

Hydrogen sulfide suppress the pathological alterations of endocrine glands induced by Gamma irradiation and cyclophosphamide

Ashraf Kassem¹; Azza Hassan¹; Eman Fayez Said Taha² and Ahmed H Osman^{*}

¹Department of Pathology, Faculty of Veterinary Medicine, Cairo University, Egypt.

²Health Radiation Research Department, National Centre for Radiation Research and Technology (NCRRT), Egyptian Atomic Energy Authority, Cairo, Egypt.

Received Date : July 04, 2022
Accepted Date : Aug 06, 2022
Published Date : Aug 25, 2022
Archived : www.jcmimagescasereports.org
Copyright : © Ahmed Hassan Osman
2022

***Corresponding Author:** Ahmed Hassan Osman, Professor and head of Pathology department, Faculty of Veterinary Medicine, Cairo University, Egypt.
Email: ahosman2007@hotmail.com

Abstract

Background: Hydrogen sulfide (H₂S) is a gaseous mediator and is usually recommended to have antioxidant, anti-inflammatory, anticancer and antiapoptotic consequences. In the endocrine system, H₂S can act at the thyroid, adrenal gland, and gonad through the hypothalamus-pituitary axis, in addition to at the pancreas there by collaborating withinside the law of many hormones of the body, and the hypothalamus pituitary axis can, in turn, adjust the manufacturing of (H₂S). Chemotherapy (Cyclophosphamide) can cause damage of thyroid, pituitary, pancreatic and adrenal glands. Radiation-induced thyroid, hypothalamus, and anterior pituitary gland disorders. Radiation-induced injury of the endocrine pancreas is known to increase the risk of diabetes mellitus. Testis tissue, radiosensitive organ, has a variety of cells that differ in their degree of sensitivity. In this manner, cancer patient needs for protective agent against different side effects of anticancer therapy especially in case of combination between chemo and radio therapies.

Presentation of the hypothesis: H₂S can guard endocrine organs and adjust hormone secretion via anti-oxidative effect and ion channel regulation. H₂S performs a function with inside the endocrine system to guard pancreatic cells, adjust insulin secretion, hold the characteristic of the adrenal cortex, sell the discharge of catecholamine, and adjust the secretion of pituitary hormones.

Implications of the hypothesis: Administrations of H₂S has protecting and therapeutic results on endocrine glands in opposition to numerous results of chemo and radio therapies. H₂S is a promising molecule for the development of new medications.

Keywords: Hydrogen sulfide; endocrine glands; pathophysiology; cyclophosphamide; Gamma irradiation.

Introduction

Over the closing several decades, hydrogen sulfide (H₂S) as obtained hobby as a manufacturer new signaling molecule, with physiological and pathophysiological roles in human issues affecting vascular biology, immune capabilities, cellular survival, metabolism, longevity, development, and strain resistance. Apart from its considered competencies in oxidative stress and inflammation, new proof has emerged revealing that H₂S consists of out physiological competencies thru focused on proteins, enzymes, and transcription factors via a post-translational modification known as per-sulfidation [1].

In this study, results of H₂S donor sodium hydrosulfide (Na

HS) on cyclophosphamide and gamma irradiation which associated to pathological changes of endocrine glands [2]. Hydrogen sulfide (H₂S) has been verified to be generated withinside the endocrine and reproductive organs and elicits numerous actions. H₂S modulates insulin secretion in pancreatic islets. Adipose tissues could grant H₂S, which regulates the close by insulin sensitivity and vascular responsiveness. Moreover, it acts at the hypothalamic-pituitary-adrenal axis and is concerned in stress responses [3]. Cyclophosphamide induces derangement in spermatogenesis and motives atrophy of seminiferous tubules and testosterone level depletion [4]. Thyroid gland damage [5] and vacuolation of interstitial spaces in the pituitary gland [6]. Pancreatic acini revealed focal slight hy-

Citation: Ahmed H Osman. Hydrogen sulfide suppress the pathological alterations of endocrine glands induced by Gamma irradiation and cyclophosphamide. *J Clin Med Img Case Rep.* 2022; 2(4): 1225.

dropic degeneration, in addition to moderate oedema and congestion. Besides, the Langerhans islets had been markedly strange and lowered in size, with atrophic and shrunken cells [7]. Adrenal glands, that are concerned withinside the frame's response to stress phenomena, demonstrated hypertrophy after high dose of cyclophosphamide, with body weight relative developing and glucocorticoids hormones immoderate secretion [8]. Endocrine late effects of irradiation may additionally be direct, ensuing in hypofunction of endocrine glands or indirect ensuing in metaplasia and most cancers [9]. Radiotherapy and chemotherapy are the handiest and systematic strategies for most cancers treatment which observed with unsafe consequences will commonly reduce the high-satisfactory of existence for victims with most cancers and may additionally purpose discontinuation of therapy [10].

H₂S CHEMISTRY

H₂S is normally used referring to the complete sulfide species. Although H₂S has appropriate solubility in water, it is nevertheless very unstable in solution. It is easy oxidized in the presence of oxygen, forming oxidized sulfide species such as sulfite (SO₃²⁻), sulfate (SO₄²⁻), thiosulfate (S₂O₃²⁻), polythionates (SnOn + ²⁻), and polysulfides (S_x²⁻), as well as other oxidized polysulfide species [11]. Hydrogen sulfide (H₂S), an everyday lethal, poisonous fuel with the odor of rotten eggs, is diagnosed as one of the three gasotransmitters in mammals, which additionally consist of nitric oxide (NO) and carbon monoxide (CO) [12,13]. Hydrogen sulfide performs relies upon on the precise circum- stance, its concentration, and the interplays with different signaling molecules, particularly NO and CO [14]. H₂S does not form hydrogen bonds and is lipophilic, allowing it to pass through biological membranes and act as a paracrine signaling molecule [15].

Synthesis and Metabolism of H₂S Inthe Endocrine System

At present, three primary recognized enzymes produce H₂S in organisms: cystathionine b-synthase (CBS), cystathionine g-lyase (CSE), and 3-mercaptopyruvate sulfur transferase (3-MST) [2]. The substrate of (CBS) and (CSE) is L-cysteine [4], whilst 3-MST can catalyze 3-mercaptopyruvate to produce H₂S [5, 6]. Recently, it has been discovered that a form of human methanethiol oxidase-selenium binding protein 1 (SELENBP1) can convert methanethiol into H₂O₂, formaldehyde, and H₂S. In adipocytes, H₂S can be produced by using SELENBP1 and is associated to the expression of CBS, CSE, and 3-MST [9]. But the amount of the enzyme that produces H₂S varies in unique tissues and organs. In endocrine glands and endocrine organs, the RNA and protein expression levels of CBS are the easiest in the pancreas, specifically in acinar cells [10]. Moreover, the CBS is hardly ever dispensed in other endocrine glands such as thyroid, parathyroid, adrenal gland, and pituitary gland [11]; however, abnormally improved in thyroid carcinoma [16]. The expression of CSE is the amplest in the liver, however low in the thyroid, pancreas, testis, ovary, and other endocrine glands [12, 14]. 3-MST is especially expressed in endocrine tissues (thyroid, parathyroid, adrenal), pancreas, gonad (tes-

tis and ovary) [17]. H₂S can have an effect on the secretion of many hormones and participate in the prevalence and improvement of endocrine diseases [18], however the impact of H₂S on the physique may additionally be biphasic [2], that is, the impact of too excessive or too low attention is the opposite, so to make certain the everyday physiological feature of the body, it is integral to preserve the attention of H₂S at an appropriate concentration. It is properly recognized that H₂S can be regulated in two ways: synthesis and consumption. In phrases of synthesis, H₂S is synthesized broadly speaking thru enzymatic and nonenzymatic pathways (such as discount of sulfur-containing compounds) and, in a few cases, launched by way of sure sulfur stored in cells [19]. The half-life of H₂S in vivo is very quick (a few seconds to a few minutes) [20,21]. Nowadays, it in the main inhibits the exercise of synthase or increases the donor in vitro, which brings awesome difficulties to the find out about of H₂S.

Biology of H₂S

H₂S is regularly produced thru the anaerobic bacterial breakdown of natural substrates in the absence of oxygen, such as in swamps and sewers (anaerobic digestion). It additionally consequences from inorganic reactions in volcanic gases, herbal gas, and some properly waters. Digestion of algae, mushrooms, garlic, and onions is believed to launch H₂S via chemical transformation and enzymatic reactions [17]. Human physique produces small quantities of H₂S and makes use of it as a signaling molecule [19]. In mammals, three enzymes are concerned in sulfur-containing amino acid metabolism and hence accountable for the in vivo manufacturing of H₂S. Two of them are pyridoxal-50-phosphate (PLP)-dependent enzymes: cystathionine β-synthase (CBS) and cystathionine γ-lyase (CSE). CBS is expressed predominantly in the central frightened device (CNS) [22].

H₂S DONORS

Inorganic sulfide salts, such as sodium sulfide (Na₂S) and hydrosulfide (Na HS) [20], diallyl disulfide (DADS) [21], Lawes son's reagent and its analogs [23], Thiol activated H₂S donors [13]. Structures of herbal meals releasing H₂S on digestion are proven in Consuming mushrooms, garlic, and onions, which include chemical substances and enzymes accountable for the transformation of the sulfur compounds, is accountable for H₂S pro- duction in the human intestine [19]. Naturally occurring donors, presently handy H₂S-releasing compounds can be divided into two groups: naturally taking place donors and artificial donors. Among the herbal source, allium household and cruciferous greens are identified to be prosperous in organosulfur compounds [24]. Among Cruciferae, veggies such as broccoli, watercress, mustard, and garden cress are wealthy in isothiocyanates, such as sulforaphane (SFN, notably existing in broccoli), allyl isothiocyanate (AITC, pretty current in black mustard), benzyl isothiocyanate (BITC, relatively current in backyard cress), 4-hydroxybenzyl isothiocyanate (HBITC, quite current in white mustard), and erucin (ERU, basically current in broccoli and rocket) [25].

Hydrogen Sulfide-Releasing Therapeutics

The literature proof suggests that hydrogen sulfide possesses the following activities: anti-inflammatory [26], anti-tumor [27], ion channel regulation [28, 29], cardiovascular protection [30] and antioxidation [31]. However, the actual function that hydrogen sulfide performs relies upon on the precise circumstance, its concentration, and the interplays with different signaling molecules, in particular NO and CO [14]. This is an important location of lookup in growing hydrogen sulfide-based therapeutics [32]. There are additionally a range of therapeutics on world markets that can generate H₂S in vivo and the place there is at least some proof of H₂S contributing to their therapeutic benefits. Indeed, quite a few advisable outcomes of H₂S for chemoprevention reducing oxidative stress, and attenuating irritation [29,33]. In rodent models, it has been proven to limit the formation of quite a number of cancers (colon, bladder, blood, liver, kidney, pancreas, lung, and mammary) [33]. There is proof that the anticancer outcomes may additionally be mediated via activation of Nfr2 [34], a signaling pathway recognized to be activated by means of H₂S [35].

Biological Effects of H₂S In The Endocrine Disorders Induced By Chemo And Radiotherapy

H₂S Promotes Pancreatic dysfunction:

Cyclophosphamide toxicity influences many organs, inclusive of the pancreas, and is exotic through glutathione depletion, lipid peroxidation, altered DNA profile, pro-inflammatory response, and apoptosis [36]. In cyclophosphamide intoxicated mice, the pancreatic acini confirmed focal average hydropic degeneration, as nicely as average edema and congestion. Besides, the Langerhans islets have been markedly irregular and decreased in size, with atrophic and shrunken cells [7]. Radiation-induced harm of the endocrine and exocrine pancreas. Diabetes mellitus (DM) has been located in preceding retrospective studies [16,37]. DM has been pronounced in 1.03–8.3% of survivors. In youth and younger adults who have acquired complete physique irradiation (TBI) or stomach irradiation, there is a threat of diabetes, which is associated to the whole dose of radiation administered to the tail of the pancreas the place the islets are concentrated [16].

The essential roles of H₂S in the pancreas are defending pancreatic β cells and regulating insulin secretion. H₂S may also guard pancreatic β cells in the following three methods consist of reduce the manufacturing of ROS; inhibit the expression of thioredoxin binding protein-2-a redox protein related with diabetes that promotes apoptosis and enlarge the content material of GSH, all of which minimize the harm of oxidative stress [38]. On the contrary, an excessive awareness of H₂S induces apoptosis of pancreatic β cells [39]. Insulin secretion is affected by means of many factors. It is recognized that the attention and oscillation of Ca²⁺, KATP channel is associated to H₂S.

H₂S can no longer solely inhibit the entry of Ca²⁺ from the plasma membrane into cells, to minimize insulin secretion, however additionally promote the launch of Ca²⁺ in mitochondria and amplify insulin secretion [40]. It is properly known that an excessive awareness of glucose can promote insulin secretion [41].

H₂S Promotes Thyroid dysfunction

Cyclophosphamide can purpose thyroid injury and alternate thyroid function [5]. There are several pathophysiological mechanisms for thyroid dysfunction that have been defined as (1) version due to lively hypothalamic pituitary axis involvement; (2) altered synthesis or clearance of thyroid hormone binding proteins detected in sure malignancies or brought about by means of most cancers cure that modifications complete however no longer free awareness of thyroid hormones; (3) alteration of thyroid hormones metabolism, which can happen in chronically unwell most cancers patients [42]. A discount of thyroid hormones may want to reason a proliferation arrest in G0-G1 in most cancers' cells, with viable influences on their sensitivity to chemotherapy agents [43].

Gamma radiation-induced thyroid problems are customary problems after radiotherapy (RT) in sufferers with head and neck cancer. The thyroid gland is a predominant endocrine organ producing thyroid hormones for keeping metabolism. Injury to the thyroid gland due to radiation can also result in momentary thyroiditis and hypothyroidism (HT). Moreover, post-RT-HT is related with the accumulative radiation dose to the thyroid gland [44]. Radiotherapy-induced thyroid abnormalities continue to be under-estimated and beneath reported. These sequelae might also encompass principal or central hypothyroidism, thyroiditis, Graves' disease, euthyroid Graves' ophthalmopathy, benign adenomas, multinodular goiter and radiation- brought on thyroid carcinoma [45]. External-beam radiotherapy induces more than a few thyroid disorders, such as important hypothyroidism (6–89%), Hashimoto's thyroiditis (0.7–48%), benign adenoma (0.6–3%), silent thyroiditis (0.6–3%), Graves' ailment (0.1–2%), Graves' ophthalmopathy (0.2–1.3%), and thyroid most cancers (0.35%) [45]. The most frequent thyroid illnesses following publicity to ionizing radiation are hypofunction and thyroid nodules. However, the expanded incidence of thyroid cancers may additionally be worried in these diseases [46].

The pathophysiological description of radiation-induced thyroid injury is associated to inhibition of follicular epithelial feature and subsequent innovative alteration of the endothelium, the impact will increase through time [47]. Even although oxidative reactions take region in all tissues and organs, the thyroid gland constitutes such an organ in which oxidative approaches are vital for thyroid hormone synthesis. It is estimated that massive quantity of reactive oxygen species, specifically of H₂O₂, are shaped in the thyroid below the physiological situation. Yet, with extra oxidative mistreatment triggered with the aid of IR, multiplied injury to macromolecules occurs, probably main to one-of-a-kind thyroid diseases, most cancers included [48].

Both anaplastic and papillary thyroid cancers are inhibited by means of hydrogen sulfide, in anaplastic cancer, hydrogen sulfide reasons the accumulation of reactive oxygen species (ROS), which inhibits cell survival and will increase apoptosis. Besides that, it promotes phone DNA injury and reasons the cell cycle to end in the G2/M phase [49]. Hydrogen sulfide inhibited cell growth in papillary carcinoma [50].

Uncertainty of the Effect of H₂S on Testicular damage

Cyclophosphamide (CYP) reasons reproductive toxicity, consisting of azoospermia, oligospermia, histological adjustments in the epididymis and testis, diminished weight of reproduc-

tive organs, and impaired fertility and growth in people and experimental animals. Because of the excessive frequency of cell division in the cells of the seminiferous epithelium, the testis is extraordinarily touchy to chemotherapeutic drugs [51]. Cyclophosphamide handled male rats confirmed low serum attention of testosterone collectively with low serum FSH and LH. CYP motives extensive minimize in recreation of testicular steroidogenic enzymes which are the key enzymes for biosynthesis of testosterone [52]. Decreased reproductive organ weights, oligo-, azoo- and teratozoospermia, low stages of testosterone and LH, atrophied seminiferous tubules, degenerated spermatogenic cells and apoptosis are some of the damaging consequences caused via Cyclophosphamide [53]. The expand in free radicals in cells can set off lipid peroxidation by means of oxidative breakdown of polyunsaturated fatty acids in the membranes of these cells. Obviously, peroxidation of sperm lipids destroys the shape of the lipid matrix in the plasma membranes, and it is related with fast loss of intracellular ATP, main to axonemal damage, lowered sperm viability and, in severe cases, even whole inhibition of spermatogenesis [54]. In addition to typical endocrinological imbalances such as ovarian failure, atypical sperm manufacturing decreased fertility and outcome, lowered implantation, and malformed or increase retarded fetuses [55].

Testis tissue, radiosensitive organ, has a range of cells that range in their diploma of radiosensitivity. The spermatogonia are very radiosensitive and kill at doses much less than three Gy in differentiation period. The infertility following irradiation is induced due to apoptosis of spermatogonia alternatively of turning into differences [18]. The germinal epithelium is very touchy to radiation-induced injury with adjustments in spermatogonia following doses as low as 0.1 Gy and everlasting infertility after fractionated doses of two Gy and above [56]. The deleterious consequences of irradiation in organ structures are on the whole mediated thru the era of ROS and cause lipid peroxidation in the cell membrane, thereby inducing DNA injury in immature germ cell's, DNA injury prompted by means of irradiation in premeiotic germ cells is detectable in essential spermatocytes and is nevertheless existing in mature spermatozoa [57]. H_2S regulates testosterone secretion with the aid of influencing luteinizing hormone (LH) secretion, indicating that H_2S may also play an essential position in testosterone secretion [58]. It blanketed the testis from chemotherapeutic drug-induced oxidative stress and irritation and elevated the recreation of antioxidant enzymes [59]. The expression of CBS and CSE are located in rat testes, however they are differentially expressed; CSE is ordinarily expressed in Sertoli cells and immature spermatogonia, whilst CBS is basically allotted in Leydig cells, Sertoli cells, and germ cells [60].

Effect of H_2S on adrenal gland

Adrenal glands, which are concerned in the body's response to stress phenomena, exhibit organ hypertrophy after a quick length of cyclophosphamide administration, with physique weight relative growing and glucocorticoids hormones extreme secretion [8]. The impact of experimental irradiation of rats in the long-term duration is accompanied by accumulation of MDA in spleen, adrenal glands, lymph nodes of the small intestine [61]. The principal mechanism of the radiation-induced regular tissues injury is the response of radiation with water in vivo to generate immoderate reactive oxygen species

(ROS) which is a most important inducer of apoptosis. Increasing proof suggests that (ROS), malondialdehyde (MDA), superoxide dismutase (SOD), and glutathione peroxidase (GSH-PX) are oxidative-stress-related. Antioxidants or free radical scavengers can also provide safety towards radiation-induced harm to hepatic and different tissues [44]. Hydrogen sulfide can have an effect on the adrenal cortex and chromaffin cells in the adrenal gland. Inhibiting cystathionine- β -synthase (CBS)/ cystathionine- γ - lyase (CSE) can purpose mitochondrial oxidative stress and dysfunction in the adrenal cortex [33].

The Role of H_2S in hypothalamus and pituitary gland

Chemotherapy (Cyclophosphamide) has been linked to hypothalamic-pituitary dysfunction in childhood most cancers survivors, manifesting as deficiencies of man or woman hormones or, in some cases, a couple of hormone deficiencies [62]. It has additionally been mentioned that this anticancer agent might also have a poor influence on talent [63]. vascular complications, seizures, and peripheral neuropathies synthesis by way of impairing neurogenesis and synaptic plasticity in hypothalamus. The majority of this neurotoxicity may want to be attributed to the improvement of oxidative stress [64]. Cyclophosphamide prompted a hemorrhagic lesion and congestion of veins and capillaries, parenchymal cells manifested scanty and hypertrophied, and vacuolation of interstitial areas in the pituitary gland [65].

Damage to the hypothalamic pituitary axis has been diagnosed as an aspect impact of radiation remedy to the base of skull. Effects brought about via non-lethal events at the cellular level are referred to as "stochastic effects." These have no dose threshold, and their incidence is associated to radiation dose; however, the severity of the impact is no longer dose related. In contrast, outcomes mediated by means of cell killing and ordinary tissue dysfunction are known as "deterministic" (or "non-stochastic") effects. These toxicities frequently have a dose threshold, and each the incidence and severity of the impact are dose associated [66]. Radiotherapy is a frequent reason of hypothalamic-pituitary dysfunction in most cancers' sufferers [67]. The hypothalamus and pituitary gland are localized shut collectively and engage by using the hypothalamic-pituitary-adrenal and-gonadal axis. Several researches confirmed predominant characteristic loss of the hypothalamus. Growth hormone (GH) deficiency is the most familiar endocrine dysfunction after hypothalamus-pituitary gland irradiation, taking place at tremendously low doses [68].

The hypothalamic-pituitary-target organ axis is a complicated organic structure, which performs an essential function in endocrine regulation. H_2S is disbursed in more than one sites in the hypothalamus, pituitary-target organ, making it an essential molecule in regulating hormone secretion, which is no longer solely twin however additionally bi-directional, that is, H_2S now not solely promotes the secretion of certain hormones however can additionally inhibit them. The complex effect of H_2S on the endocrine system may be caused by its action on different organs [69]. Hydrogen Sulfide is concerned in several physiological and pathological approaches in the body, together with dilating blood vessels (regulating blood pressure), defending tissue from ischemia-reperfusion injury, anti-inflammatory, carcinogenesis, or most cancers inhibition, and regulating hormonal metabolism through the hypothalamus and pancreas [69]. The learn about determined that hormones

produced by way of the hypothalamus-pituitary axis have an effect on H₂S synthesis. Thyroid hormone (TH) and growth hormone (GH) alter H₂S manufacturing in the liver with the aid of the TH receptor b1 and the GH receptor, respectively. In mechanism, TH inhibits CSE expression whilst GH inhibits H₂S manufacturing by substrate availability manipulate with the aid of autophagy [70].

The implication of the hypothesis

H₂S can protect endocrine organs against side effects of chemotherapy and radiotherapy in addition to regulate hormone secretion through anti-oxidative stress and ion channel regulation.

References

1. Comas F and J Moreno-Navarrete. The Impact of H₂S on Obesity-Associated Metabolic Disturbances. *Antioxidants*, 2021; 10: 633.
2. Özatik FY, et al. Protective and therapeutic effect of Hydrogen sulfide on hemorrhagic cystitis and testis dysfunction induced with Cyclophosphamide. *Turkish Journal of Medical Sciences*, 2021; 51(3): 1531-1543.
3. Zhu XY, H Gu and X Ni. Hydrogen sulfide in the endocrine and reproductive systems. *Expert Review of Clinical Pharmacology*, 2011; 4(1): 75-82.
4. Ekeleme-Egedigwe CA, et al. Antioxidant potential of garlic oil supplementation prevents cyclophosphamide-induced oxidative testicular damage and endocrine depletion in rats. *Journal of Nutrition & Intermediary Metabolism*, 2019; 18: 100109.
5. Shi Y, et al. Study on the status of thyroid function and thyroid nodules in chinese breast cancer patients. *Oncotarget*, 2017; 8(46): 80820.
6. Ayoka OA, et al. Neuro-endocrine effects of aqueous extract of *Amaranthus viridis* (Linn.) leaf in male Wistar rat model of cyclophosphamide-induced reproductive toxicity. *Toxicology reports*, 2016; 3: 608-619.
7. Khodeer DM, et al. Protective effects of evening primrose oil against cyclophosphamide-induced biochemical, histopathological, and genotoxic alterations in mice. *Pathogens*, 2020; 9(2): 98.
8. Hermenean A, A. Ardelean and C Crăciun. ADRENAL GLANDS MORPHO-FUNCTIONAL DAMAGES AT WISTAR RATS AFTER CHEMOTHERAPY ADMINISTRATION. *Studia Universitatis Vasile Goldis Seria Stiintele Vietii (Life Sciences Series)*, 2008; 18.
9. Zacharin M. Longterm endocrine effects of cancer. *International Journal of Pediatric Endocrinology*, 2013; 2013(1): 1-2.
10. Wild CP, et al. Cancer prevention Europe. *Molecular oncology*, 2019; 13(3): 528-534.
11. DeLeon ER, GF Stoy, and KR Olson. Passive loss of hydrogen sulfide in biological experiments. *Analytical biochemistry*, 2012; 421(1): 203-207.
12. Vandiver MS and SH Snyder. Hydrogen sulfide: a gasotransmitter of clinical relevance. *Journal of molecular medicine*, 2012; 90(3): 255-263.
13. Wang R. Physiological implications of hydrogen sulfide: a whiff exploration that blossomed. *Physiological reviews*, 2012; 92(2): 791-896.
14. Paul BD and SH Snyder. H₂S signalling through protein sulfhydration and beyond. *Nature reviews Molecular cell biology*, 2012; 13(8): 499-507.
15. Cuevasanta E, et al. Solubility and permeation of hydrogen sulfide in lipid membranes. *PloS one*, 2012; 7(4): e34562.
16. Friedman DN, et al. Radiation dose and volume to the pancreas and subsequent risk of diabetes mellitus: a report from the childhood cancer survivor study. *JNCI: Journal of the National Cancer Institute*, 2020; 112(5): 525-532.
17. Wallace JL. Hydrogen sulfide-releasing anti-inflammatory drugs. *Trends in pharmacological sciences*, 2007; 28(10): 501-505.
18. Barazzuol L, RP Coppes and P van Luijk. Prevention and treatment of radiotherapy-induced side effects. *Molecular oncology*, 2020; 14(7): 1538-1554.
19. Fiorucci S, et al. The emerging roles of hydrogen sulfide in the gastrointestinal tract and liver. *Gastroenterology*, 2006; 131(1): 259-271.
20. Hughes MN, MN Centelles, and KP Moore. Making and working with hydrogen sulfide: the chemistry and generation of hydrogen sulfide in vitro and its measurement in vivo: a review. *Free Radical Biology and Medicine*, 2009; 47(10): 1346-1353.
21. Huang YS., et al. Diallyl disulfide inhibits the proliferation of HT-29 human colon cancer cells by inducing differentially expressed genes. *Molecular medicine reports*, 2011; 4(3): 553-559.
22. Miles EW and JP Krau. Cystathionine β-synthase: structure, function, regulation, and location of homocystinuria-causing mutations. *Journal of Biological Chemistry*, 2004; 279(29): 29871-29874.
23. Ozturk T, E Ertas, and O Mert. Use of Lawesson's reagent in organic syntheses. *Chemical Reviews*, 2007; 107(11): 5210-5278.
24. Benavides GA, et al., Hydrogen sulfide mediates the vasoactivity of garlic. *Proceedings of the National Academy of Sciences*, 2007; 104(46): 17977-17982.
25. Corvino A, et al. Trends in H₂S-Donors Chemistry and Their Effects in Cardiovascular Diseases. *Antioxidants*, 2021; 10(3): 429.
26. Li L, and P Moore. Putative biological roles of hydrogen sulfide in health and disease: a breath of not so fresh air? *Trends in pharmacological sciences*, 2008; 29(2): 84-90.
27. Shrotriya S, et al. Diallyl Trisulfide Inhibits Phorbol Ester-Induced Tumor Promotion, Activation of AP-1, and Expression of COX-2 in Mouse Skin by Blocking JNK and Akt Signaling. *Cancer research*, 2010; 70(5): 1932-1940.
28. Zhao W, et al. The vasorelaxant effect of H₂S as a novel en-

- ogenous gaseous KATP channel opener. *The EMBO journal*, 2001; 20(21): 6008-6016.
29. Liu L, et al. Effects of H₂S on myogenic responses in rat cerebral arterioles. *Circulation Journal*, 2012; 76(4): 1012-1019.
30. Lisjak M, et al. A novel hydrogen sulfide donor causes stomatal opening and reduces nitric oxide accumulation. *Plant Physiology and Biochemistry*, 2010; 48(12): 931-935.
31. Osborne NN, et al. Glutamate oxidative injury to RGC-5 cells in culture is necrostatin sensitive and blunted by a hydrogen sulfide (H₂S)-releasing derivative of aspirin (ACS14). *Neurochemistry international*, 2012; 60(4): 365-378.
32. Zheng Y, et al. Hydrogen sulfide prodrugs—a review. *Acta Pharmaceutica Sinica B*, 2015; 5(5): 367-377.
33. Wang K, H Peng, and B Wang. Recent advances in thiol and sulfide reactive probes. *Journal of cellular biochemistry*, 2014; 115(6): 1007-1022.
34. Peng, B. and M. Xian, Hydrogen Sulfide Detection Using Nucleophilic Substitution–Cyclization-Based Fluorescent Probes. *Methods in enzymology*, 2015; 554: 47-62.
35. Peng H, et al. A fluorescent probe for fast and quantitative detection of hydrogen sulfide in blood. *Angewandte Chemie*, 2011; 123(41): 9846-9849.
36. Amiri FT, et al. Anti-apoptotic and Antioxidant Effect of Cerium Oxide Nanoparticles on Cyclophosphamide-Induced Hepatotoxicity. *Erciyes Medical Journal/Erciyes Tip Dergisi*, 2018; 40(3).
37. Friedman DN, et al. Radiation dose and volume to the pancreas and subsequent risk of diabetes mellitus: A report from the Childhood Cancer Survivor study. *American Society of Clinical Oncology*. 2018.
38. Kaneko Y, et al. Glucose-induced production of hydrogen sulfide may protect the pancreatic beta-cells from apoptotic cell death by high glucose. *FEBS letters*, 2009; 583(2): 377-382.
39. Yang G. [H. sub. 2] S as a physiologic vasorelaxant: hypertension in mice with deletion of cystathionine [gamma]-lyase. *Nature Reviews Drug Discovery*, 2009; 8(1).
40. Lu A, et al. ATP-sensitive K⁺ channels and mitochondrial permeability transition pore mediate effects of hydrogen sulfide on cytosolic Ca²⁺ homeostasis and insulin secretion in β-cells. *Pflügers Archiv-European Journal of Physiology*, 2019; 471(11): 1551-1564.
41. Yang, W., Yang G, Jia X, Wu L, Wang R. Activation of K ATP channels by H₂S, 2005; 2: 519-531.
42. Ashif Khan, M., D. Bhurani, and N.B. Agarwal, Alteration of thyroid function in Indian HER 2-negative breast cancer patients undergoing chemotherapy. *Asian Pacific Journal of Cancer Prevention*, 2015; 16(17): 7701-7705.
43. Huang J, et al. Implication from thyroid function decreasing during chemotherapy in breast cancer patients: chemosensitization role of triiodothyronine. *BMC cancer*, 2013; 13(1): 1-12.
44. Liu CH, et al. Hypothyroidism and risks of cerebrovascular complications among patients with head and neck cancer after radiotherapy. *BMC neurology*, 2021; 21(1): 1-8.
45. Jereczek-Fossa BA, et al. Radiotherapy-induced thyroid disorders. *Cancer treatment reviews*, 2004; 30(4): 369-384.
46. Feldt-Rasmussen U and ÅK Rasmussen. Autoimmunity in differentiated thyroid cancer: significance and related clinical problems. *Hormones*, 2010; 9(2): 109-117.
47. Simões-Pereira J, MS Vieira, and MC Pereira. Latency period until the development of thyroid cancer in young patients submitted to radiotherapy: Report of 10 cases. *Case reports in oncology*, 2014; 7(3): 810-814.
48. Dalton TP, HG Shertzer, and A Puga. Regulation of gene expression by reactive oxygen. *Annual review of pharmacology and toxicology*, 1999; 39(1): 67-101.
49. Zheng J, et al. Diallyl trisulfide induces G2/M cell-cycle arrest and apoptosis in anaplastic thyroid carcinoma 8505C cells. *Food & function*, 2019; 10(11): 7253-7261.
50. Xu S, et al. Diallyl trisulfide, a H₂S donor, inhibits cell growth of human papillary thyroid carcinoma KTC-1 cells through a positive feedback loop between H₂S and cystathionine-gamma-lyase. *Phytotherapy Research*, 2020; 34(5): 1154-1165.
51. Salimnejad R, JS Rad, and DM Nejad. Protective Effect of Ghrelin on Oxidative Stress and Tissue Damages of Mice Testes Followed by Chemotherapy With Cyclophosphamide. *Crescent Journal of Medical and Biological Sciences*, 2018. 5(2): p. 138-43.
52. Bakhtiary Z, et al. Ethyl pyruvate ameliorates the damage induced by cyclophosphamide on adult mice testes. *International journal of fertility & sterility*, 2016; 10(1): 79.
53. Elangovan N, et al. Cyclophosphamide treatment causes impairment of sperm and its fertilizing ability in mice. *Toxicology*, 2006; 222(1-2): 60-70.
54. Türk G, et al. Improvement of cisplatin-induced injuries to sperm quality, the oxidant-antioxidant system, and the histologic structure of the rat testis by ellagic acid. *Fertility and Sterility*, 2008; 89(5): 1474-1481.
55. Khorwal G, R Chauhan, and M Nagar. Effect of cyclophosphamide on liver in albino rats: A comparative dose dependent histomorphological study. *International Journal of Biomedical and Advance Research*, 2017; 8(3): 102-107.
56. Osterberg EC, et al. Current practices in fertility preservation in male cancer patients. *Urology annals*, 2014; 6(1): 13.
57. Ergur BU, et al. Protective effect of erythropoietin pretreatment in testicular ischemia-reperfusion injury in rats. *Journal of pediatric surgery*, 2008; 43(4): 722-728.
58. Oi Y, et al. Garlic supplementation increases testicular testosterone and decreases plasma corticosterone in rats fed a high protein diet. *The Journal of nutrition*, 2001; 131(8): 2150-2156.
59. Azarbarz N, et al. Assessment of the effect of sodium hydrogen sulfide (hydrogen sulfide donor) on cisplatin-induced testicular toxicity in rats. *Environmental Science and Pollution Research*, 2020; 27(8): 8119-8128.
60. Kimura H. Hydrogen sulfide as a biological mediator. *Antioxidants & redox signaling*, 2005; 7(5-6): 778-780.

61. Ilderbayeva GO, and DM Suleymeneva. Free Radical Pathology of the Body in the Long-Term Period under Combined Exposure to Gamma Radiation and Emotional Stress in the Experiment. *International Journal of Environmental and Science Education*, 2016; 11(17): 9873-9881.
62. Rose SR, et al. Hypothalamic dysfunction after chemotherapy. *Journal of Pediatric Endocrinology and Metabolism*, 2004; 17(1): 55-66.
63. Ramchandani D, et al. Protective effect of curculigo orchoides extract on cyclophosphamide-induced neurotoxicity in murine model. *Toxicology international*, 2014; 21(3): 232.
64. Khandelwal N, and SK Abraham. Protective effects of common anthocyanidins against genotoxic damage induced by chemotherapeutic drugs in mice. *Planta medica*, 2014; 80(15): 1278-1283.
65. Ayoka OA, et al. Neuro-endocrine effects of aqueous extract of *Amaranthus viridis* (Linn.) leaf in male Wistar rat model of cyclophosphamide-induced reproductive toxicity. *Toxicology reports*; 2016; 3: 614.
66. Vakilian S, et al. Examination of the Dose-Effect Relationship of Radiation-Induced Hypopituitarism: Results of a Case-Control Study. *Advances in Radiation Oncology*, 2021; 6(4): 100693.
67. Bast RC, and JF Holland. *Holland-Frei Cancer Medicine*, PMPH-USA. 2020; 8.
68. Voshart DC, et al. Regional Responses in Radiation-Induced Normal Tissue Damage. *Cancers*, 2021; 13(3): 367.
69. Chen HJ, et al. Role of hydrogen sulfide in the endocrine system. *Frontiers in Endocrinology*, 2021; 12.
70. Hine C, et al. Hypothalamic-pituitary axis regulates hydrogen sulfide production. *Cell metabolism*, 2017; 25(6): 1320-1333. e5.