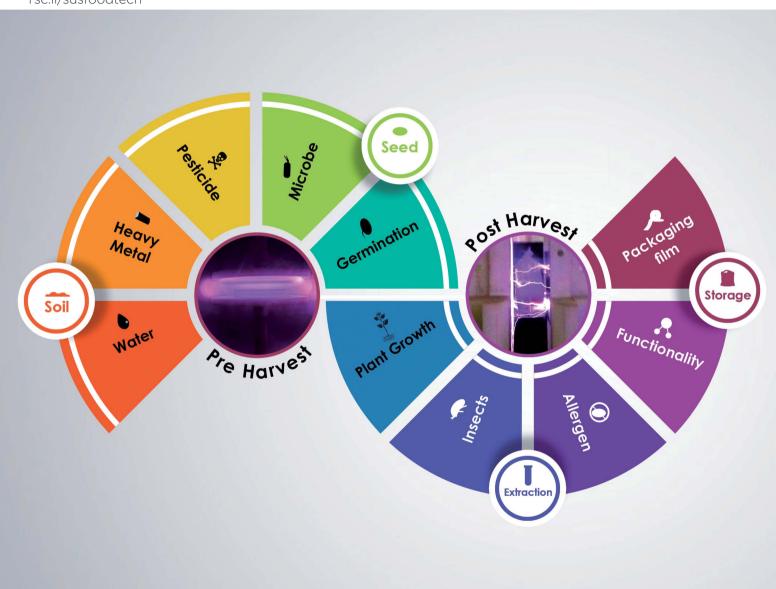
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# Sustainable Food Technology

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### **REVIEW ARTICLE**

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# Plasma processing: a sustainable technology in agri-food processing

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Globalization and rapid urbanization have led to tremendous improvement in the agriculture and food processing sectors to fulfil the food demands. In this context, managing food product safety and quality throughout the agri-food chain (pre-harvest to post-harvest) becomes vital to avoid food spoilage and increase production. Numerous innovative interventions have been investigated to achieve these goals; however, no single technology can be applied at all processing stages and may require different technologies. Nevertheless, cold plasma is a multifaceted solution for most pre and post-harvest issues, including soil/water contamination, microbial spoilage, insect infestation, and prolonged seed dormancy. In addition, the recent applications of plasma to food shape transformation is an evidence of the versatility of this technique in agri-food processing. Advantages, such as on-site production, residue, and toxic-free treatment, make the plasma process more sustainable. Reactive species, UV photons, and electrons are this plasma treatment's major compounds, giving them the peculiar and unique property to tackle most pre and post-harvest challenges. This review provides comprehensive possibilities for utilizing plasma technology throughout the agri-food chain. Various plasma systems have been developed, but their potential is limited to the lab scale. Research on large-scale applications can utilize cold plasma in future.

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#### Introduction 1.

Agriculture and food processing sectors need progressive growth in production and value addition to feed the increasing world population. Nevertheless, the required food production has to be achieved with the available resources without compromising the product quality. Thus, we need a technology that could bring progressive growth in food supply and address food security and quality-related issues. In agri-food processing,



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irrigation and soil preparation are the two significant steps; any issues associated with these steps affect the entire supply chain and reduce product quality and yield. However, due to industrialization and urbanization, effluents are continuously introduced into environment and contaminate soil and water with heavy metals and cause phytotoxicity.1,2 Further, medical and related waste containing antimicrobial contaminants also affect plant growth based on their type and nature.3 Therefore, removing these contaminants from soil and water before the farming stage could increase agricultural yield and food safety.4-7 Nevertheless, the seed germination rate and adaptability to different environments also affect crop yield. In this regard, selecting a suitable seed treatment would also increase seed tolerance against salinity and drought stress.8,9 However, the real challenge arises at the post-harvesting stage, where the harvested fresh edible agricultural produce needs to be treated

to avoid quality deterioration concerning microbes, 10-12 enzymes, 13 insects, and their secondary toxic contaminants. 14 Furthermore, in later stages, most agricultural products reach industries where they will undergo various unit operations before they are stored and distributed to consumers. 15 However, these product rheological properties, microbial/pest safety, functional properties, extract quality, and quantity would be low until these products are treated with suitable processing methods.16-18 Hence, while selecting the processing method/ treatment, we should focus on its ability to address all these issues, irrespective of the product physical nature and intended final use.

Regarding processing, non-thermal technologies are preferred over thermal technologies to avoid undesirable product quality changes. In food processing, non-thermal treatments such as pulsed light, ultrasound, UV, ozone,



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Professor Oliver Schlüter is a spokesman of the ATB program area "healthy foods", vice-head of the Department of Horticultural Engineering, and head of the ATB microbiology lab. His research work focuses emerging technologies in production primary food edible(including insects), minimal processing of food, and non-destructive quality and safety monitoring. He has over

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areas of plasma, developing technologies in both thermal and nonthermal plasma. His current research is on the generation and characterization of plasma-activated water and its applications. He has 6 granted U.S. patents, authored over 40 peer-reviewed articles, 35 conference proceedings, and 2 book chapters.



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impact of process control parameters and physicochemical characteristics of these technologies on biological materials. He has over 250 peer-reviewed research publications. Dr Tiwari is also a fellow of the Institute of Food Science & Technology (UK) and Royal Society of Chemistry (UK).

pulsed electric field, and high-pressure processing technologies are considered alternatives for thermal treatments. As the name suggests, these non-thermal food processing techniques are performed nearly at room temperature to produce safe, nutritive products and retain freshness without inducing thermal damage in the product, 19-21 and cold plasma technology is also one among them. Plasma is created by supplying high energy to gas molecules. The supplied energy dissociates molecules into atoms, and a further increase in the energy breaks those gas atoms into wholly or partially charged ions. Irving Langmuir coined the name plasma for this ionized gas in the early 20th century.22 It is the fourth state of matter and can be generated under atmospheric or vacuum conditions.<sup>23</sup> Photons, electrons, ions, atoms, free radicals, and excited or unexcited molecules are present in this state. Based on energy given to the plasma system and energy transferred to the molecules in plasma, the plasma is classified as equilibrium (thermal) and nonequilibrium (low temperature) plasma.24 One of the critical advantages of this novel technology is that it can be used for both constructive and destructive applications. Constructive approaches include functional modification,25 seed germination, plant growth,26 and quality improvement of extracts,27 while destructive applications include decontamination,28 disinfestation,29 and pesticide degradation.30 This review explains up-to-date research findings of plasma-assisted agrifood processing studies and the mechanisms involved in each process. Further, the article focuses on pre and post-harvest plasma applications specific to the commodities obtained from different plant parts (leaves, roots, fruits, vegetables, nuts, and spices).

# Mechanism of plasma in various applications of agri-food processing

### 2.1. Microbial destruction and inactivation mechanism

Though various plasma-chemical reactions are initiated during plasma generation, O<sub>3</sub>, O<sup>+</sup>, OH<sup>-</sup>, and H<sub>2</sub>O<sub>2</sub> are the primary species responsible for microbial inactivation.31 These RS breaks peptidoglycan (PG) bonds of the cell wall and causes destruction wherein the PG structural bonds like C-O, C-N, and C-C are destroyed by reactive oxygen species.32 It is also suggested that the intense bombardment of the radicals causes surface lesions leading to etching, severely affecting Gramnegative bacteria.33

Firstly, the RS oxidizes lipids, enzymes, and cytoskeletal proteins in the cell membrane and damages the cell wall. In comparison, DNA damage is caused by high-energy UV photons released from excited atoms or molecules at 220 to 280 nm. They form thymine dimers and break plasmid DNA's single and double strands that affect cell replication and other functions.34,35 In addition, RS in plasma increases the solution's acidity and further inactivates microbes.36

### 2.2. Chemical and toxin removal

Interaction of pesticides with plasma RS results in the oxidation of these chemical compounds and degrades them. Apart from oxidation, other intermediate processes also occur during the degradation and finally end in mineralization. 30,37 Pesticide and other organic or inorganic pollutant removal follows different pathways, where pollutants are converted into simple non-toxic components such as CO2, H2O, inorganic carbons, and other organic or inorganic components based on treatment time and reactive species nature. Oxidation, isomerization, and H<sub>2</sub>O/CO<sub>2</sub> removal are some essential reactions that take place during this conversion.38

### 2.3. Insect mortality

The nerve toxin effect caused by plasma ROS on the neuromuscular systems of insects induces mortality. These species enter through the respiratory pathway of insects and affect the insect behaviour and immune and circulatory systems.39-41 Mechanisms for insect mortality vary concerning the insect life stages. Ramanan et al.29 reported the different mechanisms involved in the insect's mortality of egg, larva, and adult stages.

- 2.3.1. Eggs. Mechanisms, such as disruption of embryogenesis due to ROS, eggshell cell membrane disruption due to electrostatic accumulation, and delay or lack of hatching due to anoxic or hypoxic conditions, are the significant causes of
- 2.3.2. Larva. The ROS creates severe oxidative stress in the larva and damages the cuticle and epidermis of the insect. In addition, oxidation also leads to the melanization of hemolymph and deformation of the body, causing the death of larvae.
- 2.3.3. Adult. Electrostatic excitation on insects' membranes due to high voltage discharge affects the nerves and neuromuscular system, which causes anoxia conditions (primary mode) and leads to insect death. Another mode of action (secondary mode) is the destruction of C-H bonds present in the cuticular lipid layer of the insect's surface by ROS. Damages in the insect surface layer cause dehydration and death.

### 2.4. Seed germination

Many factors influence the germination rate of plasma-treated seeds. However, the exact reasons are not clear. However, a few acceptable reasons are etching, the opening of seed coats, increased seed wettability, deposition of small bioactive molecules on the seed coat, and decontamination of seed surface microbes.42,43

### 2.5. Functionality modification

Plasma treatment alters the nature of food constituents and their property. ROS reacts with sulfur and aromatic amino acids in the protein, resulting in oxidation. Sometimes oxidation (carbonylation) also assists enzyme inactivation. In addition, the treatment also changes the secondary structure of proteins and alters their functionality. Further, ROS attacks methyl groups near the double bond regions and oxidizes them in lipids. Meanwhile, carbohydrates, aldose, and ketose form formic acids during oxidation.38

This review discusses the applications of plasma treatment on agri-food processing in various stages to enhance the agricultural production rate, reduce spoilage & waste, and improve the quantity and quality of the final product.

# 3. Plasma applications in pre-harvest stages

Plasma is used in different stages of pre and post-harvest processing of the agri-food chain (Fig. 1), where the pre-harvest operation includes soil irrigation, soil preparation, sowing, planting, *etc.*, and continues till harvesting (Fig. 2). Each stage has a noticeable impact on final product quality and yield. Hence, plasma technology has been used to obtain the desirable change during pre and post-harvest operations.

#### 3.1. Plasma treatment of water

Water is one of the vital sources for agricultural practices and is usually contaminated by microbes, heavy metals, and pharmaceutical compounds. 44-46 Removing these contaminants can improve agricultural product quality and overall production. The reactive species present in plasma can oxidize and remove the contaminants from water or effluent. 47 Furthermore,

plasma-activated water (PAW) can also be used to suppress microbial activity<sup>48</sup> and to remove chemical contaminants from water.<sup>49</sup> The interaction of reactive species with water during plasma treatment and PAW production varies based on the nature of RS (Fig. 3). The degradation of organic and inorganic compounds is shown in eqn (1).

$$\label{eq:chemical compounds + RS } \begin{subarray}{l}{l} \hline \begin{subarray}{l}{l} Chemical compounds + RS \rightarrow simple carbon/nitrogen \\ \hline \begin{subarray}{l}{l} CO_2 + H_2O \\ \hline \end{subarray} \begin{subarray}{$$

Though degradation follows different pathways, the end product will likely fall under any of those components mentioned in eqn (1). For instance, pesticide degradation (diuron) by OH<sup>-</sup> species oxidation releases CO<sub>2</sub> and H<sub>2</sub>O after producing organic acids.<sup>50</sup> Similarly, after producing different intermediate components, dimethoate degraded as non-toxic PO<sub>4</sub><sup>3-</sup> during plasma treatment.<sup>51</sup> In addition to oxidation, mineralization, removal of alkyl groups, CO<sub>2</sub>, H<sub>2</sub>O, halogens, and isomerization could take place during the degradation process. This degradation mechanism is often similar to colourant and off-odour-producing compounds.<sup>38</sup>

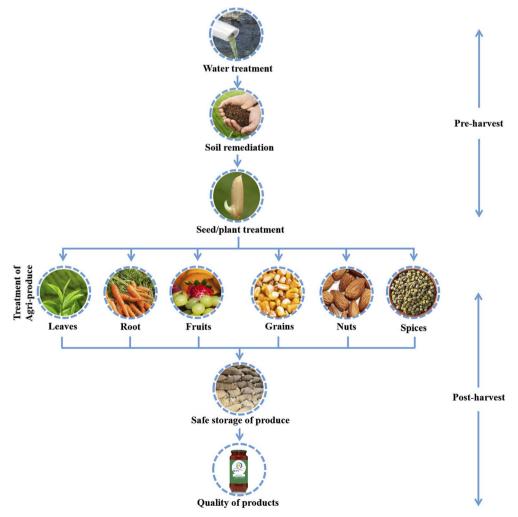
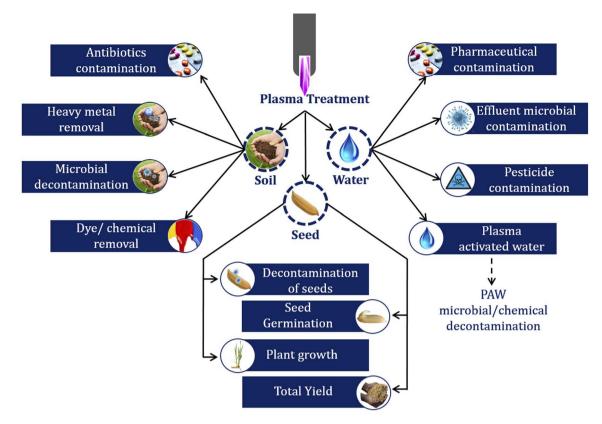
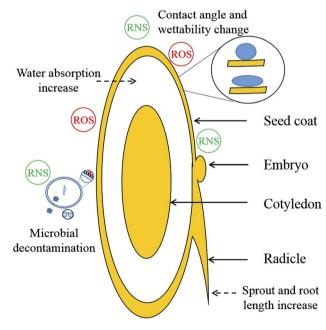


Fig. 1 Application of plasma in different stages of agri-food chain.



Plasma application in pre-harvest stages of agri-food chain.



Effect of plasma treatment on seeds.

Plasma treatment has the potential to degrade individual dyes and their mixtures (i.e., alizarin yellow + orange II + methylene blue) from water.52 Iervolino et al.53 investigated the effect of plasma on different water pollutants by varying

treatment time, power, and gas and identified the lower resistance of methylene blue (MB) and ceftriaxone against plasma (degraded within 5 min). While phenol (15 min), paracetamol (15 min), and caffeine (25 min) took more time for degradation. During this process, MB was almost entirely converted into CO<sub>2</sub>. Instead of focusing on degradation alone, Tampieri et al.54 studied the degradation and mineralization efficiency of plasma-treated rhodamine B, phenol, and metolachlor. For low pollutant concentrations, the highest mineralization efficiency of 59% and 20% was obtained for phenol and metolachlor in 30 min, respectively. To understand the combined effect of plasma with any other non-thermal method, Bradu et al.47 combined ozone treatment with plasma to remove organic pollutants from water. Ozone combined with plasma reduced the treatment time by half to remove 50% chlorophenoxyacetic herbicide (2,4-D) with doubled removal reaction rate. This study explains the role of OH<sup>-</sup> species on 2,4-D oxidation and the importance of catalytic ozonation for faster pollutant removal and mineralization. Similar to other organic pollutants, Sarangapani et al.55 reported pesticide removal concerning power and treatment time. Higher power and exposure time increased pesticide degradation into simple chemical groups due to the oxidization of active O<sub>3</sub> and OH<sup>-</sup> species. Other than organic pollutants and pesticides, microbes can also contaminate water. So, Pavlovich et al.56 reported the effect of plasma on Escherichia coli present in water and found that ozone concentration alone had a higher correlation with the antimicrobial activity of the treatment than pH and other species-generated

Table 1 Plasma applications in pre-harvest stages

Sample	Plasma characteristics	Significant results	Reference
(1) Plasma treatment on water (a) Effluent decontamination			
Dimethyl phthalate	Self-pulsing discharge (SPD) and multipin corona discharge (MCD) Input – 30 kV, 12 mA	$\ensuremath{MCD}$ was more effective in the degradation of dimethyl phthalate than $\ensuremath{SPD}$	184
Methylene blue, ceftriaxone, phenol, paracetamol and caffeine	nL min <sup>-1</sup> ) • Input power – 12 to 38 kV	Degradation and mineralization were higher when $\rm O_2$ was used at 20 kV with a 0.18 nL $\rm min^{-1}$ flow rate	53
Rhodamine B, phenol and metolachlor	Treatment time – 5 to 25 min Atmospheric plasma reactor Gas – air (flow rate – 100 mL min $^{-1}$ )	30 min. Low concentrations were removed completely Rhodamine B and metolachlor were removed in 15 and 20 min,	54
	Input – 5.9 $\pm$ 0.7 W Treatment time – up to 30 min	respectively	47
Chlorophenoxyacetic herbicide (2,4-D)	Pulsed corona discharge with zonation reactor  • Gas – oxygen (0.3 L min <sup>-1</sup> )  • Input power – 11 to 31 W  • Treatment time – up to 60 min	Plasma with ozone provides >99.8% removal of 2,4-D in 30 min of treatment time	47
Pesticide (dichlorvos, malathion, endosulfan)	DBD plasma reactor  • Gas – atmospheric pressure plasma  • Input power – 60, 70, and 80 kV <sub>RMS</sub>	Degradation was maximum for dichlorvos (78.98%), malathion (69.62%), and endosulfan (57.71) at 80 kV $_{\rm RMS}$ after 8 min	185
Bacterial culture mixed water	• Treatment time – 0, 2, 4, 6, and 8 min DBD plasma	E. coli – vortex (5 s) increases the microbial reduction up to 5 log	56
	• Gas – ambient air • Input power – 0.02 to 0.4 W cm <sup>-2</sup> • Treatment time – 30 to 300 s	in 120 s at 0.2 W cm $^{-2}$ Increased power has less effect on microbial reduction	
Polluted water (atrazine, chlorfenvinfos, 2,4- dibromophenol, and lindane)	DBD batch reactor $(R_1)$ and co-axial thin-film DBD reactor $(R_2)$ • Gas – helium at atmospheric pressure (5 L min <sup>-1</sup> ) • Input power – 30 W	$R_{\rm 2}$ was found to be efficient more efficient. For example, 15 min of treatment degraded the pollutant while the byproducts degradation was low	57
Pentoxifylline (in water)	• Treatment time – 0 to 15 min DBD plasma with coaxial configuration	92.5% pentoxifylline reduction after 90 min treatment	45
	• Gas – oxygen (flow rate – 600 SCCM) • Input power – 1.2 W • Treatment time – up to 120 min	Chemical reduction increases with increased $\mathrm{O}_3$ consumption	
Poultry wash water	Pulsed-plasma gas-discharge  • Gas – sulfur hexafluoride (SF <sub>6</sub> )/air (flow rate – 10 L min <sup>-1</sup> )  • Input power – 23.5 kV  • Treatment time – 30 s	13	46
Methylene blue (MB) water	DBD plasma with coaxial configuration	Plasma treatment of 30 min with $O_2$ gas removed 95% MB dye	186
	<ul> <li>Gas – oxygen and air (300, 600, and 900 SSCM)</li> <li>Input power – up to 1 W</li> <li>Treatment time – up to 90 min</li> </ul>	Ozone concentration influences the removal rate	
(b) PAW treatment Plasma treated water (PTW)	DBD plasma Gas – air Input – 11 W Electrode gap – 3 mm Treatment time – 1 to 10 min	Mortality rate of PTW, tap water, and reference sample after 24 h was around 85, 8 and 2% for mealybug	40

Table 1 (Contd.)

Sample	Plasma characteristics	Significant results	Reference
PTW	• Gas – atmospheric air • Input power – 3.14 W	PTW used sausage with nitrite (782 ppm) that can provide a curing effect	58
Plasma-activated water (PAW)	<ul> <li>Treatment time – 120 min</li> <li>Indirect DBD plasma</li> <li>Gas – atmospheric air</li> </ul>	Anti-microbial declined during storage After seven days, 2.4 logs and no reduction were observed in 3 h and 15 min treated PAW, respectively	59
PAW	<ul> <li>Input power - 5 W</li> <li>Treatment time - 20 min</li> <li>Atmospheric non-thermal quenched plasma</li> <li>Gas - atmospheric air (flow rate -</li> </ul>	Acidification by nitrites in PAW produces a lethal effect on ${\it Hafnia}$ alvei	60
PAW	550 L h <sup>-1</sup> )  • Treatment time − 5 min  Atmospheric pressure gliding arc plasma  • Gas − atmospheric air	S. cerevisiae – 3 log reduction after 30 min Without solid substrate H. alvei, Leuc. mesenteroides, Staph.	48
	• Input power – 1.2 W • Treatment time – 5 min	Epidermidis were eradicated after 30 min	
(2) Plasma treatment on soil remo	ediation		
. ,		54.2% of PNP was degraded after $50~s$ discharge treatment	187
Fluorene	Needle-plate pulsed corona discharge plasma Input – 30 kV	78.7% fluorene degradation was achieved after 60 min When washing is done to remove oxidation products, 99%	64
Non-aqueous phase liquid	Treatment time – 60 min Electrode gap – 20 mm Ex situ DBD plasma cylinder-to-	degradation achieved in 45 min  High energy density eradicates NAPL irrespective of its initial	65
(NAPL)-mixed soils	plane reactor • Gas − atmospheric air (flow rate − 1 L min <sup>-1</sup> )	concentration of it	
Dye-polluted soil (acid scarlet GR)	• Input power $\sim$ 25 W (energy density 675 to 10 125 J g $^{-1}$ ) • Treatment time – 2.5 to 33 min Plane-to-plane DBD plasma • Gas – atmospheric air • Input power – 15.6 to 19.6 kV (3.51	(17.6 kV) after 25 min treatment	66
<i>p</i> -Nitrophenol polluted soil (PNP)	0 1	Soil depth increase reduces PNP degradation	70
	0.5 L min <sup>-1</sup> ) • Input power – 23 kV	Quartz sand containing PNP degraded more than sand and sandy soil. Clay soil has the lowest degradation rate	
Contaminated soil	• Treatment time – 45 min Ozonizer combined DBD plasma with TiO <sub>2</sub> -based surface discharge • Gas – oxygen (flow rate – 0.5 dm³ min <sup>-1</sup> ) • Input power – 23 kV	The microbial load of ozone treated soil reduced from $5.7\times 10^6$ to $1.7\times 10^2$	72
Chloramphenicol polluted soil	<ul> <li>Treatment time – 45 min</li> <li>Atmospheric pressure DBD plasma</li> <li>Gas – oxygen, ozone, air, argon,</li> </ul>	The presence of oxygen improves the degradation compared to air Higher input voltage and flow rate increase the degradation	67
	nitrogen (flow rate – 0.15 to 1.5 L min <sup>-1</sup> )	efficiency	

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Table 1 (Contd.)

Sample	Plasma characteristics	Significant results	Reference
	• Input power – 16.4, 18.4, and 20.4		
	kV • Treatment time – 0, 5, 10, 15, 20,		
	and 25 min		
<i>p</i> -Nitrophenol polluted soil (PNP)		Plasma alone caused 78.1% of PNP degradation, while PDPTC	68
	<ul><li>catalytic (PDPTC)</li><li>Gas – atmospheric air (flow rate –</li></ul>	caused 88.8% degradation at 20 kV (10 min)	
	0.5 L min <sup>-1</sup> )		
	• Input power – 20 kV		
Pentachlorophenol (PCP) in soil	• Treatment time – 30 min Pulsed corona discharge plasma	An increase in voltage from 12 to 18 kV increased PCP	69
remuemorophenor (r or) m son	• Gas - oxygen, air, argon, nitrogen	degradation from 64 to 90% (60 min)	03
	(flow rate – up to 8 L min <sup>-1</sup> )		
	<ul> <li>Input power - 12 to 18 kV</li> <li>Treatment time - up to 60 min</li> </ul>		
Pentachlorophenol (PCP)	Pulsed corona discharge plasma	11.6, 13.6, 15.6 kV, and 17.6 kV degraded PCP up to 62, 77, 83,	71
	(PCDP)	and 87%, respectively, after 60 min	
	<ul> <li>Gas – air (flow rate – 3 L min<sup>-1</sup>)</li> <li>Input power – 11.6 to 17.6 kV</li> </ul>		
	• Treatment time – up to 60 min		
Kerosene polluted soil	Cylinder-to-plane DBD plasma	An energy level of 960 J g <sup>-1</sup> degrades the kerosene by 90%	188
	<ul> <li>Gas – air (flow rate – 1 L min<sup>-1</sup>)</li> <li>Input power – 15 to 20 kV</li> </ul>		
	• Treatment time – up to 60 min		
Contaminated soil	DBD plasma	Ozone produced during the process reduced the microbial count	62
	• Gas – oxygen and air (flow rate – 1.	and pH Increased the mineral content of the soil	
	2 L min <sup>-1</sup> )		
	• Input power – up to 30 W		
	• Treatment time – 60 min		
(3) Plasma treatment of seeds/pla			
Rice seedling	Non-equilibrium atmospheric pressure plasma	15% increase in grain yield when the plant was treated in direct plasma during the early period and no change in the later stage	
	Plasma activated ringer's lactate	passing data outly period and no change in the accessage	
	solution		
	Treatment period of solution – twice a week		
	Input – 60 Hz, 9 kV		
Maize grains	Diffuse coplanar surface barrier	Wettability increased – the amount of water imbibition was	190
	Input – 400 W with air, oxygen, and	higher 90 and more had a negative effect on growth and production	
	nitrogen	indices	
	Treatment time – 0, 30, 60, 90, 120,		
Buckwheat seeds	180, and 300 s Low-pressure radiofrequency	A shorter exposure time of up to 45 s was safer for the application	191
	system	•	
	Input - 30 W	The longest exposure of 120 s affected the fungal colonisation	
	Treatment time – 15, 30, 45. 60, 90, and 120 s		
Carrot seed	DBD	3 min had a great quantity of flavonoids, phenolics, and plant	192
	Input – 12.5 kV and 50 Hz	extracts had high radical scavenging activity	
Pea seeds (Pisum sativum L.)	Treatment time – 1 to 4 min DCSBD plasma unit with a silver	DNA damage – 240 s treatment reduces the damage by 28 to	90
	•	30.5%, 3.5%, and 21% for zeocin, distilled water, and hydrogen	
	material	peroxide treated sample, respectively	
	<ul> <li>Gas – atmospheric air (flow rate –</li> <li>3 L min<sup>-1</sup>)</li> </ul>		
	• Input power – 400 W		
Wheat	• Treatment time – 60 to 300 s	Plagma treatment of 15 and 00 a in angent 1 trops (1 tr	01
Wheat		Plasma treatment of 15 and 90 s increased HSFA4A by six and four folds after 3 h in the root	91

Table 1 (Contd.)

Sample	Plasma characteristics	Significant results	Reference
	Land on W	However, for the shoot after 6 h, a ten-fold increase was observed in the 30 s treatment	
ı	Input – 80 W Treatment time – 15, 30, 60, and 120 s		
Wheat	Atmospheric DBD plasma Gas – dry air Input – 13 kV	After treatment, GP and GR in drought stress increased from 39.3 to 50.0% and 62.7 to 80.0%, respectively	92
Radish (Raphanus sativus)	Treatment time – 4 min Double DBD reactor with plate-to- plate configuration for seed treatment	Seed GR was 40, 60, and 100% for control, PAW-15 and PAW-30, respectively $$	42
		In tap water (TW), growth was higher for control, P10, and P20 tomato plant than PAW watered plants	
	Input – 57 mJ per pulse (max) Treatment time – 10 (P10) and 20 (P20) min Double DBD reactor with cylindrical configuration – water treatment Gas – synthetic air (flow rate – 1 L min <sup>-1</sup> ) Input – 7 mJ per pulse (max) Treatment time – 15 (PAW-15) and	More growth in PAW-30 watered plants than TW	
Rapeseed ( <i>Brassica napus</i> L.)	• Input power – direct current of 20 kV (1.5 A)	Germination was 7.7% higher after 1 min of treatment The microbial load of the treated sample was maintained lesser than the control sample (1–2 log reduction)	86
Wheat seed	• Treatment time – 0 to 3 min DBD plasma Gas – air (flow rate 1.5 L min <sup>-1</sup> ) Input – 1.5 W Treatment time – 0, 1, 4, 7, 10 and 13 min	GP, GR, and GI increased in 4 min treatment from 62.5 to 77.5%, 88.0 to 95.3% and 36.7 to 41.0% compared to control	74
Broccoli seed (Brassica oleracea	CDPJ generation system  Gas – atmospheric air	Plasma reduced microbes in seed and sprout Less exposure increased germination rate, sprout length, and weight	75
Peanut ( <i>Arachis hypogaea</i> L.) seed	<ul> <li>Gas – helium (150 Pa pressure)</li> <li>Input power – 60, 80, 100, 120, and 140 W</li> </ul>	After 15 s treatment at 80, 100, and 120 W, germination potential increased by 128, 128, and 150%. Finally, at 120 W germination rate was found to be maximum	78
Wheat seeds ( <i>Triticum aestivum</i> L. cv. Eva)	<ul> <li>Treatment time – 15 s</li> <li>DCSBD – cold atmospheric pressure plasma (CAPP)</li> <li>Gas – atmospheric air</li> <li>Input power – up to 100 W cm<sup>-3</sup></li> </ul>	Water uptake increased with soaking time and dosage  Bacteria and fungi load on seeds reduced with treatment time	76
Wheat seeds (Triticum aestivum)	<ul> <li>Treatment time - 10 to 600 s</li> <li>Atmospheric pressure surface discharge plasma reactor with glass plate</li> <li>Atmospheric air (flow rate - 1</li> </ul>	1 ( )	77
Peas (Pisum sativum 'Salamanca')	L min <sup>-1</sup> ) • Input power – 2.7 W • Treatment time – 5, 15 and 30 min CAPP – surface dielectric-barrier air-discharge • Atmospheric ambient air • Input power – 9 kV <sub>pp</sub>	5 and 10 min CAPP treatment increased germination to 42 and 50%, respectively, after 24 h Germination rate reached a maximum (5 min treatment) of 65% after 48 h	82

Table 1 (Contd.)

Sample	Plasma characteristics	Significant results	Reference
	• Treatment time – 1 to 10 min		
Bean ( <i>Phaseolus vulgaris</i> )	Cold radio frequency air plasma	Water absorption increased in plasma treatment. hence,	85
		germination time reduce in plasma-treated samples	
	$10^{-2} \text{ Pa}$		
	• Input power – 20 W		
	• Treatment time – plasma		
	treatment (2 min), vacuum		
Doe goods	treatment (3 min) Coplanar surface discharge – low-	Sand commination in proceed from 77.5 to 0.50/ often 100 c	0.4
Pea seeds	temperature plasma	Seed germination increased from 77.5 to 95% after 120 s treatment	84
	• Input – 370 W	treatment	
	• Time – 60 to 600 s		
Pre germinated rice ( <i>Oryza</i>	Plasma jet with quartz tube covered	Control – 97% germination	87
sativa L.)	inner electrode		
,	<ul> <li>Argon and oxygen</li> </ul>	Treated - 93 and 91% after 5 s at 10 W power with 8, 5 mm	
		distances, respectively	
	• Input power – 10–14 W	High $\gamma$ -aminobutyric acid (GABA) content achieved in 96 h pregermination	
	• Treatment time – 5 to 10 s		
	• Distance – 5, 8 mm		
Soybean (Glycine max (L.) Merr)	CDPJ generation system	Maximum germination (G) potential, G rate, G index, and vigour	81
	4	index were observed after 80 W treatment	
	• Dry atmospheric air (2.5 m s <sup>-1</sup> )	Water uptake increased by 14.03%	
	• Input power – 0 to 120 W		
Transata (Calanna	• Treatment time – 15 s	Commingation not entirely and note in succeed has 0 and 110/	0.0
Tomato (Solanum		Germination potential and rate increased by 8 and 11%	80
Lycopersicum L.) seed	<ul><li>Helium (150 pa)</li><li>Input power – 80 W</li></ul>	Bacterial wilt severity was reduced by 25% in the treated sample	
	• Treatment time – 15 s		
Wheat (Triticum spp.)	Cold helium plasma	Germination potential and the rate increased by 6, and 6.7%,	26
vinede (17.000mm SPPI)	• Helium (150 Pa pressure)	respectively, after 80 W treatment	
	• Input power – 60, 80, 100 W	J,	
	• Treatment time – 15 s		
Poppy seed (P. somniferum L.)	Panasonic – microwave generating	Germination was 115% in 3 min treatment	89
	RF plasma		
	Input – 500 W	Plasma treated seedlings were longer than the control seedlings	
	• Time – 180 to 5400 s		
	• Gas – O <sub>2</sub> (50 mL min <sup>-1</sup> ) and Ar (50 ml min <sup>-1</sup> )		
Maize seed (Zea mays L.)	DCSBD plasma treatment	Plasma treatment for 60 s increased the root length, root fresh, and dry weight by 21, 10, and 14%, respectively	83
	• Atmospheric air	Soluble protein content in the root increased after 60 s treatment	
	• Input power – 370 W		
	• Treatment time – 60 and 120 s		
Paulownia tomentosa seeds	Glow discharge plasma treatment	Air plasma treatment at 50 W, 100 W and 200 W produced maximum germination at 15, 5, and 1 min, respectively	88
	• Air and argon (200 mTorr)	When glass plates covered the seeds, germination was reduced	
	• Input power – 50, 100 and 200 W		
	• Treatment time – 1 to 40 min		

acids. This study concluded that 5 s plasma treatment was enough to reduce  $E.\ coli$  to a not detectable level and suggested the importance of mixing on the antimicrobial effect of plasma species. In order to evaluate the effect of plasma on the removal of different pollutant mixes, Hijosa-Valsero  $et\ al.^{57}$  conducted a study where atrazine, chlorfenvinfos, 2,4-dibromophenol, and lindane pollutants were mixed with water and treated in DBD batch reactor  $(R_1)$  and co-axial thin-film DBD reactor  $(R_2)$ .

Results showed a decrease in the degradation efficiency of chemicals when their concentration increased, and even after pollutant degradation, byproduct removal took more treatment time. Similarly, pentoxifylline and its intermediates removal were analyzed by Magureanu *et al.*,<sup>45</sup> and the author reported that removal was high at a lower concentration of pollutant (100 mg L<sup>-1</sup>), higher power, high pulse rate, and higher frequency. Intermediate components produced during this

Table 2 Plasma applications in post-harvest stages

Sample	Plasma chamber	Purpose of study	Research goal	Reference
(1) Plasma treatment	of different agricultural produces			
<i>(a) Plasma treatment (</i> Baby spinach	on leaves High voltage atmospheric cold plasma	Inactivation of	Significant inactivation after 24 h through 14	102
	Input – 30–130 kV	Salmonella enterica and E. coli O157:H7	days post-treatment After 7 days of refrigerated storage – 2.6 log CFU per g – 2 min	193
Date palm leave	Treatment time – 2 min and 5 min Radiofrequency plasma Input – 80, 100, 120 W, gas pressure – 0.95 torr	Surface modification	3.5 log CFU per g – 5.0 min Wettability of leaf surface increased Surface roughness was increased due to the removal of the waxy layer and impurities	194
Green tea leaves	Treatment time – 1, 5, and 8 min DBD	Antioxidant activity	15 W and 15 min, the TPC and antioxidant activity increased by 41.14% and 41.06%	195
	Input - 5, 10 and 15 W		Catechin increased by 103.12%	
	Treatment time – 5, 10, 15 min Dielectric barrier discharge (DBD)	Microbial inactivation	<i>L. monocytogenes</i> – 2.20 log CFU per cm <sup>2</sup> reduction after 30 min	94
,	<ul><li>Atmospheric air</li><li>Input power – 15 kV</li></ul>		E. coli O157:H7 – 1.35 log MPN per cm <sup>2</sup> reduction after 15 min	
	Treatment time – 15 to 30 min			
	Plasma jet with the copper electrode and Pyrex tube • 99.99% argon (flow rate – 1 L min <sup>-1</sup> )	Microbial inactivation	E. coli – complete removal in black and green tea after 3 and 4 min, respectively Coliform – inactivated after 5 min in both	95
	• Input power – 10 kV		samples Yeast and mould – inactivated after 7 min in both the samples	
	• Distance – 1.5 cm		both the samples	
	• Treatment time – 0 to 7 min	a characteristic and a contraction	Total do main transfer and do college a	0.2
(Lactuca sativa L. var. longifolia)	Atmospheric pressure plasma with nickel coated steel needle array  • Argon (flow rate – 455.33 standard	Microbial inactivation	E. $coli$ – 10 min treatment at 12.83 kV reduces the microbial count to 1.5 logs	93
	cm³ min <sup>-1</sup> )  • Treatment time – 30 s to 10 min			
	• Input power −3.95 to 12.83 kV			
	Atmospheric pressure plasma jet • Argon (flow rate – 20 L min <sup>-1</sup> ) • Input power – 10, 20, 30, and 40 W	Microbial inactivation	The effect of plasma on microbes was higher (3.6 log reduction) at lower level surface contamination (10 <sup>4</sup> CFU per mL) after 15 s	97
		Bioactive compounds	treatment at 20 W Chlorogenic acid, caffeic acid luteolin diosmetin content were reduced as time	98
	<ul> <li>Input power – up to 35 W</li> <li>Treatment time – 0 to 120 s (overall)</li> </ul>		increased at 30 W plasma treatment	
Lamb's lettuce ( <i>Valerianella locusta</i> )	Low-pressure oxygen glow discharge plasma	Flavonoid content	Flavonoid content increased in freeze-dried leaves after 120 s exposure	99
	<ul> <li>Oxygen (0.5 mbar)</li> <li>Input power - 75 and 150 W</li> <li>Treatment time - 20 to 300 s</li> </ul>			
(b) Plasma treatment		actional to Literature	The Mark the state of the state	0.2
(Daucus carota L.)	oated steel needle array • Argon (flow rate – 455.33 standard	Microbial inactivation	E. coli – inactivation was less than 0.5 log	93
	cm³ min <sup>-1</sup> ) • Treatment time – 30 s to 10 min • Input power – 3.95 to 12.83 kV			
(c) Plasma treatment o	on fruits			105
•	DBD Input – 7% and 14% duty cycle	Decontamination	14% duty cycle and 20 min effective 1.46 log CFU per g – total aerobic mesophilic bacteria	196
	Treatment time – 5, 10, and 20 min DBD	Enzymatic browning	2.75 log CFU per g – yeast and mould Residual activity of litchi peroxidase decreased	197

Table 2 (Contd.)

Sample	Plasma chamber	Purpose of study	Research goal	Reference
Blueberry	Atmospheric cold plasma Input – 12 KV, 5 kHz	Improving antioxidant activity and microbial inactivation	Reduced the decay rate of blueberries Antioxidant activity increased but maintained at a low ROS level	198
Cavendish banana	Treatment time – 0, 30, 60, 90 s DBD	Post-harvest crown rot	60 s is the best exposure time 0.5 min optimum duration	199
	Input – 15 kV, compressed air		Colletotrichum musae inhibition – 51.89% for 0.5 min	
Tomatoes	Treatment time – 0.5, 1, 2, and 3 min DDP with aluminium-coated electrodes	Pesticide degradation	Chlorpyrifos reduced up to 89.18% at 5 W in 6 min	37
	• Gas – atmospheric air (50–75 mmHg)		Lycopene (84.86 to 15.38 mg kg <sup>-1</sup> ), beta carotene (22.82 to 6.61 mg kg <sup>-1</sup> ) and firmness were reduced	
	• Input power – 2 to 5 W			
Blueberries	Treatment time – 4 to 6 min Circular aluminium electrodes with 2 and 10 mm thick perspex dielectric material at ambient temperature (25 $\pm$ 2 °C) • Atmospheric air	· ·	Plasma treatment of 5 min at 80 kV degraded pesticides (75% of boscalid and 80% of imidacloprid)	104
	<ul> <li>Input power - 60 and 80 kV</li> <li>Treatment time - 0, 2, and 5 min</li> </ul>			
Strawberries	DBD atmospheric CP with Al electrode and polypropylene dielectric material (in pack treatment)	Microbial inactivation	$G_1$ and $G_2$ produced 3.1, 3.4 and 3.7, 3.3 log reduction of mesophiles, yeast/mould, respectively	100
	<ul> <li>G<sub>1</sub> - O<sub>2</sub>: N<sub>2</sub>: CO<sub>2</sub> of 65: 16: 19 ratio</li> <li>G<sub>2</sub> - N<sub>2</sub>: O<sub>2</sub> of 90: 10 ratio</li> <li>Treatment time - 5 min</li> <li>Input power supply 60 kV<sub>rms</sub> voltage (50 kg)</li> </ul>			
Cocktail tomatoes	Hz) Atmospheric pressure plasma with nickel	Microbial inactivation	E. coli – 12.83 kV input reduced microbial	93
(Lycopersicon lycopersicum)	• Argon (flow rate – 455.33 standard cm <sup>3</sup> min <sup>-1</sup> )		count to 1.7 log after 10 min	
	• Treatment time – 30 s to 10 min • Input power – 3.95 to 12.83 kV			
(d) Plasma treatment Peanut	on nuts  Novel CAP with rotary jet system	Inactivation of fungi	Aspergillus flavus – not detected at 180 W for	200
	Input – 180 W, 200 W	Ü	7.5 min and 200 W for 5 min  Aspergillus niger – not detected at 180 W for	
	Treatment time – 5, 7.5, 10 min		10 min and 200 W for 5 min Aflatoxin concentration after 29 days of	
Pistachio nut	DBD • Air plasma system	Decontamination of aflatoxin	storage – 16.5 ppb 4 log reduction after 180 s of treatment Maximum reduction of 52.42% of AFB 1 after	201
	• Input – 130 W, 20 kHz, and 15 KV • Treatment time – 15, 30, 60, 90, 120, 150, 180 s		180 s	
Pistachio nut	Cold plasma jet Input – 10, 15 and 20 kV and agon/gas	Pest management	Plasma exposure time – 14.04 min, voltage – 19.99 kV and Ar/air ratio – 51.65% caused	202
Dried walnut kernels	ratio – 0, 50 and 100 Radiofrequency low-pressure cold plasma • Air plasma	Microbial load reduction	increased mortality of <i>Plodia interpunctella</i> The highest log reduction occurred at 50 W 1.09 log CFU per g – total viable count, 0.97 log CFU per g – coliform	203
	• Input – 20, 30, 40 and 50 W		0.89 log CFU per g – mold	
Cashew nut	• Treatment time – 10, 15 and 20 min Low-pressure plasma-glow discharge Air plasma	Allergenicity	Did not affect the allergenicity Improved lipid extractability	204
Almond slices	Input – 80 W and 20 kHz Atmospheric argon plasma Input – 17 V, 2.26 A current Treatment time – 5, 10, 15, and 20 min	Surface disinfection	Anacardic acid content was higher 20 min treatment was effective 2.29 log CFU per g – total count 1.81 log CFU per g – yeast and mould 2.72 log CFU per g – S. aureus	205

Table 2 (Contd.)

Sample	Plasma chamber	Purpose of study	Research goal	Reference
Whole peanut (WP) and defatted peanut		Allergen reduction	The protein intensity of the Ara h 1 band was unaffected by plasma treatment	115
flour (DPF)	Input – 80 kV Gas – air		With increased time, Ara h 1 band intensity for IgE binding decreased for WP and DPF	
	Treatment time – 0 to 60 min		Antigenicity was reduced by 44 and 9.3% for DPF and WP after 60 min	
Tiger nut milk	DBD Input – 1.22 A and 30 V	Microbial inactivation	Undetectable microflora in 12 min pH reduction after 8 and 12 min protein	206
Peanut ( <i>Arachis</i> hypogaea L.)	Treatment time – 2, 4, 6, 8, and 12 min Coplanar – multi hollow surface DBD unit	Quality characteristics	decreased beyond 4 min Oleic and linoleic acid content was reduced from 43.47 to 35.74% and 32.56 to 24.49% for fresh and treated peanuts	114
	• Atmospheric air with 20 to 30% humidity (flow rate – 0.5 to 20 L min <sup>-1</sup> )		Input power, treatment time, and gas mixer used in the study directly affected peanut quality	
	• Input power – 10 to 40 W Treatment time – 1 to 15 min			
Unpeeled almonds	DCSBD plasma Input – 350 W Gas – dry air, $O_2$ , $N_2$ , $CO_2$ and $CO_2$ /Ar mix $(90\% + 10\%)$	Microbial decontamination	Air, $O_2$ , and $N_2$ plasma reduced microbial count by 5, 4.8 and 2 logs after 15 min	107
	Gas flow rate – 0.8 L min <sup>-1</sup> Treatment time – 15 min			
Hazelnuts	Atmospheric pressure fluidized bed plasma (APFBP) reactor Input – 460 to 655 W	Aflatoxin removal	1st reactor: <i>A. flavus</i> and <i>A. parasiticus</i> count reduced by more than 4 log in 5 min at 655 W 2nd reactor: both microbes count reduced by	
	Gas – dry air and $N_2$ (flow rate – 3000 L $h^{-1}$ ) Treatment time – 1 to 5 min		more than 3 log A similar trend was followed in both reactors when $N_2$ gas was used	
	1st reactor – 49 mm diameter, 2nd reactor – 65 mm diameter			
Walnut fruits (J. Regia L.)	Plasma jet with a copper electrode and Pyrex tube • Argon (1 L min <sup>-1</sup> ) • Input power – 15 kV DC supply	Microbial decontamination	The fresh and dried walnut microbial load was removed after 10 and 11 min treatment, respectively, for all walnut types	109
	• Treatment time – 3 to 11 min			
Hazelnuts	DBD plasma  Input – 0.4 to 2 kW	Aflatoxin removal	Pure $N_2$ or $N_2$ with $O_2$ (0.1%) provided higher toxin removal  After 12 min exposure, residual AFB <sub>1</sub> and AFs	
	•		in nuts were 29.1 and 30.4% at 1150 W	
	Gas – pure $N_2$ or $N_2$ and air mix (flow rate – 120 L min <sup>-1</sup> with 7 bar) Treatment time – 1 to 12 min			
Hazelnuts	Distance – 50 mm APFBP reactor	Microbial	At 655 W (25 kHz) within 1 min exposure, 2 log	110
	Input – 460 to 655 W Gas – dry air (flow rate – 3000 L h <sup>-1</sup> ) Treatment time – 1 to 5 min	decontamination	reduction was achieved, and the natural flora was reduced by 3.45 log in 2 min	
Almonds	Gliding arc plasma • Input – 590 W • Gas – air and nitrogen (60 psi) • Time – 0, 10, and 20 s	Microbial decontamination	Plasma reduced 1.34 log CFU per mL of $\it E.~coli$ O157:H7 C9490 after 20 s treatment with 6 cm distance	
Hazelnut, peanut, and pistachio nut	• Distance – 2, 4, 6 cm Low-pressure CP (LPCP) sterilization unit	Microbial decontamination	In hazelnuts, $SF_6$ plasma reduces <i>D</i> -value (1.1 min) compared to air plasma (4.2 min)	113
	• Air or sulfur hexafluoride (SF <sub>6</sub> )		5 times more log reduction was observed in $SF_6$ than in air plasma (5 min)	
	<ul> <li>Input power - 300W</li> <li>Treatment time - 5 to 20 min</li> </ul>			
Almonds	DBD plasma Input – 16 to 30 kV	Microbial decontamination	4 to 5 log reduction in $E$ . $coli$ was observed in all almond varieties after 30 min	112

Table 2 (Contd.)

Sample	Plasma chamber	Purpose of study	Research goal	Reference
	Discharge gap – 10 mm			
(e) Plasma treatment Saffron	on spices Low-pressure cold plasma technology Input – 70, 90, and 110 W Treatment time – 5, 10, 15, and 30 min	Decontamination	110 W for 30 min – high microbial reduction 3.52 log CFU per g – TVC 4.62 log CFU per g – coliforms 2.38 log CFU per g – mold	207
Black pepper corn	Low-pressure cold plasma technology	Decontamination	4.12 log CFU per g – yeast 2.50 W for 20 min showed the highest decontamination	208
	Input - 150 W and 250 W		0.88 log CFU per g – aerobic plate	
Black pepper	Treatment time – 10 min and 20 min DSCBD – DBD	Decontamination	3.66 log CFU per g – yeast and mould <i>B. subtilis</i> count reduced by 5.06 log after 5 min exposure	117
	Input – 400 W		B. subtilis spores, E. coli and S. Enteritidis D – values were 142, 47 and 58 s	
	Gas – ambient air			
Saffron	Treatment time – 60 to 300 s Plasma jet	Bioactive compound	Safranal content was reduced by 14% and 21%	118
our or	Input – 8 and 12 kV	Broaten't compound	at 8 kV and 12 kV, respectively However, isophorone (4 and 8%) and 4- ketoisophorone (2 and 6%) increased by 8 min Ar plasma at 8 kV and 12 kV, respectively	
	Gas - Ar with 5, 10, and 20% O2 (flow rate		, , ,	
Red pepper flake	<ul> <li>1 L min<sup>-1</sup>)</li> <li>Treatment time – 6 and 8 min</li> <li>Microwave integrated CP treatment with high (HMCPT) and low microwave density</li> </ul>		1.8 and 0.6 log per cm <sup>2</sup> reduction was observed in spore count in HMCPT and	119
	(LMCPT) Input – 900 W		LMCPT Based on particle size, the microbial reduction was more for flakes (1.4 and 2.7 log per cm <sup>2</sup> ) than particles (0.8 and 1.2 log per cm <sup>2</sup> ) in LMCPT and HMCPT	
	Gas – He (flow rate – 1 std L $min^{-1}$ with 0.7 kPa)		Estati i una ristati i	
Onion ( <i>Allium</i> cepa L.) powder	Treatment time – 20 min Microwave-powered CP treatment with HMCPT and LMCPT	Microbial reduction	B. cereus spores reduction was more in HMCPT (2.1 log per cm <sup>2</sup> ) than LMCPT (1.2 log per cm <sup>2</sup> )	120
	Input – 400 to 900 W		For microbial reduction, vacuum air drying of the sample was superior to hot air drying	,
	Gas – He (flow rate – 1 L min <sup>-1</sup> at 0.7 kPa) Treatment time – 10 to 40 min			
Onion powder	DBD – CP treatment Input – 9 kV Treatment time – 20 min	Microbial reduction	E. coli and L. monocytogenes count reduced by more than 1 log after 20 min at 9 kV, while more than 2 log reduction was observed in S. Enteritidis	121
Whole black pepper	Radiofrequency $(G_1)$ and microwave $(G_2)$ generated plasma jet	Microbial reduction	Entertitis $G_2$ plasma was more effective against aerobics and spores than $G_1$ plasma	28
	<ul> <li>G<sub>1</sub> used argon (10 L min<sup>-1</sup>)</li> <li>G<sub>2</sub> used air (18 L min<sup>-1</sup>)</li> <li>Input power - 30 W (G<sub>1</sub>), 1.2 kW (G<sub>2</sub>)</li> <li>Treatment time - 15 min (G<sub>1</sub>) and (G<sub>2</sub>)</li> <li>30 min</li> </ul>			
Black pepper	Atmospheric pressure plasma jet (APPJ)  • Argon plasma (flow rate – 20 L min <sup>-1</sup> )  • Argon (0.5 L min <sup>-1</sup> ) + CO <sub>2</sub> (20 L min <sup>-1</sup> )  plasma 0.5 (argon) + 20 (CO <sub>2</sub> ) 0  • Air plasma (20 L min <sup>-1</sup> )  • Argon + H <sub>2</sub> O plasma (20 L min <sup>-1</sup> )  • Input power – 280 V (8 A)	Microbial reduction	Argon and $CO_2$ gas mixer plasma treatment inactivated <i>B. subtilis</i> after 5 min of treatment	

Table 2 (Contd.)

Sample	Plasma chamber	Purpose of study	Research goal	Reference
	Fluid bed APPJ treatment  • Air and argon (flow rate – 20 and 14 L min <sup>-1</sup> , respectively)  • Treatment time – 0 to 80 s	Microbial reduction	Microbial reduction of 1.5 and 5 log reduction of $Salmonella$ was observed after 20 and 80 s of treatment	
	RF – low-pressure cold plasma	Microbial reduction	$\rm H_2O_2$ usage (60 min) reduced microbial count by 1 to 3 logs	124
	Input – 300 and 400 W		After 60 min, Ar plasma caused higher microbial reduction than air, $O_2$ and $N_2$ plasma (for all microbes)	
	Gas – $O_2$ , $N_2$ , air, and Ar (0.3 and 9 mbar) Treatment time – 15 to 60 min			
(Capsicum	Microwave powered CP	Microbial reduction	20 min plasma treatment at 900 W reduced Aspergillus flavus ( $N_2$ gas at 667 pa) by 2.5 logs	125
, -	• Air and argon (267 to 26 680 pa)		Bacillus cereus required heating for inactivation	
	<ul> <li>Input - 300 to 900 W</li> <li>Treatment time - 0 to 80 s</li> </ul>			
(2) Plasma treatment (a) Grain safety and q	t for safe storage, extraction and quality en	nhancement		
	DBD	Degradation of	50 kV and 5 min degraded DON by 83.99%	209
	Input – 30, 40, and 50 kV	deoxynivalenol	55 KV and 5 mm degraded DON by 65.39%	209
	Treatment time – 4, 8, 12, and 16 min	aconginitatenoi		
	Atmospheric cold plasma	Increased storage period to study the occurrence	In vishal – 40 W, 15 and 20 min – grain damage – 1 to 3%	210
	Input – 40, 50, 60 W Treatment time – 10, 15, and 20 min	of Callosobruchus chinensis	In other samples, the damage was 0%	
Canola grain, canola meal and barley grains	ACP DBD and ACP-jet	Degradation of Zearalenone	ACP-DBD – 3 min degradation was 91.6%, 83.2%, and 64.8% for canola grain, canola meal, and barley grain	211
	• ACP – DBD – input – 0–34 kV, 1 A and 300 W, 3500 Hz, duty cycle – 70% and output pulse – 10 µs		ACP jet – 85% Ar + 15% $O_2$ – high degradation	
	• ACP-jet – 1500 Hz, voltage – 0–22 kV, 70% duty cycle, 10 µs			
	And 0–0.025 A with 75% Ar + 25% $N_2$ • Treatment time – 0.5, 1, 3, 5, and 15 min			
·	Atmospheric cold plasma (DBD) • Input – 0–34 kV, 1 A, and 300 W, 3500 Hz, duty cycle – 70% and output	Removal of deoxynivalenol	6 min – 48.9% and 10 min – 54.4%	212
	pulse – 10 μs			
	• Treatment time – 0, 2, 4, 6, 8, and 10 min DBD plasma jet		<i>T. castaneum</i> mortality for Ar, Ar + $N_2$ (20 sccm) and Ar + $N_2$ (80 sccm) gas mixture were 90, 70 and 46%, and for <i>T. confusum</i> 93%, 63% and 57%, respectively. A similar trend was observed for $O_2$ admixture	126
	Input – 90–130 W Gas – Ar (3000 sccm), $O_2$ (1 to 8 sccm) with Ar, $N_2$ (0 to 80 sccm) with Ar			
	Treatment time – 10 to 15 min CP with DBD		Long-time exposure reduced microbes on the	127
	• Atmospheric air		surface Wheat germination was affected due to the long exposure time	
	• Input power – 80 kV		iong exposure time	
	• Treatment time – 5 and 20 min			
Wheat germ	DPD plasma		Lipase activity was 27.11 and 25.03% at 20 and 24 kV plasma voltage after 25 min	129
	Input voltage – 20 and 24 kV		Lipoxygenase activity was 55.18 and 49.98% for the same condition	

Table 2 (Contd.)

<b></b>			
Treatment time – 5 to 35 min			
Distance - 1.5 and 2 cm			
CP with DBD		Total isolated colonies were reduced to 27	131
1		from 250 after 10 s treatment	
• Input power – 80 kV			
• Treatment time – 3, 10, and 30 s			
DBD plasma with		10 kV pulse voltage removed 3 times more	130
polymethylmethacrylate		endospores than 6 kV treatment	
Gas – Ar (flow rate – 2.8 nm)			
Input – 6 to 10 kV			
<u> </u>			
Low-pressure plasma circulating fluidized		700 and 900 W with 10% $O_2$ mix reduced 1.91	132
bed reactor		and 2.59 log microbes after 10 min	
Gas – Ar and $O_2$ (15 norm. L min <sup>-1</sup> )			
Input - 700 and 900 W			
Time – up to 73.5 s			
Low-pressure CP		Microbial contamination reduced below 1%	133
		for Aspergillus spp. and Penicillum spp.	
• Treatment time – 30 s to 30 min			
Bell jar type of plasma apparatus	Pest control	60 W and 30 min showed complete inhibition	213
		-	
• •	Insect inactivation	The maximum mortality rate was found for	29
- , , , ,			
		100 / 0)	
	Insect control	Complete mortality was achieved by increased	134
	miscer control		101
• Input power = 500 to 3000 V			
		distance for Tribottum custumeum addits	
	Insect control	An increase in exposure time and input power	125
•	msect control		133
-		· ·	
		mortality of 1. castaneum	
Electrode distance – 2 to 6 min			
ACP	Functionality study		214
		1 0	
Input – 50 kV and 60 kV		ē	
		technological properties	
Pin-to-plate ACP	•		215
	modification		
Input – 170, 200, and 230 V			
		increased from 9.65 g $g^{-1}$ to 14.11 g $g^{-1}$ water	
		absorption – 6.39 g g $^{-1}$ to 7.66 g g $^{-1}$	
Treatment time - 5, 10, and 15 min		Swelling power – 7.28 g $g^{-1}$ to 8.79 g $g^{-1}$	
		Water holding – 2.93 g $g^{-1}$ to 3.48 g $g^{-1}$	
Multipin electrical discharge atmospheric	Functional property		216
cold plasma	modification	functional properties	
Input – 13 and 24 W		Water absorption, oil absorption capacity,	
		swelling, and solubility increased	
Treatment times - 10, 20, and 30 min			
, ,			
Radiofrequency CP treatment	Functional modification		137
Gas – Ar and CO <sub>2</sub> (flow rate of 10 and 25		Solvent retention of different flour types	
	• Atmospheric air • Input power – 80 kV • Treatment time – 3, 10, and 30 s DBD plasma with polymethylmethacrylate Gas – Ar (flow rate – 2.8 nm) Input – 6 to 10 kV Time – up to 10 min Low-pressure plasma circulating fluidized bed reactor Gas – Ar and O <sub>2</sub> (15 norm. L min <sup>-1</sup> ) Input – 700 and 900 W Time – up to 73.5 s Low-pressure CP • Air gases or SF <sub>6</sub> (300 torr) • Input power – 300 W • Treatment time – 30 s to 30 min  Bell jar type of plasma apparatus Input – 40 and 60 W Treatment time – 20, 25, and 30 min DBD CP treatment • Atmospheric air (1 mbar) • Input power – 1 to 10 kV • Treatment time – 2 and 20 min DBD atmospheric CP • Atmospheric air (flow rate – 1 L min <sup>-1</sup> ) • Input power – 500 to 3000 V • Treatment time – 0.5 to 7 min DBD plasma Input – 500 to 3000 V Atmospheric air (1 mbar) Treatment time – 0.5 to 7 min Electrode distance – 2 to 6 min ACP Input – 50 kV and 60 kV Treatment time – 5 and 10 min Pin-to-plate ACP  Input – 170, 200, and 230 V  Treatment time – 5, 10, and 15 min Multipin electrical discharge atmospheric cold plasma Input – 13 and 24 W  Treatment times – 10, 20, and 30 min Radiofrequency CP treatment Input – 120 W	OP with DBD  Atmospheric air  Input power – 80 kV  Treatment time – 3, 10, and 30 s  DBD plasma with polymethylmethacrylate Gas – Ar (flow rate – 2.8 mm) Input – 6 to 10 kV Time – up to 10 min Low-pressure plasma circulating fluidized bed reactor Gas – Ar and O <sub>2</sub> (15 norm. L min <sup>-1</sup> ) Input – 700 and 900 W Time – up to 73.5 s Low-pressure CP  Air gases or SF <sub>6</sub> (300 torr)  Input power – 300 W  Treatment time – 30 s to 30 min  Bell jar type of plasma apparatus Input – 40 and 60 W Treatment time – 20, 25, and 30 min  DBD CP treatment  Atmospheric air (1 mbar)  Input power – 1 to 10 kV  Treatment time – 2 and 20 min DBD atmospheric CP  Atmospheric air (flow rate – 1 L min <sup>-1</sup> ) Input power – 500 to 3000 V  Treatment time – 0.5 to 7 min DBD plasma Input – 500 to 3000 V  Atmospheric air (1 mbar) Treatment time – 0.5 to 7 min Electrode distance – 2 to 6 min  ACP  Functionality study  Input – 50 kV and 60 kV  Treatment time – 5 and 10 min Pin-to-plate ACP  Functionality modification  Input – 170, 200, and 230 V  Treatment time – 5, 10, and 15 min  Multipin electrical discharge atmospheric Functional property cold plasma Input – 13 and 24 W  Treatment times – 10, 20, and 30 min  Radiofrequency CP treatment Input – 120 W  Functional modification Input – 120 W	CP with DBD  Atmospheric air  Input power – 80 kV  Treatment time – 3, 10, and 30 s  DBD plasma with polymethylmethacrylate Gas – Ar (10m rate – 2, 8 m) Input – 6 to 10 kV Time – up to 10 min Low-pressure plasma circulating fluidized bed reactor Gas – Ar and O <sub>2</sub> (15 norm. 1, min <sup>-1</sup> ) Input – 700 and 900 W Time – up to 173.5 s Low-pressure CP Air gase or SFe, (300 torr) Input – 700 and 900 W Treatment time – 30 s to 30 min  BBell jar type of plasma apparatus Input – 40 and 60 W Treatment time – 20, 25, and 30 min DBD CP treatment DBD CP treatment DBD Atmospheric air (1 mbar) Input nee – 10 to 10 min DBD atmospheric CP  Atmospheric air (1 mbar) Input – 500 to 3000 V  Atmospheric air (1 mbar) Input – 500 to 3000 V  Atmospheric air (1 mbar) Input – 50 to 3000 V  Atmospheric air (1 mbar) Input – 50 to 3000 V  Atmospheric air (1 mbar) Input – 50 to 3000 V  Atmospheric air (1 mbar) Input – 50 to 3000 V  Atmospheric air (1 mbar) Input – 50 to 3000 V  Atmospheric air (1 mbar) Input – 50 to 3000 V  Atmospheric air (1 mbar) Input – 50 to 3000 V  Atmospheric air (1 mbar) Input – 10 jo ky and 60 kV  Treatment time – 5 and 10 min  Bell pin type or blasma apparatus Input – 50 kV and 60 kV  Treatment time – 5 and 10 min  Bell pin type or blasma apparatus Input – 50 kV and 60 kV  Treatment time – 5 and 10 min  Bell pin type or blasma apparatus Input – 170, 200, and 230 V  Treatment time – 5, 10, and 15 min  Treatment time – 5, 10, and 15 min  Treatment time – 5, 10, and 15 min  Treatment time – 5, 10, and 30 min  Input – 170, 200, and 230 V  Treatment time – 1, 20, and 30 min  Treatment time – 1, 20, and 30 min  Input – 170, 20, and 230 V  Treatment time – 1, 20, and 30 min  Treatment time – 1, 20, and 30 min  Treatment time – 1, 20, and 30 min  Treatment time – 5, 10, and 15 min  ACP  Treatment time – 5, 10, and 15 min  Treatmen

Table 2 (Contd.)

Sample	Plasma chamber	Purpose of study	Research goal	Referenc
	Treatment time – 1 h (2 $\times$ 30 min)			
Rice starch	Bell jar CP reactor	Functional property	Amylose content, pH, turbidity, and starch	138
	• Atmospheric air (0.15 mbar)	modification	hydrolysis percentage were reduced At maximum power level and time, GT is reduced	
	• Input power – 40 to 60 W		reduced	
	• Treatment time – 5 and 10 min			
Wheat flour	Continuous atmospheric pressure cold	Fatty acid profile	FFA and phospholipid were reduced after 60	139
	plasma		and 120 s treatment at 20 V	
	Input – 15 and 20 V		Pasting properties were unaffected	
	Treatment time – 60 and 120 s			
,	d DBD plasma ozone treatment	Functional property	Higher transmittance and swelling power for	140
ong rice)		modification	treated rice pastes and grains, respectively	
	Input – 60 and 70 kV		Syneresis increased for both flours	
	Gas – atmospheric air			
Crain nos (Disum	Treatment time – 5 and 10 min	Eunational property	DDI galubility in area god with plasma	1.41
Grain pea (Pisum	Surface DBD air plasma	Functional property modification	PPI solubility increased with plasma	141
ativum)	• Input – 8.8 kV <sub>PP</sub> • Gas – air	mounicauoli	treatment time up to 191% (10 min) in distilled water. PPF solubility reduced from 71	
	• Time – up to 10 min		to 33% for the same treatment time	
Wheat flour (soft,	DBD atmospheric CP	Functional property	Peak time and peak integral of soft and hard	25
nard)	• Atmospheric air (flow rate – 1 L min <sup>-1</sup> )		flour increased significantly when voltage and	-
,	• Input power – 60 and 70 kV		treatment time exceeded 60 kV and 5 min	
	• Treatment time – 5 and 10 min			
Wheat flour	Counter flow - cold plasma generated	Functional property	Wet gluten (34.2 to 32.1%), dry gluten (9.7 to	142
(	Gas – $O_3$ at 1000 ppm (flow rate – 2.5	modification	7.9%) and gluten softening (13 to 10 mm)	
	$L \min^{-1}$ )		values decreased after 45 min, while	
	,			
	Treatment time – 30 and 45 min		sedimentation (56 to 61 cm <sup>3</sup> ) and bread-	
			making strength index (50 to 56) values	
	Treatment time – 30 and 45 min		sedimentation (56 to 61 cm <sup>3</sup> ) and bread- making strength index (50 to 56) values increased compared to control	
` ,	Treatment time – 30 and 45 min  nt on minimally processed and processed for	ruits products quality	making strength index (50 to 56) values	
(a) Minimally proces	Treatment time – 30 and 45 min  nt on minimally processed and processed fi		making strength index (50 to 56) values increased compared to control	217
(a) Minimally proces Fresh-cut	Treatment time – 30 and 45 min  nt on minimally processed and processed fi  ssed fruit products  DBD	ruits products quality  Quality and flavour	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria	217
a) Minimally proces Fresh-cut	nt on minimally processed and processed fi seed fruit products DBD Input – 40 kV		making strength index (50 to 56) values increased compared to control	217
a) Minimally proces Fresh-cut antaloupe	nt on minimally processed and processed fi seed fruit products DBD Input – 40 kV Treatment time – 90 s	Quality and flavour	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage	
a) Minimally proces Fresh-cut cantaloupe	nt on minimally processed and processed fi seed fruit products DBD Input – 40 kV		making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity	217
a) Minimally proces Fresh-cut cantaloupe	nt on minimally processed and processed fi seed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor	Quality and flavour Enzyme activity	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage	
a) Minimally proces resh-cut antaloupe	nt on minimally processed and processed fi seed fruit products DBD Input – 40 kV Treatment time – 90 s	Quality and flavour Enzyme activity	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%	
a) Minimally proces resh-cut antaloupe	nt on minimally processed and processed finds fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  • Input voltage: 4.8 to 6.9 kV and 12 to 22	Quality and flavour Enzyme activity	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased	
a) Minimally proces Presh-cut cantaloupe Banana slices	nt on minimally processed and processed fissed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  • Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz	Quality and flavour Enzyme activity	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased  2.1 log CFU per g reduction in total aerobic	218 219
a) Minimally proces Presh-cut cantaloupe Banana slices	nt on minimally processed and processed finds fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma	Quality and flavour  Enzyme activity  Surface decontamination	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased	218 219
a) Minimally proces Presh-cut rantaloupe Banana slices	nt on minimally processed and processed finds fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV	Quality and flavour  Enzyme activity  Surface decontamination	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased  2.1 log CFU per g reduction in total aerobic	218 219
a) Minimally proces Presh-cut cantaloupe Banana slices Carrot discs	nt on minimally processed and processed finds fruit products DBD Input - 40 kV Treatment time - 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min	Quality and flavour  Enzyme activity  Surface decontamination	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased  2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV	218
a) Minimally proces Fresh-cut cantaloupe Banana slices Carrot discs Fresh-cut pitaya	nt on minimally processed and processed finds fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic	218 219
a) Minimally proces Presh-cut Presh-cut Presh-cut Presh-cut discs Presh-cut pitaya	nt on minimally processed and processed finds fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety	218
a) Minimally proces resh-cut antaloupe sanana slices carrot discs	nt on minimally processed and processed finds fruit products DBD Input - 40 kV Treatment time - 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased,	218
a) Minimally proces Presh-cut Pantaloupe Banana slices Carrot discs Presh-cut pitaya	nt on minimally processed and processed finds fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor Input voltage: 40, 50, 60, and 70 kV	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety	218
a) Minimally proces Fresh-cut cantaloupe Banana slices Carrot discs Fresh-cut pitaya fruit	nt on minimally processed and processed for seed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity	218 219 220
a) Minimally proces Presh-cut Pantaloupe Banana slices Carrot discs Presh-cut pitaya ruit	nt on minimally processed and processed fixed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity  65 kV for 1 min was effective in achieving	218
a) Minimally proces Fresh-cut cantaloupe Banana slices Carrot discs Fresh-cut pitaya fruit	nt on minimally processed and processed fixed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor  Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor  Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma Input voltage: 45 kV, 65 kV	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity	218 219 220
a) Minimally proces Presh-cut Presh-cut Pantaloupe Banana slices Presh-cut pitaya Pruit Presh-cut pear	nt on minimally processed and processed fixed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor  Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor  Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma Input voltage: 45 kV, 65 kV Treatment time: 1 and 5 min	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity  Microbial inactivation	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity  65 kV for 1 min was effective in achieving a shelf-life of 7 days	<ul><li>218</li><li>219</li><li>220</li><li>221</li></ul>
a) Minimally proces Presh-cut Presh-cut Pantaloupe Banana slices Presh-cut pitaya Pruit Presh-cut pear Presh-cut apples	nt on minimally processed and processed fixed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor  Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor  Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma Input voltage: 45 kV, 65 kV Treatment time: 1 and 5 min Double barrier discharge type gas plasma	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity  Microbial inactivation	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity  65 kV for 1 min was effective in achieving a shelf-life of 7 days  Browning area – 17% reduction in RD and	218 219 220
a) Minimally proces Presh-cut Presh-cut Pannana slices Presh-cut pitaya Pruit Presh-cut pear Presh-cut apples Pink Lady®, Fuji,	nt on minimally processed and processed for seed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor  Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor  Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma Input voltage: 45 kV, 65 kV Treatment time: 1 and 5 min Double barrier discharge type gas plasma with three brass electrodes	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity  Microbial inactivation	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity  65 kV for 1 min was effective in achieving a shelf-life of 7 days  Browning area – 17% reduction in RD and 50% in other samples (30 min). Whereas	218 219 220 221
a) Minimally proces Fresh-cut Fresh-cut Fresh-cut Fresh-cut pitaya Fresh-cut pear Fresh-cut apples Pink Lady®, Fuji, Modì®, Red	nt on minimally processed and processed for seed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor  Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor  Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma Input voltage: 45 kV, 65 kV Treatment time: 1 and 5 min Double barrier discharge type gas plasma with three brass electrodes Atmospheric air	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity  Microbial inactivation	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity  65 kV for 1 min was effective in achieving a shelf-life of 7 days  Browning area – 17% reduction in RD and 50% in other samples (30 min). Whereas 60 min treatment gave 86% and 58%	<ul><li>218</li><li>219</li><li>220</li><li>221</li></ul>
a) Minimally proces Presh-cut Pantaloupe Banana slices Carrot discs Presh-cut pitaya Presh-cut pear Presh-cut apples Pink Lady®, Fuji, Modì®, Red	nt on minimally processed and processed fixed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma Input voltage: 45 kV, 65 kV Treatment time: 1 and 5 min Double barrier discharge type gas plasma with three brass electrodes Atmospheric air Input power – 150 W	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity  Microbial inactivation  Enzymatic browning	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity  65 kV for 1 min was effective in achieving a shelf-life of 7 days  Browning area – 17% reduction in RD and 50% in other samples (30 min). Whereas	<ul><li>218</li><li>219</li><li>220</li><li>221</li></ul>
a) Minimally proces Fresh-cut Fresh-cut Fresh-cut Fresh-cut pitaya Fresh-cut pear Fresh-cut apples Pink Lady®, Fuji, Modì®, Red	nt on minimally processed and processed fixed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor  Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor  Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma Input voltage: 45 kV, 65 kV Treatment time: 1 and 5 min Double barrier discharge type gas plasma with three brass electrodes Atmospheric air Input power – 150 W Treatment time – 30 (15 + 15) and 60 (30	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity  Microbial inactivation  Enzymatic browning	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity  65 kV for 1 min was effective in achieving a shelf-life of 7 days  Browning area – 17% reduction in RD and 50% in other samples (30 min). Whereas 60 min treatment gave 86% and 58%	<ul><li>218</li><li>219</li><li>220</li><li>221</li></ul>
a) Minimally proces	nt on minimally processed and processed fixed fruit products DBD Input – 40 kV Treatment time – 90 s DBD reactor  Input voltage: 4.8 to 6.9 kV and 12 to 22 kHz Treatment time: 0.6 to 2.6 min DBD reactor Gas: atmospheric pressure plasma Input voltage: 60 kV, 80 kV, and 100 kV Treatment time: 0, 1, 2, 3, 4, 5 min DBD reactor Input voltage: 40, 50, 60, and 70 kV Treatment time: 1, 3, 5, and 7 min Atmospheric cold plasma Input voltage: 45 kV, 65 kV Treatment time: 1 and 5 min Double barrier discharge type gas plasma with three brass electrodes Atmospheric air Input power – 150 W	Quality and flavour  Enzyme activity  Surface decontamination  Accumulation of phenolics and antioxidant activity  Microbial inactivation  Enzymatic browning	making strength index (50 to 56) values increased compared to control  CP treatment inhibited the growth of bacteria and mould during 10 days storage  Polyphenol oxidase and peroxidase activity decreased to 70% and 100%  Total phenol and flavonoid content increased to 50%  Vitamin B6 increased 2.1 log CFU per g reduction in total aerobic mesophiles and yeast and mould at 5 min and 100 KV  60 kV for 5 min reduced the total aerobic bacteria and increased safety  The content of individual phenolics increased, resulting in increased antioxidant activity  65 kV for 1 min was effective in achieving a shelf-life of 7 days  Browning area – 17% reduction in RD and 50% in other samples (30 min). Whereas 60 min treatment gave 86% and 58%	218 219 220 221

Table 2 (Contd.)

Sample	Plasma chamber	Purpose of study	Research goal	Reference
			Extracted polyphenol did not affect the cell	
	• Input power 150 W		viability	
	<ul> <li>Input power – 150 W</li> <li>Treatment time – 30 (15 + 15) and 120</li> </ul>			
	(60 + 60) min			
resh-cut apples	Dielectric barrier discharge (DBD)	Polyphenol content	Polyphenol oxidase activity - reduced by 12,	144
	chamber with brass electrodes		32, and 58% after 10, 20, and 30 min	
	• Atmospheric air		treatment	
	• Input power – 9 V (15 kV measured potential difference)			
	• Treatment time – 10, 20 and 30 min (5 +	-		
	5, 10 + 10 and 15 + 15 min each major			
h) Dungagad fruit nu	slice side)			
<i>b) Processed fruit pr</i> Apple juice	OBD and glow discharge plasma	Browning enzyme	Both systems inactivated polyphenol oxidase	222
чры јакс	DDD and glow discharge plasma	activity, antioxidant	and peroxidase and increased total phenolic	222
		capacity, and total	content and antioxidant activity	
	DBD	phenolic content	DBD – better at increasing total phenolic and	
	Input 50 000 Hz		antioxidant capacity	
	Input – 50–900 Hz Glow discharge			
	Input – air plasma flow rate: 10–30			
	mL min <sup>-1</sup>			
	Treatment time – 10–30 min	P! 11 111 C	Market and a laborated and a l	222
Cashew apple juice	ACP	Bio accessibility of vitamin C	Malic acid and phenylalanine increased for ACP 700	223
		vitaliiii G	Increased the vitamin C bio accessibility at	
			ACP 700	
Camu–camu juice	DBD	Bioavailability of	Higher excitation frequency increased the	224
	Input – 24 kV and frequency – 200, 420,	vitamin C	availability of ascorbic acid Degradation of anthocyanins and peroxidase	
	583, 698, and 960 Hz		enzyme at a higher frequency	
Kiwi turbid juice	DBD	Microbial inactivation	• 12 kV, volume – 18 mL and discharge time –	225
		and quality changes	1 min was the optimum condition	
	Input – 10, 25, and 40 kV		• Polyphenol content – 0.18 mg g <sup>-1</sup> , chlorophyll content – 3.47	
	Treatment time – 1, 3 and 5 min		• Sterilization rate – 18.03%	
	Volume – 10, 15, and 20 mL			
Siriguela juice	Low-pressure plasma processing	Bioactive component	Enhanced the bioactive component content	226
	Nitrogen gas flow rate – 10–30 mL min <sup>-1</sup>		Highest pigment increase – 10 min – 10 mL min <sup>–1</sup>	
	Treatment time – 5 to 15 min		Highest antioxidant activity – 15 min – 20	
			mL min <sup>-1</sup>	
Blueberry juice	Cold plasma jet	Bioavailability of	2 min exhibited high anthocyanin	227
	Input – 11 kV, 1000 Hz and oxygen gas	vitamin C	Long treatment decrement of vitamin C	
	concentration – 0, 0.5 and 1 Treatment time – 2, 4, and 6 min		1% concentration of O <sub>2</sub> resulted in the	
	2, 1, and 5 mm		reduction of Bacillus by 7.2 logs CFU per g	
apple juice	Dielectric barrier discharge (DBD) plasma	Microbial inactivation	4.0 log CFU per mL at 30 W (40 s)	146
	chamber with 2 mm electrode distance			
	from sample surface at 23 $\pm$ 2 $^{\circ}$ C temperature			
	Atmospheric air		4.2 log CFU per mL at 40 W (40 s)	
	• Input power – 30, 40, 50 W		4.34 log CFU per mL at 50 W (30 s)	
Banana starch	• Treatment time – 0 to 40 s	Starch modification	No abances in amulass and resistant -t	150
	Corona electrical discharge (CED) plasma Input – 30, 40, and 50 kV	i staten modification	No changes in amylose and resistant starch contents. The crystalline to amorphous ratio	130
	Treatment time – 3 min		increased after CED plasma treatment	
Sour cheery nectar	APPJ setup		E. coli – 3.34 log reduction at 650 W after 120 s	
Гоmato juice	• Atmospheric dry air (flow rate – 3000 L	Microbial inactivation	E. coli – 1.43 log reduction at 650 W after 120 s	3
Apple juice	h <sup>-1</sup> )  • Input power − 650 W		<i>E. coli</i> – 4.02 log reduction at 650 W after 120 s	1
Appie juice Orange juice	• Treatment time – 0 to 120 s		<i>E. coli</i> – 4.02 log reduction at 650 W after 120 s	

Reference

Sample

Table 2 (Contd.)

Plasma chamber

Sample	Plasma chamber	Purpose of study	Research goal	Reference	
White grape juice	High voltage atmospheric pressure CP with plexiglass and polypropylene dielectric barrier • Atmospheric dry air (Ziploc bag) • Input power – 80 kV (peak) • Treatment time – 1 to 4 min	Microbial inactivation	S. cerevisiae population reduced to 7.4 logs (80 kV for 4 min)	149	
(4) Plasma treatmen	at on essential oil extraction and quality				
Hyssop	DBD	Changes in essential oil content during storage	Trichomes were vulnerable to plasma treatment	228	
	Input - 17, 20, and 23 kV		Increased treatment duration reduced the EC content		
	Treatment time - 1, 5 and 10 min				
Fennel and	DBD	Essential oil yield	19 kV and 10 min yield 1.89% (v/w) for fennel	229	
spearmint leaves	Input – 17–23 kV		and 1.81% (v/w) for spearmint leaves		
	Treatment time – 5–15 min				
Lemon peel	Low-pressure DBD cold plasma	Essential oil yield	Plasma treatment for 10 min at 2.5 kV	152	
	• Gas – atmospheric air (1 mbar)		increased the extraction yield by 149.34%		
	• Input – 1 to 2.5 kV				
	• Treatment time – 10 min				
Lemon verbena	Low-pressure CP connected with vacuum	Essential oil yield	Essential oil (EO) content increased by 36.7% 153		
(Lippa citriodora Kunth.)	pump		during the first 1 min of treatment and later		
	Nitrogen, argon, oxygen		decreased due to higher exposure time		
0	• Treatment time – 0, 1, 3, and 5 min	Parametel - 21 - 1-14	pl to the liber library and the laws a	2=	
Sweet basil (Ocimum		Essential oil yield	Plasma treated basil EO produced more	27	
basilicum)	<ul><li>Air/helium</li><li>Treatment time - 5 s</li></ul>		antioxidant activity (94.82%) than untreated (90.64%) basil at a 250 $\mu g$ mL <sup>-1</sup> concentration		
Laman naal		Ecceptial oil viold			
Lemon peel	DBD plasma treatment	Essential oil yield	Maximum EO yield was obtained in the argon 15 plasma (flow rate – 15 mL min <sup>-1</sup> ) after 1 min		
	• Argon, oxygen, nitrogen, or air (10–20		Argon plasma of $1 \times 1 \times 1$ cm size lemon peel		
	mL min <sup>-1</sup> )		produces maximum EO extraction efficiency		
	• Input power – 30 kV (peak)				
	• Treatment time – 1 to 15 min				
Clove oil	Plasma jet with pyrex glass tube at	Essential oil yield	microbial inhibition (A. niger, Penicillium sp.,	151	
	atmospheric pressure				
	• Argon (flow rate of 10 L min <sup>-1</sup> )		and Rhizopus sp.) was reduced to 20 times for		
			clove oil and 8 to 9 times for eugenol after		
			10 min at 40 W		

Purpose of study

Research goal

degradation were entirely removed after 120 min. Industrial wastewater with higher microbial contamination might also affect the water quality of agri-farms. So, industrial wastewater needs decontamination studies, and Rowan et al.46 conducted a study on poultry wash water to investigate microbial decontamination. In this study, other than RS and UV photons, nitric and carbonic acids formed during plasma treatment resulted in pH reduction (acidified) that helped in microbial destruction. These components resulted in the microbial reduction of 8 logs (less than or equal) to complete removal after 30 s of treatment, and the sensitivity of treatments varied for different microbes (Bacillus cereus endospores < Listeria monocytogenes < Salmonella enterica Typhimurium < Salmonella enterica Enteritidis < E. coli < Campylobacter jejuni < Campylobacter coli).

Apart from using plasma units for treating different water samples, it is also possible to carry the plasma species in water by which the treatments can be done. Ten Bosch et al.40 reported the effect of plasma-treated water on insect mortality

(Mealybugs). Though Plasma Treated Water (PTW) and Classically Acidified Water (CAW) had no significant difference, the CAW mortality rate was higher than PTW at the same pH due to the complexity of RS present in PTW. This PTW can also be used as an alternative for sodium nitrite in sausage curing with similar sensory, microbial, and peroxide values as that of control.58 Even though plasma-activated water (PAW) possesses antimicrobial properties, its effectiveness reduces during the storage period. Traylor et al.59 reported that PAW prepared with 15 min and 3 h dosage intervals were found to have a similar antimicrobial effect of more than 5 logs. However, after 30 min, the effectiveness of PAW prepared with 15 min exposure time reduced from 5.6 logs to 2.4 logs, while the other was stable for 2 days. Similarly, the lethal effect of stable RS on Hafnia alvei was reported, and among NO-, NO2-, and H2O2 species, acidification of PAW by NO2 species acted as a primary factor in microbial reduction.60 Kamgang-Youbi et al.48 reported the influence of pH reduction and substrate absence on

saccharomyces destruction and found that due to its cell size, initial population, and adhesiveness of substrate surface, their resistance was higher against plasma treatment.

Table 1 shows the different plasma generation methods and significant results achieved in water treatments. However, further research must be explored to characterize the degradation process's end product. Since the future industrialization of PAW usage may produce a massive quantity of used PAW, there is a need for research on its recyclability and toxicity.

### 3.2. Plasma treatment for soil remediation

Industrial and agricultural-related wastes such as heavy metals, pesticides, petroleum, and its products cause soil contamination, which could affect humans and the ecosystem. 61 These contaminants' concentration in soil has to be reduced to avoid losses in production and cross-contamination in agricultural produces. Plasma technology has already gained popularity at the industrial level as an emerging technology for different treatments. Hence, this technology can be used in soil decontamination.62

In plasma assisted soil remediation, factors such as soil thickness and its type affect the treatment voltage, frequency, soil thickness, and air flow rate. The effectiveness of the method was identified against different types of soil (sandy soil and loam soil).63

Soil pollutant removal varies depending upon the nature of the soil as well as the pollutant. Zhan et al.64 studied the influence of treatment parameters such as time and washing, soil properties such as pH, moisture, and initial concentration of pollutants on fluorene removal from soil. The study showed a significant increase in fluorene degradation (from 33.8 to 57.9%) while reducing the soil moisture content from 20 to 0.6%. In addition, the efficiency increased from 60.6 to 71% when the pH of the soil rose from 3.0 to 9.0. Washing, increasing treatment time, and reducing initial pollutant concentration are reported to be critical factors for pollutant removal. Aggelopoulos et al.65 studied the non-aqueous phase liquid (NAPL)-mixed soil decontamination with high and low energy density plasma treatment to avoid this influence of initial pollutant concentration. When the ex situ DBD plasma treatment energy density was around 10 000 J g per soil, NAPL was obliterated, irrespective of the initial concentration. However, low energy density was affected by its increased concentration. Lu et al. 66 included the effect of frequency, gas flow, and input power in removing acid scarlet GR dye from the soil. It was found that an increase in DBD plasma input power (3.51 to 5.72 W), frequency (200–300 Hz), and gas flow rate (0.5– 1.0 L min<sup>-1</sup>) improved dye degradation. At the same time, a higher air flow rate (1.5 L min<sup>-1</sup>) reduced efficiency and O<sub>3</sub> and OH<sup>-</sup> reactive species were reported as critical factors in this degradation. However, to find the effectiveness of different gas sources on degradation other than flow rate, Lou et al. 67 used O2 and air for chloramphenicol degradation, where the efficiency of O<sub>2</sub> (41%) was higher than air (26%). For better efficiency, optimum moisture content ( $\leq 10\%$ ) and Fe<sup>0</sup> addition were needed, and that can increase reactive species and discharge

channels, respectively. Similarly, Wang et al.68 used a pulsed discharge plasma-TiO<sub>2</sub> catalytic (PDPTC) reactor to enhance the reactor performance for organic component removal. In this study, TDPTC (55.1%) removed higher organic components than the plasma reactor performed without TiO<sub>2</sub> (42.9%).

For pentachlorophenol (PCP) degradation, Wang et al. 69 used an optimum flow rate (3 L min<sup>-1</sup>) and a high O<sub>2</sub> environment to increase the degradation rate. However, it was found that prolonged exposure of pollutants (4 to 96 h) to soil reduced the degradation by 13.4% as the pollutant gets into the soil granules deeply. So, apart from process parameters and other soil properties, the depth of the soil also decides the degradation efficiency. To understand this, Wang et al. 70 studied p-nitrophenol (PNP) degradation at a different depth from the surface of the soil using plasma. From this study, it was found that an increase in soil depth reduces the degradation rate from 77.9 (0-2 mm depth) to 52.8% (10-12 mm depth), and the removal of PNP in moist clay soil (44.1%) was higher than dry soil (11.3%) with increased discharge voltage. Similarly, Wang et al.71 studied the effect of the size of the soil granule on the degradation of pentachlorophenol (PCP) and found higher degradation in 20 mesh size granular soil (87%) than in 10 mesh size granular soil (72%). In addition to granule size, the high pH of the soil also assists the PCP degradation. Unlike other studies, Stryczewska et al.72 focused on microbial decontamination and plasmainduced soil property change. Here, the soil pH and temperature changes occurred with changes in O3 concentration and treatment time, resulting in decontamination. A similar study was carried out with different electrode configurations, and this study reported the importance of screw or pyramid-shaped electrodes on higher O<sub>3</sub> production. O<sub>3</sub> and NO<sup>-</sup> produced in this process altered the soil conductivity (34 to 79 mS m<sup>-1</sup>) and microbial population.62 Later, Redolfi et al. (2010) studied the RS oxidation effect on kerosene-mixed soil to analyze the pollutant residues in the plasma exhaust gas and its bioavailability. Results indicated that only a negligible amount of organic components came along with the gas outlet, while most of them were retained in the soil and did not convert from a solid to a gas state. Hence the bioavailability of kerosene byproducts increased in the soil.

Extended research is needed to understand the effect of plasma treatment on soil mineral content, residue toxicity, and their effects on plant growth and crop yield after treatment. Apart from the decontamination of soil, the effect of plasma on desirable organisms present in it also needs to be addressed.

### 3.3. Plasma treatment of seeds/plants

Improving germination rate and agricultural yield is the primary concern in increasing food production,73 and plasma treatment has the potential to improve these two aspects of agriculture processing by altering the germination rate (GR), germination potential (GP), vigor index (VI), water absorbance, contact angle, wettability, and growing tolerance of seed. At optimum process conditions, these changes can influence the overall production of food produce by favouring seed and plant growth.

3.3.1. Germination and plant growth. Since germination is a crucial factor in deciding seed quality, efforts were made to increase the germination behaviour of seeds through plasma technology. Li et al.74 reported an increasing trend in GP, GR, GI, and VI of wheat seeds up to the treatment time of 4 min, while the root and shoot lengths had a positive response during germination. Kim et al.75 treated broccoli seeds in a corona discharge plasma jet (CDPI) to know the additional benefits of this treatment. They found microbial reduction on the seed surface along with increased germination rate and growth; meanwhile, the sensory and physicochemical parameters were unaffected. Different factors are involved in seed germination (Fig. 3), and water absorption is one among them, Zahoranová et al.76 highlighted this in a wheat seed germination study and found an increase in water absorption of each seed at 2 h (6.41 to 9.60 mg) and 8 h (12.53 to 16.07 mg) of soaking times with respect to the control. In addition to this, GR (21%), VI-I (28%), VI-II (36%), and surface microbial reduction (2 logs) also increased. Along with germination and water absorption, Dobrin et al.77 reported the average increase in root length (32.89 to 36.49 cm), sprout length (4.26 to 4.34 cm), root weight (0.78 to 1.06 g), and root to shoot ratio (0.88 to 1.2) of treated wheat seeds after 15 min treatment. Though most research shows a positive impact on germination and physical characteristics, morphological changes and yield of crops during the growth periods were unidentified. So, Ling et al.78 decided to work on peanut (Arachis hypogaea L.) seed with helium plasma and recorded the increase in yield, plant height, stem diameter, shoot and root dry weight, area, thickness, water, and nitrogen content of leaf at 120 W power level. Not only seeds but plants such as basil showed more growth in height, around 10 mm, than the control.27 Apart from plasma treatment, PAW also helped in the growth of seeds, and Sivachandiran42 reported about a 28% to 45% increase in the stem length of radish after PAW watering. For tomato and pepper plants, plasma and PAW combined treatment showed positive plant growth with healthy leaves. Similarly, a synergistic effect was observed in the germination of mung bean seeds when they were treated with plasma and PAW. The germination percentage reached up to 100% (from 94%) when plasma and PAW (CO2) were used; however, the PAW generated using O<sub>2</sub> [PAW(O<sub>2</sub>)] inactivated more microorganisms than the other PAW.79

Apart from germination enhancement, plasma also has the ability to change the minerals, pigments, enzymes, and other nutritional compositions of the seeds. For example, in capacitively coupled plasma (CCP) treated tomato seeds, calcium (7.73%) and boron (11.53%) contents were increased; 80 whereas in soybean, nutrient fractions such as soluble protein and soluble sugar and enzymes such as peroxidase (POD) and phenylalanine ammonia-lyase (PAL) activities increased after CDPJ treatment.81 In helium plasma-treated wheat (Triticum spp.), chlorophyll content increased by 9.8% more than that in the control.26 However, photosynthesis efficiency and flavonoid content in peas are reduced due to UV-C production.82 Similarly, time-dependent reduction of dehydrogenase (27%) and catalase activities (75%) were observed in the roots of germinated maize

seeds.83 In comparison, structural changes were observed with increased indolyl acetic acid (13.7%) in peas.84 For beans, germination time (44 to 40 h) and exotesta contact angle were reduced after plasma treatment.85

In contrast to other research works, Puligundla et al.86 analyzed the negative impact of plasma over dosage on rapeseed germination and suggested the optimum treatment time for better germination since 1 min treatment showed better results in this study. However, the study on pre-germinated brown rice by Sookwong et al.87 showed a reduction in germination rate after the treatment and explained the dependency of treatment time and distance. Along with other dependent variables, gas used in treatment also has a considerable impact on seed germination, and studies by Živković et al.88 on paulownia tomentosa seed proved that by showing maximum germination at 15 min for air plasma, unlike argon plasma. It also noted that the impact of plasma on seeds could not be carried out for a long time since poppy seeds treated at 500 W plasma showed higher GR on the first day of germination but reduced on the sixth day. i.e., 13% reduction was observed in 3 min treated sample.89

3.3.2. Plasma-induced seed tolerance. In recent times, plasma-induced seed tolerance has been getting researchers' attention, which led to experiments on different stress parameters such as drought, heavy metals, toxins, and salinity against plasma treatment. For example, Kyzek et al.90 reported the adaptive response of plasma-treated pea seeds (Pisum sativum L.) against zeocin and found an increased tolerance against drought and heavy metals due to the interaction of RNS with seeds. However, the most common problem of salinityinduced stress was investigated by Iranbakhsh et al.91 for wheat, and this study showed an increase in heat shock factor (HSF) A4A and salinity tolerance after plasma treatment. Further, it increased the wheat plant's immunity, shoot weight, and enzyme activity (peroxidase and phenylalanine ammonialyase). In addition to saline tolerance, drought tolerance also increased for wheat due to the production of abscisic acid. Subsequently, proline (12.7%) and soluble sugar (16.4%) contents also increased with the treatment.92

Seed treatment results and their effects vary based on the plasma characteristics used in the respective treatments (Table 1). Many studies have shown a significant increase in the germination of seeds, but detailed studies are required to identify the effect on the quality and yield of the final product. In addition, the negative impact of plasma treatment on seeds needs to be addressed with the evident mode of action, which could provide the limitations of plasma treatment for seeds (Table 2).

# Plasma applications in postharvest stages

### 4.1. Plasma treatment of different agriculture produces

Some of the edible products, such as leaves, roots, fruits, nuts, and spices, are either contaminated by microbes or will have allergens. Plasma treatment can decrease the contaminant level

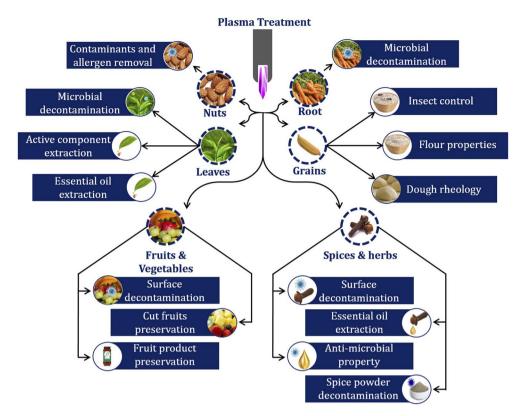


Fig. 4 Plasma application in post-harvest stages of agri-food chain.

and increase the active component percentage in food products. However, based on the surface morphology of different commodities, the effect may change and usually produces a smooth surface that will have more microbial removal rate than the unevenly surfaced one, and this could be due to the improper penetration of plasma species into the porous structures.<sup>93</sup> Decontamination and disinfestation of grains, higher

UV Changes in cell Wall potential ROS Cell wall ROS Lipid RNS Damage oxidation RNS Protein modification Cell wall **Protein** Layers damage Leakage 🌎

Fig. 5 Effect of plasma treatment on microbial cell.

extraction of EO, and functional modification of flours are some advantages of plasma treatments that will be discussed in this section. However, like any other treatment, plasma produces minor undesirable changes that will also be covered here. Different post-harvest applications of plasma are shown in Fig. 4.

**4.1.1.** Leaves. Interaction of plasma species with microbes causes itching, and thus inactivation occurs, while DNA damages caused by UV photons further enhance this effect (Fig. 5). So, microbial contamination on freshly harvested leaves can be reduced by plasma treatment. Pasquali et al.94 reported the microbial decontamination of radicchio leaf (Cichorium intybus L.) after DBD plasma treatment and found a significant removal of microbial load. However, the sample stored after treatment had less chroma (C\*) value at 4 °C. In order to improve the inactivation efficiency, different gas mixes were suggested, and the Ar-O2 gas mix is one among them. In romaine lettuce, Ar and O2 gas mix gave better decontamination efficiency without altering the texture of the produce.93 However, this gas combination is not ideal for decontaminating all microbes since they have different physical and morphological characteristics. For example, black and green tea were treated with an argon plasma jet, where E. coli and coliform were removed within 5 min, while yeast and mould took more time. Further, it also increased the total phenolic content (TPC) of black tea (10.77 to 11.38 g GAE/100 g) and green tea (14.94 to 16.02 g GAE/100 g), but caffeine and colour values were not affected significantly.95 Similarly, microwave plasma reduced

the microbial (viable) count of dried peppermint, stinging nettle, and lemongrass leaves by up to 1.6 logs within 7.5 min of treatment time.96 Though decontamination of leaves was the primary goal of these studies, Baier et al.97 found a permanent reduction in photosynthetic efficiency  $(F_v/F_m)$  of corn leaves when a higher power level was used. Nevertheless, in the case of lamb's lettuce, the interaction of reactive species caused the reduction in bioactive phenolic components and not in the photo- or thermo-desorption processes. Further reductions such as protocatechuic acid (16%), chlorogenic acid (29%), and caffeinic acid (35%) were observed during the treatment, along with some structural changes.98 Similar reductions were observed in flavonoids, phenolic acids, and the contact angle of lamb's lettuce.99

4.1.2. Roots. Surface morphology is one of the critical factors in deciding the decontamination efficiency of plasma treatment, Bermúdez-Aguirre et al.93 reported that the surface morphology of baby carrots restricted the penetration of plasma into the porous surface areas, and hence the microbial log reduction was found to be less.

4.1.3. Fruits. Freshly harvested fruit surfaces can either be contaminated by microbes or by pesticides. For microbial removal, similar to Amini and Ghoranneviss's95 study, strawberries were treated in DBD atmospheric cold plasma at 60 kV<sub>rms</sub> voltage in high oxygen (G1) and high nitrogen (G2) environments. The results showed that G2 was more efficient against mesophiles, whereas G1 was effective on yeast/mould. To maintain more firmness and  $L^*$  value,  $G_1$  was preferred since it produced more ozone ( $G_1$  - 2800 ppm) than  $G_2$  (600 ppm).100 For cocktail tomatoes, along with E. coli inactivation, an increase in  $a^*$  value was observed with respect to the treatment time. However, at a fixed treatment time of 10 min at 12.83, 11.18, and 6.90 kV power levels, a slight reducing trend of a\* value was observed.93 However, to achieve the desired microbial reduction from fruit surfaces, it is essential to select a suitable feed gas. A recent study revealed that using air plasma on red currants was only able to reduce fungal population (1.28) log) and not aerobic mesophilic. At the same time, using nitrogen plasma induced up to 1.11 log microbial reduction in

aerobic mesophilic without causing any lethal effect on yeast and moulds.101 Among the various plasma types, cold atmospheric pressure plasma (CAPP) is commonly employed for decontamination of these kinds of fruit and berries.102 In addition to plasma treatment and PAW, plasma activated air (PAA) also helps in the decontamination of fresh fruits and vegetables (apples cucumbers, tomatoes, and carrots). Interestingly, the commodity with a rough surface (carrot) achieved a higher microbial reduction of up to 5 logs within 5 min of PAA treatment compared to smooth surfaced apple and tomato samples ( $\sim$ 3 log reduction). <sup>103</sup>

As pesticides are used extensively in agricultural commodities, their residues remain on the surface of the commodities even after primary and secondary processing. Hence, Ranjitha Gracy et al. 37 applied plasma treatment to tomatoes to validate the chlorpyrifos reduction in DBD plasma with different dosage levels. The results showed around 90% pesticide reduction due to the conversion of the phosphorothiol group (P=S) into the phosphoryl group (P=O). After the treatment, considerable changes in firmness, total phenolic content, and colour index were observed. Earlier, Sarangapani et al. 104 studied the same with blueberries contaminated with boscalid and imidacloprid pesticides and found at least a 75% reduction in both pesticides. Along with this, ascorbic acid content increased from 8.91 to 14.01 mg/100, while time and power level increases reduced the total flavonoids, anthocyanin, and TPC. Different interactions of plasma with water are shown in Fig. 6.

Apart from plasma treatment, PAW also helps in fruit processing. For example, PAW water can clean the fungicide contaminated tomatoes and can cause up to 85.3% and 79.47% reduction in the chlorothalonil and thiram contents. 105 Further, they are also capable of reducing microbial contamination of fruits and vegetables.106

4.1.4. Nuts. Nuts are high-quality oil and nutritive food source; hence, it is susceptible to microbial contamination. In order to remove the contaminants from the surface, it needs to be processed without significant quality deterioration. However, plasma treatment provides surface decontamination with no or minimal quality changes. In unpeeled almonds, air plasma

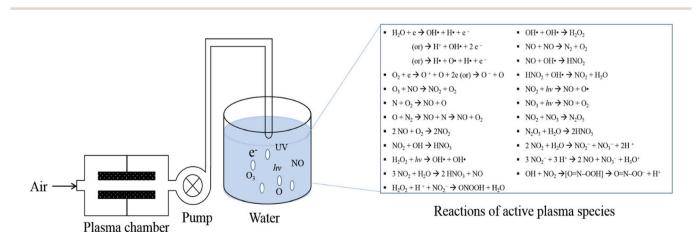


Fig. 6 Interaction of reactive species with water in plasma treatment and PAW generation.

increased the inactivation of salmonella; however, air mixed with N2 caused browning. 107 A similar decontamination study on hazelnuts showed Aspergillus spp. Reduction even after 30 days of storage due to the presence of RS.108 The effect of moisture content and different cultivars on microbial removal of nuts were experimented with walnut fruits in a plasma jet using argon gas and observed that microbial removal was high in dried walnuts than the fresh ones for the given treatment time. It is due to the low water activity, moisture content, and rugged texture of dried walnuts. After treatment, no further microbial growth was observed till 30 days (4 °C) except in a few (Shahmirzad and Taleghan) walnut types. 109 Apart from product variables, the nature of the microbes present in nuts also has varying responses to the treatment. In Dasan et al.110 report on hazelnuts, Aspergillus flavus reduced up to 4.5 log per g, whereas A. parasiticus reduction was 4.19 log per g under the same conditions. The difference in morphology, toxic production, and geographical range of these microbial species could not be the factor for the difference in microbial removal. However, molecular research is needed to get more information about this effect. To evaluate the feed gas effect, Niemira111 reported the microbial reduction in almonds with air and nitrogen plasma and found higher microbial removal in air-treated almonds than the nitrogen treated due to ROS in air plasma. With this, almonds' decontamination can also be increased by higher treatment time, input voltage, and frequency. 112 Likewise, hazelnut, peanut, and pistachio nut were treated in lowpressure air or sulfur hexafluoride (SF<sub>6</sub>) cold plasma. Though SF<sub>6</sub> showed higher microbial reduction than air plasma, aflatoxin removal was higher (50%) in air plasma treatment than SF<sub>6</sub> (20%) after 20 min treatment. Peanuts and pistachio nuts were not entirely sterilized after 20 min SF<sub>6</sub> treatment, and hazelnut microbial load reduced linearly till 5 min of SF<sub>6</sub> treatment.113

Some quality changes are also associated with nuts during treatment. For instance, peanuts were treated with coplanar DBD plasma; though antioxidant capacity was not changed due to short exposure to ROS, the TPC increased from 200.23 to 341.15 mg (GAE/100 g) after 25 W treatment for 8 min (0.5 L min<sup>-1</sup>) since phenols protect cells against the damaging effects UV and ROS. However, lipid oxidation and moisture reduction were observed due to the oxidation of strong ROS species.114 Another problem with peanuts is the Ara h 1 allergens present in them; however, it was reduced in the whole peanut (WP) and defatted peanut flour (DPF) through binding epitope modifications caused by protein and lipid functionality changes after the treatment.115 Aflatoxins are commonly found in nuts, which is a hazardous toxin, though it was removed about 70% from hazelnut (Aflatoxin B1-AFB1 and total Aflatoxin-AFs) by DBD plasma treatment at 1150 W. Changes in the lactone ring were considered to be the reason for degradation and this effect was increased with power and time.116

4.1.5. Spices. Mošovská et al. 117 experimented with plasma on black pepper and found that B. subtilis spores were more resistant than E. coli and S. Enteritidis. After treatment, minimal surface damage and moisture loss with unchanged sensory properties were observed. Amini et al.118 reported that the ester and volatile compounds of saffron in response to the plasma treatment. Safranal was degraded by plasma treatment and increased (from 21% to 33% at 12 kV) by adding O2 (5%) to the plasma gas composition. In contrast, isophorone and 4ketoisophorone increased by O2 gas mixer. Plasma treatment reduced crocin esters, and an O2 gas mixer enhanced the effect in Ar plasma.

Kim et al. 119 reported decontamination of red pepper flakes from B. cereus in high microwave density (HMCPT) and low microwave density (LMCPT) CP treatment. It was observed that IR drying and flaked samples were suited for microbial reduction. Higher sample water activity  $(a_{w})$  produced a more lethal effect on microbial reduction. Kim et al. 120 treated onion powder in microwave-powered plasma. Microbial inactivation was high in HMCPT than in LMCPT. In addition, the treatment drying method influenced microbial reduction. Onion Aw reduced from 0.26 to 0.16 and 0.12 in LMCPT and HMCPT. Plasma treatment with low temperature (4 °C) storage prevented microbial growth. Quercetin concentration and DPPH activity followed a reducing trend in HMCPT samples, and for the control sample, an increasing trend was observed during storage at both temperatures (4 °C and 25 °C). Won et al. 121 reported the effect of He-plasma on onion preservation. In plasma treatment, increased E. coli, L. monocytogenes and S. Enteritidis inactivation by about 2 log per cm<sup>2</sup> when  $a_{\rm w}$ increased from 0.4 to 0.8 was noted, and also, the increase in particle size positively influenced inactivation.

Hertwig et al.28 reported the effect of radio frequency (G1) and microwave (G2) generated plasma jet on black pepper. G1 plasma treatment for 15 min caused 0.7 logs and 0.6 log inactivation of aerobic and spore count, whereas G2 plasma treatment caused 1.7 logs and 1.4 inactivation of an aerobic and spore count for the same time interval. The piperine content of both plasma-treated samples was reduced slightly, and G2 plasma was effective against S. enterica. Takemura et al. 122 reported the microbial reduction in black pepper in plasma treatment. Argon and CO2 mixed plasma treatment reduced microbial load better than other combinations. The growth retardation of untreated samples was less than air and argon + CO<sub>2</sub> mixer plasma. Similarly, Sun et al. 123 reported APPJ plasma treatment of black pepper on surface microbial reduction. A sample stored at a higher temperature (37 °C) and low RH (33%) reduced the initial microbial count more than that stored at a low temperature and RH. During the treatment, peppercorn surface temperature reached above 120 °C after 80 s exposure. Combined storage condition and 80 s plasma treatment reduced an average of about seven microbial log counts. Grabowski et al. 124 treated black pepper with plasma for decontamination. Spore & non-spore forming aerobic bacteria and anaerobic spore-forming bacteria were removed during the treatment. Increased treatment time increased microbial reduction but resulted in water loss. Lumps on pepper were observed after treatment except in O<sub>2</sub> and H<sub>2</sub>O<sub>2</sub> plasma. Kim et al.125 reported the effect of microwave-powered CP on the decontamination of A. flavus from red pepper powder. The treatments performed at higher power levels reduced the water activity of pepper due to evaporation, which further increased

the ROS concentration. Colour change was insignificant, and He and O<sub>2</sub> gas mix with heating up to 90 °C inactivated B. cereus spores.

Research has shown a considerable microbial reduction in most agricultural products and enhancement in active components. However, specific issues, such as colour change, weight loss, and bioactive quality losses, must be addressed by selecting proper plasma characteristics and treatment time. Apart from that, the mode of action of RS or UV produced from plasma treatment on the food components needs to be explained to understand the reason for positive and negative impacts on quality.

### 4.2. Plasma treatment for safe storage, extraction, and quality enhancement of agricultural products

Removal of microbes and pests during storage is essential to increase the quality of goods. Fumigation and other thermal treatments result in either residue formation or quality deterioration. Plasma treatment facilitates microbial removal and disinfestation without affecting product quality. Meanwhile, it also inactivates the enzymes that reduce the product quality or shelf life. In some cases, plasma treatment improves the functional properties of food constituents.

**4.2.1.** Grain safety and quality. Carpen et al. 126 reported the effect of nitrogen and oxygen gas mixture with Ar plasma on the mortality of Tribolium castaneum and T. confusum in wheat. When O<sub>2</sub> was added with Ar plasma at the concentration levels of 2.8 and 8.4 seem, mortality reduced from 88% to 65% and 1.3% for T. castaneum and 96% to 51% and 0% for T. confusum, respectively. The reduction in mortality was assumed to be due to temperature reduction caused by gas mixing. Similarly, when wheat and barley were subjected to plasma treatment to remove microbial contamination, bacteria and fungi in barley were reduced to 2.4 logs and 2.1 logs after 20 min (24 h retention), while in wheat, it was reduced to 1.5 logs and 2.5 logs after 20 min for bacteria and fungi, respectively. It was found that germination was increased after 5 min treatment with 2 h retention in wheat, and the contact angle was reduced.127 Apart from insects, microbes and their secondary metabolites also affect grain quality. Hence, Durek et al.128 studied the importance of plasma treatment and its processing variable (feed gas: CO<sub>2</sub> + O<sub>2</sub>, CO<sub>2</sub>; time: 1-3 min) effect on Aspergillus niger and Penicillium verrucosum contaminated barley. Though both microbe's counts were reduced in all plasma treatments, the P. verrucosum's produced mycotoxin (ochratoxin A) content reduced only in the  $CO_2 + O_2$  plasma treatment (49.0 ng g<sup>-1</sup> to 27.5 ng g<sup>-1</sup> after 1 min and 23.8 ng g<sup>-1</sup> after 3 min). When  $CO_2$ alone was used on barley, it increased the mycotoxin content upon storage.

Tolouie et al. 129 reported the plasma effect on wheat germ enzyme activity to enhance its shelf life. Lipase and lipoxygenase activities were reduced more in 24 kV plasma treatment, while antioxidant activity and total phenols were unchanged. Lipase and lipoxygenase activities were recovered by 1.18 and 6.52 U g<sup>-1</sup>, respectively, after 30 days. Butscher et al. 130 reported bacterial inactivation of plasma treatment in wheat grain where water content was reduced by 10.48 to 9.35% gluten, and the falling number was unaffected. Kordas et al. 131 reported around ten times reduction in the fungal colonies of wheat seeds when exposed to up to 10 s of plasma treatment, and it also resulted in more than 98% germination energy. However, the root and leaf length decreased during the same treatment time. Butscher et al.132 reported the effect of power level and O2 concentration on microbial destruction in wheat. Higher O2 levels and power levels positively affect microbes, while the extensograph and amylograph parameters such as elasticity, strain resistance, energy, P/L ratio, and gelatinization increased after plasma treatment. Selcuk et al.133 reported the surface structure of grains and legumes on microbial disinfection of air and SF<sub>6</sub> plasma. Cooking quality and water absorbance had no significant effect due to treatments. However, the seed germination rate was maintained above 85% after treatment. Wheat disinfection was 99% in SF<sub>6</sub> plasma treatment.

4.2.2. Flour safety and quality. Ramanan et al.29 reported the effect of CP on the mortality of T. castaneum at different life stages (egg, larva, and adult). Low voltage levels and less electrode distance were found not to affect mortality. However, higher voltage (3000 V) mortality was affected by distance. T. castaneum mortality prediction model showed no significant difference with experimental mortality rate values (egg, larva, adult) at a 95% confidence level. At 2500 V, maximum mortality was found at a 3.7 cm electrode distance. Eggs, larvae, and adults had 93.33%, 93.33%, and 100% mortality, whereas the predicted model showed 100%, 80%, and 100% mortality, respectively. Similarly, Radhakrishnan et al. 134 demonstrated T. castaneum disinfestation using CP. Refined wheat flour treated with plasma produces 100% mortality at 1750 V for 7 min and 3 min with 4 cm and 2 cm electrode distances. While, at 3000 V voltage and 4 cm electrode distance, even 3 min treatment

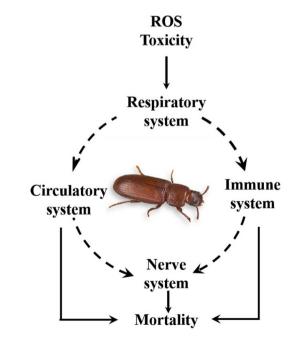


Fig. 7 Mortality of plasma treated insect.

achieved the same mortality. It was also found that there were no significant changes in the colour of the flour after treatment. Likewise, for maida flour, mortality of T. castaneum after plasma treatment was reported by Mahendran et al.  $^{135}$  Results indicated that 1750 V at 2 cm for 3 min, 1750 V at 4 cm for 7 min, and 3000 V at 4 cm for 3 min treatments produced 100% mortality. Different mechanisms of insect mortality are shown in Fig. 7. Apart from disinfestation, plasma treatment also assisted in reducing the microbial load (7.22 CFU per g to 4.73 CFU per g) of tenebrio flour and altered the flour properties such as solubility, water and oil binding capacity.  $^{136}$ 

Studies have found the effect of plasma treatment on rheological properties and changes in flours. Held  $et~al.^{137}$  reported the flour and dough properties of hard and soft wheat. Secondary structure contribution to protein ( $\beta$ -turns,  $\alpha$ -helices, random and  $\beta$ -sheet) was significantly affected after plasma treatment. Peak maximum hard, soft and intermediate flour time increased after plasma treatment while extensibility values decreased. Thirumdas  $et~al.^{138}$  reported the changes in rice starch after CP treatment. At 60 W power level, gelatinization temperature (GT) reduced after 10 min of treatment. At the same time, the pasting temperature was reduced for all treatments. An increase in peak viscosity, storage modulus (except 60 W to 5 min) and loss modulus (except 60 W to 5 min) was

found. Bahrami *et al.*<sup>139</sup> studied the effect of plasma on the functional properties of wheat flour. It was found that protein fractions in flour changed with a higher dosage, while non-polar lipids and glycolipids fractionation had no difference after treatment, and the dough became strong due to protein and lipid modifications. When short and long rice flours were treated with plasma, it showed an increase in transmittance, swelling power, and gel syneresis. Though rice protein was not affected by plasma treatment, the pasting properties of long rice flour increased due to starch molecules' cross-linkage by ozone oxidation. In contrast, after treatment, amino acids were reduced.<sup>140</sup>

Pea protein-rich flour (PPF), pea starch-rich four (PSF), pea testa flour (PTF), and pea protein isolate (PPI) fractions of pea flour were treated in surface DBD plasma, which led to a mass loss of 2.1, 1.2, 1.3, and 1.1% for PPF, PSF, PTF, and PPI, respectively. Colour changes were increased for PPF and PPI and reduced for PSF when exposure time increased. PPF and PTF water and fat binding increased as treatment time increased to 10 min. Misra *et al.* Feported plasma's effect on wheat flour's rheological properties. Peak time and peak integral of soft and hard flour increased significantly when voltage and treatment time exceeded 60 kV and 5 min. Ozone concentration increased with respect to time and input voltage

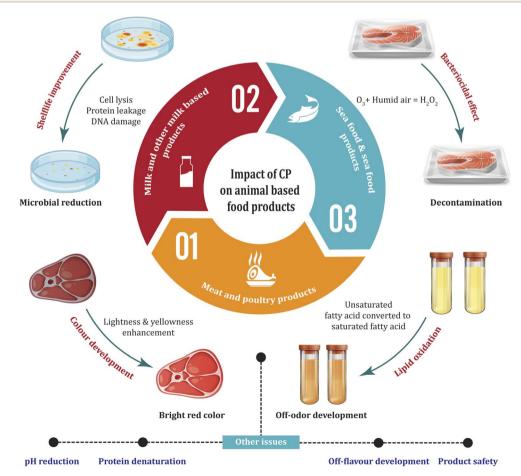


Fig. 8 Impact of cold plasma on animal based food products.

increased.  $\alpha$ -helix,  $\beta$ -turn +  $\beta$ -sheet percentage were reduced at 60 kV for 5 min exposure in both samples. However, they increased at 70 kV after 10 min time in soft wheat flour.

Plasma-induced flour functionality changes also resulted in end product quality changes. Menkovska et al. 142 reported the effect of plasma on wheat flour and dough quality changes. An increase in farinograph values (except dough softening) was observed with respect to treatment time. Similarly, alveograph values increased (except elasticity) with respect to time. Shape formation ratio, specific and total loaf volumes increased from 0.47 to 0.58 h per day, 3.79 to 4.55 cm<sup>3</sup>, and 560 to 657 cm<sup>3</sup>, respectively, after 45 min of treatment due to gas retention and crumb cell increase.

Few research studies have focused on the mode of action of ROS on insects, while the effect of UV and RNS on insects is unclear. In addition, PTW and CAW have not produced the same mortality, even though they produced similar pH. Thus, the actual reason behind the mortality of insects in PTW is unclear. Industrial use of PTW has many practical issues since it hydrates/increases the moisture content of produce.

# 4.3. Processed and minimally processed fruits products

The primary reason for minimal processing is to preserve the food with minimal quality changes. However, as non-thermal processing, plasma can help retard browning and microbial activity without adding preservatives and retains the quality of the product. Browning in cut fruit surface is the major problem in preservation. In the case of apples (Pink Lady®, Fuji, Modì®, Red Delicious (RD)), the browning area reduced significantly after plasma treatment due to less residual PPO activity of 10% and 50% in Fuji and Modi, respectively, after 60 min. This could be due to the changes in the secondary structure of proteins and modification of the amino acid side chain of enzymes. 143 Along with browning control, other parameters such as soluble solids, dry matter, acidity, firmness, and rupture strength were increased;144 however, antioxidant capacity and total phenol index (TPI) were reduced in apples.145 When the juice was extracted from apples, more than 4 logs of microbial reduction were achieved at 30 W (40 s), 40 W (40 s), and 50 W (30 s) without TSS change. However, the treatment affected total phenolic content (41.7 to 32.4 GAE mg/100 g), antioxidant capacity, pH, and colour values.146 These data confirmed the study of Surowsky et al.147 in which a plasma jet was applied to reduce Citrobacter freundii loads in apple juice by about 5 log cycles after a plasma exposure of 480 s using argon and 0.1% oxygen plus a subsequent storage time of 24 h. Similar effects were found in sour cheery nectar and other juices (tomato, apple, and orange) with a minimum of 1 log reduction in E. coli at 650 W after 120 s treatment time. However, in this case, TPC increased by more than 14% except in orange juice (9.52%). 148 However, for white grape juice along with S. cerevisiae inactivation, phenols (720.62) to 445.02 GAE μg per mL), flavonoid (265.21 to 231.04 CE μg per mL), DPPH (88.16 to 82.24), and antioxidant capacity (679.35 to 637.61) were reduced after plasma treatment. The degradation of aromatic rings in phenols due to RS causes a reduction in

total phenolic content. Thus it is also reflected in DPPH and antioxidant capacity reduction.149

Wu et al. 150 reported the effect of corona electrical discharge (CED) plasma on banana starch property changes. DSC's onset temperature increased to 60.0, 60.5, and 61.2 °C from 57.2 °C at 30, 40, and 50 kV cm<sup>-1</sup> intensities due to the cross-linkages created by RS in the polymeric chain of starch granules. This effect was seen in peak and conclusion temperatures increase and gelatinization enthalpy reduction. In terms of pasting behaviour, peak viscosity reduced from 100.4 (control) to 44.4 RVU (50 kV cm<sup>-1</sup>) while pasting temperature reached 92.1 °C from 74.4 °C (control) due to higher crystallinity and swelling resistance of starch.

Though plasma can reduce the microbial load of food products, it also causes certain quality deteriorations in it. Microbial reduction can be correlated with the changes in the acidity of the plasma-treated sample, but matrix effects have to be considered for clear identification of the inactivation mechanisms, and a focus on evaluating the correlation between the plasma treatment parameter and the subsequent product quality changes is needed.

#### 4.4. Essential oil extraction and quality

Essential oils naturally have anti-bacterial properties; when treated with plasma, the properties can be enhanced and reduce the quantity of EO required to perform the same effect. While the treatment also affects the EO components due to higher oxidation in a few studies.

EO obtained from plasma-treated sweet basil produced more antioxidant activity with higher eugenol content, while the treated seed growth after a month was 40 mm higher than the control.<sup>27</sup> Similarly, an increase in the antimicrobial property of clove oil was achieved after plasma jet processing. Due to this, the minimum concentration required for microbial inhibition (A. niger, Penicillium sp., and Rhizopus sp.) was reduced for clove oil and eugenol.151 Other than improving EO quality, the extraction yields also increased in microwave pretreated lemon peel at different plasma power levels (1.0 kV, 1.5 kV, and 2.0 kV), and it was reported that the increase in lemon peel EO yield is due to the rupture that happened on the oil glands by etching, and it was confirmed by SEM images. 152 Similarly, Ebadi et al. 153 reported the same in lemon verbena (Lippa citriodora Kunth.) without any pretreatment after short-time plasma exposure and found an increase in spathulenol (8.1%) and globulol (7.3%) content after 5 min treatment. Earlier, Kodama et al. 154 studied the lemon peel with different gases and peel sizes to understand the effect of process and product variables on extraction. The results found that DBD used with Ar gas induced damage in lemon peel oil glands and increased the EO extraction in the initial stages of treatment. However, the surface area increase (size reduction) reduced the limonene,  $\gamma$ -terpinene, and  $\beta$ pinene concentrations in EO due to overexposure. Similarly, plasma treatment induced both negative and positive effects in the bioactive compounds of essential oil extracted from turmeric powder. However, the treatment reduced the microbial count from turmeric powder by 1.5 logs. 155

### Effect on the food products of animal origin

### 5.1. Dairy products

Milk and its products are highly perishable and often contaminated by bacteria such as Salmonella spp., Streptococcus, Coliforms, Enterococcus, Bacillus, etc. Subjecting the milk to plasma treatment generates free radicals inside the sample and offers antibacterial properties. However, the antibacterial nature of the milk relies on the plasma treatment intensity. Therefore, DBD plasma is the most studied plasma treatment for milk and milk products (fat-free- dry powder and cheese). The treatment was also effective against most milk-contaminating microorganisms (Ex: E. coli, L. monocytogenes, and Salmonella typhimurium). Further, the plasma treatment produced superior quality products compared to pasteurized ones. However, its effectiveness is not superior to UHT treatment. 156,157

 $H_2O_2$  reactive species are the major cause for the increase of milk's antibacterial properties following plasma treatment. However, few studies suggest direct damage in the microbial cell walls and protein leakage due to plasma species. 158-160 Nevertheless, based on the plasma discharge type, the mechanism of microbial inactivation will vary. Specifically, the microbes present in the liquid of the thin layer will observe more direct cell wall damage than the ones present in the larger volumes. The indirect plasma species effect in the high-volume liquid samples is the plasma species's high reactivity and dissolving property. Therefore, the secondary reactive species generated in the liquid samples will have a greater influence on the microbes when using a larger volume of samples.

Plasma treatment tends to reduce the pH of liquid samples due to the generation of H<sup>+</sup> ions. However, in milk samples, the phosphate and milk casein buffering effect limits the pH change and causes only a mild reduction. Short-time plasma treatments did not affect the milk and milk products' colour. However, long exposure resulted in the reduction of yellowness and lightness. Further, the treatment also increases the saturated fatty acid content in the milk by oxidizing the unsaturated fatty acids. Apart from that, lipid oxidation also increases in milk products due to the oxidation of plasma species. Similarly, the reactive species oxidize sulfonate and nitrate the side chains of amino acids.156

### 5.2. Seafood

Cold plasma is effective against most spoilage-causing microorganisms such as L. monocytogenes, Staphylococcus aureus, and Enterobacteriaceae. However, depending on the packing conditions, the treatment effectiveness varies for different seafood products. For example, using CO2 and O2 gas inside the package of the seafood product enhances CP-assisted microbial reduction. In addition, the natural humidity inside the packaging material allows the plasma reactive species interaction and subsequently provides a bactericidal effect to the microbes post the plasma treatment.161

During plasma treatment, the hydrogen in the sample dissociates into H<sup>+</sup> ions and reduces the pH of the sample. The

increase in acidity subsequently triggers protein breakdown. In addition, reactive species also denature proteins through oxidation and fragmentation. The oxidation process causes the proteins to form aggregates with cross-linkages. Like other lipid sources, the seafood lipids oxidize due to plasma treatment and induce off-flavor upon prolonged exposure. However, CP treatment can retort the activity of several enzymes (α-amylase, lipase, alkaline phosphatase, peroxidase, and lipoxygenase) that spoil seafood. Furthermore, CP improved the colour value of the seafood samples by increasing the  $L^*$  and b values through lipid oxidation and pigment production (Fig. 8).161

### 5.3. Meat and poultry

Salmonella spp., E. coli O157:H7, S. aureus, L. monocytogenes, and Campylobacter jejuni contaminate most meat and poultry products. These contaminants can be removed effectively using CP treatment. In addition, studies have shown the decontamination efficiency of cold plasma in many types of meat and poultry products (i.e., pork, beef, poultry meat, and egg). The mechanism of action is similar to that discussed in the earlier sections. However, the quality characteristics vary based on the product's nature.162

At minimal exposure level, CP treatment does not induce any colour changes. However, at longer exposure times, the  $L^*$ (reduce), a (increase), and b (increase) values will change significantly. However, the treated sample partially recovered from the colour degradation during the storage period. Similarly, most of the plasma-treated meat and poultry samples observed a minimal effect on the off-odor and water content. However, it can be avoided by minimizing the treatment duration. When it comes to pH, inert gas CP treatment did not cause any considerable changes in meat or poultry products. However, excess nitrogen reactive species reportedly reduced the pH of the CP-treated samples. Nevertheless, the lipids in meat and poultry products also are oxidized due to the reactive plasma species. Further, the treatment also results in the degradation (oxidation) of proteins due to the acidic compounds (H<sub>2</sub>O<sub>2</sub>) produced by CP treatment.162

# Sensory and consumer perception of plasma treated-food products

Though non-thermal technologies preserve the freshness of food products, some techniques impair the sensory qualities. For example, the oxidation of lipid compounds during plasma treatment results in the production of off-flavor-producing compounds. On the other hand, CP treatment also improved the sensory qualities of certain food products. For instance, cut cantaloupe exhibited better sensory characteristics after 40 kV plasma treatment (90 s). Whereas 15 kV CP treated dairy beverage did not cause any sensory attribute changes to the product. However, products like tender coconut had a negative impact (chemical odour) on the sensory quality due to CP (49.4 kJ kg<sup>-1</sup>) treatment. 163 Therefore, understanding the CP effect on the sensory properties of different food products will be a breakthrough in plasma research. Nevertheless, consumer

perception of the plasma-treated products also needs to be counted for the success of plasma-treated food products. In this regard, consumers' perceptions of food products can be analyzed through the brain-computer interface method to obtain unbiased results from the consumers. 164 Therefore, Coutinho et al. 165 surveyed the perception of chocolate milk with 1085 participants. The study revealed that consumers are more concerned about the CP-treated product price and feel the plasma-treated product will be costlier than the conventionally processed products.

### Other novel applications

### 7.1. Waterless (hydroponic) plant cultivation

Plasma activated water not only assists plant growth in stress (saline, drought, or heavy metal) conditions but also in soilless conditions. Generally, the nitrogen required for plant growth is supplied directly from the soil or from soil microbes. Hence, when the plants are grown in hydroponic conditions, substrates like coir peats, rice husk, etc., are used to support the plant root and provide nutrients. However, PAW itself contains RNS that act as a nitrogen source for plants and eliminate the requirement of substrates. In addition, ROS species also produce H<sub>2</sub>O<sub>2</sub> in PAW, which helps in controlling infections on the plant roots.166 Nevertheless, the PAW effect on plants may not always be positive, as few studies mentioned a negative effect on plant growth (Lettuce); however, after PAW supply, the secondary metabolites (Epicatechin, rutin, quercetin, total phenolic) were found to be high in both roots and leaves. 167,168 However, a recent study utilized the plasma jet generated PAW solely for fulfilling the nitrogen requirement of corn and lettuce plants and assisted in seed germination and plant growth. 169,170

### 7.2. Cold plasma as green fertilizer

The rapid climate change and increased population will have an impact on food shortage in the future. Therefore, increased crop production is needed, which is done with the utilization of chemical fertilizers for agricultural crops. Hence a sustainable technology with less energy consumption and reduced chemical fertilizer use are of interest to researchers and producers to meet the demands.171 Cold plasma has the potential to be used as a green fertilizer in the future due to its increased efficiency and eco-friendly production. As the plasma treatment will lead to the production of reactive oxygen species and reactive nitrogen species in water, it acts as a fertilization liquid facilitating plant development. The H<sub>2</sub>O<sub>2</sub> formed during the plasma treatment initiated the abscisic acid hormone and gibberellin, which is responsible for seed germination and dormancy. Similarly, the NO<sub>3</sub> generated in PAW has a direct impact on the growth rate of the plant, as nitrate is a crucial element for plant growth. One of the major advantages of PAW as a green fertilizer is the presence of nutrients as ions rather than salts, which helps in easy absorption.170 Matra et al.172 have found that the PAW spray generated from the gliding arc plasma generator enhanced the amount of nitrogenous fertilizer in treated water. Similarly, Subramanian et al.173 have reported that PAW

generated from DBD has a higher specific energy of nitrogen, which is desirable for agricultural purposes. Stoleru et al. 168 have also concluded that the nitrites and nitrates from PAW worked as a fertilizer and enhanced the growth of plants. Though there are many studies on PAW as green fertilizer, the appropriate quantity of water used and concentration of nitrates could be a major breakthrough in this application.

### 7.3. Shape transformation

3D printing and sessile drop drying are the two commonly used additive manufacturing practices to produce shape-changing 2D food structures (xerogels). Both these techniques produce flat 2D food structures that can transform their existing structure into a defined 3D structure under external stimuli (water, oil, drying, or pH) contact.174-177 However, to achieve the desired shape-shift, flat 2D structures need to be coated with food-grade constraint materials (i.e., ethyl cellulose). These constraints act as a barrier between stimuli and food surfaces and limit the interaction in the coated areas. However, the uncoated areas interact with stimuli and induce stress gradients between coated and uncoated areas. As a result, the relative expansion increase causes the food structure to fold or bend in a specific pattern to produce a desirable 3D structure. 178-180 Nevertheless, surface wettability (i.e., water absorption and oil absorption) and the binding nature of food surface with constraint material decide the success of these shape-transforming foods. Both these factors can be improved by treating these 2D foods in plasma. Plasma etching increases the surface area of food, thereby its binding behaviour and water/oil absorption.30,180 Hence, to utilize the advantages of plasma treatment in shape transformation, Gupta et al.181 treated wheat xerogels using glow discharge plasma (power: 7.32 W, duration: 5 min) and then coated them with ethyl cellulose in a defined pattern. Due to plasma treatment, the swelling gradient created between coated and uncoated areas of xerogels increased and resulted in better shape transformation when immersed in 90 °C hot water. Similar results were obtained in Stephen et al. 182 study on oiltriggered shape transformation, where the author treated corn xerogels in cold plasma (voltage: 1 kV, duration: 5 min) prior to cellulose acetate linear strip coating. As a result, the 2D structure curled into a spiral shape within 2 s in hot (220 °C) coconut oil due to the high relative extension and constraint material binding. Likewise, Cheeyattil et al. 183 obtained flower shape and samosa shape in the barley flour xerogel through oleomorphic shape shifting. However, same xerogel can be used to obtain water and oil-based shape transformation. Therefore, Jaspin et al.179 tried obtaining complicated flower shapes from plasmatreated flat xerogels using hot water and oil stimuli and succeeded in both attempts.

### Conclusion

Plasma treatment significantly affected the agri-food chain in terms of maintaining the quality and safety of products, increasing germination, extraction efficiency, plant growth, and removing hazardous contaminants from water, soil, and agricultural produces. The interaction of plasma with food products, microbes, allergens, toxins, enzymes, insects, and other constituents varied depending on process variables such as gas composition, gas flow rate, power level, frequency, treatment time, and type of plasma chamber; product variables such as cultivar type, moisture, and other compositions; and other variables such as nature of pesticide, microbes, chemical contaminants, and their initial concentrations. Different mechanisms were involved in different plasma applications, such as microbial inactivation by surface etching and DNA modification, chemical removal by oxidation, allergen removal by protein and lipid functionality modification, germination by surface modification (wettability and surface etching), seed tolerance by heat shock protein production, disinfestation by nerve toxin effect, higher EO extraction by oil gland damages, browning retardation by changes in the secondary structure of proteins and modification of amino acid side chain, etc. However, many other reasons, such as species and cultivardependent efficiency change of treatment in microbes and agricultural commodities, extract quality improvement, uncertain quantity changes in food properties (i.e., increase and decrease in TPC, colour changes, etc.), the effect of RS on consumption to human health and plant growth needs to be explained in further research studies. Also, plasma treatment's large-scale application in these areas is still not stated. Laboratory experiments were mostly done for a lesser quantity of samples. Cost, efficiency, safety, and productivity will be the concern when plasma treatments are used for large-scale operations. Understanding plasma treatment and RS mode of action can increase the chance of productive utilization of plasma treatment by avoiding deterioration during treatment. Some studies have mentioned the long-term effect of RS on product quality and safety.

### Conflicts of Interest

The authors also declare that there are no conflicts of interest.

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