Oxidative Stress Associated with Chilling Injury in Immature Fruit: Postharvest Technological and Biotechnological Solutions

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Abstract: Immature, vegetable-like fruits are produced by crops of great economic importance, including cucumbers, zucchini, eggplants and bell peppers, among others. Because of their high respiration rates, associated with high rates of dehydration and metabolism, and their susceptibility to chilling injury (CI), vegetable fruits are highly perishable commodities, requiring particular storage conditions to avoid postharvest losses. This review focuses on the oxidative stress that affects the postharvest quality of vegetable fruits under chilling storage. We define the physiological and biochemical factors that are associated with the oxidative stress and the development of CI symptoms in these commodities, and discuss the different physical, chemical and biotechnological approaches that have been proposed to reduce oxidative stress while enhancing the chilling tolerance of vegetable fruits.

Keywords: chilling injury; immature fruit; oxidative stress

1. Postharvest Technology and Physiology of Immature Fruits

1.1. Immature Fruits

A number of immature vegetable-like fruits are produced by dicotyledonous species of great economic importance, within the families Cucurbitaceae (pumpkins, cucumbers, zucchini, bitter gourds and luffa), Solanaceae (eggplants, bell peppers), Fabaceae (peas, broad beans), and Malvaceae (okra). These are herbaceous and annual species of subtropical or tropical origin, contributing with fibre, vitamins and other non-nutritive but beneficial additions to the human diet.

Thus, eggplant fruit is rich in polyphenols, including hydroxycinnamic acid and its derivative chlorogenic acid, which have a potent antioxidant capability [1]. Eggplants and other immature fruits are also excellent sources of natural pigments and other antioxidant constituents such as chlorophyll a, chlorophyll b, and ascorbate [2].

Cucumbers have a very high percentage of water and very few calories, along with potential antidiabetic, hypolipidemic and antioxidant activity, mainly based on their high polyphenol content [3], up to 10 mg/100 g of flavonols and nearly 60 mg/100 g of proanthocyanidins [4]. Moreover, cucumber cucurbitacins exhibit anti-cancer activity as well as having purgative, anti-inflammatory and cosmetic pharmacological applications [5]. These medicinal properties are highly valued, both for their therapeutic activity within Indian and Chinese folk medicine and for use in the cosmetic industry [6].
In zucchini and other types of summer and gourd squash, the content of antioxidant compounds is of increasing interest for breeders, who have tried to increase the content of carotenoids and other nutritional compound and to improve external appearance and organoleptic characteristics [7,8].

Bitter gourd is notable for its medicinal properties. It is traditionally used in India, Korea, China and other Asiatic countries for the treatment of diabetes, its most common traditional use, but it is also used for the treatment of several illness such as dysmenorrhea, rheumatism, psoriasis and other [9]. Its medicinal properties are based on its high content of phenolic and saponin compounds, which are associated with antioxidant activity [10,11], and other compounds such as cucurbitane-type triterpenoids, and cucurbitane-type triterpene glycoside and insulin-like peptide [12–14]. The immature fruit of bitter gourd contains a high vitamin C content and it is a good source of vitamin A and some minerals such as iron [10,11,15].

Fresh luffa fruits are used as vegetables, but their seeds are used in traditional medicine in China due to antipyretic and anthelmintic properties [16]. The constituents of luffa fruits show antioxidant activity, due to their high vitamin C content, plus carotenoids and phenolic compounds such as catechins, flavonoids and anthocyanins [17]. These compounds show preventive effects on several types of cardiovascular complaints and cancer [18]. Du et al. [17] revealed that luffa fruits are rich in phenolic substances with high antioxidant potential such as cinnamic acid derivates and the flavonoid glycosides. Moreover, the luffa fruit is a recognized antibacterial agent due to its high content in tannins [19].

1.2. Harvest Time/Optimal Developmental Stage

The quality and the postharvest life of immature fruits are conditioned both by the developmental stage of immature fruits and the choice of harvest time [20]. Several harvesting indices (HI), including fruit size and shape, colour, texture, glossiness, among others, have been developed to assess the harvest time of fruit and vegetables [21]. The mature fruit HI are related to maximum growth and full ripening, a process required for the fruits to acquire their optimal organoleptic properties [22]. In immature fruits, however, the key parameter of the HI is fruit size, which is mainly chosen on the basis of consumer and market demands.

Since vegetable fruits are green at the edible stage, before seeds are fully enlarged and hardened, they can be harvested at various stages of development [23]. Zucchini HI, for example, are mainly based on the size and colour required by the marked (with an average length of about 20 cm), and the fruit being harvested just before hardening and darkening of fruit peel occurs and before undesirable seeds start to develop [24]. Firmness, colour and external glossiness are also common parameters used to define the HI of eggplants, cucumbers or bell peppers [25].

Cucumbers are harvested at nearly full size, but always before the seeds become hardened, at the moment when jelly-like material fills the seed cavity. The quality of a cucumber is also based on fruit glossiness, firmness and dark green peel [26].

Eggplants are harvested before reaching full size; however, there is a wide range of ontogeny stages at which they could be marketed [3] and the quality is primarily based on globular or elongated shape, firmness and a dark purple peel. Okra pods, on the other hand, are harvested while they are still immature; that is, having accumulated enough mucilage, but before becoming fibrous. This generally occurs within two to six weeks after anthesis [27].

1.3. Postharvest Physiology

Despite their great commercial and cultural importance, vegetable fruits are very high perishable commodities. Their epidermis is not yet fully developed and they are harvested at a developmental phase in which storage compounds have not accrued. Their high respiration rates, associated with high rates of dehydration and metabolism, lead to a rapid spoilage during storage [28]. The short commercial life of these fruits is therefore conditioned by their developmental stage at harvest time and their particular physiology during their postharvest storage conditions. Moreover, these products,
due their tropical or subtropical origin, are susceptible to chilling injury (CI) when stored at low but non-freezing temperatures.

Three main factors are known to control the postharvest physiology of immature fruits: the respiration rate, the production of ethylene, and weight loss associated with dehydration. The respiration rate of immature fruit is much higher than that of mature fruit. While tomatoes, melons and mangoes have respiration rates ranging from 16 to 46 mg CO$_2$ kg$^{-1}$ h$^{-1}$ at 10 °C, cucumbers, summer squash and okra have rates ranging from 30 to 143 mg CO$_2$ kg$^{-1}$ h$^{-1}$ at the same temperature [25]. Respiration can be controlled by several factors such as temperature and atmospheric composition.

The speed of metabolic processes is increased by a factor of two for every 10 °C; therefore, respiration rates can be slowed down by reducing the temperature of fruit, although vegetable fruits are not adaptable to long-term storage because they are susceptible to CI [29]. The optimal storage temperature for immature fruit oscillates between 10 and 13 °C, as CI are induced at temperatures below 8–9 °C, this limits its long-distance distribution and marketing. Moreover, most vegetable fruits are susceptible to dehydration and mechanical injury [30]. Since mechanical injury also affects the fruit respiration rate, correct postharvest management that includes not only the control of the temperature but also the avoidance of mechanical damage is necessary to maintain the quality of the fruit [31].

The plant hormone ethylene is also important in controlling fruit physiology during its postharvest conservation period. Since they are immature at harvest, the fruits in question behave as non-climacteric and produces therefore a low rate of ethylene at harvesting and during storage [32,33]. In mature fruit the most conspicuous response to ethylene is related to ripening, but in green vegetables and immature fruits exposure to low levels of ethylene causes yellowing of green tissues, thus reducing the postharvest life of the commodities.

There are several factors promoting ethylene production, including mechanical injury, decay, insect damage and some types of stress such as low or high temperature. Megías et al. [34] demonstrated that cold-induced ethylene in zucchini is not produced during the cold storage period but after transferring the fruit to 20 °C. Also, the production level depended upon both the duration of cold storage and the period of conditioning time at room temperature [34,35]. It is indeed remarkable that, even after cold storage, ethylene production is dependent upon the level of sensitivity of the fruit to cold. Thus, sensitive cultivars of zucchini presented higher ethylene production, while less sensitive ones showed a lower level [36,37].

This cold-induced ethylene production has also been observed in other immature fruits such as cucumbers, eggplants and other climacteric and non-climacteric fruits such as citrus, kiwis and pears, among others [38–42]. It is always associated with an upregulation of ethylene biosynthesis genes [37,43–47].

Dehydration is also a major cause of fruit wilting and softening, and negatively affects nutritional quality. The dermal system of fruits and vegetable (cuticle, stomata, epidermal cells, trichomes) is responsible for regulation of water loss. This system is developed during the growth and development stages of the fruit so that it can fulfil its function when the fruit is mature [48]. The partial development of the dermal system makes immature fruits highly susceptible to water loss. The first symptom associated with fruit dehydration is wilt, but in cucumbers it has been demonstrated that dehydration also upregulates the activity of the cell-wall-degrading enzymes like poligalacturonase and pectinesterase, which finally leads to fruit softening and shrivelling [49]. Moreover, water loss increases ethylene production, which, as Lurie et al. [50] and Kubo et al. [49] pointed out, could explain the accelerated senescence observed in bell peppers and cucumbers.

Massolo et al. [51] found that eggplant fruit treated with 1-methylcyclopropene (1-MCP) maintained fruit firmness concomitantly with a reduction in water loss, this probably being caused by a delay produced by 1-MCP in senescence of the calyx.
2. Chilling Injury and Oxidative Stress

2.1. Chilling Injury (CI)

The storage of fruit under cold conditions is a generalized technology used to avoid rapid
decay and maintain quality. Low temperature slows down many of the processes responsible for
the deterioration and loss of quality in vegetable fruits [52]. However, many fruits and vegetables,
including immature fruits, are susceptible to chilling injury (CI). Their storage at cold but non-freezing
temperatures triggers a number of CI symptoms that irreversibly reduce the external and internal
quality of the product [53].

Chilling injury refers to a syndrome that involves several physiological events, as well as the
characteristic and recognizable symptoms of cold-stored fruit. The type and extent of this syndrome
varies with the species, cultivar, cold storage conditions and other factors, including farming conditions.
For example, while in eggplants a characteristic expression of chilling injury includes a net browning
of the pulp [51], in cucumbers and zucchinis the damage may include surface pitting, dehydration
and large sunken areas, as well as discoloration. There is, therefore, no one single symptom or type of
damage that is common to all products and would allow for the assessment of CI in all commodities.

Nevertheless, there are instances of several symptoms appearing together that can be recognized
as the consequence of cold storage. Some CI changes occur at cellular level, including alterations of
membrane structure, cell plasmolysis and increased electrolyte leakage [54,55]. Others changes imply
alterations in the metabolism, including higher levels of ethylene production, and the accumulation of
abnormal compounds such as malonyldialdehyde (MDA) as a resulted of either anaerobic respiration
or oxidative damage [24,37].

One of the most notable CI-associated alterations, however, affects the external appearance of
the fruit and include damage to the fruit surface, including pitting, large sunken areas, discoloration,
translucent water-soaked spots and water-soaked areas and deep lesions that can reach the
subepidermal tissues [55]. These macroscopic changes are commonly used to assess the extent of cold
damage [24,36,46,56].

The most common CI symptoms in immature fruit include weight and firmness loss, electrolyte
leakage, and appreciable damages to the fruit surface. In cucumbers, watery pitting is the main
symptom of CI [57]. Ultrastructural analysis of this damage has revealed that they are associated with
cracks in the cuticle and the sinking of epidermal cells near the stomata, which together lead to a raised
transpiration rate, as well as a decay due to the growth of necrotrophic fungi [55,58]. Another specific
symptom of CI in cucumbers is discoloration due to chlorophyll degradation that takes place at low
temperature [59].

In the case of zucchini, pitting is the main chilling symptom. Microscopical analysis of this pitting
has shown that the surface depressions are caused by cell death and cell collapse, associated with
solubilisation of pectin and cell wall degradation [60]. When the symptom of pitting is visible, chilling
decay is irreversible; Balandrán-Quintana et al. [61] suggested that respiration rate may be an indicator
of chilling damage before visible symptoms appear and injury becomes irreversible.

At the mature green or breaker stage, before the fruit has completely ripened, bell peppers are also
susceptible to cold storage, although the degree of susceptibility depends on the cultivar [62,63]. Green
bell peppers show similar symptoms to those described as appearing in cucumbers and zucchini, along
with shrivelling, resulting from fruit moisture loss and also seed browning [62,64]. These are also major
chilling symptoms in eggplants [65,66]. Browning is caused by oxidation of phenolic compounds by
the enzyme polyphenol oxidase (PPO), which seems to be dependent on low temperature. This enzyme
catalyses the o-hydroxylation of monophenols to o-diphenol and further oxidation of o-diphenols to
o-quinones, which react with amino acids or proteins, generating brown pigmentation [67,68].

Okra is moderately susceptible to CI, with the most common symptoms of chilling injury being
water-soaked lesions, pitting, discoloration, appearance of mould or mildew and increased decay
(even after removal to warmer temperatures, as during marketing). The sensitivity to CI varies
according to okra cultivars and phenological stage of pods. Young mucilaginous pods are more sensitive than larger pods. Usually the symptoms appear after two days of cold storage if the temperature is as low as 2 °C. The green pods turn a brownish olive-green, while yellow cultivars turn brown or brownish-red [69–71].

Luffa fruits are highly perishable and their postharvest life is no more than one week at room temperature. During the process of decay, the fruits appear yellow and are wilting. Luffa is highly sensitive to chilling injury and the most visible symptoms include discoloration, watery brown or black spots on and under the epidermis, and a higher level of decay including severe decomposition, especially if the fruit is being rewarmed after cold storage [72,73]. Table 1 shows the chilling injury symptoms in some vegetable-like fruits.

Table 1. Chilling injury symptoms in some immature fruits.

| Immature Fruit | Symptoms                                                                 | Threshold Temperature (°C) | References  |
|----------------|--------------------------------------------------------------------------|----------------------------|-------------|
| Cucumber       | Surface pitting, increased yellowing and disease susceptibility, water-soaked areas of the flesh | 10–12                      | [26,38,55]  |
| Eggplant       | Surface pitting and scald, browning of the flesh and seeds               | 8–12                       | [51]        |
| Bell pepper    | Surface pitting, water-soaked areas, seed browning, and decay            |                            | [62,74]     |
| Okra           | Discoloration; water-soaked areas; surface pitting; exuding lesions, and decay by mould or mildew, calyx discoloration | 7–10                       | [27,75]     |
| Zucchini       | Surface pitting, large sunken areas, dehydration, discoloration         | 7–10                       | [24,36]     |
| Bitter gourd   | Pitting that coalesced to form large sunken dark brown pits; surface discoloration; internal tissue breakdown; decay | 8–10                       | [76]        |

2.2. Oxidative Stress

2.2.1. Overview

The environmental and physical changes that the produce undergoes before and after harvest may induce the production of reactive oxygen species (ROS), which are the cause of oxidative damage during the postharvest of immature fruit and vegetables, inducing decay of the product and loss of quality. Oxidative stress starts with an elevated production of ROS as a result of numerous processes such as photosynthesis, respiration, photorespiration and oxidative burst, which occur at different cellular locations and within different cellular organelles [77,78]. ROS include superoxide anion, hydrogen peroxide, hydroxyl radical, nitric oxide, and peroxynitrite. If the production of ROS increases dramatically, as occurs under environmental stress, hydroxyl radical reacts with membrane lipids, inducing peroxidation, which leads to membrane degradation. Malondialdehyde (MDA) is a product of this lipid peroxidation, and is used as an indicator of stress in some tissues [79].

During immature fruit postharvest, oxidative stress avoidance is important for the maintenance of fruit quality. Two strategies have been developed to avoid or tolerate oxidative stress, generating a response that includes metabolic changes at biochemical and molecular level [80]. These changes include the induction of the enzymatic and non-enzymatic mechanisms of the antioxidant defence: superoxide anions are detoxified by the enzyme superoxide dismutase (SOD), which produces hydrogen peroxide that can be scavenged by catalase (CAT) and several peroxidases (POD) such as thioredoxin peroxidase and glutathione peroxidase, as well as the enzymes belonging to the Foyer–Halliwell–Asada cycle [78]. In this cycle, ascorbate peroxidase (APX) uses ascorbate for the
reduction of hydrogen peroxide. The oxidized ascorbate is recycled by monodehydroascorbate reductase (MDHAR) and dehydroascorbate reductase (DHAR), in this case oxidizing glutathione that is reduced by glutathione reductase (GR) [81,82]. As for non-enzymatic antioxidants, there are several metabolites in plants such as carotenoids, phenolic compounds, ascorbate and glutathione that can be used as ROS scavengers. In tissues with a high rate of respiration such as developing fruit, in addition to ROS scavenging mechanisms the mitochondrial alternative oxidase (AOX) provides an alternative electron flow. This activity can reduce the production of ROS in situations where the electron transport chain is saturated or suffers some damage [83].

Another way to prevent oxidative stress is the activation of metabolic pathways involved in the maintenance of redox status, such as the GABA shunt pathway [84]. In Arabidopsis, mutants disrupted in the enzyme of the GABA shunt succinic semialdehyde dehydrogenase were unable to scavenge hydrogen peroxide [85].

When the generation of ROS exceeds the capacity of the plant tissue to maintain cellular redox homeostasis, oxidative stress appears [86]. This stress in fruit and vegetables can be detected either directly as accumulation of ROS, MDA, and the apparition of brown pigments added to enhanced electrolyte leakage, or indirectly as changes in enzymatic and non-enzymatic antioxidant systems [87]. With regard to acclimatisation to abiotic stress, several phytohormones play a main role in the activation of the response. It has been reported that an intricate interplay between phytohormones and ROS [88], added to elevated levels of ROS, also modulate the levels and functions of phytohormones [80].

2.2.2. Low Temperature and Oxidative Stress in Vegetable Fruit

The storage of vegetable fruit at low temperature is necessary in order to reduce fruit respiration, thus avoiding postharvest losses. However, cold storage generates CI in many vegetable fruits, such as cucumbers, zucchini and eggplants, by inducing ROS production [3,36,89]. It was proposed that lipid peroxidation was associated with CI, after the study by Parkin and Kuo [79] in which chilling-induced lipid degradation in cucumber fruit was detected. The plastids were suggested as the site of chilling-induced peroxidation. By monitoring electrolyte leakage and changes in thylakoid lipids in cold-stored cucumber fruit, Hariyadi and Parkin [90] proved the existence of oxidative stress during cold storage, and confirmed the sensitivity of plastid membranes to chilling.

Fruit are structures with higher respiratory activity; however, low-temperature storage may induce an uncoupling in the respiratory chain, giving rise to ROS. In plants exposed to low temperature a sharp increase in their cyanide-sensitive respiration has been reported, mainly due to an induction of AOX activity [83]. AOX is a very efficient way of diverting the electron flow and reducing the generation of superoxide anion, hydrogen peroxide, and hydroxyl radical. Low temperature also reduces antioxidant enzymatic activity in fruit of chilling sensitive cultivars, thereby decreasing their ability to cope with elevated ROS production [36,62]. In addition, treatments used to reduce the damage caused by cold storage in sensitive fruits lead to a reduction of oxidative stress by increasing the antioxidant defence [24,91–94].

In cucumbers, a decrease of the non-enzymatic antioxidant glutathione, α-tocopherol, and ascorbate has been found during cold storage [90]. More recently, ROS accumulation has been assessed in cold-stored cucumber fruit, detecting a gradual increase in hydrogen peroxide content and the rate of superoxide anion production during storage [89,95].

In this species, the chilling tolerance depends on the cultivar [59] and is affected by fruit maturity. Fruits at earlier developmental stages are more susceptible to chilling stress, associated with a higher MDA content and electrolyte leakage, than more mature fruit [96,97]. At earlier developmental stages, cold-stored cucumber fruit shows a lower transcription levels of the main antioxidant enzymes (SOD, CAT, POD, MDHAR, DHAR, GR) even after rewarming, which may be the cause higher oxidative damage [97]. At later stages of development, however, the fruit displays a higher ascorbate content, which improves antioxidant capacity and allows longer cold storage [96].
Many studies have been undertaken to determine the defence mechanisms of the zucchini fruit against CI [37,46,60,94,98–101]. Whilst analysing the postharvest condition of several cultivars of zucchini fruit, it has been demonstrated that chilling tolerance in this species is also cultivar-dependent [35,36]. Cold storage induces the accumulation of hydrogen peroxide in both exocarp and mesocarp tissues of this fruit [102]. In addition, a strong positive correlation has been detected in exocarp tissue between hydrogen peroxide and MDA content and CI, whereas CAT activity and CI symptom correlate negatively in this same tissue [36].

Soluble sugars also play an important role in the ROS balance under conditions of stress, either by acting as ROS scavengers, as osmoprotectants or by stabilizing cellular membranes [103,104]. The fruit of a chilling-tolerant cultivar of zucchini accumulated higher levels of soluble sugars under cold stress, associated with an increased antioxidant defence [36,99]. Moreover, the induction of the GABA shunt pathway in cold-stored zucchini fruit has been considered as a contribution to the acquisition of chilling tolerance by increasing the production of reducing agents and energy providers [91,92,100]. It can therefore be deduced that an enhanced antioxidant defence and energy status improves the quality and cold tolerance of zucchini fruit.

In green bell peppers, it has been reported that low-temperature storage causes a decrease in SOD and CAT activities in sensitive cultivars at the most susceptible stages of ripeness [62]. This CAT response could be related to the ultrastructure changes that take place as a consequence of chilling damage, such as plastid degradation and the disappearance of peroxisomes [105]. In this study, a proteomic analysis proved that one of the main processes affected by chilling in pepper fruit is redox metabolism. Among the antioxidant responses in peppers was a significant increase in the induction of AOX in cold-stored fruit [106]. Specifically, this work reported the accumulation of two AOX, one of them (CaAOX1) being an unprocessed transcript due to altered RNA splicing that takes place at 0 °C. Although CaAOX1 is expressed, it is accumulated both during cold storage and during treatments that improve chilling response, induce earlier and higher expression levels of this gene. Another chilling symptom in bell peppers is seed browning, which has also been also associated with certain stages of maturity of this fruit [64]. These authors have reported lower of levels of CAT and POD but higher Superoxide Dismutase (SOD), Lipoxygenase (LOX), Phenylalanine Ammonia Lyase (PAL), Polyphenol Oxidase (PPO) activity in seeds during cold-sensitive stage, as well as increased MDA and saturated fatty acids content.

Eggplants are affected by CI when stored at temperatures below 10 °C, but their sensitivity differs among cultivars, and depends on: the size at harvest; the developmental stage and environmental conditions. However, the differences in chilling susceptibility within the normal eggplant harvest stages are not clear yet. Zaro et al. [3] showed, in two cultivars, that during the early stage of development (baby eggplant) the chilling damage was less than in later stages; these latter containing a higher content in antioxidants. In the darker cultivars of eggplant, the epidermis is rich in anthocyanins, such as delphinidin-3-(p-coumaroyl rutoside)-5-glycoside (nasunin) and/or delphinidin 3-rutoside (D3R), both of which contribute to hydroxyl radical and superoxide anion radical scavenging activities [107]. However, these darker cultivars are very susceptible to browning when stored for prolonged periods or at low temperatures. Indeed, the levels of phenols have been positively correlated with the incidence of browning [108]. Concellon et al. [66] reported an increase in membrane and organelle disruption during the cold storage of eggplants. This loss of cell compartmentalization makes phenols accessible to PPO. In a similar way, calyx browning correlated positively with PPO activity [109].
3. Postharvest Treatments that Reduce Oxidative Stress and Chilling Injury

Different postharvest technologies have been developed to avoid or delay the development of chilling injury during the postharvest storage of immature fruits. The current market trend is to avoid synthetic chemicals, whilst encouraging the use of sustainable postharvest technologies based on physical treatment (temperature, relative humidity and modified atmosphere packaging) and biochemical treatments using natural products and growth regulators (Table 2).

3.1. Physical Approaches

3.1.1. Temperature Treatments

Postharvest heat treatments prior to low-temperature storage have been used to prevent or mitigate the action of CI in certain immature fruits, including cucumbers [110,111], peppers [112] and zucchini [46,60].

Hot water dipping (HWD), intermittent warming (IW) or hot water rinsing and brushing (HWRB) have prolonged the shelf life of vegetables and fruits [113]. These short-term bathing techniques, along with intermittent warming (IW), induce heat shock protein (HSP) genes [113,114], polyamines (PA) production [93,112] and the natural antioxidant defence system within the fruit itself [115,116]. This maintains the quality of some immature fruit commodities [45,117–120]. A recent report in zucchini has also shown that warm-water treatments are effective for maintaining fruit firmness while preventing the proliferation of fruit microflora [121].

Postharvest preconditioning treatments at moderate temperatures before cold storage allow a progressive adaptation of the fruit to chilling temperatures. In zucchini, it has been reported that temperature-preconditioning treatments at 15 °C for two days, before storage at the chilling temperature of 4 °C, reduces CI symptoms by preventing the deterioration of the cell membrane and improving the antioxidant status of the preconditioned fruit [60,122,123].

These treatments were able to reduce lipid peroxidation and ion leakage, which are indicative of a lack of membrane integrity, while diminishing H$_2$O$_2$ content, thereby inducing the activities of antioxidant enzymes such as ascorbate peroxidase (APX) and catalase (CAT), and increasing ATP pool and proline accumulation [60]. The reduction of pitting and fruit weight in temperature-preconditioning zucchini fruits was also accompanied by a reduction of both cold-induced ethylene and the fruit respiration rate in the more chilling-susceptible cultivars [46], suggesting that ethylene is also a modulator of CI in this immature fruit.

In the case of cucumbers, a preconditioning treatment at 10 °C encouraged chilling tolerance associated with a reduced accumulation of MDA and ROS and with induced activity of SOD, APX, and CAT, together with a higher content of the ROS scavengers ascorbic acid and glutathione [124].

3.1.2. The Use of Controlled and Modified Atmospheres

Fruit respiration rate and fruit dehydration are two physiological aspects that limit the shelf life of fresh fruits during their postharvest storage period [125–127]. The use of controlled atmosphere (CA), modified atmosphere packaging (MAP), and edible coatings alters the gaseous atmosphere where the main active gases (O$_2$, CO$_2$, ethylene and H$_2$O vapour) surrounding the fruit [128–133]. Thus, fruit quality was better maintained under CA and MAP, this being associated with reduced oxidative stress in the refrigerated immature fruits of different species such as cucumbers, peppers and zucchini squash.

Although rarely used during storage of immature fruit commodities, there are some reports demonstrating the positive effects of CA on cucumbers and zucchini [94,134–136]. Fahmy and Nakano [134] reported that cucumber fruit stored under CA with low O$_2$ reduced weight loss, changes in skin colour, electrolyte leakage and MDA accumulation. Low O$_2$ storage atmosphere were also responsible for reducing CI in zucchini squash, maintaining higher levels of the polyamines spermidine and spermine [137]. Exposure to 40% CO$_2$ prior to chilling storage at 2 °C were found to be effective in
reducing CI in zucchini squash, while reducing abscisic acid, putrescine, spermidine and spermine content [136]. Under CA with a high level of O₂, however, zucchini fruit slightly induced AOX, as well as SOD, APX and CAT antioxidant activity together with total phenolics, suggesting the involvement of alternative respiratory pathway during chilling stress [94].

3.1.3. The Effects of Different Types of Wrapping

The use of plastic coverings with differing levels of permeability to active gases produces a balance in CO₂ production and O₂ consumption that passively modifies the atmosphere of freshly-packaged produce. Due to its comparatively low cost, MAP has been widely used in the industry for the maintenance of fruit quality and the extension of product shelf life, in both mature and immature fruits [53,75,119,128,138–140].

Individual shrink wrapping (ISW) is a form of MAP that prolongs the shelf life and maintains the harvest freshness of fruits and vegetables. The main advantages of ISW include a reduction in CI, together with reductions in fruit dehydration, deformation and decay, which prevent secondary infection. In cucumbers and peppers, ISW using polyethylene film has been proved to be useful for extending fruit shelf life and reducing weight loss while retaining freshness, colour and firmness [135,141–143].

Shrink packaging in combination with HWRB treatment reduced weight loss, softening, decay incidence and CI in green peppers, while reducing LOX activity and electrolyte leakage [114,118]. Shrink-wrap film was also used to extend the shelf life of zucchini stored at 4 °C, reducing the loss of both fruit weight and firmness as well as reducing CI symptoms. This coincided with a reduction of cold-induced ethylene production, a downregulation of ethylene biosynthesis and signalling genes, a decrease in fruit respiration rate and the accumulation of oxidative stress metabolites such as H₂O₂ and MDA [24].

3.1.4. The Effects of Edible Coatings

Edible coatings also preserve the quality of vegetable fruits by reducing water loss and CI symptoms during postharvest storage, and by activating the antioxidant system that reduces membrane damage and prevents flesh browning. Alone or in combination with essential oils from diverse origins, edible coatings are being increasingly used for their non-toxic, biodegradable and biocompatible properties, reducing thereby the need for disposable non-degradable packaging materials. The application of ceramide lessens CI symptoms in pepper fruits stored at 4 °C, delays the degradation of chlorophylls and soluble proteins, suppresses the accumulation of MDA by maintaining the integrity of the cell membrane, and enhances the activity of the antioxidant enzymes POD, CAT, and APX [144]. Chitosan-based coatings were also observed to improve the shelf life of cucumbers [145,146], zucchini [121], bell peppers, [147–150] eggplants [151] and sponge gourds [152].

3.2. Chemical Approaches

Given that oxidative stress and CI symptoms are controlled by growth regulators and metabolites inducing signal transduction mechanisms, many postharvest technologies are based on treatments using hormones, hormone inhibitors, or metabolites involved in stress responses. These include ethylene inhibitors, brassinosteroids, abscisic acid (ABA), salicylic acid, jasmonic acid and polyamines. Others are based on natural non-toxic products.

3.2.1. Ethylene (ET)

ET regulates a wide range of biochemical, physiological and developmental processes that can damage climacteric and non-climacteric fruits during postharvest storage [153–155]. The use of ET biosynthesis and response inhibitors has been extensively implemented in the industry to extend postharvest self life and to delay senescence and thereby decay of the fruit.
The gaseous ET analogue 1-methylcyclopropene (1-MCP) can irreversibly bind to ethylene receptors, thus avoiding subsequent ethylene response [156,157]. Postharvest treatments with 1-MCP are effective, stable over time and non-toxic, but their effectiveness is highly variable and depends upon the species and cultivar employed, the treatment conditions and the stage of ripening of the fruit. It has been demonstrated that 1-MCP successfully prevents deterioration of fruit quality by delaying ripening or senescence in many fruits, but also by inducing chilling tolerance. This has been reported in the immature fruits of eggplant, peppers and zucchini [37,51,158].

In zucchini, 1-MCP delays the onset of CI symptoms such as pitting and fruit weight loss in the fruit of the most chilling-susceptible cultivars, which coincides with a reduction in the respiration rate and cold-induced ET and includes a downregulation of ET biosynthesis and signalling genes [37]. These results indicate that ethylene is not only a response of the fruit to cold damage, as occurs upon rewarming, but could also play a regulatory role in the onset of CI in the non-climacteric zucchini fruit.

Postharvest technologies based on ethylene removal, such as that of catalyst titanium dioxide (TiO$_2$), and the ethylene absorbents CaCl$_2$ and KMnO$_4$, have been employed to prolong the shelf life of fruit produce [159]. A new palladium-based ethylene adsorbing technology has also been developed, which scavenges ethylene for the control of fruit ripening [159]. Nevertheless, none of these treatments have been so far used for the postharvest management of immature fruits.

3.2.2. Brassinosteroids (BRs)

BRs have also been reported to be effective in delaying CI symptoms in immature fruits. Wang et al. [160] showed that BRs mitigate the action of CI in green bell peppers, accompanied by a reduction in electrolyte leakage, a reduction MDA content, and enhanced the activity of antioxidant enzymes such as CAT, APX, and GR. In the same way, Gao et al. [161] have found that BR alleviated CI and flesh browning in eggplant fruit maintaining membrane integrity and moisture content, by declining in the accumulation of H$_2$O$_2$ and by inducing total phenolics and PAL, PPO, and POD activities.

3.2.3. Abscisic Acid (ABA)

ABA is accumulated in plants in response to many stress factors, including drought, salinity and cold [162,163]. The accumulation of ABA prevents cellular dehydration by promoting stomata closure and the accumulation of osmoprotectant molecules such as soluble sugars and proline [164,165]. Moreover, ABA can induce the antioxidant defence response which reduces the effects of the oxidative stress that appear during dehydration [166], and can promote the induction of genes that are involved in cold and dehydration tolerance. These genes are known as COR (Cold Regulated) and ABRE (ABA Responsive Element Binding) genes [167–169].

In vegetable fruits, we have recently tested that the application of ABA induces cold tolerance in the chilling-sensitive zucchini cultivar Sinatra, while treatment with the ABA inhibitor sodium tungstate reduces the chilling tolerance of the chilling-tolerant cultivar Natura (Carvajal et al., unpublished).

3.2.4. Salicylic Acid (SA) and Jasmonic Acid (JA)

SA and JA, as well as their methyl esters, known as methyl salicylate (MeSA) and methyl jamonate (MeJA), are natural and safe signalling molecules that were applied to fruits and vegetables to enhance CI tolerance and to maintain the postharvest quality of fresh produce [170]. In luffa fruits (sponge gourd), the application of SA alleviates postharvest CI by decreasing electrolyte leakage, reducing the accumulation of MDA and total phenolics, enhancing the activity of antioxidant enzymes such as SOD, CAT and APX, and inhibiting the activity of phenylalanine ammonia-lyase (PAL) and polyphenol oxidase (PPO) [73]. In order to facilitate salicylic acid biological function, efforts have been made to synthesize salicylic-acid-grafted chitosan. The so called salicyloyl chitosan was utilized because of its superior water solubility and adhesiveness to fruit surface [171]. Zhang et al. [116] reported that the
salicyloyl chitosan coating alleviated chilling injury and weight loss in cucumber fruits, which was associated with reduced respiration rate, lower electrolyte leakage, reduced MDA accumulation, and higher activity of the antioxidant enzymes SOD, CAT, APX and GR.

JA and MeJA are also regulators of stress responses [172,173]. The induction of JA genes in Arabidopsis thaliana [174], and rice [175] conferred enhanced chilling tolerance. In species with immature fruits, it has been reported that MeJA treatment alleviates CI and reduces ethylene production in cold-stored eggplant fruits [176], but was ineffective in reducing CI in zucchini squash [177]. In green bell peppers, MeSA and MeJa enhance CI tolerance by inducing the expression of the AOX gene, thereby maintaining the balance between ROS production and general antioxidant system activity [178]. In cucumbers, MeJA induces chilling tolerance by inhibiting H\textsubscript{2}O\textsubscript{2} generation, and enhancing both CAT activity and CAT gene expression [179].

3.2.5. Polyamines (PAs)

PAs are a group of phytohormone-like aliphatic amine compounds, major types of which are the triamine spermidine (Spd), the tetramine spermine (Spm), and their diamine precursor, putrescine (Put). They produce pleiotropic effects in plants, including defence against various abiotic stresses. Since PA and ET share the same precursor, S-adenosylmethionine (SAM), high levels of PA would provoke a reduction in ET levels followed by an increase in CI tolerance as observed in peppers [110,112] and in zucchini [98,99,122,180].

In zucchini, the enhanced tolerance to CI in putrescine-treated fruits was associated with reduced levels of hydrogen peroxide and malondialdehyde, which could be related to changes in LOX activity [180]. These advantageous changes appear to occur by activating the antioxidant system via the accumulation of ascorbate and ferric reducing/antioxidant power (FRAP), and by inducing the activities of APX, CAT and GR antioxidant enzymes [92]. The response of cold stored zucchini fruit to PA treatments could be also due to other physiological responses that protect cells from oxidative damage, including \gamma\textsubscript{-}aminobutyric acid (GABA) degradation via the GABA shunt pathway [180] and via the accumulation of soluble sugars including glucose, fructose and raffinose, as has been observed in putrescine-treated zucchini fruits under cold storage [92,99].

PAs also enhance the chilling tolerance of cucumber plants through modulating antioxidative system in the leaf [181]. There is, however, no data concerning the effect of these plant growth regulators on the oxidative status and CI tolerance of the cucumber fruit.

3.2.6. Nitric Oxide (NO)

Nitric oxide (NO) is an important signalling molecule involved in many plant physiological processes [182]. It has been reported that NO protects plant cells against oxidative stress by reducing ROS accumulation, and has been used as a postharvest treatment for maintaining fruit quality and alleviating CI symptoms. In cucumbers, the treatment with NO was effective in alleviating CI symptoms, while reducing oxidative stress damage due to a decrease in O\textsubscript{2}\textsuperscript{-} and H\textsubscript{2}O\textsubscript{2} and by activating the antioxidant enzymes CAT, SOD, APX and POD [95,179]. NO fumigation was also applied to extend the postharvest life of green beans [183].

3.2.7. Other Chemical Treatments

Calcium treatments have been shown to retain the firmness and quality of vegetables and fruit as Ca contributes to the structural integrity of plant cell walls and membranes [184]. De Bruin et al. [121] reported that in zucchini fruit treated with CaCl\textsubscript{2} flesh firmness was effectively maintained during 14 days of storage, in contrast to the control fruit, which showed a decline in flesh firmness after the first seven days of storage.
6-Benzylaminopurine (6-BA) is closely related to plant stress tolerance. Little research has been carried out on the postharvest cold tolerance of vegetable fruits. Chen and Yang [89] have shown that 6-BA alleviated CI in cucumber fruit through improving antioxidant enzyme activity and total antioxidant capacity, and by maintaining higher levels of ATP content and energy charge. Fruits treated with 6-BA maintained higher levels of chlorophyll, ascorbic acid, total phenolics and total antioxidant capacity. Furthermore, under chilling stress, this treatment reduced the increase in membrane permeability and lipid peroxidation, delayed the increase in both \( \text{O}_2^- \) and \( \text{H}_2\text{O}_2 \) production, and increased the activities of SOD, CAT, APX and GR.

Massolo et al. [185] have stated the effect of postharvest cytokinin (CK) treatment on refrigerated, round, soft-rind squash. CK-treated fruit showed slower deterioration and dehydration, and remained firmer than the control. The treatments prevented phenolic-compound accumulation and decreased pectin solubility. The treated squash had higher levels of tightly-bound polyuronides, indicating a substantial delay in cell wall dismantling. CK sprays also reduced neutral sugar solubility from pectin-rich fractions [185].

Glycine betaine (GB), a stress-ameliorating compound, has been used to reduce CI in sweet peppers [186]. The authors suggested that GB increases chilling tolerance by inducing the activity and expression of POD, CAT APX and GR antioxidative enzymes and genes, thus alleviating the potential injury resulting from CI. The amelioration of CI in sweet peppers by GB was associated with a reduction in electrolyte leakage, MDA content, and lipid peroxidation.

### 3.3. Breeding and Biotechnological Approaches

Plant breeding and biotechnology approaches are strategies that show potential for decreasing oxidative stress while enhancing CI tolerance in fruits. Numerous examples of transgenic plants showing reduced CI are described in the literature, although a few of them refer to the postharvest quality of immature fruits.

Breeding for tolerance to postharvest CI has rarely been used for enhancing the postharvest quality of immature fruits. The observed genetic variability of CI among commercial zucchini hybrids [36], and the greater variation among traditional local varieties in gene banks [35], opens up the possibility to achieve new zucchini cultivars by combining different sources of CI tolerance in zucchini.

With this objective in mind, Megías et al. [35] have recently described a study of the mode of inheritance of a source of cold tolerance in zucchini, finding a cosegregation between cold-induced ethylene production and CI index and weight loss, suggesting that cold-induced ethylene can be used as a marker to select postharvest cold tolerance in segregating populations.
Table 2. Postharvest treatments alleviating oxidative stress and CI in immature fruits.

| Postharvest Technologies | Technology                        | Species       | Effects                                                                 | References                  |
|--------------------------|-----------------------------------|---------------|-------------------------------------------------------------------------|----------------------------|
|                          |                                   | Cucumber      | Reduced electrolyte leakage, chilling-induced ethylene, and ACS and ACO activity | [117]                      |
|                          |                                   |               | Reduced electrolyte leakage and MDA, and enhanced PLD and LOX activity  | [110,111]                  |
| Heat treatment           |                                   | Green bell pepper | Reduced CI, electrolyte leakage and LOX activity                        | [114]                      |
|                          |                                   |               | Reduced weight loss, softening, decay and CI                           | [118]                      |
|                          |                                   | Zucchini      | Induced HSP genes, maintained flesh firmness                           | [113]                      |
| Physical                 |                                   | Eggplant      | Retarded CI, reduced spermidine                                         | [119]                      |
| Temperature preconditioning treatment |                          | Cucumber      | Increased soluble solids, ascorbic acid, and MDA, $O_2^-$ and $H_2O_2$, induced activity of antioxidant enzymes, and the scavengers AsA and glutathione | [124]                      |
|                          |                                   | Zucchini      | Alleviated CI and weight loss, reduced $H_2O_2$, MDA and ascorbic acid content, and induced activity of antioxidant enzymes | [46,91,93]                 |
| Controlled atmospheres, CO$_2$ and O$_2$ treatments |                          | Cucumber      | Alleviated chilling injury, weight loss and changed in peel colour, maintenance of electrolyte leakage and MDA | [134]                      |
|                          |                                   | Zucchini      | Reduced CI, increased levels of spermidine, spermine and total phenolics, induced activities of alternative oxidase, SOD, APX and CAT | [94,136,137]               |
| Controlled atmospheres, Use of plastic covers |                          | Green bell pepper | Reduced CI and weight loss, maintained of ACC, Put, and ABA levels, reduced ascorbic acid content | [139,140]                 |
|                          |                                   | Okra          | Reduced weight loss and ascorbic acid content, increased titratable acidity | [75]                       |
|                          |                                   | Eggplant      | Retarded chilling injury, decreased spermidine levels                  | [119]                      |
|                          |                                   | Cucumber      | Reduced weight loss, decay and fruit deformation, maintenance of freshness, colour and firmness | [141,142]                 |
|                          |                                   | Green bell pepper | Reduced CI, weight loss, membrane leakage and LOX activity, induction of HSP from the HSP70 family | [114,118,140,143]         |
|                          |                                   | Zucchini      | Reduced ethylene production and ethylene gene expression, reduced $H_2O_2$ and MDA | [24]                       |
| Ceramide coating         |                                   | Green bell pepper | Maintenance of membrane integrity, reduced MDA, enhanced activity of POD, CAT, and APX | [144]                      |
Table 2. Cont.

| Postharvest Technologies | Technology          | Species     | Effects                                                                 | References |
|--------------------------|--------------------|-------------|------------------------------------------------------------------------|------------|
| Physical                 | Chitosan coating   | Zucchini    | Reduced CI, preservation of flesh firmness                             | [121]      |
|                          |                    | Cucumber    | Reduced CI, electrolyte leakage and MDA accumulation, increased content of soluble solids, chlorophyll and ascorbic acid, SA, and induced activity of SOD, CAT, APX and GR | [116]      |
|                          |                    | Sponge gourd| Delayed PPO, increased content of ascorbic acid and total phenolics    | [152]      |
| Chemical                 | 1-MCP              | Green bell pepper | Delaying senescence associated with enhanced antioxidant enzyme activities | [158]      |
|                          |                    | Zucchini    | Reduced fruit weight loss, respiration rate and cold-induced ethylene, reduced expression of ethylene genes | [37]       |
|                          |                    | Eggplant    | Reduced weight loss and browning, and reduced of PAL, PPO and POD activity, and total phenolics | [51]       |
|                          | Brassinosteroids   | Green bell pepper | Decreased electrolyte leakage and MDA content, enhanced CAT, APX, and GR activities | [160]      |
|                          |                    | Eggplant    | Maintenance of membrane integrity and moisture and reduced flesh browning, reduced phenolic accumulation and repressed PAL, PPO, and POD activities | [161]      |
|                          | ABA                | Zucchini    | Delayed development of CI symptoms                                    | [188]      |
|                          | Salicyloyl chitosan coating | Cucumber | Higher total soluble solids, chlorophyll and ascorbic acid content, reduced electrolyte leakage and MDA, and induction of SOD, CAT, APX and GR | [116]      |
|                          | SA and MeSA        | Sponge gourd| Higher antioxidant activity reduce MDA, enhanced SOD, CAT, APX activities | [73]       |
|                          | MeSA and MeJA      | Green bell pepper | Increased expression of AOX gene                                       | [106,178]  |
|                          |                    | Cucumber    | Reduced H$_2$O$_2$ accumulation enhanced catalase activity             | [179]      |
|                          |                    | Eggplant    | Reduced ethylene production                                           | [176]      |
|                          | MeJA               | Cucumber    | Induced chilling tolerance by inhibiting H$_2$O$_2$ generation and CAT activity | [179]      |
|                          | PAs                | Green bell pepper | Reduction of ethylene production                                       | [112]      |
|                          |                    | Zucchini    | Induction of APX, CAT and GR activities, increased content of ascorbate, FRAP, glucose, fructose and raffinose, and reduced LOX activity | [98,99,122,180] |
| Nitric oxide             | Cucumber           | Reduced lipid peroxidation, O$_2^-$ and H$_2$O$_2$ accumulation, and enhanced CAT, SOD, APX and POD activities | [95,176]  |
|                          |                    | Green beans | Shell life extension                                                   | [183]      |
| 6-BA                     | Cucumber           | Slower deterioration and dehydration, phenolic compound accumulation, and decreased pectin and sugar solubility, delayed cell wall disassembly | [185]      |
| CK                       | Zucchini           | Slower deterioration and dehydration, phenolic compound accumulation, and decreased pectin and sugar solubility, delayed cell wall disassembly | [185]      |
| GB                       | Green bell pepper  | Slower deterioration and dehydration, phenolic compound accumulation, and decreased pectin and sugar solubility, delayed cell wall disassembly | [185]      |
Several biotechnological approaches have been developed to induce tolerance to oxidative stress by enhancing enzymatic and non-enzymatic mechanisms of ROS detoxification. The accumulation of some antioxidant compounds including ascorbate, glutathione, tocopherols and tocotrienols, polyphenols, carotenoids, flavonoids and anthocyanins protect plants against oxidative stress, but the most used ROS scavenging mechanisms in transgenic plants were those that overexpress genes encoding for the enzymes involved in \( \text{O}_2^- \) and \( \text{H}_2\text{O}_2 \) detoxification: SOD, PX and CAT (Table 3). In recent years, post-transcriptional mechanisms based on non-coding regulatory micro RNA (miRNA) have been also used to induce tolerance towards multiple stresses, including oxidative and abiotic stresses [189]. Most of these biotechnological approaches have been developed and tested in model species, but some of them were also tested and shown to be effective in immature fruit species (Table 2).

Lee et al. [190] reported the generation of a transgenic cucumber plant that overexpress a gene from cassava (mSOD1) under a fruit specific promoter, which was used to produce an anti-ageing cosmetic from the cucumber fruit. Similarly, the overexpression of the tomato Cu/Zn SOD gene in peppers increased regeneration efficiency while reducing \( \text{H}_2\text{O}_2 \) accumulation [191]. The overexpression of APX was a means of inducing tolerance to oxidative stress in cucumbers [192] and bell peppers [193], while APX genes from eggplants, peppers and sponge gourds were found to be effective in conferring flooding and drought tolerance and in reducing oxidative injury in Arabidopsis and rice [193–195]. The cucumber NITRIC OXIDE ASSOCIATED 1 gene (CsNOA1) is induced by chilling stress, and its constitutive overexpression in cucumbers led to a greater accumulation of soluble sugars and starch, plus an upregulation of Cold-regulatory C-repeat binding factor 3 (CBF3) expression, and a lower CI index in cucumber seedlings [179]. The overexpression of other genes, including pepper LOX and S-ADENOSYL METHIONINE DECARBOXYLASE (SAMDC) genes, reduced lipid peroxidation in membrane cells and reduced ROS, meanwhile inducing higher tolerance to Arabidopsis biotic and abiotic stresses [196–198].

Transcription factors and upstream regulators involved in abiotic and oxidative stress responses have been manipulated in transgenic plants with immature fruits. The Arabidopsis CBF1 gene driven by the inducible promoter RD29A enhanced chilling stress tolerance in cucumbers, concomitantly with an increase in activity of antioxidant enzymes SOD and CAT and a higher free proline content and relative water content in leaves [199]. In the same way, transgenic cucumbers overexpressing the ICE1 gene are more tolerant to chilling stress, showing enhanced expression of cold-responsive genes, a greater concentration of sugar and proline, and a reduced accumulation of MDA [200]. In eggplants, the overexpression of the R2R3 MYB transcription factor gene SmMYB1 resulted in transgenic eggplants that accumulate higher concentrations of anthocyanin in leaves, petals, stamens, and also in the fruit peel and flesh. The GM seedlings exhibited a higher antioxidant activity concomitant with a greater tolerance to freezing and better recovery under rewarming conditions [201]. The ectopic expression of pepper RECEPTOR-LIKE KINASE 1 (CaRLK1) and ankyrin repeat domain zinc finger transcription factor gene CaKRI demonstrated that they function, respectively, as negative and positive regulators of ROS accumulation, resulting in plants that were respectively more sensitive and tolerant to oxidative stress [202,203].

Most of the biotechnological studies detailed above have focused on the vegetative organs of plants, few having been conducted on the fruit. Future studies are needed, therefore, so as to address the effects of these antioxidant transgenes on ROS scavenging systems and chilling tolerance in the immature fruit of marketable species.
Table 3. Transgenic plants of immature fruit species with enhanced tolerance to oxidative stress.

| Species        | Transgene                                      | Effect of Overexpression/Silencing                                    | References |
|---------------|-----------------------------------------------|---------------------------------------------------------------------|------------|
| Cucumber      | *Cassava SOD*                                 | Anti-oxidative cucumber as a functional cosmetic material            | [190]      |
|               | *Cucumber Nitric Oxide Synthase Associated 1 gene CsNOS1* | Chilling tolerance of cucumber seedling                              | [179]      |
|               | *Arabidopsis CBF1*                            | Protection against chilling stress in cucumber leaves                | [199]      |
|               | *Cucumber Mitogen-activated protein kinase gene CsNMAPK* | ROS scavenge and osmotic adjustment in cucumbers under salt stress  | [204]      |
|               | *Cucumber Inducer of CBF expression 1 (ICE1)*  | Enhanced chilling tolerance                                          | [200]      |
|               | *Cucumber APX*                                | Alteration in ascorbate and glutathione redox states and increased sensitivity to ozone-induced oxidative stress in tobacco | [192]      |
| Eggplant      | *Eggplant APX (SmAPX)*                        | Greater resistance to flooding and less oxidative injury in *Arabidopsis* and rice | [193,194] |
|               | *Eggplant SmMYB1 gene encoding a R2R3 MYB transcription factor* | Increasing anthocyanin content in different organs, including fruit peels, and enhanced tolerance to seedling freezing stress | [201]      |
| Bell pepper   | *Pepper LOX gene CaLOX1*                      | Reduction of lipid peroxidation and H₂O₂, associated with high salinity and drought stress tolerance, and defence against pathogens in *Arabidopsis* | [196,197] |
|               | *Pepper PX gene CaPO2*                        | Drought and oxidative stress tolerance in *Arabidopsis*             | [195]      |
|               | *Pepper S-adenosylmethionine decarboxylase gene SAMDC* | Reduced ROS production and drought tolerance in *Arabidopsis*        | [198]      |
|               | *Capsicum annumRECEPTOR-LIKE KINASE 1 (CaRLK1)* | Suppress plant cell death by increasing concentration of superoxide anion | [203]      |
| Tomato        | *Cu/Zn SOD*                                   | Increased resistance to oxidative damage and improved shoot regeneration in pepper | [191]      |
|               | *Pepper ankyrin repeat domain zinc finger transcription factor gene CaKRI* | Reduced levels of free oxygen radicals. Enhanced tolerance to salinity and oxidative stress in tomato | [63]       |
|               | *Pepper ascorbate peroxidase-like 1 gene CaPOA1* | Enhances tolerance to oxidative stress and pathogens in tobacco       | [205]      |
| Sponge gourd  | *Luffa cylindrica ASCORBATE PEROXIDASE (LcAPX)* | Enhanced flood tolerance in transgenic *Arabidopsis* plants by reducing H₂O₂ and MDA | [193]      |

4. Conclusions and Perspectives

Immature vegetable-like fruits are highly perishable commodities. They are in a phase of development in which the metabolism, and therefore respiration rate and transpiration, is very high. Although cold storage decreases their metabolism, these fruits are very susceptible to CI, which is associated with an increase in the production of ROS and the induction of oxidative stress. To avoid such stress, a series of coordinated protective mechanisms are triggered in the fruit, including enzymatic and non-enzymatic antioxidant defence mechanisms, the activation of an alternative electron transport by mitochondrial Alternative Oxidase, and the induction of metabolic pathways that maintain the redox balance.

Several physical, chemical and biotechnological approaches have been developed in immature fruit to reduce post-harvest losses caused by cold damage, many of which are aimed at reducing ROS production and avoiding oxidative stress. In addition to their effectiveness in preventing CI symptoms, new postharvest technologies should be increasingly sustainable, promoting the use of natural products and avoiding chemical compounds that may affect food safety and environmental
preservation. In this sense, plant breeding is offered as a sustainable solution that should be taken more and more into account for the development of new varieties that are increasingly tolerant to oxidative stress and CI. Moreover, the biotechnological strategies should not only be used to improve our understanding of oxidative stress regulation, but also to provide tolerant plant material for future breeding programs.

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References

1. Singh, B.N.; Singh, B.R.; Singh, R.L.; Prakash, D.; Singh, D.P.; Sarma, B.K.; Upadhyay, G.; Singh, H.B. Polyphenolics from various extracts/fractions of red onion (Allium cepa) peel with potent antioxidant and antimutagenic activities. Food Chem. Toxicol. 2009, 47, 1161–1167. [CrossRef] [PubMed]
2. Blanco-Díaz, M.T.; Font, R.; Martínez-Valdivieso, D.; del Río-Celestino, M. Diversity of natural pigments and phytochemical compounds from exocarp and mesocarp of 27 Cucurbita pepo accessions. Sci. Hortic. 2015, 197, 357–365. [CrossRef]
3. Zaro, M.J.; Chaves, A.R.; Vicente, A.R.; Concellón, A. Distribution, stability and fate of phenolic compounds in white and purple eggplants (Solanum melongena L.). Postharvest Biol. Technol. 2014, 92, 70–78. [CrossRef]
4. Mukherjee, P.K.; Nema, N.K.; Maity, N.; Sarkar, B.K. Phytochemical and therapeutic potential of cucumber. Fitoterapia 2013, 84, 227–236. [CrossRef] [PubMed]
5. Nema, N.K.; Maity, N.; Sarkar, B.; Mukherjee, P.K. Cucumis sativus fruit-potential antioxidant, anti-hyaluronidase, and anti-elastase agent. Arch. Dermatol. Res. 2011, 303, 247–252. [CrossRef] [PubMed]
6. Aburjai, T.; Natsh, F.M. Plants used in cosmetics. Phyther. Res. 2003, 17, 987–1000. [CrossRef] [PubMed]
7. Konopacka, D.; Seroczyńska, A.; Korzeniewska, A.; Jesionkowska, K.; Niemirowicz-Szczytt, K.; Płocharski, W. Studies on the usefulness of Cucurbita maxima for the production of ready-to-eat dried vegetable snacks with a high carotenoid content. LWT Food Sci. Technol. 2010, 43, 302–309. [CrossRef]
8. Blanco-Díaz, M.T.; del Río-Celestino, M.; Martínez-Valdivieso, D.; Font, R. Use of visible and near-infrared spectroscopy for predicting antioxidant compounds in summer squash (Cucurbita pepo ssp. pepo). Food Chem. 2014, 164, 301–308. [CrossRef] [PubMed]
9. Grover, J.; Yadav, S. Pharmacological actions and potential uses of Momordica charantia: A review. J. Ethnopharmacol. 2004, 93, 123–132. [CrossRef] [PubMed]
10. Wu, S.-J.; Ng, L.-T. Antioxidant and free radical scavenging activities of wild bitter melon (Momordica charantia Linn. var. abbreviata Ser.) in Taiwan. LWT Food Sci. Technol. 2008, 41, 323–330. [CrossRef]
11. Tan, S.P.; Parks, S.E.; Stathopoulos, C.E.; Roach, P.D. Greenhouse-grown bitter melon: Production and quality characteristics. J. Sci. Food Agric. 2014, 94, 1896–1903. [CrossRef] [PubMed]
12. Kubola, J.; Siriamornpun, S. Phenolic contents and antioxidant activities of bitter gourd (Momordica charantia L.) leaf, stem and fruit fraction extracts in vitro. Food Chem. 2008, 110, 881–890. [CrossRef] [PubMed]
13. Joseph, B.; Jini, D. Antidiabetic effects of Momordica charantia (bitter melon) and its medicinal potency. Asian Pac. J. Trop. Dis. 2013, 3, 93–102. [CrossRef]
14. Zhang, L.-J.; Liaw, C-C.; Hsiao, P-C.; Huang, H.-C.; Lin, Z.-H.; Hsu, F.-L.; Kuo, Y.-H. Cucurbitane-type glycosides from the fruits of Momordica charantia and their hypoglycaemic and cytotoxic activities. J. Funct. Foods 2014, 6, 564–574. [CrossRef]
15. Alam, M.A.; Uddin, R.; Subhan, N.; Rahman, M.M.; Jain, P.; Reza, H.M. Beneficial role of bitter melon supplementation in obesity and related complications in metabolic syndrome. J. Lipids 2015, 2015, 1–18. [CrossRef] [PubMed]
16. Jamuna, S.; Karthika, K.; Paulsami, S. Phytochemical and pharmacological properties of certain medicinally important species of Cucurbitaceae family—A review. J. Res. Biol. 2015, 5, 1835–1849.
38. Wang, C.Y.; Adams, D.O. Chilling-induced ethylene production in cucumbers (Cucumis sativus L.). *Plant Physiol.* 1982, 69, 424–427. [CrossRef] [PubMed]
39. Hyodo, H.; Fukasawa, R. Ethylene production in kiwifruit. *J. Jpn. Soc. Hortic. Sci.* 1985, 54, 209–215. [CrossRef]
40. McCollum, T.G.; McDonald, R.E. Electrolyte leakage, respiration, and ethylene production as indices of chilling injury in grapefruit. *HortScience* 1991, 26, 1191–1192.
41. Lafuente, M.T.; Zacarías, L.; Martínez-Téllez, M.A.; Sanchez-Ballesta, M.T.; Dupille, E. Phenylalanine ammonia-lyase as related to ethylene in the development of chilling symptoms during cold storage of citrus fruits. *J. Agric. Food Chem.* 60(20), 49, 6020–6025. [CrossRef]
42. Chen, J.; Zhao, Y.; Chen, X.; Peng, Y.; Hurr, B.M.; Mao, L. The Role of ethylene and calcium in programmed cell death of cold-stored cucumber fruit. *J. Food Biochem.* 2014, 38, 337–344. [CrossRef]
43. Zacarías, L.; Lafuente, M.T.; Marcos, J.F.; Saladie, M.; Dupille, E. Regulation of ethylene biosynthesis during cold storage of the chilling-sensitive Fortune mandarin fruit. In *Biotechnology of the Plant Hormone Ethylene III*; Vendrell, E., Klee, H., Pech, J.C., Romojaro, F., Eds.; IOS Press: Amsterdam, The Netherlands, 2003; pp. 112–117.
44. Sapitnitskaya, M.; Maul, P.; McCollum, G.T.; Guy, C.L.; Weiss, B.; Samach, A.; Porat, R. Postharvest heat and conditioning treatments activate different molecular responses and reduce chilling injuries in grapefruit. *J. Exp. Bot.* 2006, 57, 2943–2953. [CrossRef] [PubMed]
45. Maul, P.; McCollum, G.; Guy, C.L.; Porat, R. Temperature conditioning alters transcript abundance of genes related to chilling stress in “Marsh” grapefruit flavedo. *Postharvest Biol. Technol.* 2011, 60, 177–185. [CrossRef]
46. Mégias, Z.; Martínez, C.; Manzano, S.; Barrera, A.; Rosales, R.; Valenzuela, J.L.; Garrido, D.; Jamilena, M. Cold-induced ethylene in relation to chilling injury and chilling sensitivity in the non-climacteric fruit of zucchini (*Cucurbita pepo* L.). *LWT Food Sci. Technol.* 2014, 57, 194–199. [CrossRef]
47. Lado, J.; Rodrigo, M.J.; Cronje, P.; Zacarías, L. Involvement of lycopene in the induction of tolerance to chilling injury in grapefruit. *Postharvest Biol. Technol.* 2015, 100, 176–186. [CrossRef]
48. Glenn, G.M.; Chiou, B.S.; Imam, S.H.; Wood, D.F.; Orts, W.J. Role of cuticles in produce quality and preservation. In *Produce Degradation: Pathways and Prevention*; Lamikanra, O., Imam, S., Eds.; CRC Press: Boca Raton, FL, USA, 2005; pp. 20–53.
49. Kubo, Y.; Xue, Y.; Nakatsuka, A.; Mathooko, F.M.; Inaba, A.; Nakamura, R. Expression of a water stress-induced polygalacturonase gene in harvested cucumber fruit. *J. Jpn. Soc. Hortic. Sci.* 2000, 69, 273–279. [CrossRef]
50. Lurie, S.; Shapiro, B.; Ben-Yehoshua, S. Effects of water stress and degree of ripeness on rate of senescence of harvested bell pepper fruit. *J. Am. Soc. Hortic. Sci.* 1986, 111, 880–885.
51. Massolo, J.F.; Concillón, A.; Chaves, A.R.; Vicente, A.R. 1-Methylcyclopropene (1-MCP) delays senescence, maintains quality and reduces browning of non-climacteric eggplant (*Solanum melongena* L.) fruit. *Postharvest Biol. Technol.* 2011, 59, 10–15. [CrossRef]
52. Rao, C.G. Engineering for Storage of Fruits and Vegetables: Cold Storage, Controlled Atmosphere Storage, Modified Atmosphere Storage; Rao, C.G., Ed.; Academic Press: Cambridge, MA, USA, 2015.
53. Sevilla, L.; Sanchez-Ballesta, M.T.; Romojaro, F.; Flores, F.B. Physiological, hormonal and molecular mechanisms regulating chilling injury in horticultural species. Postharvest technologies applied to reduce its impact. *J. Sci. Food Agric.* 2009, 89, 555–573. [CrossRef]
54. Kratsch, H.A.; Wise, R.R. The ultrastructure of chilling stress. *Plant Cell Environ.* 2000, 23, 337–350. [CrossRef]
55. Fernández-Trujillo, J.P.; Martínez, J.A. Ultrastructure of the onset of chilling injury in cucumber fruit. *J. Appl. Bot. Food Qual.* 2006, 80, 100–110.
56. Sayyari, M.; Babalar, M.; Kalantari, S.; Martínez-Romero, D.; Guillen, F.; Serrano, M.; Valero, D. Vapour treatments with methyl salicylate or methyl jasmonate alleviated chilling injury and enhanced antioxidant potential during postharvest storage of pomegranates. *Food Chem.* 2011, 124, 964–970. [CrossRef]
57. Zhao, Y.; Chen, J.; Tao, X.; Zheng, X.; Mao, L. The possible role of BAX and BI-1 genes in chilling-induced cell death in cucumber fruit. *Acta Physiol. Plant.* 2014, 36, 1345–1351. [CrossRef]
58. Tatsumi, Y.; Maeda, K.; Murata, T. Morphological changes in cucumber fruit surfaces associated with chilling injury. *J. Jpn. Soc. Hortic. Sci.* 1987, 56, 187–192. [CrossRef]
59. Hakim, A.; Purvis, A.C.; Mullinix, B.G. Differences in chilling sensitivity of cucumber varieties depends on storage temperature and the physiological dysfunction evaluated. *Postharvest Biol. Technol.* 1999, 17, 97–104. [CrossRef]
60. Carvajal, F.; Palma, F.; Jamilena, M.; Garrido, D. Cell wall metabolism and chilling injury during postharvest cold storage in zucchini fruit. *Postharvest Biol. Technol.* 2015, 108, 68–77. [CrossRef]
61. Balandrán-Quintana, R.R.; Mendoza-Wilson, A.M.; Gardea-Béjar, A.A.; Vargas-Arispuro, L.; Martínez-Téllez, M. A Irreversibility of chilling injury in zucchini squash (*Cucurbita pepo* L.) could be a programmed event long before the visible symptoms are evident. *Biochem. Biophys. Res. Commun.* 2003, 307, 553–557. [CrossRef]
62. Lim, C.S.; Kang, S.M.; Cho, J.L.; Gross, K.C. Antioxidizing enzyme activities in chilling-sensitive and chilling-tolerant pepper fruit as affected by stage of ripeness and storage temperature. *J. Am. Soc. Hortic Sci.* 2009, 134, 156–163.
63. Lim, C.S.; Kang, S.M.; Cho, J.L.; Gross, K.C.; Woolf, A.B. Bell Pepper (*Capsicum annuum* L.) fruits are susceptible to chilling injury at the breaker stage of ripeness. *HortScience* 2007, 42, 1659–1664.
64. Boonyaritthongchai, P.; Srilaong, V.; Wongs-Aree, C.; Techavuthiporn, C. Effect of methyl jasmonate on polyphenol oxidase. *J. Agric. Food Chem.* 2000, 48, 695–700. [CrossRef]
65. Pérez-Gilbert, M.; García-Carmona, F. Characterization of catecholase and cresolase activities of eggplant polyphenol oxidase. *J. Agric. Food Chem.* 2007, 55, 385–386. [CrossRef]
66. Concellón, A.; Añón, M.C.; Chaves, A.R. Effect of low temperature storage on physical and physiological characteristics of eggplant fruit (*Solanum melongena* L.). *LWT Food Sci. Technol.* 2007, 40, 389–396. [CrossRef]
67. Tomás-Barberán, F.A.; Espín, J.C. Phenolic compounds and related enzymes as determinants of quality in fruits and vegetables. *J. Sci. Food Agric.* 2001, 81, 853–876. [CrossRef]
68. Concellón, A.; Añón, M.C.; Chaves, A.R. Characterization and changes in polyphenol oxidase from eggplant fruit (*Solanum melongena* L.) during storage at low temperature. *Food Chem.* 2004, 88, 17–24. [CrossRef]
69. Perkins-VEazie, P.M.; Collins, J.K. Cultivar, packaging, and storage temperature differences in postharvest shelf life of okra. *HortTechnology* 1992, 2, 350–352.
70. Cantwell, M.; Suslow, T. *Perishables Handling #107;* Department of Plant Sciences, University of California, Davis: Davis, CA, USA, 2001.
71. Boonyaritthongchai, P.; Srlaong, V.; Wongs-Aree, C.; Techavuthiporn, C. Effect of methyl jasmonate on reducing chilling injury symptom and maintaining postharvest quality of okra (*Hibiscus esculentum* L.). *Acta Hortic.* 2013, 1012, 1119–1124. [CrossRef]
72. Zong, R.; Cantwell, M.; Morris, L.L. Postharvest handling of Asian specialty vegetables under study. * Calif Agric.* 1993, 47, 27–29.
73. Han, C.; Zuo, J.H.; Wang, Q.; Dong, H.Z.; Gao, L.P. Salicylic acid alleviates postharvest chilling injury of sponge gourd (*Luffa cylindrica*). *J. Integr. Agric.* 2017, 16, 735–741. [CrossRef]
74. Purvis, A.C. Diphenylamine reduces chilling injury of green bell pepper fruit. *Postharvest Biol. Technol.* 2002, 25, 41–48. [CrossRef]
75. Babarinde, G.O.; Fabunmi, O.A. Effects of packaging materials and storage temperature on quality of fresh okra (*Abelmoschus esculentus*) fruit. *Agric. Trop. Subtrop.* 2009, 42, 151–156.
76. Wang, L.; Li, Q.; Cao, J.; Cai, T.; Jiang, W. Keeping quality of fresh-cut bitter gourd (*Momordica charantia* L.) at low temperature of storage. *J. Food Process. Preserv.* 2007, 31, 571–582. [CrossRef]
77. Del Río, L.A.; López-Huertas, E. ROS generation in peroxisomes and its role in cell signaling. *Plant Cell Physiol.* 2016, 57, 1364–1376. [CrossRef] [PubMed]
78. Mittler, R.; Vanderauwera, S.; Giolery, M.; van Breusegem, F. Reactive oxygen gene network of plants. *Trends Plant Sci.* 2004, 9, 490–498. [CrossRef] [PubMed]
79. Parkin, K.L.; Kuo, S.J. Chilling-induced lipid degradation in cucumber (*Cucumis sativa* L. cv Hybrid C) fruit. *Plant Physiol.* 1989, 90, 1049–1056. [CrossRef] [PubMed]
80. Choudhury, F.K.; Rivero, R.M.; Blumwald, E.; Mittler, R. Reactive oxygen species, abiotic stress and stress combination. *Plant J.* 2016, 90, 856–867. [CrossRef] [PubMed]
81. Asada, K. Production and scavenging of reactive oxygen species in chloroplasts and their functions. *Plant Physiol.* 2006, 141, 391–396. [CrossRef] [PubMed]
82. Foyer, C.H.; Noctor, G. Ascorbate and glutathione: The heart of the redox hub. *Plant Physiol.* 2011, 155, 2–18. [CrossRef] [PubMed]
83. Vanlerberghe, G.C.G. Alternative oxidase: A mitochondrial respiratory pathway to maintain metabolic homeostasis during abiotic and biotic stress in plants. Int. J. Mol. Sci. 2013, 14, 6805–6847. [CrossRef] [PubMed]
84. Bouché, N.; Fromm, H. GABA in plants: Just a metabolite? Trends Plant Sci. 2004, 9, 110–115. [CrossRef] [PubMed]
85. Bouché, N.; Fait, A.; Bouchez, D.; Moller, S.G.; Fromm, H. Mitochondrial succinic-semialdehyde dehydrogenase of the γ-aminobutyrate shunt is required to restrict levels of reactive oxygen intermediates in plants. Proc. Natl. Acad. Sci. USA 2003, 100, 6843–6848. [CrossRef] [PubMed]
86. Hodges, D.M.; Lester, G.E.; Munro, K.D.; Toivonen, P.M.A. Oxidative stress: Importance for postharvest quality. HortScience 2004, 39, 924–929.
87. Toivonen, P.M.A. Postharvest storage procedures and oxidative stress. HortScience 2004, 39, 938–942.
88. Raja, V.; Majeed, U.; Kang, H.; Andrabi, K.I.; John, R. Abiotic stress: Interplay between ROS, hormones and MAPKs. Environ. Exp. Bot. 2017, 137, 142–157. [CrossRef]
89. Chen, B.; Yang, H. 6-Benzylaminopurine alleviates chilling injury of postharvest cucumber fruit through modulating antioxidative system and energy status. J. Sci. Food. Agric. 2013, 93, 1915–1921. [CrossRef] [PubMed]
90. Hariyadi, P.; Parkin, K.L. Chilling-induced oxidative stress in cucumber fruits. Postharvest Biol. Technol. 1991, 1, 33–45. [CrossRef]
91. Carvajal, F.; Palma, F.; Jamilena, M.; Garrido, D. Preconditioning treatment induces chilling tolerance in zucchini fruit improving different physiological mechanisms against cold injury. Ann. Appl. Biol. 2015, 166, 340–354. [CrossRef]
92. Palma, F.; Carvajal, F.; Jamilena, M.; Garrido, D. Putrescine treatment increases the antioxidative response and carbohydrate content in zucchini fruit stored at low temperature. Postharvest Biol. Technol. 2016, 118, 68–70. [CrossRef]
93. Wang, C.Y. Effect of temperature preconditioning on catalase, peroxidase, and superoxide dismutase in chilled zucchini squash. Postharvest Biol. Technol. 1995, 5, 67–76. [CrossRef]
94. Zheng, Y.; Fung, R.W.M.; Wang, S.Y.; Wang, C.Y. Transcript levels of antioxidative genes and oxygen radical scavenging enzyme activities in chilled zucchini squash in response to superatmospheric oxygen. Postharvest Biol. Technol. 2008, 47, 151–158. [CrossRef]
95. Yang, H.; Wu, F.; Cheng, J. Reduced chilling injury in cucumber by nitric oxide and the antioxidant response. Food Chem. 2011, 127, 1237–1242. [CrossRef] [PubMed]
96. Qian, C.; He, Z.; Zhao, Y.; Mi, H.; Chen, X.; Mao, L. Maturity-dependent chilling tolerance regulated by the antioxidative capacity in postharvest cucumber (Cucumis sativus L.) fruits. J. Sci. Food. Agric. 2013, 93, 626–633. [CrossRef] [PubMed]
97. Qian, C.; Mi, H.; Zhao, Y.; He, Z.; Mao, L. Effect of maturity stage on the gene expression of antioxidative enzymes in cucumber (Cucumis sativus L.) fruits under chilling stress. J. Integr. Agric. 2013, 12, 1495–1500. [CrossRef]
98. Martinez-Téllez, M.; Ramos-Clamont, M.; Gardea, A.; Vargas-Arispuro, I. Effect of infiltrated polyamines on polygalacturonase activity and chilling injury responses in zucchini squash (Cucurbita pepo L.). Biochem. Biophys. Res. Commun. 2002, 295, 98–101. [CrossRef]
99. Palma, F.; Carvajal, F.; Lluch, C.; Jamilena, M.; Garrido, D. Changes in carbohydrate content in zucchini fruit (Cucurbita pepo L.) under low temperature stress. Plant Sci. 2014, 217, 78–86. [CrossRef] [PubMed]
100. Palma, F.; Carvajal, F.; Jamilena, M.; Garrido, D. Contribution of polyamines and other related metabolites to the maintenance of zucchini fruit quality during cold storage. Plant Physiol. Biochem. 2014, 82, 161–171. [CrossRef] [PubMed]
101. Wang, C.Y.; Buta, J.G. Methyl jasmonate reduces chilling injury in Cucurbita pepo through its regulation of abscisic acid and polyamine levels. Environ. Exp. Bot. 1994, 34, 427–432. [CrossRef]
102. Gualanduzzi, S.; Baraldi, E.; Braschi, I.; Carnevali, F.; Gessa, C.E.; De Santis, A. Respiration, hydrogen peroxide levels and antioxidative enzyme activities during cold storage of zucchini squash fruit. Postharvest Biol. Technol. 2009, 52, 16–23. [CrossRef]
103. Couée, I.; Sulmon, C.; Gouesbet, G.; El Amrani, A. Involvement of soluble sugars in reactive oxygen species balance and responses to oxidative stress in plants. J. Exp. Bot. 2006, 57, 449–459. [CrossRef] [PubMed]
104. Nishizawa, A.; Yabuta, Y.; Shigeoka, S. Galactinol and raffinose constitute a novel function to protect plants from oxidative damage. *Plant Physiol.* 2008, 147, 1251–1263. [CrossRef] [PubMed]
105. Sánchez-Bel, P.; Egea, I.; Sánchez-Ballesta, M.T.; Martínez-Madrid, C.; Fernández-Garcia, N.; Romojaro, F.; Olmos, E.; Estrella, E.; Bolarin, M.C.; Flores, F.B. Understanding the mechanisms of chilling injury in bell pepper fruits using the proteomic approach. *J. Proteom.* 2012, 75, 5463–5478. [CrossRef]
106. Fung, R.W.M.; Wang, C.Y.; Smith, D.L.; Gross, K.C.; Tian, M. MeSA and MeJA increase steady-state transcript levels of alternative oxidase and resistance against chilling injury in sweet peppers (*Capsicum annuum* L.). *Plant Sci.* 2004, 166, 711–719. [CrossRef]
107. Kaneyuki, T.; Noda, Y.; Traber, M.G.; Mori, A.; Packer, L. Superoxide anion and hydroxyl radical scavenging activities of vegetable extracts measured using electron spin resonance. *IIJMB Life* 1999, 47, 979–989. [CrossRef]
108. Prohens, J.; Rodríguez-Burruezo, A.; Raigón, M.D.; Nuez, F. Total phenolic concentration and browning susceptibility in a collection of different varietal types and hybrids of eggplant: Implications for breeding for higher nutritional quality and reduced browning. *J. Am. Soc. Hortic. Sci.* 2007, 132, 638–646.
109. Fan, L.; Shi, J.; Zuo, J.; Gao, L.; Lv, J.; Wang, Q. Methyl jasmonate delays postharvest ripening and senescence in the non-climacteric eggplant (*Solanum melongena* L.) fruit. *Postharvest Biol. Technol.* 2016, 120, 76–83. [CrossRef]
110. Mao, L.; Pang, H.; Wang, G.; Zhu, C. Phospholipase D and lipoxygenase activity of cucumber fruit in response to chilling stress. *Postharvest Biol. Technol.* 2007, 44, 42–47. [CrossRef]
111. Mao, L.-C.; Wang, G.-Z.; Zhu, C.-G.; Pang, H.-Q. Involvement of phospholipase D and lipoxygenase in response to chilling stress in postharvest cucumber fruits. *Plant Sci.* 2007, 172, 400–405. [CrossRef]
112. González-Aguilar, G.A.; Fortiz, J.; Cruz, R.; Baez, R.; Wang, C.Y. Methyl jasmonate reduces chilling injury and maintains postharvest quality of mango fruit. *J. Agric. Food Chem.* 2000, 48, 515–519. [CrossRef] [PubMed]
113. Lurie, S. Postharvest heat treatments. *Postharvest Biol. Technol.* 1998, 14, 257–269. [CrossRef]
114. Bar-Yosef, A.; Alkalai-Tuvia, S.; Perzelan, Y.; Aharon, Z.; Liic’, Z.; Lurie, S.; Fallik, E. Effect of shrink packaging in combination with rinsing and brushing treatment on chilling injury and decay of sweet pepper during storage. *Adv. Hortic. Sci.* 2009, 23, 225–230.
115. Sato, Y.; Murakami, T.; Funatsuki, H.; Matsuba, S.; Saruyama, H.; Tanida, M. Heat shock-mediated APX gene expression and protection against chilling injury in rice seedlings. *J. Exp. Bot.* 2001, 52, 145–151. [CrossRef] [PubMed]
116. Zhang, Y.; Zhang, M.; Yang, H. Postharvest chitosan-g-salicylic acid application alleviates chilling injury and preserves cucumber fruit quality during cold storage. *Food Chem.* 2015, 174, 558–563. [CrossRef] [PubMed]
117. McCollum, T.G.; Doostdar, H.; Mayer, R.T.; McDonald, R.E. Immersion of cucumber fruit in heated water enhances fruit quality during refrigerated storage of eggplants. *J. Agric. Food Chem.* 2001, 49, 4700–4705. [CrossRef] [PubMed]
118. Ilić, Z.S.; Trajković, R.; Pavlović, R.; Alkalai-Tuvia, S.; Perzelan, Y.; Fallik, E. Effect of heat treatment and individual shrink packaging on quality and nutritional value of bell pepper stored at suboptimal temperature. *Int. J. Food Sci. Technol.* 2012, 47, 83–90. [CrossRef]
119. Rodríguez, S. del C.; López, B.; Chavesz, A.R. Effect of different treatments on the evolution of polyamines during refrigerated storage of eggplants. *J. Agric. Food Chem.* 2001, 49, 4700–4705. [CrossRef]
120. Li, N.; Parsons, B.L.; Liu, D.; Mattoo, A.K. Accumulation of wound-inducible ACC synthase transcript in tomato fruit is inhibited by salicylic acid and polyamines. *Plant Mol. Biol.* 1992, 18, 477–487. [CrossRef] [PubMed]
121. De Bruin, W.; Rossouw, W.; Korsten, L. Comparison of safe alternative dipping treatment to maintain quality of zucchini. *J. Food Qual.* 2016, 39, 109–115. [CrossRef]
122. Yi Wang, C. Combined treatment of heat shock and low temperature conditioning reduces chilling injury in zucchini squash. *Postharvest Biol. Technol.* 1994, 4, 65–73. [CrossRef]
123. Wang, H.; Qian, Z.; Ma, S.; Zhou, Y.; Patrick, J.W.; Duan, X.; Jiang, Y.; Qu, H. Energy status of ripening and postharvest senescent fruit of litchi (*Litchi chinensis* Sonn.). *BMC Plant Biol.* 2013, 13, 55. [CrossRef] [PubMed]
124. Wang, B.; Zhu, S. Pre-storage cold acclimation maintained quality of cold-stored cucumber through differentially and orderly activating ROS scavengers. *Postharvest Biol. Technol.* 2017, 129, 1–8. [CrossRef]
125. Kader, A.A. Biochemical and physiological basis for effects of controlled and modified atmospheres on fruits and vegetables. *Food Technol.* 1986, 40, 99–100.
126. Kader, A.A. Atmosphere modification. In Postharvest Physiology and Pathology of Vegetables; Bartz, J.A., Brecht, J.K., Weichmann, J., Eds.; Marcel Dekker: New York, NY, USA, 2003; p. 733.
127. Kader, A.A.; Saltveit, M.E. Respiración y gas exchange. In Postharvest Physiology and Pathology of Vegetables; Bartz, J.A., Brecht, J.K., Weichmann, J., Eds.; Marcel Dekker: New York, NY, USA, 2003; pp. 7–29. ISBN 0-8247-0687-0.
128. Kader, A.A.; Zagory, D.; Kerbel, E.L.; Wang, C.Y. Modified atmosphere packaging of fruits and vegetables. Crit. Rev. Food Sci. Nutr. 1989, 28, 1–30. [CrossRef] [PubMed]
129. Scully, A.D.; Horsham, M.A. Active packaging for fruits and vegetables. In Intelligent and Active Packaging for Fruits and Vegetables; Wilson, C.L., Ed.; CRC Press: Boca Raton, FL, USA, 2008; pp. 7–29, ISBN-13: 978-0849391668.
130. Yahia, E.M. Introduction. In Modified and Controlled Atmospheres for the Storage, Transportation and Packaging of Horticultural Commodities; Yahia, E.M., Ed.; CRC Press, Taylor & Francis Group: Boca Raton, FL, USA, 2009; pp. 1–16, ISBN-13: 978-1420069570.
131. Mangaraj, S.G.T. Modified atmosphere packaging—An ideal food preservation technique. J. Food Sci. Technol. 2009, 46, 399–410.
132. Kanellis, A.K.; Tonutti, P.P. Biochemical and molecular aspects of modified and controlled atmospheres. In Modified and Controlled Atmospheres for the Storage, Transportation and Packaging of Horticultural Commodities; Yahia, E.M., Ed.; CRC Press, Taylor & Francis Group: Boca Raton, FL, USA, 2009; pp. 553–567, ISBN-13: 978-14200695702009.
133. Mishra, B.; Khatkar, B.S.; Garg, M.K.; Wilson, L.A. Permeability of edible coatings. J. Food Sci. Technol. 2010, 47, 109–113. [CrossRef] [PubMed]
134. Fahmy, K.; Nakano, K. Optimal design of modified atmosphere packaging for alleviating chilling injury in cucumber fruit. Environ. Control Biol. 2014, 52, 233–240. [CrossRef]
135. Wang, C.Y.; Qi, L. Modified atmosphere packaging alleviates chilling injury in cucumbers. Postharvest Biol. Technol. 1997, 10, 195–200. [CrossRef]
136. Serrano, M.; Pretel, M.T.; Martinez-Madrid, M.C.; Romojoaro, F.; Riquelme, F. CO₂ treatment of zucchini squash reduces chilling-induced physiological changes. J. Agric. Food Chem. 1998, 46, 2465–2468. [CrossRef]
137. Wang, C.Y.; Ji, Z.L. Effect of low-oxygen storage on chilling injury and polyamines in zucchini squash. Sci. Hortic. 1989, 39, 1–7. [CrossRef]
138. Zagory, D.; Kader, A.A. Modified atmosphere packaging of fresh produce. Food Technol. 1988, 42, 70–77.
139. Serrano, M.; Martinez-Madrid, M.C.; Pretel, M.T.; Riquelme, F.; Romojoaro, F. Modified Atmosphere Packaging minimizes increases in putrescine and abscisic acid levels caused by chilling injury in pepper fruit. J. Agric. Food Chem. 1997, 45, 1668–1672. [CrossRef]
140. Sahoo, N.R.; Bal, L.M.; Pal, U.S.; Sahoo, D. A Comparative study on the effect of packaging material and storage environment on shelf life of fresh bell-pepper. J. Food Meas Charact. 2014, 8, 164–170. [CrossRef]
141. Rao, D.S.; Rao, K.G.; Krishnamurthy, S. Extension of shelf life of cucumber by modified atmosphere packaging (MAP) and shrink wrapping. Indian Food Pack. 2000, 54, 65–71.
142. Dhall, R.K.; Sharma, S.R.; Mahajan, B.V.C. Effect of shrink wrap packaging for maintaining quality of cucumber during storage. J. Food Sci. Technol. 2012, 49, 495–499. [CrossRef] [PubMed]
143. Mahajan, B.V.C.; Dhillon, W.S.; Sidhu, M.K.; Jindal, S.K.; Dhaliwal, M.S.; Singh, S. Effect of packaging and storage environments on quality and shelf-life of bell pepper (Capsicum annuum). Indian J. Agric. Sci. 2016, 86, 738–742.
144. Shi, J.Y.; Gao, L.P.; Zuo, J.H.; Wang, Q.; Wang, Q.; Fang, L. Effect of exogenous ceramide treatment on chilling tolerance in green bell pepper fruits. Mod. Food Sci. Technol. 2016, 32, 164–170.
145. Ghaouth, A.; Arul, J.; Ponnampalam, R.; Boulet, M. Use of chitosan coating to reduce water loss and maintain quality of cucumber and bell pepper fruits. J. Food Process. Preserv. 1991, 15, 359–368. [CrossRef]
146. Mohammadi, A.; Hashemi, M.; Hosseini, S.M. Postharvest treatment of nanochitosan-based coating loaded with Zataria multiflora essential oil improves antioxidant activity and extends shelf-life of cucumber. Innov. Food Sci. Emerg. Technol. 2016, 33, 580–588. [CrossRef]
147. Abbasi, N.A.; Ashraf, S.; Ali, I.; Butt, S.J. Enhancing storage life of bell pepper by UV-C irradiation and edible coatings. Pak. J. Agric. Sci. 2015, 52, 403–411.
148. Gholamipour, F.K.; Kamari, S.; Ghasemnezhad, M.; Ghazvini, R.F. Effect of chitosan coating on weight loss and postharvest quality of green pepper (Capsicum annuum L. Fruits. Acta Hortic. 2010, 877, 821–826. [CrossRef]

149. Beaulieu, J.C.; Park, H.S.; Mims, A.G.B.; Kuk, M.S. Extension of green bell pepper shelf life using oilseed-derived lipid films from soapstock. Ind. Crops Prod. 2009, 30, 271–275. [CrossRef]

150. Lerdthanangkul, S.; Krochta, J.M. Edible coating effects on postharvest quality of green bell peppers. J. Food Sci. 1996, 61, 176–179. [CrossRef]

151. de Souza, P.A.; Arroucha, E.M.M.; de Souza, A.E.; da Costa, A.R.; de Ferreira, G.S.; Bezerra Neto, F. Conservação pós-colheita de berinjela com revestimentos de fúcsia de mandioca ou filme de PVC. Hortic. Bras. 2009, 27, 235–239. [CrossRef]

152. Han, C.; Zuo, J.; Wang, Q.; Xu, L.; Zhai, B.; Wang, Z.; Dong, H.; Gao, L. Effects of chitosan coating on postharvest quality and shelf life of sponge gourd (Luffa cylindrica) during storage. Sci. Hortic. 2014, 166, 1–8. [CrossRef]

153. Abeles, F.B.; Morgan, P.W.; Saltveit, M.E. The role of ethylene in agriculture. In Ethylene in Plant Biology, 2nd ed.; Abeles, F.B., Morgan, P.W., Saltveit, M.E., Eds.; Academic Press: Cambridge, MA, USA, 1992; pp. 264–296. [CrossRef]

154. Barry, C.S.; Giovannoni, J.J. Ethylene and fruit ripening. J. Plant Growth Regul. 2007, 26, 143–159. [CrossRef]

155. Gao, H.; Kang, L.; Liu, Q.; Cheng, N.; Wang, B.; Cao, W. Effect of 24-epibrassinolide treatment on the cold acclimation process of Arabidopsis thaliana (L.) Heynh. Plant Mol. Biol. 2010, 877, 821–826. [CrossRef]

156. Blankenship, S.M.; Dole, J.M. 1-Methylcyclopropene: A review. Postharvest Biol. Technol. 2003, 28, 1–25. [CrossRef]

157. Watkins, C.B. The use of 1-methylcyclopropene (1-MCP) on fruits and vegetables. Biotechnol. Adv. 2006, 24, 389–409. [CrossRef] [PubMed]

158. Cao, S.; Yang, Z.; Zheng, Y. Effect of 1-methylcyclopene on senescence and quality maintenance of green bell pepper (Capsicum annuum L.) Fruits. Acta Hortic. 2010, 877, 821–826. [CrossRef]

159. Smith, A.W.J.; Poulston, S.; Rowsell, L.; Terry, L.A.; Anderson, J.A. A New Palladium-based ethylene scavenger to control ethylene-induced ripening of climacteric fruit. Platin. Met. Rev. 2009, 53, 112–122. [CrossRef]

160. Wang, Q.; Ding, T.; Gao, L.; Pang, J.; Yang, N. Effect of brassinolide on chilling injury of green bell pepper in storage. Sci. Hortic. 2012, 144, 195–200. [CrossRef]

161. Gao, H.; Kang, L.; Liu, Q.; Cheng, N.; Wang, B.; Cao, W. Effect of 24-epibrassinolide treatment on the metabolism of eggplant fruits in relation to development of pulp browning under chilling stress. J. Food Sci. Technol. 2015, 52, 3394–3401. [CrossRef] [PubMed]

162. Swamy, P.M.; Smith, B.N. Role of abscisic acid in plant stress tolerance. Curr. Sci. 1999, 76, 1220–1227.

163. Zhu, J.-K. Salt and drought stress signal transduction in plants. Annu. Rev. Plant Biol. 2002, 53, 247–273. [CrossRef] [PubMed]

164. Cheng, W.H.; Endo, A.; Zhou, L.; Penney, J.; Chen, H.-C.; Arroyo, A.; Leon, P.; Nambara, E.; Asami, T.; Seo, M. A unique short-chain dehydrogenase/reductase in Arabidopsis glucose signaling and abscisic acid biosynthesis and functions. Plant Cell 2002, 14, 2723–2743. [CrossRef] [PubMed]

165. Saito, S.; Hirai, N.; Matsumoto, C.; Ohigashi, H.; Ohta, D.; Sakata, K.; Mizutani, M. Arabidopsis CYP707A1 encode (+)-abscisic acid 8′-hydroxylase, a key enzyme in the oxidative catabolism of abscisic acid. Plant Physiol. 2004, 134, 1439–1449. [CrossRef] [PubMed]

166. Jiang, M.; Zhang, J. Cross-talk between calcium and reactive oxygen species originated from NADPH oxidase in abscisic acid-induced antioxidant defence in leaves of maize seedlings. Plant Cell Environ. 2003, 26, 929–939. [CrossRef] [PubMed]

167. Kurkela, S.; Franck, M. Cloning and characterization of a cold-and ABA-inducible Arabidopsis gene. Plant Mol. Biol. 1990, 15, 137–144. [CrossRef] [PubMed]

168. Lang, V.; Palva, E.T. The expression of a rab-related gene, rab18, is induced by abscisic acid during the cold acclimation process of Arabidopsis thaliana (L.) Heynh. Plant Mol. Biol. 1992, 20, 951–962. [CrossRef] [PubMed]
169. Mantyla, E.; Lang, V.; Palva, E.T. Role of abscisic acid in drought-induced freezing tolerance, cold acclimation, and accumulation of LT178 and RAB18 proteins in *Arabidopsis thaliana*. *Plant Physiol.* 1995, 107, 141–148. [CrossRef] [PubMed]

170. Asghari, M.; Aghdam, M.S. Impact of salicylic acid on post-harvest physiology of horticultural crops. *Trends Food Sci. Technol.* 2010, 21, 502–509. [CrossRef]

171. Hu, Q.; Wang, T.; Zhou, M.; Xue, J.; Luo, Y. In vitro antioxidant-activity evaluation of gallic-acid-grafted chitosan conjugate synthesized by free-radical-induced grafting method. *J. Agric. Food Chem.* 2016, 64, 5893–5900. [CrossRef] [PubMed]

172. Wasternack, C. Jasmonates: An update on biosynthesis, signal transduction and action in plant stress response, growth and development. *Ann. Bot.* 2007, 100, 681–697. [CrossRef] [PubMed]

173. Ziosi, V.; Bonghi, C.; Bregoli, A.M.; Trainotti, L.; Biondi, S.; Sutthiwat, S.; Kondo, S.; Costa, G.; Torrigiani, P. Jasmonate-induced transcriptional changes suggest a negative interference with the ripening syndrome in peach fruit. *J. Exp. Bot.* 2008, 59, 563–573. [CrossRef] [PubMed]

174. Hu, Y.; Jiang, L.; Wang, F.; Yu, D. Jasmonate regulates the inducer of CBF expression-C-repeat binding factor/DRE binding factor1 cascade and freezing tolerance in *Arabidopsis*. *Plant Cell.* 2013, 25, 2907–2924. [CrossRef] [PubMed]

175. Du, H.; Liu, H.; Xiong, L. Endogenous auxin and jasmonic acid levels are differentially modulated by abiotic stresses in rice. *Front. Plant Sci.* 2013, 4, 397. [CrossRef] [PubMed]

176. Carrera, R.; Zapata, S.; García, A.; Aguado, E.; Rebollos, M.M.; Manzano, S.; Jamilena, M.; Valenzuela, J.L. Efectos de los tratamientos de Metil Jasmonato y Ácido Salicilico en la calidad poscosecha y daños por frío de frutos de berenjena. *Actas Port. Hortic.* 2016, 28, 239–244.

177. Zapata, S.; Carrera, R.; Manzano, S.; García, A.; Aguado, E.; Garrido, D.; Rebollos, M.M.; Manzano, S.; Valenzuela, J.L.; Valenzuela, J.L. Efectos de los tratamientos de Metil Jasmonato y Ácido Salicilico en la reducción del daño por frío en calabacin. *Actas Port. Hortic.* 2016, 28, 232–238.

178. Purvis, A.C. Role of the alternative oxidase in limiting superoxide production by plant mitochondria. *Physiol. Plant.* 1997, 100, 165–170. [CrossRef]

179. Liu, Y.; Yang, X.; Zhu, S.; Wang, Y. Postharvest application of MeJA and NO reduced chilling injury in cucumber (*Cucumis sativus*) through inhibition of H$_2$O$_2$ accumulation. *Postharvest Biol. Technol.* 2016, 119, 77–83. [CrossRef]

180. Palma, F.; Carvajal, F.; Ramos, J.M.; Jamilena, M.; Garrido, D. Effect of putrescine application on maintenance of zucchini fruit quality during cold storage: Contribution of GABA shunt and other related nitrogen metabolites. *Postharvest Biol. Technol.* 2015, 99, 131–140. [CrossRef]

181. Zhang, W.; Jiang, B.; Li, W.; Song, H.; Yu, Y.; Chen, J. Polyamines enhance chilling tolerance of cucumber (*Cucumis sativus L.*) through modulating antioxidative system. *Sci. Hortic.* 2009, 122, 200–208. [CrossRef]

182. Wink, D.A.; Mitchell, J.B. Chemical biology of nitric oxide: Insights into regulatory, cytotoxic, and cytoprotective mechanisms of nitric oxide. *Free Radic. Biol. Med.* 1998, 25, 434–456. [CrossRef]

183. Soegiarto, L.; Wills, R.B.H. Short term fumigation with nitric oxide gas in air to extend the postharvest life of broccoli, green bean, and bok choy. *Hortotechnology* 2004, 14, 538–540.

184. Izumi, H.; Watada, A.E. Calcium treatment to maintain quality of zucchini squash slices. *J. Food Sci.* 1995, 60, 789–793. [CrossRef]

185. Massolo, J.F.; Lemoine, M.L.; Chaves, A.R.; Concellón, A.; Vicente, A.R. Benzyl-aminopurine (BAP) treatments delay cell wall degradation and softening, improving quality maintenance of refrigerated summer squash. *Postharvest Biol. Technol.* 2014, 93, 122–129. [CrossRef]

186. Wang, Q.; Ding, T.; Zuo, J.; Gao, L.; Fan, L. Amelioration of postharvest chilling injury in sweet pepper by glycine betaine. *Postharvest Biol. Technol.* 2016, 112, 114–120. [CrossRef]

187. Liu, L.; Wei, Y.; Shi, F.; Liu, C.; Liu, X.; Ji, S. Intermittent warming improves postharvest quality of bell peppers and reduces chilling injury. *Postharvest Biol. Technol.* 2015, 18, 18–25. [CrossRef]

188. Wang, C.Y. Effect of abscisic acid on chilling injury of zucchini squash. *J. Plant Growth Regul.* 1991, 10, 101–105. [CrossRef]

189. Banerjee, A.; Roychoudhury, A.; Krishnamoorthi, S. Emerging techniques to decipher microRNAs (miRNAs) and their regulatory role in conferring abiotic stress tolerance of plants. *Plant Biotechnol. Rep.* 2016, 10, 185–205. [CrossRef]
190. Lee, H.-S.; Kwon, E.-J.; Kwon, S.-Y.; Jeong, Y.-J.; Lee, E.-M.; Jo, M.-H.; Kim, H.-S.; Woo, I.-S.; Shinmyo, A.; Yoshida, K. Transgenic cucumber fruits that produce elevated level of an anti-aging superoxide dismutase. *Mol. Breed.* 2003, 11, 213–220. [CrossRef]

191. Chatzidimitriadou, K.; Nianiou-Obeidat, I.; Madesis, P.; Perl-Treves, R.; Tsafaritis, A. Expression of SOD transgene in pepper confer stress tolerance and improve shoot regeneration. *Electron. J. Biotechnol.* 2009, 12, 7–8. [CrossRef]

192. Sanmartin, M.; Drogoudi, P.D.; Lyons, T.; Pateraki, I.; Barnes, J.; Kanellis, A.K. Over-expression of ascorbate oxidase in the apoplast of transgenic tobacco results in altered ascorbate and glutathione redox states and increased sensitivity to ozone. *Planta* 2003, 216, 918–928. [CrossRef] [PubMed]

193. Chiang, C.-M.; Chen, C.-C.; Chen, S.-P.; Lin, K.-H.; Chen, L.-R.; Su, Y.-H.; Yen, H.-C. Overexpression of the ascorbate peroxidase gene from eggplant and sponge gourd enhances flood tolerance in transgenic *Arabidopsis*. *J. Plant Res.* 2017, 130, 373–386. [CrossRef] [PubMed]

194. Chiang, C.M.; Chen, L.F.O.; Shih, S.W.; Lin, K.H. Expression of eggplant ascorbate peroxidase increases the tolerance of transgenic rice plants to flooding stress. *J. Plant Biochem. Biotechnol.* 2015, 24, 257–267. [CrossRef]

195. Choi, H.W.; Hwang, B.K. The pepper extracellular peroxidase *CaPO2* is required for salt, drought and oxidative stress tolerance as well as resistance to fungal pathogens. *Plant Physiol.* 2012, 235, 1369–1382. [CrossRef] [PubMed]

196. Hwang, I.S.; Hwang, B.K. The pepper 9-lipoxygenase gene *CaLOX1* functions in defense and cell death responses to microbial pathogens. *Plant Physiol.* 2010, 152, 948–967. [CrossRef] [PubMed]

197. Lim, C.W.; Han, S.-W.; Hwang, I.S.; Kim, D.S.; Hwang, B.K.; Lee, S.C. The pepper lipoxygenase *CaLOX1* plays a role in osmotic, drought and high salinity stress response. *Plant Cell Physiol.* 2015, 56, 930–942. [CrossRef] [PubMed]

198. Wi, S.J.; Kim, S.J.; Park, K.Y. Constitutive S-adenosylmethionine decarboxylase gene expression increases drought tolerance through inhibition of reactive oxygen species accumulation in *Arabidopsis*. *Planta* 2014, 239, 979–988. [CrossRef] [PubMed]

199. Gupta, N.; Rathore, M.; Goyary, D.; Khare, N.; Anandhan, S.; Pande, V.; Ahmed, Z. Marker-free transgenic cucumber expressing *Arabidopsis* *CBF1* gene confers chilling stress tolerance. *Biol. Plant.* 2012, 56, 57–63. [CrossRef]

200. Liu, L.; Duan, L.; Zhang, J.; Zhang, Z.; Mi, G.; Ren, H. Cucumber (*Cucumis sativus* L.) over-expressing cold-induced transcriptome regulator ICE1 exhibits changed morphological characters and enhances chilling tolerance. *Sci. Hortic.* 2010, 124, 29–33. [CrossRef]

201. Zhang, Y.; Chu, G.; Hu, Z.; Gao, Q.; Cui, B.; Tian, S.; Wang, B.; Chen, G. Genetically engineered anthocyanin pathway for high health-promoting pigment production in eggplant. *Mol. Breed.* 2016, 36, 54. [CrossRef]

202. Seong, E.-S.; Choi, D.-I.; Cho, H.-S.; Lim, C.-K.; Cho, H.-J.; Wang, M.-H. Characterization of a stress-responsive ankyrin repeat-containing zinc finger protein of *Capsicum annuum* (CaKR1). *BMB Rep.* 2007, 40, 952–958. [CrossRef]

203. Yi, S.Y.; Lee, D.J.; Yeom, S.-I.; Yoon, J.; Kim, Y.-H.; Kwon, S.-Y.; Choi, D. A novel pepper (*Capsicum annuum*) receptor-like kinase functions as a negative regulator of plant cell death via accumulation of superoxide anions. *New Phytol.* 2010, 185, 701–715. [CrossRef] [PubMed]

204. Xu, H.; Sun, X.; Wang, X.; Shi, Q.; Yang, X.; Yang, F. Involvement of a cucumber MAPK gene (*CsNMAPK*) in positive regulation of ROS scavengence and osmotic adjustment under salt stress. *Sci. Hortic.* 2011, 127, 488–493. [CrossRef]

205. Sarowar, S.; Kim, E.N.; Kim, Y.J.; Ok, S.H.; Kim, K.D.; Hwang, B.K.; Shin, J.S. Overexpression of a pepper ascorbate peroxidase-like 1 gene in tobacco plants enhances tolerance to oxidative stress and pathogens. *Plant Sci.* 2005, 169, 55–63. [CrossRef]

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