Environmental Toxins and Human Cancer

Kyaw Myint Oo Professor, Rector (Retired), University of Dagon Member of the Executive Committee, Myanmar Academy of Arts and Science Ministery of Education, Yangon

1. Introduction

Cancer is caused by changes to certain genes that alter the way our cells function. Some of these genetic changes occur naturally when DNA is replicated during the process of cell division. But others are the result of environmental exposures that damage DNA. One research study concludes that as many as 35% of all cancers are driven by Environmental toxins. A British Government White Paper concluded that across 4000 common compounds used in-home, two thirds were toxic, and one third were probably carcinogenic.

Many well-known scientists, public health officials and physicians have been sounding alarms about the links between environmental toxins and human cancer for years now. The incidence of certain cancers, particularly thyroid cancer and leukemia, may very well rise in the Japanese population most heavily exposed to radiation from the Fukushima accident. Environmental toxins pose potentially grave threats to our health, and accidents only compound these threats both locally and for people all over the world who breathe air, eat food, and drink water. People can avoid some cancer-causing exposures, such as tobacco smoke and the sun's rays. But others are harder to avoid, especially if they are in the air we breathe, the water we drink, the food we eat, or the materials we use to do our jobs.

Scientists are studying which exposures may cause or contribute to the development of cancer. Understanding which exposures are harmful, and where they are found, may help people to avoid them. The chemicals in our water, air, and food, the materials in our home, and non ionizing radiation present cancer risks. But that doesn't mean that we are defenseless. Researchers have identified several mechanisms by which most cancer-producing toxins disrupt our body's defense systems. Compelling evidence reveals how we can defend against these carcinogenic mechanisms.

2. Sources of Cancer-Inducing and Cancer-Promoting Toxins

While it is impossible to avoid all cancer-causing environmental toxins, it is important to be aware of some of the most prominent sources. Researchers have compiled a list of common toxins broken down by their environmental sources. As you'll see from this list, these can be found in sources we interact with on a daily basis, including our food, water, plastic, cell phones, and even sunlight (Table 1).

- Aflatoxins are toxic chemicals produced by *Aspergillus* fungi growing on grains and peanuts, particularly those stored improperly. Chronic exposure induces cancer by multiple mechanisms.
- **Polycyclic aromatic hydrocarbons** are chemical structures composed of carbon, hydrogen, and occasionally other atoms. They are products of fossil fuel combustion, particularly

petrochemicals, and are a major source of cancer-causing chemicals in polluted air. Air pollutants (polycyclic aromatic hydrocarbons, tobacco smoke).

- **Bisphenol A (BPA)** is one of the highest-volume toxic chemicals found worldwide. It is used in making all kinds of plastics and resins, including water bottles and food containers.
- Arsenic. People are exposed to elevated levels of inorganic arsenic through drinking contaminated water, using contaminated water in food preparation and irrigation of food crops, industrial processes, eating contaminated food and smoking tobacco. Long-term exposure to arsenic in drinking water can cause cancer in the skin, lungs, bladder and kidney. It can also cause other skin changes such as thickening and pigmentation.
- **PCBs** are fat-soluble substances to which everyone is exposed through ingesting animal fats, inhalation, or dermal contact. Exposure to PCBs is associated with an increased risk of certain cancers of the digestive tract, liver and skin. PCB exposure is also associated with reproductive deficiencies, such as reduced growth rates, retarded development, and certain neurological effects which may or may not persist beyond infancy.
- **Heavy metals** (including cadmium, arsenic, nickel, lead, and mercury) are naturally occurring components of the earth's crust. Human exposure results from mining, smelting, and petroleum manufacturing, all of which release heavy metals into the air, water, and soil.
- **Pesticides and herbicides**, especially those containing organic chemicals bonded to chlorine or bromine, are found in agricultural settings, where they make their way into the food chain. Sadly, even after the highly toxic dichlorodiphenyltrichloroethane (DDT) was banned, risks still abound, both because of persistent DDT in the environment and because newer compounds intended to replace DDT (such as methoxychlor) are turning out to have their own cancer-inducing properties.
- **Tobacco smoke** is a serious health hazard causing more than 41,000 deaths per year. It can cause or make worse a wide range of damaging health effects in children and adults, including lung cancer, respiratory infections and asthma. Tobacco smoke causes approximately 7,330 deaths from lung cancer and 33,950 deaths from heart disease each year.
- **Dioxins** are commonly found in foods of animal origin (meat, dairy, and fish, depending on the country of origin).
- **Phthalates** are chemicals used to soften plastics. They are found in a wide variety of products, including bottles, shampoo, cosmetics, lotions, nail polish, and deodorant. Pre-natal exposure to phthalates is associated with adverse genital development and can significantly reduce masculine behavior in boys. Women with high exposure to phthalates while pregnant report significantly more disruptive behavior in their children, while other research has found phthalate exposure can lead to thyroid dysfunction in adults.
- **Triclosan** is an antibacterial and antifungal agent found in some consumer products, including toothpaste, soaps, detergents, toys, and surgical cleaning treatments. Its efficacy as an antimicrobial agent, the risk of antimicrobial resistance, and its possible role in disrupted hormonal development remains controversial.
- **Heterocyclic amines** are chemicals that form when meat is cooked at high temperatures (e.g., grilled or broiled).
- **Cyprodinil** is a fungicide. Cypronidil causes eye and skin irritation, and it may cause an allergic skin reaction. The major target organs of cyprodinil are the liver and kidney.
- Ultraviolet radiation is a natural component of sunlight, but serves as a powerful source of many of the changes that lead to cancer.

• **Electromagnetic field radiation**, especially the kind produced by cellular phones and their transmitting stations, are only now emerging as potential environmental threats. Such radiation is associated with DNA damage, potentially leading to cancer.

This is by no means an exhaustive listing of cancer-related environmental toxins. Toxins are ubiquitous, particularly in our highly industrialized society. They are, therefore, nearly impossible to avoid, but as we have read, we know that we are not helpless. We can arm ourselves with knowledge about natural products capable of offsetting much of the increased cancer risk posed by environmental toxins.

Basic		
Mechanism	Toxins	
Increased	• Aflatoxin A1	
DNA damage	• All politicants (polycyclic aromatic hydrocarbons, tobacco smoke)	
	 Arsenic 	
	• Bisphenol A (BPA)	
	Cadmium chloride	
	Microwave radiation	
	Mobile phone radiation	
	Nickel salts	
	Pesticides	
	Ultraviolet light	
Activation by	Aflatoxin B1	
liver enzymes	• Dioxin	
	 Heterocyclic amines (from cooking meat) Nicotine 	
	 Polycyclic aromatic hydrocarbons 	
	(e.g., benzo[a]pyrene)	
Suppress	Dioxin	
immune	• Mercury	
surveillance	• Mycotoxins (e.g., aflatoxins,	
	fumonisins, and deoxynivalenol)	
	 Perfluorinated hydrocarbons 	
	Tobacco smoke	
	Ultraviolet light	

Table 1. Toxins that induce or promote cancers	
(International Agency for Research on Cancer, 201	1)

Endocrine	• BPA
disruption	• Cyprodinil (a fungicide related to
	polycyclic aromatic hydrocarbons)
	• Dioxin
	• Heavy metals (arsenic, cadmium, lead,
	mercury)
	• Methoxychlor
	• Phthalates
	• PCBs
	• Polycyclic aromatic hydrocarbons
	Triclosan
Loss of	Aflatoxins
apoptosis	• Heavy metals
1 1	Pesticides/herbicides
	• UV light
	č

3. How Toxins Produce Cancer and Powerful Cancer Prevention from Nutrients

The heavily industrialized nature of our modern world constantly exposes us to toxic, cancer inducing, and cancer promoting influences. Chemical toxin as well as various sources of radiation can initiate cancer through DNA damage and promote tumor development through mechanisms involving liver enzyme systems, suppressed immunity, disruption of your hormones, and hijacking the cells' normal death-inducing programs. You can't realistically evade all of the factors that can cause cancer in your life, but you can leverage modern scientific knowledge to your benefit. Many nutrients are available with known cancer-fighting benefits, including those that directly counteract the major processes by which toxic compounds and radiation promote malignancies.

Despite the vast number and diversity of cancer types, there are a relatively small number of events that typically occur in the progression from healthy cell to malignancy. Toxic environmental chemicals, electromagnetic fields, and ionizing radiation may initiate and/or promote malignancy, operating along a number of mechanisms. Important five mechanisms include:

- DNA damage,
- Liver detoxification impairment,
- Immune impairment,
- Endocrine disruptors, and
- Loss of apoptosis.

3.1. DNA Damage

The first way toxins may lead to cancer is by breaking DNA strands. Damage to DNA is a major initiating factor in cellular transformation to cancer. DNA damage can be caused by toxins that break

DNA strands (such as pro-oxidant chemicals or ionizing radiation). Such damage can induce mutations in the DNA that trigger cancer. That is why nutrients that prevent DNA damage, or ones that promote its **repair**, are so potent in protecting against cancer. Given the role of oxidative stress in causing such damage, **nutrients that reduce DNA damage** (i.e. **vitamins C** and **E** and the trace mineral **selenium**), are often considered as a first line of defense.

Surprisingly, **probiotics**, which are normally associated with improved gastrointestinal function, have been found to be effective at reducing DNA damage specifically in the colon. This may help to prevent **colon cancer**, the third cause of cancer-related deaths in the US. And, while sunlight exposure can boost vitamin D levels, such exposure also raises DNA skin damage, but **vitamin D supplementation** can protect against DNA damage throughout the body. Nearly a dozen nutrients have been found to prevent DNA damage, resulting in a positive impact on cancers of the prostate, colon, breast, skin, liver, and more.

Powerful Cancer Prevention from nutrients that reduce DNA damage. It is easy to become fearful about the vast numbers of toxic chemicals and other influences all around us, lurking to produce catastrophic cancers. Fortunately, there are solutions in the form of specific nutrients with powerful cancer-preventing effects. Compelling scientific studies show that specific nutrients counteract major processes by which toxic compounds and radiation promote malignancies. Table 2 lists a few of the nutrients known for their DNA-protective effect.

(International ingeney for Research on Suncer, 2011)		
Nutrient	Cancers Affected	
Carotenoids (lutein, astaxanthin, lycopene)	Prostate, colon	
Coenzyme Q10	Colon, head-and-neck	
Fish oil (omega-3 fats)	Colon, skin	
Genistein (from soy)	Leukemia	
Plant polyphenols	Multiple	
Probiotics (Lacto-bacillus rhamnosus)	Colon	
Quercetin	Head-and-neck	
Selenium	Colon, breast	
Vitamin C	Breast, colon, skin	
Vitamin D	Colon, skin	
Vitamin E	Liver, breast	

Table 2. Nutrients that reduce DNA damage (International Agency for Research on Cancer 2011)

3.2. Liver Detoxification Systems

The second way environmental toxins cause cancer is through their detrimental impact on liver detoxification systems. **Liver detoxification systems** play a major role in managing ingested toxins because blood from the digestive tract goes to the liver before being pumped around to the remainder of the body.

The liver has two major detoxification pathways: Phase I and Phase II. In **Phase I** enzymes convert toxic chemicals into compounds that may be more toxic than the parent compound. Unfortunately, if the toxic load is too heavy, it can cause overactivity of Phase I enzymes, which can

have the reverse effect of converting relatively harmless substances into potential DNA-damaging carcinogens. Making matters worse, the worst offenders of overactive Phase I enzymes are substances some people encounter on a daily basis, including alcohol, saturated fats, and exhaust fumes, among others.

In **Phase II** detoxification, the liver adds another substance to the toxic chemical in order to make it more water soluble. This allows your body to excrete the toxin through bile or urine, helping remove the potentially carcinogenic substance from the body. For these reasons, cancer-preventive nutrients that influence liver metabolism are generally those that *regulate* toxin-enhancing Phase I reactions, promote toxin-neutralizing Phase II reactions, or, in many cases, do both. Nutrients that regulate these liver detoxification systems come largely from dietary plants and their extracts.

Several nutrients have this dual action on liver enzymes, including curcumin, folic acid, and garlic, among others. Research suggests this may have a positive impact on preventing some of the most common and deadly cancers. List of nutrients that prevent cancer by regulating the liver's detoxification enzymes are shown in Table 3.

Nutrient	Impact on Liver	Cancers Affected
	Enzymes	
Chlorophyllin	Inhibit Phase I;	Liver, colon,
	boost Phase II	prostate
Curcumin	Inhibit Phase I; boost	Breast, colon,
	Phase II	prostate, pancreas
Folic acid	Inhibit Phase I;	Breast, pancreas
	boost Phase II	
Garlic	Inhibit Phase I;	Breast, liver,
	boost Phase II	prostate
Genistein	Boost Phase II	Colon
Isothiocyanates	Inhibit Phase I;	Liver, colon,
(sulforaphane, PEITC) from	boost Phase II	breast, prostate
cruciferous vegetables		
Plant flavonoids	Inhibit Phase I;	Multiple
(i.e. chrysin, genistein,	boost Phase II	
quercetin)		
Silymarin (milk thistle)	Boost Phase II	Liver

Table 3.	Nutrients that prevent cancer by regulating the liver's
	detoxification enzymes. (International Agency for Research
	on Concor 2011)

3.3. Immune Surveillance

The third way environmental toxins can cause cancer is through their impact on immune surveillance. **Immune surveillance** refers to the immune system's continual search for cells bearing signs that they have become cancerous. A number of environmental toxins can suppress immune surveillance, raising the risk that a malignant cell will slip under the radar, form a tumor, and successfully spread to other parts of the body. **Nutrients that enhance immune surveillance** are only

now being recognized as powerful contributors to the body's lifelong fight against cancer. These nutrients boost those components of the immune system that are responsible for recognizing the unique tumor "markers" displayed on the surface of malignant cells, and then destroying those cells.

Nutrient	Immune Mechanism	Cancers Affected	
Enzymatically modified	Increases natural killer cell	Leukemia, Multiple	
rice bran	activity	Myeloma, Liver	
Cistanche extracts	Increases naïve T-cells,	Colon	
	increased expression of		
	transforming growth factor		
	beta, decreases		
	inflammation		
Grape seed	Induction of	Skin	
proanthocyanidins	immunoregulatory		
	cytokines; stimulation of		
	tumor-destroying T cells		
Green tea polyphenols	Activation of tumor-killing	Colon, skin, lung,	
(EGCG)	T-lymphocytes and natural	prostate, breast	
	killer cells; induction of		
	immuno-regulatory		
	cytokines		
Probiotics (lactic acid	Decrease inflammation;	Colon, skin	
bacteria)	increase immunoregulatory		
	cytokines; increased		
	interferon-gamma		
	production		
Reishi mushroom	Enhanced proliferation of	Liver, lymphoma,	
(Ganoderma lucidum)	tumor-killing T-	lung	
	lymphocytes, antibody-		
	producing B-lymphocytes,		
	and natural killer cells		
Resveratrol	Sensitizes tumor cells to	Leukemia	
	killing by cytokine-induced		
	killer cells; enhances		
	cytokine-induced killer cell		
	activity		
Silymarin	Induction of	Skin	
	immunoregulatory		
	cytokines; stimulation of		
	tumor-destroying T cells		
Vitamin D	Reduced inflammation	Colon	

Table 4. Nutrients that boost immune surveillance.(International Agency for Research on Cancer, 2011)

Nutrients that enhance immune surveillance may stimulate growth and proliferation of **tumor-detecting lymphocytes**, promote a vigorous attack on tumor cells by so-called "**natural killer cells**," and/or stimulate **antibody production**, which aids in immobilization and destruction of malignant cells. Table 4 lists nutrients capable of activating one or more components of the immune system in order to destroy developing cancers.

3.4. Endocrine Disruptors

The fourth way environmental toxins can cause cancer is through their impact on endocrine disruptors. **Endocrine disruptors** are chemicals that interact with sex hormones and/or their receptors to promote cancer development. Not surprisingly, **nutrients that inhibit endocrine disruptors** show promise in preventing hormone-dependent cancers such as those of the breast, uterus, and prostate. Although scientists don't yet fully know how these nutrients work to inhibit endocrine disruptors, it may involve enhanced excretion or reduced absorption of toxins from the intestinal tract.

	Toxin Inhibited	Cancers
Nutrient		Affected
Chlorella pyrenoidosa	Dioxin (a polychlorinated	Breast, prostate
(More efficiently obtained	biphenyl [PCB]),	
as chlorophyllin)	perfluorinated compounds	
	(PFCs)	
Folic acid	Bisphenol A (BPA),	Breast, prostate
	phthalates	_
Genistein	BPA, phthalates	Breast, prostate
Probiotics (Lactobacillus,	BPA	Breast, prostate
Bifidobacterium)		
Vitamin C	Heavy metals (lead,	Liver, lung,
	copper, iron)	prostate
Vitamin E	Heavy metals (lead,	Liver, lung,
	copper, iron)	prostate

Table 5. Nutrients capable of inhibiting endocrine-disrupting pollutants(International Agency for Research on Cancer, 2011)

Table 5 lists nutrients capable of inhibiting endocrine-disrupting pollutants. In addition, there are also a number of **plant flavonoids** (i.e. chrysin, genistein, quercetin) that are effective against endocrine disrupting toxins. They appear to reduce the activity of estrogen-producing enzymes such as aromatase, thereby reducing overall sex hormone predominance and starving hormone-dependent tumors of their vital growth factors.

3.5. Loss of Apoptosis

Another way environmental toxins are associated with cancer is through inducing a loss of apoptosis, or programmed cell death. **Loss of apoptosis** refers to the "immortality" typical of cancer cells. Normal body cells are programmed to die off when appropriate. Cancer cells have lost this ability (often as a result of DNA damage), which allows them to reproduce essentially without limit. A number

of chemical toxins, particularly *aflatoxin*, a potent inducer of liver damage, can switch off the gene responsible for producing apoptosis, which results in cancer promotion.

Nutrients that restore cells' natural ability to die by apoptosis represent the final category in our listing of nutrients that help fight against cancers caused by environmental toxins. These nutrients typically act by modifying various signaling pathways. This means that they can activate genes that become suppressed when cells become cancerous, including genes that normally support the graceful death of a cell that is no longer useful or poses a threat. By restoring the natural self-destruction program initiated by apoptosis genes, these nutrients put a sharp roadblock in the way of a developing tumor. This allows other anticancer mechanisms such as immune surveillance to clear the remainder of the battlefield. Nutrients known to promote apoptosis include coffee extract, quercetin, pine bark extract, and selenium. Research shows they have a positive impact on bladder, colon, and ovarian cancers, among others. Detailed list of nutrients that promote or restore apoptosis capabilities in malignant cells are shown in Table 6.

Nutrient	Cancers Affected
Chlorophyllin	Bladder
Coffee extract	Colon
Curcumin	Leukemia, colon
Emblica officinalis (amla; Indian	Ovary
gooseberry)	
Green tea extract (EGCG)	Leukemia, lymphoma, head-
	and-neck
Lycopene	Prostate
Phenyl isothiocyanate (PEITC) from	Bladder, lung
cruciferous vegetables	
Pine bark extract (Enzogenol)	Leukemia
Prebiotics (fermentable fiber, which	Colon
produces butyrate, induces apoptosis)	
Probiotics (Lactobacillus salivarius)	Oral
Propolis	Colon
Quercetin	Ovary
Red clover isoflavones	Prostate
Rosemary (carnosol)	Prostate, colon, skin, breast,
	kidney, liver
Sarsaparilla (Smilax glabra)	Multiple
Selenium	Colon, lung, prostate
Soy isoflavones (genistein, daidzein)	Prostate

Table 6. Nutrients that promote or restore apoptosis(International Agency for Research on Cancer, 2011)

4. Conclusion

We are awash in a sea of toxins and invisible radiation that constantly promotes malignant transformation of our cells, leading to persistently high rates of cancer. Despite the seemingly immeasurable amount of environmental toxins, there are five mechanisms through which they typically work to promote cancer. This allows us to identify nutrients that have cancer-fighting properties that work specifically against these mechanisms. Just as the chemicals that cause cancer do so by multiple mechanisms, natural products offer multiple, overlapping, and complementary approaches to cancer prevention.

By becoming familiar with the major cancer-inducing and cancer-promoting toxic influences in your world, you can then develop a supplement regimen that covers all five mechanisms by which we know that toxins and radiation induce cancerous changes. By choosing carefully from among the nutrients listed in this article, you can establish a solid cancer-fighting base in your own body—one that works with your natural defenses to defeat cancers before they get established.

Environmental toxins can cause serious health effects when exposure is allowed to accumulate. Problems usually result from prolonged or excessive exposure. While it is impossible to completely eliminate exposure, a few simple steps will go a long way towards protecting you and your family:

- 1. Filtering home tap water and not storing water in plastic bottle.
- 2. Not using plastic plates to heat food in a microwave oven.
- 3. Reduce use of canned foods and eat mostly fresh or frozen foods.
- 4. Use baby bottles that are BPA free (or better yet use glass bottles) and look for toys labeled BPA free.
- 5. Use PVC-free containers. Buy plastic wrap and bags made from polyethylene and use glass containers. If you do use plastic containers, do not heat or microwave them.
- 6. Choose phthalate-free toys. Many large toymakers have pledged to stop using phthalates, but be sure to look for toys made from polypropylene or polyethylene.
- 7. Purchase phthalate-free beauty products. Avoid nail polish, perfumes, colognes, and other scented products that list phthalates as an ingredient.
- 8. Eating food grown without pesticides or chemical fertilizers.
- 9. Avoiding processed, charred and well-done meats.
- 10. Reducing cell phone usage.
- 11. Reducing exposure to radiation from medical sources by discussing with healthcare providers whether medical tests or procedures (such as CT-scans) that use radiation are really necessary.
- 12. Get your home air and water checked for radon.

There is no need to freak out over occasional exposure to environmental toxins. Just look for simple ways to reduce your everyday exposure. Make changes slowly, one at a time, in a manageable way, and you will decrease your risk with minimal stress.

References

- Abnet CC. Carcinogenic food contaminants. *Cancer Invest*. 2007 Apr-May;25(3): 189-196.
- Aktar MW, Sengupta D, Chowdhury A. Impact of pesticides use in agriculture: their benefits and hazards. *Interdiscip Toxicol*. 2009 Mar;2(1):1-12.
- Ananthaswamy HN, Pierceall WE. Molecular mechanisms of ultraviolet radiation carcinogenesis. *Photochem Photobiol*. 1990 Dec;52(6):1119-36.
- Balabanic D, Rupnik M, Klemencic AK. Negative impact of endocrine-disrupting compounds on human reproductive health. *Reprod Fertil Dev.* 2011;23(3):403-16.
- De Iuliis GN, Newey RJ, King BV, Aitken RJ. Mobile phone radiation induces reactive oxygen species production and DNA damage in human spermatozoa in vitro. *PLoS One*. 2009;4(7):e6446.
- Erkekoglu P, Kocer-Gumusel B. Genotoxicity of phthalates. *Toxicol Mech Methods*. 2014 Dec;24(9):616-26.
- Hamid AS, Tesfamariam IG, Zhang Y, Zhang ZG. Aflatoxin B1-induced hepatocellular carcinoma in developing countries: Geographical distribution, mechanism of action and prevention. *Oncology Letters*. 2013;5(4):1087-92
- Hardell L, Sage C. Biological effects from electromagnetic field exposure and public exposure standards. *Biomed Pharmacother*. 2008 Feb;62(2):104-9.
- Hung LJ, Chan TF, Wu CH, Chiu HF, Yang CY. Traffic air pollution and risk of death from ovarian cancer in Taiwan: fine particulate matter (PM2.5) as a proxy marker. *J Toxicol Environ Health A*. 2012;75(3):174-82.
- International Agency for Research on Cancer (2011): Global Cancer Facts and Figures. Second Edition. American Cancer Society, Inc. 57 pp.
- International Agency for Cancer Research.(IACR).(2011): Biennial Report. Lyon, France. 35pp.
- Jeng HA. Exposure to endocrine disrupting chemicals and male reproductive health. *Front Public Health*. 2014 Jun 5;2:55.
- Kim JY, Yi BR, Go RE, Hwang KA, Nam KH, Choi KC. Methoxychlor and triclosan stimulates ovarian cancer growth by regulating cell cycle- and apoptosis-related genes via an estrogen receptor-dependent pathway. *Environ Toxicol Pharmacol.* 2014 May;37(3):1264-74.
- Lee CC, Liu JY, Lin JK, Chu JS, Shew JY. p53 point mutation enhanced by hepatic regeneration in aflatoxin B1-induced rat liver tumors and preneoplastic lesions. *Cancer Lett.* 1998 Mar 13; 125(1-2):1-7
- Liu C, Gao P, Xu SC, et al. Mobile phone radiation induces mode-dependent DNA damage in a mouse spermatocyte-derived cell line: a protective role of melatonin. *Int J Radiat Biol*. 2013 Nov;89(11):993-1001.
- Lutsenko EA, Cárcamo JM, Golde DW. Vitamin C prevents DNA mutation induced by oxidative stress. *J Biol Chem.* 2002 May 10;277(19):16895-9.
- Mihai CT, Rotinberg P, Brinza F, Vochita G. Extremely low-frequency electromagnetic fields cause DNA strand breaks in normal cells. *Journal of Environmental Health Science and Engineering*. 2014;12:15.
- Modem S, Dicarlo SE, Reddy TR. Fresh Garlic Extract Induces Growth Arrest and Morphological Differentiation of MCF7 Breast Cancer Cells. *Genes Cancer*. 2012 Feb;3(2):177-86.

- Morgan LL, Miller AB, Sasco A, Davis DL. Mobile phone radiation causes brain tumors and should be classified as a probable human carcinogen (2A) (Review). *Int J Oncol*. 2015 May;46(5):1865-71.
- Oishi K, Sato T, Yokoi W, Yoshida Y, Ito M, Sawada H. Effect of probiotics, Bifidobacterium breve and Lactobacillus casei, on bisphenol A exposure in rats. *Biosci Biotechnol Biochem*. 2008 Jun;72(6):1409-15.
- Ormond G, Nieuwenhuijsen MJ, Nelson P, et al. Endocrine disruptors in the workplace, hair spray, folate supplementation, and risk of hypospadias: case–control study. *Environ Health Perspect*. 2009;117(2):303-7.
- Pericleous M, Rossi RE, Mandair D, Whyand T, Caplin ME. Nutrition and pancreatic cancer. *Anticancer Res.* 2014 Jan;34(1):9-21.
- Pestana D, Teixeira D, Faria A, Domingues V, Monteiro R, Calhau C. Effects of environmental organochlorine pesticides on human breast cancer: putative involvement on invasive cell ability. *Environ Toxicol.* 2015 Feb;30(2):168-76.
- Rauscher-Gabernig E, Mischek D, Moche W, Prean M. Dietary intake of dioxins, furans and dioxin-like PCBs in Austria. *Food Addit Contam Part A Chem Anal Control Expo Risk* Assess. 2013;30(10):1770-9.
- Reichrath J, Rass K. Ultraviolet damage, DNA repair and vitamin D in nonmelanoma skin cancer and in malignant melanoma: an update. *Adv Exp Med Biol.* 2014;810:208-33.
- Revilla E, Santa-Maria C, Miramontes E, et al. Antiproliferative and immunoactivatory ability of an enzymatic extract from rice bran. *Food Chem.* 2013 Jan 15;136(2):526-31.
- Rota M, Bosetti C, Boccia S, Boffetta P, La Vecchia C. Occupational exposures to polycyclic aromatic hydrocarbons and respiratory and urinary tract cancers: an updated systematic review and a meta-analysis to 2014. *Arch Toxicol.* 2014 Aug;88(8):1479-90.
- Singh S, Li SS. Epigenetic effects of environmental chemicals bisphenol a and phthalates. *Int J Mol Sci.* 2012;13(8):10143-53.
- Sugimura T. Nutrition and dietary carcinogens. Carcinogenesis. 2000 Mar;21(3):387-95.
- Van Vleet TR, Watterson TL, Klein PJ, Coulombe RA, Jr. Aflatoxin B1 alters the expression of p53 in cytochrome P450-expressing human lung cells. *Toxicol Sci.* 2006 Feb;89(2):399-407.
- Vetvicka V, Vetvickova J. Glucan-resveratrol-vitamin C combination offers protection against toxic agents. *Toxins (Basel)*. 2012 Nov;4(11):1301-8.
- Volkow ND, Tomasi D, Wang GJ, et al. Effects of cell phone radiofrequency signal exposure on brain glucose metabolism. *Jama*. 2011 Feb 23;305(8):808-13.
- WHO. (2008): World cancer report 2008.
- WHO. (2011): Cancer Control Program. (NCCP core self-assessment tool)
- Wicki A, Hagmann, J. (2011): Diet and cancer. Swiss medical weekly141: w13250
- Xin F, Jiang L, Liu X, et al. Bisphenol A induces oxidative stress-associated DNA damage in INS-1 cells. *Mutat Res Genet Toxicol Environ Mutagen*. 2014 Jul 15;769:29-33.

Websites

http://www.cancer.gov/cancertopics/factsheet/Prevention/HPV-vaccine

http://www.cancer.gov/cancertopics/types/disparities

http://www.naturalnews.com/021808_cancer_prevention.html#ixzz2SOzRbCKI

J. Myan. Acad. Tech. 18(1-2). 2018