nitrosamides, and related compounds are an important group of environmental carcinogens. VC decreases formation of nitrosamine formation whereas lipid-soluble derivatives of VC are also very efficient in reducing the nitrosamine content of foods. VC induces apoptosis in the cells through the disruption of mitochondrial membrane potential and by suppressing the translocation of transferring receptor from cytosol to membrane. It also attenuates the proliferation of cancer cells via prevention of growth of these cells. It arrests the growth at G1 stage which has approximate with the modulation of the activity of p53-p21Waf1/Cip1 and CDK2.

**Cardiac diseases:** Oxidative modification of low density lipoprotein (LDL) increases atherogenicity as it is taken up more quickly by macrophages, monocytes and smooth muscle cells beneath the arterial endothelium and results in increased accumulation of cholesterol ester leading to formation of foam cells and the initiation of the lesion of atherosclerosis. Other methods by which it causes atherosclerosis is by retaining monocyte-macrophages in the arterial wall and by being cytotoxic towards endothelial cells. VC has protective role against oxidation of LDL, it even preserves the endogenous antioxidants in LDL. It has been observed that VC along with DHA prevent the initiation of lipid peroxidation in LDL and also preserved the LDL-associated antioxidants like α-tocopherol, β-carotene, and lycopene, intact VC works synergistically with vitamin E, ubiquinol-10, β-carotene in LDL to suppress the peroxidation.

VC increases level of high density lipoprotein (HDL) and this association was stronger in older people and in males than in younger people and in females. In patients suffering from elevated total cholesterol levels if VC is administered in high dose, there is reduction in total cholesterol levels. Vasodilation is compromised in patients with atherosclerosis and this may lead to myocardial infarction and stroke. Literature suggests that administration of VC results in improved vasodilation in patients of coronary heart disease, angina pectoris, congestive heart failure, diabetes, high cholesterol, and high blood pressure.

**Lead Toxicity:** Lead increases the level of lipid peroxidation and brain thiobarbituric acid-reactive substances, decreases the capacity of antioxidant defence system and even decreases blood haemoglobin, glutathione peroxidise, superoxide dismutase levels and liver glutathione levels. Various clinical trials have concluded that on administration of VC alone or in combination with other agents there is decrease in lead concentration and there is lesser tissue damage which is otherwise caused by increased lead levels in the body. In a study vitamin C and vitamin B were administered to counteract the effect of high lead concentration and it was concluded that both the vitamins reduced the damage to liver cells from oxidative damage caused by lead, but their effect was dependent on their concentration. Ebuehi et al. administered vitamin C and vitamin E to evaluate their effect in attenuating hepatotoxicity and oxidative stress caused by lead and the study presented that administration of the vitamins reduced the toxic effects induced by lead. As mentioned lead intake decreases blood haemoglobin, in a study VC was given along with iron to examine their effect on anaemia caused by lead. Their intake prevented the growth depression and anaemia caused and reduced tissue lead contents, specifically in bone in which most of the accumulated lead was observed to be present. Goyer and Cherian claimed that chelating effect of VC was equivalent to that of EDTA when examined in relation to lead toxicity.

Lead causes alterations in hematopoietic system and drug metabolizing enzymes, keeping this in view Vij et al. examined the effect of VC in restoring normal physiology in male rats. They concluded that administering VC restored blood delta aminolevulinic acid dehydratase, uroporphyrinogen I synthetase and a few drug metabolizing enzymes and a significant reduction in lead concentration of blood and liver. In another study reduction in neurotoxic effects including neuronal damage and apoptotic cell death was investigated in rats. The rats were given 100mg/kg/day of VC and histopathological evaluation presented that VC reduced apoptosis in the developing hippocampus and also spares hippocampal CA1, CA3 and dentate gyrus (DG) neuron and blood levels of lead.

**Bone:** VC is a cofactor in the hydroxylation of proline and lysine. The intracellular level of VC in osteoblasts is regulated by sodium dependent transporter proteins present in the plasma membrane. VC reduces the iron prosthetic group of the hydroxylases and thus appears to play a pivotal role in maintenance of bone health. Its lower intake has been related to reduced bone mass and increase in rate of bone loss, increase in frequency of fractures and higher occurrence of osteoporosis. It is suggested that VC enhances bone health and its low level plays a significant role in development of osteoporosis. VC stimulates assembling of fibril and also the proliferation of various cell types. It plays major role in both proliferation of osteoblasts and even in their differentiation to generate bone matrix proteins. In a study effect of vitamin C concentration on osteoblasts was examined and the results presented that the proliferation of these bone forming cells increased with increase in VC and the highest proliferation was observed at level of 200µg/ml but at the concentration of 300µg/ml and above VC appeared to be toxic. It is also observed that osteoblasts increase the expression of osteonectin and osteocalcin on adding vitamin C.

A population based cohort study by Sahni et al. conducted a study to evaluate the role of VC on bone mineral density and a positive correlation was observed between the two. Rodriguez et al. described a case of regional transient osteoporosis of the foot. When the patient was evaluated radiologically severe osteopenia in the feet and bone marrow oedema was observed. The authors concluded that these finding were related to severe vitamin C deficiency. Lynch et al. observed an unusual form of osteoporosis in Johannesburg subjects and noticed a significant correlation with vitamin C deficiency.

**Ocular Tissue:** VC is present in the eye structure of many species and its concentration is higher than in other tissues. The level of VC in eye is highest in aqueous humor and lowest in retina and it has been generally considered that aqueous humor serves as the source of VC in the ocular tissues. The considered primary role of VC in the ocular tissues is to provide protection against oxidative damage and precisely the damage induced by light. Reiss et al. stated that the high concentration of ascorbate in the eye might be an adaptation of the ocular tissue to protect itself from solar radiation. VC prevents the...
riboflavin-mediated, light-induced damage to the cation pump in the lens.\textsuperscript{90,91} and even lowers the photoperoxidation of the membranes.\textsuperscript{51}

Cataracts are a leading cause of visual impairment throughout the world. Consumption of VC has been observed to show an inverse relationship with occurrence of cataract. Few studies have concluded that increased dietary vitamin C intake\textsuperscript{2,23} and increased blood levels of vitamin C\textsuperscript{64} to be associated with decreased risk of cataracts. VC has protective effect on all three types of cataracts including cortical, nuclear, and posterior subcapsular but most on posterior sub-capsular type.\textsuperscript{52,56}

**Gout:** VC supplementation is observed to provide protective effects against gout as it reduces hyperuricemia which is a precursor of gout. The protective effect of vitamin C against hyperuricemia is due to its competition with uric acid for renal reabsorption via an anion-exchange transport system in the proximal tubules.\textsuperscript{72} Enomoto et al.,\textsuperscript{24} in 2002 stated that the other mechanism behind it might be through inhibition of urate transporter 1 which is the main target for uricosurics or inhibition of Na-dependent anion co-transporter or both in the proximal tubule. Huang et al.,\textsuperscript{28} conducted a clinical trial and concluded that on administration of 500mg/day there was significant increase in glomerular filtration rate hence giving another mechanism behind uricosuric effect of VC. A study in USA stated that after supplementation with VC there was a decrease of 45% in level of uric acid with intake of 1500mg or more and about 34% lower risk for vitamin C intake between 1000 and 1499mg/day compared to no intake of vitamin C and the results obtained were independent of diet body mass index, age, hypertension, diuretic use, alcohol, and chronic renal failure.\textsuperscript{40}

**Hypervitaminosis A:** Rodahl observed that detrimental effects of hypervitaminosis A were increased in animals suffering from deficiency of vitamin C than in non-scorbutic animals.\textsuperscript{61} Wendt and Schroeder stated that administration of VC had favourable action on toxic effects of hypervitaminosis A and delayed deposition of the vitamin in the liver.\textsuperscript{52}

**Treatment**

**Diabetes:** It has been reported that leukocyte VC levels of diabetic patients is low as compared to non-diabetic on giving both kind of subjects same amount of the vitamin and it reduces the level of advanced glycation end products which are formed by irreversible glycation of proteins in oxidative conditions.\textsuperscript{47} In a study by Osganian et al.,\textsuperscript{44} intake of VC in the dose of 400mg/day or more was observed to reduce the fatal and nonfatal coronary heart disease in diabetics. Levy et al.,\textsuperscript{65} studied intake of vitamins and their impact on the progression of coronary artery atherosclerosis and the results presented that vitamin supplementation was related with improvement in coronary atherosclerosis in diabetic females having two copies of haptoglobin 1 gene whereas the condition worsened in diabetic females having two copies of haptoglobin 2 gene thus claiming that genetic factors of the individual also play a primary role. In an experiment effect of VC on diabetes induced endothelial dysfunction and generation of reactive oxygen species was examined. The results showed that reduced VC was associated with increased dihydrothromadine 123, which is an indication of increase in microvascular oxidative stress. Endothelial dysfunction was observed to be reduced or completely eliminated on administration of VC.\textsuperscript{66}

In a study effect of vitamin C along with metformin on fasting (FG) and post meal blood glucose (PGB) and glycosylated hemoglobin (GHB) in patients with type 2 diabetes was evaluated. It was observed that there was an identifiable decrease in FG and PBG with increase in VC levels. The proposed dose of VC was 1,000mg/day which was found to be simple, safe, and an effective means of preventing and decreasing chronic complications of diabetes. The proposed mechanism was its antioxidant capacity by providing protection from the oxygen species and even to β cells from getting damaged, other reason for beneficial effect of VC in diabetes is positive action of VC on vitamin E.\textsuperscript{67} Literature suggests that diabetics have increased occurrence of gastric ulceration, in a study effect of VC on this manifestation of diabetes was examined. The study concluded that VC shows protective effects on mucosa of stomach from formation of peptic ulcers and even increase secretion of gastric acid without disturbing the cytoarchitecture of the gastric mucosa.\textsuperscript{68}

**Cognitive impairment:** Oxidative damage to neuronal cell membranes and mitochondrial DNA has been proposed to cause Alzheimer’s disease (AD) whereas oxidation of LDL enhances the risk of vascular dementia.\textsuperscript{69} Due to this reason VC therapy has also been proposed in cognitive impairment. VC acts as a scavenger of free radicals in the brain and lower levels of this vitamin has been related to occurrence of AD. Charlton et al.,\textsuperscript{70} evaluated levels of VC in patients with dementia and AD. It was observed that patients suffering from AD or senile dementia had lower plasma VC levels than control subjects of same age group. It has also been stated that even in a healthy elderly, a higher level of plasma vitamin C was associated with better cognitive functioning. The authors suggested that adequate dose of VC to play its cognitive function would be 200mg/day. Perrig et al.,\textsuperscript{71} followed 446 Swiss subjects for 22years and concluded that high serum level of AA was associated with superior performance on tests of memory. Several cross sectional and case-control studies of cognitive impairment have observed that cognitive impairment is associated with lower levels of serum vitamin C.\textsuperscript{72,73}

**Pulmonary function:** VC has been shown to improve pulmonary function, Bucca et al. administered VC to patients suffering from bronchial reactive conditions caused by infections, allergy or exposure to air pollution or cigarette smoke. It was observed that VC prevented the increase in bronchial activity induced by foreign particle, due to allergic reaction or infection. Pathophysiologically, inflammation in the respiratory tissue attracts a number of neutrophil to the lung and these activated neutrophils generate oxygen free radicals. These species interact with α-1-protease thus leading to inhibition of proteolysis. VC protects α-1-protease from free radicals as well as from cigarette smoke.\textsuperscript{74} Low concentration of the vitamin causes reduction in forced expiratory volume. It attenuates bronchoconstriction caused due to inhalation of nitrogen dioxide and ozone and even reduces bronchial reactivity to histamine and methacholine.\textsuperscript{75} VC has anti-bronchospastic action in subjects with exercise induced bronchospasm.\textsuperscript{76}

VC levels in asthmatics have been observed to be reduced and it has also been stated that VC has a protective effect on asthma.\textsuperscript{77} On giving VC to asthmatics the frequency and severity of attacks reduces.\textsuperscript{78} Its administration with standard anti-asthmatics improves leukocyte motility, reduces antistreptolysin O levels and reduces immunoglobulin-E levels and titres of antibodies to the respiratory viruses.\textsuperscript{79} In a study it was observed that vitamin C was significantly associated with a lower risk of prevalent asthma, with highest association with the subgroup of youngsters exposed to cigarette smoke.\textsuperscript{80} Brittle asthma is a rare form of severe asthma in which patient suffers from repeated life threatening attacks. There is...
increased risk of mortality and morbidity in patients suffering from brittle asthma. VC has also been observed to play a pivotal role in pathophysiology of brittle asthma as well. It was observed that the nutrient intake of vitamin C was lesser in patients with brittle asthma than in other cases.91

Cancer: Few studies carried out in 1970s and 1980s have reported that on administration of large dosage of VC result in increase in the survival time and improvement of the quality of life of patients in the last stage of cancer.92 Gottlieb93 stated that VC fights back the oxidative damage induced during chemotherapy and radiation therapy thus helps in inhibition of the growth of cancerous cells. Prasad et al. found that high doses of VC not only can protect normal cells during cancer treatment but also can help fight tumours. At the level of 5 mM it causes a decrease of 50% in survival of cancer cells and the processes like apoptosis, pyknosis and necrosis are dependent on intracellular ascorbate. The destruction of cancer cells depend on concentration of hydrogen peroxide formation which is generated by ascorbate in the extracellular medium.94

In a study An et al.,95 studied effect of VC during treatment of colon cancer. It was observed that co-administration of VC with a cancer drug cisplatin increases sensitivity of the cancer cells towards the drug and even modifies tumour suppressor gene p58, thus enhancing the treatment results and further prognosis. It also increases the tumoricidal action of dacarbazine, tamoxifen, doxorubicin and paclitaxel.96,97 When VC is administered along with vitamin K, the combination enhances the therapeutic effect of six different chemotherapy agents.98 Inflammation has a pivotal role in development of tumours as it affects tumour proliferation, angiogenesis, metastasis and causes resistance to therapy. The features of inflammation related to cancer development are leukocyte infiltration, cytokine build-up, tissue remodelling and angiogenesis. The infiltrated leukocyte includes cytokines such as IL1, IL6, TNFα, TGFβ, FGF, EGF and HGF21, as well as chemokines such as CCL2 and CXCL8. Administration of VC reduces inflammation, in a study it was indicated that on intake of VC there was reduction in production of IL-2, IL-6 and TNFα. When VC is given through IVC therapy it attenuates cancer cell production and proliferation.99

Common Cold: Usage of AA in treatment of common cold was first documented in 1930s.100 Since then a large number of clinical trials have been conducted to test the hypothesis. In a Cochrane systematic review it was stated that a clinical trial observed administration of 8g of VC, the symptoms had more “short colds” of lesser severity than a day with administration of 4g. The review also mentioned that with intake of 1g daily of VC the symptoms reduced in both adults as well as children.101 Constantini et al.,91 evaluated role of VC in treatment and prevention of upper respiratory tract infections in swimmers. They concluded that VC reduced the severity and duration of the infection in males but not in females. Maggini et al.,102 co-administered vitamin C along with zinc and claimed that their co-administration reduced the duration of rhinorrhea and the severity of symptoms.

Deficiency of vitamin C

Scurvy

Scurvy, disease caused due to deficiency of vitamin C, the disease has its initials in the late bronze age, its description was written in Ebers Papyrus (1500 B.C.) and was even mentioned by Socrates (469 B.C. – 399 B.C.), he mentioned scurvy as pain in legs, gangrene of gums and loss of teeth as a symptoms complex. Later in around 1497, the disease was observed in sea explorers and in the voyage of Vasco da Gama. The disease caused death of about 2/3rd of the crew members. In 1553 voyage of Jacques Cartier suffered from the same symptoms, many members died and many got cured after taking tea prepared from the bark and leaves of spruce tree. Few more sailors and voyages explained the occurrence of scurvy in the later years but scientific explanations was given by an English navy surgeon, James Lind in “Treatise on Scurvy” in 1755.4

The term ‘scurvy’ for the disease resulting from prolonged vitamin C deficiency had origins in ‘scorbutus’ (Latin), ‘scurbur’ (French), and ‘Skorbut’ (German). Scurvy was a common problem in the world’s navies and is estimated to have affected 2million sailors. In 1747, James Lind conducted a trial of six different treatments for 12 sailors with scurvy: only oranges and lemons were effective in treating scurvy. Scurvy also occurred on land, as many cases occurred with the ‘great potato famine’ in Ireland in 1845.51

The first reported manuscript available in Medline was in 1847 by Dr. T Shatter. He described three stages of this deficiency. In the first stage he stated that the appearance was characterized by the general and usual indications of debility, feelings of weakness and disinclination to exertion, nervousness, oppression of the breathing, with a feeling of faintness, attacks of chills, superficial pains of the limbs, facial, labial, glossal and gingival paleness, with small and soft pulse.

In the second stage, there is feeling of debility, nervousness is less intense, the respiration is slightly accelerated, hearing is mildly oppressed, and attacks of faintness occasionally supervene; the general body pain is increased, spongy and swollen gingiva which bleeds easily, halitosis, occurrence of petechial spots on the limbs, general bruising of the skin, the pulse is small, feeble, and slightly accelerated, occasional attacks of fever, in few cases joints are stiff, enlarged and painful, formation of nodes on the clavicle, sternum, and tibia.

In the third and last stage, he observed depressed breathing, a sinuous discharge on coughing, swollen and painful gingiva, fetid breath, discharge of body fluid, appearance of petechiae and tendency to bleeding from the gingiva and mucous surfaces of the vagina and rectum, formation of serous effusion on the neural membranes leading to confused state of mind.103

Sources

Plant derivatives are superior sources of vitamin C.95 Since humans cannot synthesise VC, their principle source of the vitamin is dietary fruit and vegetables.96 Hampl et al.,107 conducted a survey on diet of school children and concluded that children consuming higher fruits and vegetables had higher level of VC and were more healthy. In 2004 Hung et al.98 stated that higher consumption of fruits and vegetables was associated with a comparable decrease in major chronic disease risk and further supported the recommendation of consuming five or more servings of fruits and vegetables daily.99 The content of VC in fruits and vegetables varies in a wide range and it even varies from the condition of the fruit and vegetable, raw plant product has the highest content of the vitamin, there is decrease in the content of VC with increase in storage temperature and duration. VC is sensitive to chemical and enzymatic oxidation during the processing and cooking.5 Other factors that affect the content of VC are oxygen, heat and light (Tables 1–5).108–111
| Sr. no. | Source                                | Vitamin C content (mg/100gm) |
|---------|---------------------------------------|------------------------------|
| 1       | Pepper, hot chili, green              | 242                          |
| 2       | Pepper, sweet, yellow                 | 183                          |
| 3       | Peppers, hot chili, red               | 143                          |
| 4       | Drumstick pods                       | 141                          |
| 5       | Pokeberry shoots                      | 136                          |
| 6       | Parsley, fresh                       | 133                          |
| 7       | Mustard spinach, tender green         | 130                          |
| 8       | Kale                                  | 120                          |
| 9       | Peppers, jalepano                     | 118                          |
| 10      | Vine spinach (basella)                | 102                          |
| 11      | Taro, tahitian, raw                  | 96                           |
| 12      | Broccoli leaves and flower cluster    | 93                           |
| 13      | Pepper, Hungarian                    | 92                           |
| 14      | Cauliflower, green                   | 88                           |
| 15      | Bitter gourd, leafy tips             | 88                           |
| 16      | Brussels sprouts                     | 85                           |
| 17      | Lambs quarter                        | 80                           |
| 18      | Sesbania flower                      | 73                           |
| 19      | Mustard green                        | 70                           |
| 20      | Cress, garden                        | 69                           |

**Table 1** Highest sources of Vitamin C (Vegetables - Raw) (values are in mg/100gm)
### Table 2: Highest sources of Vitamin C (Fruits - Raw) (values are in mg/100gm)

| Sr. no. | Source                                         | Vitamin C content (mg/100g) |
|---------|------------------------------------------------|----------------------------|
| 1       | Camu camu (Justi et al., 2000)                 | 2800                       |
| 2       | Acerola                                         | 1667                       |
| 3       | Guava                                           | 228                        |
| 4       | Lemon                                           | 182                        |
| 5       | Currants, European black                        | 181                        |
| 6       | Kiwifruit                                       | 92                         |
| 7       | Longans                                         | 84                         |
| 8       | Litchi                                          | 71                         |
| 9       | Orange with peel                                | 71                         |
| 10      | Jujube                                          | 69                         |
| 11      | Persimmons, native                              | 66                         |
| 12      | Pummelno                                        | 61                         |
| 13      | Papaya                                          | 60                         |
| 14      | Strawberries                                    | 58                         |
| 15      | Abiyuch                                         | 54                         |
| 16      | Clementines                                     | 48                         |
| 17      | Pineapple                                       | 47                         |
| 18      | Kumquats                                        | 43                         |
| 19      | Grapefruit, pink and red, California and Arizona| 38                         |
| 20      | Carissa (natal plum)                            | 38                         |
Table 3 Vegetable sources of Vitamin C and effect of food processing on Vitamin C content (values are in mg/100gm)

| Sr. no. | Source (vegetable) | Raw   | Canned | Cooked | Frozen |
|---------|--------------------|-------|--------|--------|--------|
| 1       | Pepper, sweet, red | 127.7 | 46.5   | -      | 58.7   |
| 2       | Kale               | 120   | -      | 41     | 39.3   |
| 3       | Broccoli           | 89.2  | -      | 64.9   | 56.4   |
| 4       | Pepper, sweet, green | 80.4 | 46.5   | 74.4   | 58.7   |
| 5       | Mustard green      | 70    | -      | 25.3   | 25.3   |
| 6       | Peas, edible-podded | 60   | -      | 47.9   | 22     |
| 7       | Cabbage, red       | 57    | -      | 34.4   | -      |
| 8       | Cauliflower        | 48.2  | -      | 44.3   | 48.8   |
| 9       | Peas, green        | 40    | 7.8    | 14.2   | 18     |
| 10      | Kidney beans       | 38.7  | -      | 35.6   | -      |
| 11      | Spinach            | 28.1  | 13.5   | 9.8    | 5.5    |
| 12      | Lima beans         | 23.4  | -      | 16.3   | 12.4   |
| 13      | Okra (lady finger) | 23    | -      | 16.3   | 12.4   |
| 14      | Radish oriental    | 22    | -      | 15.1   | -      |
| 15      | Beans pinto        | 21.7  | 0.7    | 6.1    | -      |
| 16      | Turnips            | 21    | -      | 11.6   | 4.4    |

Table 4 Fruit sources of Vitamin C and effect of food processing on Vitamin C content (values are in mg/100gm)

| Sr. No. | Source   | Raw | Dried | Frozen | Canned |
|---------|----------|-----|-------|--------|--------|
| 1       | Lemon    | 182 | -     | 31.5   | -      |
| 2       | Longans  | 84  | 28    | -      | -      |
| 3       | Jujube   | 69  | 13    | -      | -      |
| 4       | Papaya   | 60.9| -     | -      | 3.5    |
| 5       | Strawberries | 58.8| -     | 41.2   | -      |
| 6       | Pineapple| 47.8| -     | 8      | 7.4    |
| 7       | Mangoes  | 36.4| -     | -      | 15.2   |
| 8       | Gooseberry| 27.7| -     | -      | 10     |
| 9       | Raspberries | 26.2| -     | 16.5   | 8.7    |
| 10      | Blackberry| 21  | -     | 3.1    | 2.8    |

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The medicinal value and the numerous sources of vitamin C - a review

Table 5 Fruit Juice and content of Vitamin C (values are in mg/100gm)

| Sr. no. | Source                  | Raw | Frozen | Canned |
|---------|-------------------------|-----|--------|--------|
| 1       | Orange juice            | 50  | 137.9  | 30.1   |
| 2       | Grapefruit juice        | 38  | 119.8  | 29.2   |
| 3       | Tangerine juice         | 31  | 85.1   | 22     |
| 4       | Lime juice              | 30  | -      | 6.4    |
| 5       | Passion fruit, purple, juice | 29.8 | -    | -      |
| 6       | Cranberry juice         | 9.8 | -      | -      |
| 7       | Pomegranate juice       | 0.1 | -      | -      |
| 8       | Pineapple juice         | -   | 42     | 10     |
| 9       | Blackberry juice        | -   | -      | 11.3   |
| 10      | Apple juice             | -   | 2.1    | 0.9    |

Excess dosage

The toxicity of VC is low but doses crossing upper intake level may cause diarrhoea, abdominal cramps, gastrointestinal disturbances, flatus and nausea. Other than these there is increase in formation renal stones and the mechanism behind it is conversion of VC to oxalate and the property of acidifying urinary. In a study it was observed that on intake of 1000mg of supplemental VC given twice daily increased the level of excretion of oxalate by upto 22%. In patients suffering from hereditary hemochromatosis it has been seen that excessive VC intake results in increased iron overload and tissue damage, no such results have been presented in normal individuals. Other effects of VC toxicity include reduced vitamin B12 and copper levels, erosion of dental enamel, and allergic responses (Table 6).

Table 6 Recommended daily allowance

| Sr. no. | Individual            | Recommended daily allowance (RDA) |
|---------|-----------------------|-----------------------------------|
|         | **Pediatric**         |                                   |
| 1       | Birth - 6months (M & F) | 40mg                             |
| 2       | Infants 6-12months (M & F) | 50mg                        |
| 3       | Children 1-3years (M & F) | 15mg                         |
| 4       | Children 4-8years (M & F) | 25mg                        |
| 5       | Children 9-13years (M & F) | 45mg                       |
| 6       | Adolescent female 14-18years | 65mg                     |
| 7       | Adolescent male 14-18years | 75mg                     |
| 8       | Pregnant female 14-18years | 80mg                     |
| 9       | Lactating female 14-18years | 115mg                   |
|         | **Adults**            |                                   |
| 10      | Male over 18years     | 90mg                             |
| 11      | Female over 18years   | 75mg                             |
| 12      | Pregnant female over 18years | 85mg                    |
| 13      | Lactating female over 18years | 120mg                  |

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Conclusion

From the data mentioned it can be observed that Vitamin C has a long list of treating as well as preventing systemic diseases. If taken on a regular basis can be highly sanative for the human body. As mentioned, human beings are unable to synthesize Vitamin C and require an external source to fulfill the recommended daily allowance and fruits and vegetables are its richest sources. We jotted down the sources of Vitamin C along with impact of food processing on the concentration of the vitamin in the plant source and recommend increased consumption of these plant sources.

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Conflict of interest

Author declares that there is no conflict of interest.

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