

Available online freely at www.isisn.org

Bioscience Research Print ISSN: 1811-9506 Online ISSN: 2218-3973

OPEN ACCESS

REVIEW ARTICLE

Journal by Innovative Scientific Information & Services Network

BIOSCIENCE RESEARCH, 2023 20(1): 117-128.



Salman Aloufi

Department of Biotechnology, Faculty of Science, Taif University, Box 11099, Taif 21944, Saudi Arabia

*Correspondence: s.alouf @tu.edu.sa Received: 22-02-2023, Revised: 24-03-2023, Accepted: 26-03-2023 e-Published: 27-03-2023

The fruits are exposed to many infections and diseases on farms or during storage due to pathogens in the plant such as fungi. Many of these infections cause the fruit to lose many properties and impair its quality. In most cases, when infection occurs, the fruits begin to change their skin and continue until their complete death. Some studies showed that genes play a major role in resisting many pathogens in fruits, and depend on those genes in choosing good fruits, which helped to study genetic diversity among fruit types. Also, many previous studies have shown that there are some varieties that have a great genetic ability to resist the most common pathogens, which made the fruits grow and ripen properly. There are a group of hormones associated with ripening in fruits and play an important role in regulating the ripening process according to climatic conditions, the nature of the environment, and the genetic diversity of fruit varieties. Some studies have indicated that epigenetic control of gene expression (without changing the original DNA sequence) controls fruit ripening as morphological differences can occur without changing the DNA sequence. This scientific review focuses on the associations between plant hormones, transcription factors, and epigenetic modifications that control fruit ripening by discussing the latest findings of comparative genomics and system biology.

Keywords: Fruit ripening, Ethylene, phytohormone, gene resistance, DNA methylation, Epigenetic.

INTRODUCTION

The fruit is characterised by the floral plants. They exist in many forms and sorts. Nowadays, the forefather's fruit, dry and cracked, presumably emerged in the early days of the Cretans; later in cretaceous, or in early luscious fruits developed. The primary territory, mechanisms for seed distribution are the differentiation of fruits from the dried fractures to fruits and berries from fleshy stone, connected to vertebrates (Gao, et al. 2019). Fruit ripening is an intricate and well-coordinated development process. The fleshy fruits become delectable, tasty, and soft throughout ripening, attracting cattle and aiding in seed dispersal. In addition to softening, more regularly accumulated sugars, acids, colours, and volatile substances often increase animal interest and taste (Sánchez-Sevilla, et al. 2017). Moreover Fruit serves as an essential dietary supplement by providing humans with nutrients, vitamins, fibre, and antioxidants. "From an agronomic perspective, the nutritional value, flavour, quality, and shelf life of the fruit are characteristics. Important adjustments to moulding processes include coloration, wall hardness (raising sucrose, acid loss, taste, etc.), and pigmentation (loss of green coloration and increases, depending on the species and cultivar).

The following variables apply to colouring and increase in non-photosynthetic pigments (production of

volatile compounds giving a characteristic odor) (Sánchez-Sevilla, et al. 2017).

"Although Malus domestica is one of the most important and renowned fruit cultivations in the globe", farmers still confront several unresolved production, storage and marketing challenges. For at least four month in cold shops, most commercially cultivated fruit must be sold. However, mushroom Rot causes considerable loss of fruit during storage. In the low-consumption production systems (e.g., organic olives, agroforestry), fruits are especially sensitive and losses up to 20 times higher than in conventional orchards can be prevented from being effectively protected against fungicides. Increased returns are typically limited since after a shorter shelf life the product must be placed on the market (Zhou, 2019).

The amount of fungicides that are permissible in standard manufacturing systems has decreased, and in an increasing number of countries, use of fungicides after harvest is completely outlawed. Moreover, the use of the shielding gas 1-MCP (1-methylcyclopropene) is prohibited. Marketing goals highlighted the desire for high-strength apple cultivars, such as in Scandinavia (Li, et al. 2020). As a result, various deposits' relative levels of resistance and susceptibility to rot are increasing.

Differences between varieties of phenotypic and physiological features such as fruit ripening, including changes in the contents of intrinsic ethylene (IEC), fruit texture, structures of fruit skin, and chemical contents, for example sugar, as well as a variety of antifungal elements such as chlorogenic acid and quercetine may be related. This research includes information about ways of measuring damage caused by different storage rots and the influence of fruit ripening factors on storage rot. This will fortunately be beneficial for plant farmers wishing to create cultivars with enhanced resistance to warehouse red assaults.

As a well-coordinated growing process, the ripening of fruit and seeds coincide. "Thousands of genes control the gradual softening and/or lignification of pericarp layers, and the maturing process controls the accumulation of carbohydrates, acids, pigments, and volatile chemical emissions." The key to increasing yields is a better understanding of the mechanics underlying fruit ripening (Giovannoni, et al. 2017). Tomato demonstrates its involvement in the regulation of maturity of ethylene and related molecular networks, mutations that prevent a fruit maturing transition (Ito, et al. 2017). However, it remains unclear how the other plant hormones play. In this review, one will discuss the tight connectedness of plant hormones, transcription factors and epigenetic alterations to maintain precise control over the ripening process. There is a discussion of recent results in comparative genomics and system biology.

The Cause of Red Storage Fungi

Many big fungi are often attacked in both orchards and cold warehouses, generally known as storage rot. Symptoms begin first as changes in the skin and the whole fruit may continue to decay. "Apple Storage Rot includes two types of biotrophics, N. malicorticis, and N. malicorticis perennans, and Colletotrichum, which are known as C. acutatum and C. gloaosporioides, divided into several taxa while other imports are called "N. malicortia" or "N. malicortia," which may be known to include neo-fabraea (e.g. N. alba (syn. N. vagabunda)) ("pathogenic wounds") (Li, et al. 2020). Furthermore, there are various fungi that can induce apple flowering and huge necrotic patches on the fruit, known as floral red top or calyx final rot. Relative value varies considerably from year to year, including Alternaria spp., Botryosphaería obtusa or Cadophora luteo-olivacea to Fusarium spp., Mucor pyriformis, Neonectria ditissima or Phacidiopycnis washingtonensis" (Chen, et al. 2019).

Storage Rot Damage Quantification

Selecting and advancing the development of optimum varieties is based on genetic diversity among variation types (each variety generally reflects an apple-genotype or other crop produced via clone). Studies from orchards and warehouses showed that the cultivars have a significant susceptibility to some of the most common rot (Chen, et al. 2019). However, only natural infections cannot correctly measure the level of resistance, as the quantity and virulence of the inoculum might fluctuate. While a few fungi may be to blame for the symptoms in the foetus, some fungi also struggle with obvious harm. It may be difficult to classify the fungus in question, but longterm molecular study can help by supplementing the available morphological data (Gao, *et al.* 2020).

inoculations Experimental with well-defined inoculations were done for different fungi to collect adequate quantification data for a wide number of genotypes. In rot, such as P. spansum, B. cinerea, and M. wounding and inoculation fructigène, with the conidiospores of the fungus are generally used to inoculate numerous fruit (Li, et al. 2017). Damage and inoculation can also be performed at a time by use of a plastically disposable pipette, allowing one to three inoculation sites for each fruit to be treated in large quantities in a limited period. For each lesion evaluated at pre-defined time following storage the severity of the illness is generally estimated as an average (Li, et al. 2019).

Linear regression may compute the repetition rate (LEGR) if repeated action is done over several days or weeks. The lesion width = severity of the illness (S), in addition to the most widely used criterion, may also be measured as the disease effect by the rate of inoculations that cause lesions (I). Sometimes (I — S)/ 100 is based on the severity of illness. In the investigation on the wild M. sieversii growths implanted in P. expansion wounds, however, S and I were strongly linked.

Resistance mechanisms and gene resistance search

According to studies on 11 different varieties of apples colonised by *Botryosphaeria dothidea*, which causes the apple ring red in Asia, the thickness of a fruit's cuticle and the quantity of spiracles are crucial factors for biotrophic fungi that infect spirals. A flowery open cup may also provide a pathway for the decay of many deposits. Advanced recognition and signalling networks depend on the complex interactions between fungi and their hosts in order for both biotrophic and necrotrophic fungi to be able to infect their hosts (Xiao, et al. 2020). The buildup of reactive oxygen (ROS) species, known as oxidative explosion, is a frequent reaction by plants to a fungal assault.

This might have a direct detrimental impact on the pathogens through damage to the cell membrane and the mitochondrial proteins. The plant then has to eliminate the excess ROS, often by raising the amount of glutathione and ascorbate. The fungal attack takes place through generating a wider range of chemicals, "For example the host's innate immune system is defeated by phytotoxic chemicals, cells wall modificators, and protein effectors. These changes have a complex effect on several separate fruit components. The total phenol concentration therefore increased in one study, presumably due to phenylalanine lyase activation. Soon after P. expansion inoculation and then again decreased "(Xiao, et al. 2020).

In an unpublished GWA investigation of the extent of

the lesions for 180 P. vaccine expansion, principally for apple growing in Northern Europe, no significant correlation has been discovered on the ground (Tian, et al. 2019). Axiom®Apple 480K Field above LG3 Lesion decline (Figure 1). "In general, significant SNP-based associations in unrelated material require a much bigger sample dimension. This putative QTL is near to QTL, as stated by Norelli et al. for p. expansion resistance in M. sieversii was not established yet. The consequences, however, are quite fascinating, since the location of these two QTLs implies, not only in Wild-type M. sieversii but also in M. è I domestica, that there is a considerably difference in P resistance" (Liu, et al. 2017).

As genome-wide sequencing tools become more widely available, genes can now also be found utilising them. These kinds of data, along with mass segregation analysis, were used to identify resistance genes against C. fruticaola leaf spots on glomerella leaves (Liu, et al. 2017). The complete genomic comparison of the profiles of the SNP from the 'Golden Delicacies' to 'Fuji' crosses, across resistent and susceptible volumes, enabled the locus of the Rgls to be mated precisely to LG15, It is not known whether Glomerella leaf spots will be beneficial for forecasting the rotten resistance produced by the same species during apple fruit storage. Previous studies have shown a clear organ specialisation, with differences in its influence on the oxidant defencing mechanism of the host enzyme in at least certain isolates of C. fructicola (Yang, et al. 2017).

Effect of Fruit Ripening, Texture and Chemical Fruit Composition

In this review, the day on which the variety is ready for (commercial) harvest is defined as "renewal period." If many varieties are evaluated in a given trial, they generally calculate the ripening time as the amount of days between the harvest of the oldest ripening variety and the harvest. The period for renewals of recognised main varieties like Gala or Golden Delicia cultivated in every environment and adjustment by location and year if necessary can provide comparative statistics for varieties grown at various places (Zhang, 2018). "Fruit ripeness" refers on the other hand to the stage that the fruit is capable of reaching, (e.g. unripe, medium ripe, overripe). Early ripening plants (summer apples) tend to produce high menopause and ethylene and ripen fast whereas late ripening (fall apples) have less breathing and production of ethylene and are ripe more Slower (Yue, et al. 2020).

It is difficult to describe the "texture" of a fruit since it consists of so many different elements, including chewability, crunchiness, juiciness, and hardness. The resistance parameter is the one that is most frequently measured, and a straightforward, portable penetrometer can be used to do so. An automated texture analyzer has recently been used in research to examine several texture features at once. The examination of 86 types of apples with this instrument indicated a large concordance between mechanical factors and the 'resistance,' while the acoustic set was in accordance with the perception 'crack' by humans (Wang, et al. 2020).

During harvesting and after storage resistance is frequently tested as the harvest time decreases. In combination or most commonly just after an IEC climatic spike, generally with a large drop. Commercial fruits are generally picked, especially when meant for long term storage, while they are in a specific balance, i.e. before the IEC is increased (Chen, et al. 2021). The stiffness after the cold storage influences not only the initial stiffness but also the choice of apple types in current olive grovers. The loss of rigidity is called fruit softening (harshness difference between the measures) (Fan, *et al.* 2018).

The result is often divided into weeks of storage to offer the "smoothing rate" and enable comparison of the variety stored across different time periods. An extensive investigation of the Swedish and Norwegian variation collections revealed a positive correlation between ripening and harvest resistance (early harvest ripening is weaker than late ripening), and a negative correlation between ripening and soaking rates (Fan, et al. 2018). Varieties of (early ripening lose power more quickly than those that ripening early) (Latest Ripening). "However, there has been no link between the date of commercial harvest (i.e. time of ripening) and the strength of the fruit when 23 Belgian cultivars were compared" (Lü, et al. 2018).

The proportion of various compounds in cellulose and husks influences the environmental variables such as soil, pruning, fertilisation, irrigation and weather conditions. Genetically different genetically modified chemicals can also lead to a high degree of reproductive Resistance in genotypes rich with favourable substances (Fu, et al. 2017). However, a series of diverse genotypes are rather complicated to quantify the chemical content and should be evaluated throughout similar fruit ripening phases. The hydrolysis of starch during the ripening phase is responsible for considerable growth of sugars (commonly known as solutionable solids, SSCs) in the fruit body and the breathing metabolism reduces acidics (oft known as titrated acidity, BP) (Lü, et al. 2018).

The ripeness of the investigated specimens may potentially interfere with, with the fruits of the initial specimens characterised by a relatively low ascorbate at harvest and a quicker decrease of the fruits of the late ripening species in the next 10 days (Khaksar, et al. 2019). Furthermore, the chemical composition of the fruit generally varies in response; different fungal species acidify or alkalize their environment in order to aid infection and necrotrophic fruit development. Apple sugars in particular have a major role in breathing, which is increased during illness, and in consumption of energy. However, the defensive systems may be characterised by various phenolic compounds (Li, et al. 2020).

Ripening During Hormonal and Transcriptional Adolescent Regulation

Fruits are typically categorised according to their airborne activities and the ethylene production profiles throughout ripening into two physiological categories, menopausal and nonbactérial. For the correct Ripening of fruits and preventing the production and perception of this hormone, the synthesis of ethylene on climate fruit such as tomatoes, apples and bananas is crucial (Chang, 2020).

Initially focused on the dried, fragmented silica of Arabidopsis model plant was efforts to find the transcriptional legislation underpinning the formation of fruits and berries? These findings illustrated the function played in tissue specificity and dehiscence process by numerous MADS box transcription factors (Chang, 2020). "The identification of the silicon valve edge was established by the redundant SHATTERPROOF 1/2 (SHPs, members of the AGAMOUS subfamily": when relocated, the fruit became irrelevant. In spite of major structural variations between dry and fleshy fruits, further focussing particularly investigation, on tomatoes, discovered some orthologies for these Arabidopsis genes in order to govern the ripening of those MADS boxes. Some of the regulatory networks underlying fruit growth during fleshy fruit development have now been maintained (Li, et al. 2020).

"There are a number of key developments in the processes of the control of ripening, including renal, immature, colourless (Cnr), green ripening (Gr), green meat (gf), high pigmento1 (hp1), high pigment2 (hp2), and never ripe. The reindeer mutant encodes the SEPALLATA clade's partly deleted MADS-box protein whereas Cnr is an epigenetic change that changes the methylation of the SQUAMOSA promoter protein (SPB). NOR is a transcription factors member of the NAC domain" (Chang, 2020).

"A new work integrating transcriptomics, proteomics, and targeted metabolite analysis increased the expression of chain-regulated ethylene genes and expanded our understanding of the function of this hormone in controlling proteins and metabolites in tomato ripening (Khaksar, et al. 2019). These results supports the idea that either the kidney in one cascade does not operate together to control the ripeness or suggests that it is unlikely to function earlier than the kidney in genes associated to ethylene expression/ripening" (Li, et al. 2020).

More recently, a combination technique based on chromatin immunoprecipitation and transcriptome analysis has been revealed for the RIN to interact with promoters of over 200 genes and to govern the exhibition of its objectives through activation or substitution. The RIN goal genes are essential regulator controls on maturity, including CNR and NOR, as well as renowned active pathways (e.g. carotenoid accumulation, chlorophyll rupture, ethylene synthesis, and perception) (Fu, et al. 2017).

"The lack of respiratory explosions, and minimal endogenous production of ethylene compared with normal menopause fruits, fruit such as strawberries, citrus fruits and grapes were designated as nonbacterial. In potatoes, certain variations seem to be resistant to ethylen, whereas the expression of the ripening-specific genes has been stimulated by other types of pepper treated with exogenous ethylene" (Lü, et al. 2018).

Ethylene has a fairly high concentration of green fruits, drops in white fruit and then grows throughout its refining phase in strawberries, which have proved to be the major model of the non-specific fruit ripening. Interestingly, the latest rise is accompanied by a higher breathing rate which is comparable to that in early puberty of postmenopausal foetuses (Fan, et al. 2018). Different techniques were employed to better understand the role of ethylene in strawberry development. Other cell-related genes such as β -Galactosidase, pectin methylesterase or β -xylosidase were controlled by outside ethylen whereas expression in other genes, such as expanine, was not responsive to ethylene, FaEXP2 (Chen, et al. 2021).

Phytohormones have a significant impact on a complex network of transcription factors and genetic regulators that regulate fruit development, a crucial step in maximising seed dispersal. According to various ripening-defective mutants, the effects of applying exogenous hormones, and transcriptome analyses, abscisic acid (ABA) and ethylene are the primary regulators of ripening and senescence in both dry and fleshy fruits. While ethylene is the most well-known participant in the fruit's final stage of life, ABA also plays a critical regulatory role by helping to increase ethylene production and serving as a stress hormone in response to disease attack and dehydration (Forlani *et al.* 2019)

Since the nineteenth century, ethylene's effects on plants have been understood, and it is well-known as the phytohormone involved in a variety of plant growth and development processes, including the ripening of fruit. Understanding the mechanisms of climacteric fruit ripening and the part ethylene plays in this process has been crucial to increasing fruit yield and fruit quality.

Numerous studies have been conducted on the biochemistry, genetics, and physiology of ripening in economically significant fruit crops, and a wealth of knowledge is now available on everything from the ethylene biosynthesis pathway to the mechanisms of perception, signalling, and gene expression control (Chaves and Mello-Farias, 2006).

Based on the involvement of ethylene during maturation, fruits can be divided into two main types. Nonclimacteric fruits, like cherries, strawberries, and pineapple, mature without the help of ethylene. The 'climacteric crisis', which occurs at the same time as the respiratory peak during ripening in climatic fruits such tomato, avocado, melon, apple, pear, and kiwifruit, is marked by an exceptional increase in ethylene production (Abeles *et al.* 2012). Fruit undergoes a number of structural and biochemical changes during maturation that give them distinct organoleptic features, such as changes to the fruit's appearance, texture, and flavour (Seymour *et al.* 2012).

Several studies have shown that ethylene signalling elements and associated transcription factors are involved in the regulation of fruit ripening. The phytohormone ethylene plays a crucial role in climacteric fruit ripening (Liu *et al.* 2020).

During fruit development and ripening, two ethylene biosynthetic mechanisms have been identified. At the preripening stages, System 1 dominates and is characterised by autoinhibitory regulation of ethylene synthesis. When system 2 is switched on and ethylene production becomes autocatalytic, climacteric ripening begins(Liu *et al.* 2020).

Recent investigations of the transgenic and metabolomic levels of ethyleneally sensitive transgenic strawberries show that regular growth of fruits needs that Ethylene in both sections of strawberries, cornet and containers behaves differently. These results show that, although not as relevant as menopausal fruits, the ripening of strawberry fruits may be influenced by ethylene (Wang, et al. 2020).

A recent comparative study of transcriptome and metabolome (tomatoes and potatoes) showed that both species have comparable ethylene-mediated components of their signal throughout the development of climate and non-bacterial fruit. However, the regulations of these genes in peppers are very different and may indicate an altered sensitivity to ethylene or other regulators than to tomatoes (Yue, et al. 2020). The ethylène biosynthesis, the aminocyclopropane-1-carboxylic acid synthase (ACC) and the ACC oxidase genes are not activated in potatoes, contrary to the scenario observed in tomatoes. Ethylene absorptions are upwardly controlled throughout pepper ripening, for instance by genes linked to walls cells, ethylene reactivity factor 3 (ERF3) and biosynthetic genes for carotenoids (Zhang, 2018).

Other frequently regulated genes have been described between menopausal and non-menopausal fruits. "For the correct growth and ripening of strawberries, the SEPALLATA gene (SEP1/2; MADS-box) is needed. Similarly, the MADS-box SEP3 gene also shows ripening-related expression in banana categorised as menopausal fruit (Yang, et al. 2017). The expression of the MADS2 gene is also related with fruit firmness in apples, whilst the blueberries are used to regulate the production in Anthocyanine in the SQUAMOSA MADS-box orthologist of TDR4 in the tomatoes" (Liu, et al. 2017).

Current understanding of the involvement of hormones – other than ethylene – in menopausal and nonmenopausal fruit growth and ripening is limited (Yang, et al. 2017). The most plentiful form of free Auxin, Indole 3 Acetic Acid (IAA), which decreased in tomatoes, peppers, bananas, nutmegs and Strawberries was accompanied by a rise in its conjugate form of IAA Asp. Catalysed by the IAA-amino (GH3) synthase gene, the conjugation reaction. Fifteen GH3 family members have been described in tomatoes, although only two show ripening-related expression patterns (Liu, et al. 2017).

The over-expressing tomatoes of the pepper gene GH3 display ripening expectations that correspond to the idea that the IAA-to-AA-conjugated ratio may contribute to a period of ripening rather than IAA levels. No single growth regulator appears to have a beneficial role in nonclimate fruit similar to that of ethylene, although the ripening of nonsubstantial fruits is negatively affected by auxine (Tian, et al. 2019). The treatment of several ripening-specific genes with exogenous auxin has been found to decrease Strawberries expression. Auxine also seems to have an adverse function to play in the regulation of ripening via synthetic auxine therapy, which slows down the expression of a number of ripening genes (Xiao, et al. 2020).

Due to the important function of auxins in nonbacterial fruit growth and ripening, the potential role of other plant hormones such as gibberellins has been poorly regarded (GA) (Li, et al. 2019). In the case of strawberries however, topical application of GA3 to fruit ripening has discovered that the development of the red colour has been delayed significantly and genes in cell growth and cell wall disintegration have altered expression. Abscisic acid (ABA) in plants is recognised to involve phytohormonic acid in numerous growth, development, and environmental stress response aspects (Gao, et al. 2020).

ABA encourages a sugar buildup in meat fruits to control the environment and non-bacterial fruit ripening. Certain cell wall genes, such as polygalacturonase and pectin methylesterase, are downregulated as a result of the tomatization of the gene that catalyses the early step ABA (NCEd1 9x-epoxycarotenoid of synthesis dioxigenase) (Li, et al. 2017). More life. Long life. The NCED expression decrease also corresponds to a delay in strawberry ripening. ABA is said to cause strawberries and raisins to ripe. However, in raisins, GH3 promoter study found ABRE-like elements, which may imply that the ratio of ABA to Auxin is linked to early ripening, is unclear to its processes (Chen, et al. 2019).

In recent years, there has been a significant increase in our understanding of the molecular phenomena at the transcriptional, biochemical, hormonal, and metabolite levels underlying the development of climate and nonbacterial fruits (see Figures 1 and 2). However, throughout the shift from unripe to mad fruit, one still do not comprehend the evolution of the hormonal response (Li, et al. 2020). To present, most published research on regulating transcription and metabolism at spatial and temporal levels have very low resolution and are still constrained by the coverage of many molecular units of the cell. Improved technologies and statistical tools, however, enable us to further strengthen our analytical

Salman Aloufi

capabilities, in order to tackle difficulties like subcellular compartments and the contrast behaviour. In addition it will be necessary to understand the genetic control of fruit growth and ripening by making the high quality data of the fruit genome sequence available (Zhou, 2019).

Epigenetic Reshaping in Ripening

The epigenetic control of gene expression is increasingly acknowledged as a genome module (the inheritance without changing the original DNA sequence). morphological Inherited differences can arise spontaneously at a generic locus of plants, without changing the underlying DNA sequence (Sánchez-Sevilla, et al. 2017). Epigenetic alterations are possible. The form of the epigenetic regulation is DNA methylation. It is involved in transcription control, stress reactions and also plays a vital function in safeguarding the genome from transposable element activity (TE) and other repetitive sequences. DNA in the production plant is methylated to cytosine residues in three sequences (CG, CHG and CHH, where H = A, C or T) (Ito, et al. 2017).

The contribution of system profiling techniques will assist to discover and clarify the connection between epigenomic reshaping and the transcriptional regulation that takes place during the ripening process (Giovannoni, et al. 2017).

Analyse the epigenetic alterations in Arabidopsis shows that in their coding area, at least a third of the genes expressed are methylation and just 5 percent of the genes in promoter regions are methylated. But methylation promoter genes have a more selective expression of tissue (Yang, et al. 2017).

The first frequency and distribution research in tomato cytosine methylation sites is over 20 years ago. Cyticosine methylation between two tomato species and the patterns of methylation were found to be reasonably plentiful polymorphisms in which parents are stably inherited from Mendel to Mendel (Li, et al. 2020). The availability of methylation models for specific tissues and the overall 5 mC decline in frequency in the tissue development also prompted the scientists to hypothesise change in chosen alleles' methylation state during plant development (Zhou, 2019).

In *Prunus spp*, endocarp phénotypes are widely diversified, the majority of which occur in conventional reproduction. (Sánchez-Sevilla, et al. 2017) have shown that the thickness, durability and bitterness of the fruiting body varies with almond shells. Seeds of some kinds of peach, apricots and plum plums owing to open fruit body are easily susceptible to pests and illnesses. This abnormality known as the "split pit" is due to downward regulation of the biosynthetic genes of phenylpropanoids. The pit development includes environmental conditions, growing techniques and ripening period. Especially early types of fish and penny plums are often more likely to break seeds because their hole is not sufficiently reinforced to resist rapidly increasing fruit body power (Minas 2016).

As a consequence of the inadequate growth of the endocarpal layer, partly naked seeds have been denominated the solitary natural phenotype in the wild feather population (Giovannoni, et al. 2017). The stonefree genotype is very environmentally sensitive as the whole endocarp can develop throughout the years at mild temperatures in the spring, while in the colder years the very cold stone is present. The lack of endocarpic tissues implies that the whole endocarpic layer is not present in this mutant (Olubero et al. 2017).

Recently, the impact of cytosine methylation on ripening tomato fruits was demonstrated in the molecular characterization of colourless, immature phenotypes. The immature Cnr tomato mutant lacks ripe berries, has a green pericarp, and is unable to withstand the administration of exogenous ethylene (Olubero et al. 2017). The Cnr locus gene was discovered by positional cloning as an SPB protein, however no modification of the gene coding sequence could be ascribed to the immature phénotype. By contrast, at the area upstream from the start of ATG the bisulfite sequence of the Cnr allle mutant has shown hypermethylation of cytosine (Minas 2016).

The expression of the Cnr gene was significantly decreased as a result of this hypermethylation situation. This immature phenotype was caused by an inherited pattern of cytosine hypermethylation, specifically in the Cnr gene promoter (Harris, 2018). In addition, the Cnr promoter appears to have been demethylated in a particular area shortly before the start of the ripening during the formation of normal tomato (cf. Liberto). This led to the assumption that DNA methylation was used to regulate fruit ripening (Lang, 2017).

A genomic picture of the connection between the genetic ripening programme and the status of DNA methylation is shown by recent studies by (Burdon, 2017). The scientists injected a chemical inhibitor of cytosine 5-azcidine methyulation in the sites of the ulnar and in the column of tomato fruits based on earlier discoveries on the nature of the Cnr (epi) mutation. The inhibitor for methylation caused the development of ripening red areas in which the production of typical ripening genes was anticipated (phytoene synthase 1 and polygalacturonase) (Olubero et al. 2017).

Moreover, the area of the Cnr Promotor has been demethylated from the greens of the fruit in the red sectors, signalling adequate epigenetic indication of Cnr's demethylation to trigger ripening (Hilioti, et al. 2016). The authors subsequently extended their ideas on the significance of cytosesine methylation, including abnormal ripening by Cnr and renal mutations, which detailed the full sequence of methylomas from tomato leaves, unripe and ripe fruit (Ichinose, 2017).

At least three significant outcomes were found in the sequence of the whole epigenome: The degree of methylation of the first transcription sites (ITS) upstream regions steadily reduced during fruit development in wild

Salman Aloufi

foetuses, in the Cnr and rin mutants of ripening (Burdon, 2017). Which have been consistently higher in CG levels in TSS and also in Cnr, this general drop was not seen, which was similar to those reported in the leaves. Common ripening gene promoters are being gradually demethylated as wild foetuses are developing (Harris, 2018).

Additional evidence of a relationship between cytosine ripening and methylation was obtained by the identification of ChIP-Seq fruit binding sites. 292 genes with established ripening functions were included in the RIN objective set (Minas 2016). The "hot areas," continuous methylation or overlaps upstream of TSS, have been proven to represent RIN binding sites. "Analysis of the methylation state of these regions shows that they have gradually been demethylated as they transition from green fruits to red fruits, which is correlated with greater transcription levels of RIN target genes (Lang, 2017).

Previous study has demonstrated that Cnr epimutant was blocked by RIN binding to only a restricted number of promoters, indicating that RIN binding might be prevented (Harris, 2018). The major results are that I local treatment immature fruit with chemical substances of bv demethylating DNA speeds up ripening, ripening gene promoters containing RIN sites gradually desmethylate, but in the case of defective mutants of ripening stav stablely hypermethylised and RIN does not bind CNR promoters together (and possibly all hypermethylated target gene promoters). Assigning a major role in the coordination of tomato fruit ripening to the epigenome structure and the developmental dynamics" (Burdon, 2017).

The general scenario given thus far also indicates that the gradual demethylation by gene promoters associated with ripening may be a requirement for the binding of transcriptional regulators, which induces an accumulation of the transcripts associated with ripening (Minas 2016). The mechanism that promotes demethylation, however, remains mysterious with regular fruit growth and additional studies to identify the "lack of bonds" are needed. Give us an assumption that epigenetic alterations in fruit phenotypes will allow the future regular screening of crop epigenomes that will speed up the identification of ePigenetic changes due to the rising relevance (Olubero et al. 2017).

In addition to the analyses of traditional genetical variants in future development methods, testing for epigenomic structure and dynamics are assumed to exist. Epigenetic methods to improved yield can substantially change characteristics of fruit quality, particularly for qualities whose allegiance during domestication or recent severe reproductive stress has been decreased (Hilioti, et al. 2016). Future modelling work for epigenomic and small-range RNAs will give a significantly better understanding of complicated dynamics behind this well-regulated biological process in combination with the most often utilised profiling of transcripts, proteins, Enzymes

and metabolites, as illustrated in Figure 2 (Ichinose, 2017).

CONCLUSION

The major entrance sites are animal and/or human fruit wounds when necrotrophic red, for example blue mould induced by P. expansion, is kept. Therefore, the variety's resistance may be assessed appropriately by inoculating the wounds with fungal spores. Penicillium expansion has been assumed to oppose the process of fruit protection by regulating endogenous production of ethylene. It also lowers the pH of the host tissue and secretes lytic enzymes which accelerate breakdown of the cell wall. This leads to a superior performance than early ripening variants, which generally are characterised by a lower ethylene explosion environment and a greater fruit firmness when infected with bleuish moth. In general, the link between the lesion disintegration and the fruit's chemical composition is less obvious. But shows that the resistance level is affected mostly by the fundamental phenolics and by contents changes owing to infection. There is significantly less evidence available for other necrotrophs, such as Botrytis and Monilinia, but the resistance of these fungi is presumably similar to that of fruit ripening.

Colletotrichum and Neofabraea, for example, are largely penetrated by biotrophic storage rot through open pores in fruit development of the flowers or spiracles and microcracking. Whey vaccination therefore only assesses a portion of these fungi's genetic immunity. Sadly, woundfree techniques of immunisation are longer to use and harm is harder to assess. The effects of ethylene-related mechanisms on biotrophic resistance cannot yet be identified because to the small number of thorough investigations of these fungi. However, in comparison with the well researched P. expanse, the link with the ripening period and fruit rigidity is less evident for Colletotrichum and Neofabraea.

The number one wish list for plant breeders is good germplasm with high-impact and well-defined genes, preferably provided with simple DNA markers. The best development of rot-resistant cultivars would require a variety of geographical plasmas to span wide differences in ripening time. Therefore, excellent immunisation procedures need to be developed to acquire huge numbers of phenotypic data, which will provide the basis for genetic research.

CONFLICT OF INTEREST

The authors declared that present study was performed in absence of any conflict of interest

ACKNOWLEDGEMENT

Author appreciate the cooperation of department of Biotechnology, faculty of science at Taif university on this review..

Copyrights: © 2023@ author (s).

This is an open access article distributed under the terms of the **Creative Commons Attribution License (CC BY 4.0)**, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author(s) and source are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

REFERENCES

- Aragüez, I. *et al.* Eugenol production in achenes and receptacles of strawberry fruits is catalyzed by synthases exhibiting distinct kinetics. *Plant Physiol* **163**, 946–958 (2013).
- Ben, L. & Salzberg, S. L. Fast gapped-read alignment with Bowtie 2. *Nature mathods* **9**, 357–359 (2012).
- Burdon J, Pidakala P, Martin P, Billing D. Softening of 'Hayward' kiwifruit on the vine and in storage: The effects of temperature. Scientia Horticulturae. (2017); 220:176–82.https://doi.org/10.1016/j.scienta. 2017.04.004.
- Chai, Y. M. *et al.* Brassinosteroid is involved in strawberry fruit ripening. *Plant Growth Regulation* **69**, 63–69 (2013).
- Chambers, A. H. *et al.* Identification of a strawberry flavor gene candidate using an integrated genetic-genomicanalytical chemistry approach. *BMC Genomics* **15**, 217 (2014).
- Chang, H. Y., & Tong, C. (2020). Identification of Candidate Genes Involved in Fruit Ripening and Crispness Retention Through Transcriptome Analyses of a 'Honeycrisp'Population. *Plants*, *9*(10), 1335.
- Chen, H., Li, H., Lu, X., Chen, L., Liu, J., & Wu, H. (2019). Identification and expression analysis of GRAS transcription factors to elucidate candidate genes related to stolons, fruit ripening and abiotic stresses in woodland strawberry (Fragaria vesca). *International journal of molecular sciences*, *20*(18), 4593.
- Chen, S., Wu, J., Zhang, Y., Zhao, Y., Xu, W., Li, Y., & Xie, J. (2021). Genome-Wide Analysis of Coding and Non-coding RNA Reveals a Conserved miR164–NAC–mRNA Regulatory Pathway for Disease Defense in Populus. *Frontiers in Genetics*, *12*.
- Chen, W. W. et al. Requirement of CHROMOMETHYLASE3 for somatic inheritance of the spontaneous tomato epimutation colourless nonripening. Sci. Rep. 5, 9192 (2015).
- Cherian, S., Figueroa, C. R. & Nair, H. 'Movers and shakers' in the regulation of fruit ripening: a crossdissection of climacteric versus non-climacteric fruit. *J Exp Bot* **65**, 4705–4722 (2014).
- Concha, C. M. Methyl jasmonate treatment induces changes in fruit ripening by modifying the expression

of several ripening genes in Fragaria chiloensis fruit. *Plant Physiol Biochem* **70**, 433–444 (2013).

- Cumplido-Laso, G. *et al.* The fruit ripening-related gene FaAAT2 encodes an acyl transferase involved in strawberry aroma biogenesis. *J Exp Bot* **63**, 4275– 4290 (2012).
- Daminato, M., Guzzo, F. & Casadoro, G. A SHATTERPROOF-like gene controls ripening in nonclimacteric strawberries, and auxin and abscisic acid antagonistically affect its expression. *J Exp Bot* **64**, 3775–3786 (2013).
- Fan, Z. Q., Ba, L. J., Shan, W., Xiao, Y. Y., Lu, W. J., Kuang, J. F., & Chen, J. Y. (2018). A banana R2R3-MYB transcription factor MaMYB3 is involved in fruit ripening through modulation of starch degradation by repressing starch degradation-related genes and MabHLH6. *The Plant Journal*, *96*(6), 1191-1205.
- Filler Hayut, S., Melamed Bessudo, C., and Levy, A. A. (2017). Targeted recombination between homologous chromosomes for precise breeding in tomato. Nat. Commun. 8:15605. doi: 10.1038/ncomms15605
- Fu, C. C., Han, Y. C., Kuang, J. F., Chen, J. Y., & Lu, W. J. (2017). Papaya CpEIN3a and CpNAC2 cooperatively regulate carotenoid biosynthesis-related genes CpPDS2/4, CpLCY-e and CpCHY-b during fruit ripening. *Plant and Cell Physiology*, 58(12), 2155-2165.
- Gao, J., Zhang, Y., Li, Z., & Liu, M. (2020). Role of ethylene response factors (ERFs) in fruit ripening. *Food Quality and Safety*, *4*(1), 15-20.
- Gao, Y., Zhu, N., Zhu, X., Wu, M., Jiang, C. Z., Grierson, D., ... & Qu, G. (2019). Diversity and redundancy of the ripening regulatory networks revealed by the fruitENCODE and the new CRISPR/Cas9 CNR and NOR mutants. *Horticulture Research*, *6*(1), 1-10.
- Gapper, N. E.,McQuinn, R. P., and Giovannoni, J. J. (2013). Molecular and genetic regulation of fruit ripening. Plant Mol. Biol. 82, 575–591. doi: 10.1007/s11103-013-0050-3
- Gerstein, M. B. et al. Architecture of the human regulatory network derived from ENCODE data. Nature 489, 91–100 (2012).
- Giovannoni J, Nguyen C, Ampofo B, Zhong S, Fei Z. The Epigenome and Transcriptional Dynamics of Fruit Ripening. Annual Review of Plant Biology. (2017); 68:61–84. https://doi.org/10.1146/annurevarplant-042916-040906 PMID: 28226232.
- Giovannoni, J., Nguyen, C., Ampofo, B., Zhong, S. L. & Fei, Z. J. The epigenome and transcriptional dynamics of fruit ripening. Annu. Rev. Plant. Biol. 68, 61–84 (2017).
- Han, Y. *et al.* FaSnRK2.6, an ortholog of Open Stomata 1, is a Negative Regulator of Strawberry Fruit Development and Ripening. *Plant Physiol.* doi: 10.1104/pp.114.251314 (2015).
- Harris, C. J. et al. A DNA methylation reader complex that

Fruit Ripening and Various Injuries: the Crosslink between Genetics and Phytohormones- A review

Salman Aloufi

enhances gene transcription. Science 362, 1182 (2018).

- Hilioti, Z., Ganopoulos, I., Ajith, S., Bossis, I., and Tsaftaris, A. (2016). A novel arrangement of zinc finger nuclease system for in vivo targeted genome engineering: the tomato LEC1-LIKE4 gene case. Plant Cell Rep. 35, 2241–2255. doi: 10.1007/s00299-016-2031-x
- Hilioti, Z., Ganopoulos, I., Bossis, I., and Tsaftaris, A. (2014). LEC1-LIKE paralog transcription factor: how to survive extinction and fit in NF-Y protein complex. Gene 543, 220–233. doi: 10.1016/j.gene.2014.04.019
- Ichinose, M., and Sugita, M. (2017). RNA editing and its molecular mechanism in plant organelles. Genes 8:5.
- Ito, Y. et al. Re-evaluation of the rin mutation and the role of RIN in the induction of tomato ripening. Nat. Plants 3, 866–874 (2017).
- Ito, Y. et al. Re-evaluation of the rin mutation and the role of RIN in the induction of tomato ripening. Nat. Plants 3, 866–874 (2017).
- Ito, Y., Nishizawa-Yokoi, A., Endo, M., Mikami, M., and Toki, S. (2015). CRISPR/Cas9-mediated mutagenesis of the RIN locus that regulates tomato fruit ripening. Biochem. Biophys. Res. Commun. 467, 76–82. doi: 10.1016/j.bbrc. 2015.09.117
- Ito, Y., Nishizawa-Yokoi, A., Endo, M., Mikami, M., Shima, Y., Nakamura, N., et al. (2017). Re-evaluation of the rin mutation and the role of RIN in the induction of tomato ripening. Nat. Plants 3, 866–874. doi: 10.1038/s41477-017-0041-5
- Jia, H. F. *et al.* Sucrose functions as a signal involved in the regulation of strawberry fruit development and ripening. *New Phytol* **198**, 453–465 (2013).
- Jia, H. F. *et al.* Type 2C protein phosphatase ABI1 is a negative regulator of strawberry fruit ripening. *J Exp Bot* **64**, 1677–1687 (2013).
- Jia, H., and Wang, N. (2014). Targeted genome editing of sweet orange using Cas9/sgRNA. PLoS One 9:e93806. doi: 10.1371/journal.pone.0093806
- Jia, H., Orbovi'c, V., Jones, J. B., and Wang, N. (2016). Modification of the PthA4 effector binding elements in Type I CsLOB1 promoter using Cas9/sgRNA to produce transgenic Duncan grapefruit alleviating Xcc1pthA4:dCsLOB1.3 infection. Plant Biotechnol. J. 14, 1291–1301. doi: 10.1111/pbi.12495
- Jia, H., Zhang, Y., Orbovi'c, V., Xu, J., White, F. F., Jones, J. B., et al. (2017). Genome editing of the disease susceptibility gene CsLOB1 in citrus confers resistance to citrus canker. Plant Biotechnol. J. 15, 817–823. doi: 10.1111/pbi.12677
- Jinek, M., Chylinski, K., Fonfara, I., Hauer, M., Doudna, J. A., and Charpentier, E. (2012). A programmable dual-RNA-guided DNA endonuclease in adaptive bacterial immunity. Science 337, 816–821. doi: 10.1126/science.1225829
- Karlova, R., Chapman, N., David, K., Angenent, G. C.,

Seymour, G. B., and de Maagd, R. A. (2014). Transcriptional control of fleshy fruit development and ripening. J. Exp. Bot. 65, 4527–4541. doi: 10.1093/jxb/ eru316

- Kaur, N., Alok, A., Shivani, Kaur, N., Pandey, P., Awasthi, P., et al. (2018). CRISPR/Cas9-mediated efficient editing in phytoene desaturase (PDS) demonstrates precise manipulation in banana cv. Rasthali genome. Funct. Integr. Genomics 18, 89–99. doi: 10.1007/s10142-017-0577-5
- Khaksar, G., Sangchay, W., Pinsorn, P., Sangpong, L., & Sirikantaramas, S. (2019). Genome-wide analysis of the Dof gene family in durian reveals fruit ripening-associated and cultivar-dependent Dof transcription factors. *Scientific reports*, *9*(1), 1-13.
- Klap, C., Yeshayahou, E., Bolger, A. M., Arazi, T., Gupta, S. K., Shabtai, S., et al. (2017). Tomato facultative parthenocarpy results from SIAGAMOUSLIKE 6 loss of function. Plant Biotechnol. J. 15, 634–647. doi: 10.1111/pbi. 12662
- Kou, X. H., Watkins, C. B. & Gan, S. S. Arabidopsis AtNAP regulates fruit senescence. J. Exp. Bot. 63, 6139–6147 (2012)
- Kumar, R., Khurana, A., and Sharma, A. K. (2014). Role of plant hormones and their interplay in development and ripening of fleshy fruits. J. Exp. Bot. 65, 4561– 4575. doi: 10.1093/jxb/eru277
- Lang, Z. et al. Critical roles of DNA demethylation in the activation of ripeninginduced genes and inhibition of ripening-repressed genes in tomato fruit. Proc. Natl Acad. Sci. USA 114, E4511–E4519 (2017).
- Li, B., Fan, R., Yang, Q., Hu, C., Sheng, O., Deng, G., ... & Yi, G. (2020). Genome-wide identification and characterization of the NAC transcription factor family in Musa acuminata and expression analysis during fruit ripening. *International journal of molecular sciences*, *21*(2), 634.
- Li, D., Mou, W., Xia, R., Li, L., Zawora, C., Ying, T., ... & Luo, Z. (2019). Integrated analysis of high-throughput sequencing data shows abscisic acid-responsive genes and miRNAs in strawberry receptacle fruit ripening. *Horticulture research*, *6*(1), 1-13.
- Li, L. *et al.* Quantitative proteomic investigation employing stable isotope labeling by peptide dimethylation on proteins of strawberry fruit at different ripening stages. *J Proteomics* **94**, 219–239 (2013).
- Li, R., Fu, D., Zhu, B., Luo, Y., and Zhu, H. (2018). CRISPR/Cas9-mediated mutagenesis of IncRNA1459 alters tomato fruit ripening. Plant J. 160:595.
- Li, S. et al. The RIN-MC fusion of MADS-Box transcription factors has transcriptional activity and modulates expression of many ripening genes. Plant Physiol. 176, 891–909 (2018).
- Li, T., Xu, Y., Zhang, L., Ji, Y., Tan, D., Yuan, H., & Wang, A. (2017). The jasmonate-activated transcription factor MdMYC2 regulates ETHYLENE RESPONSE

Fruit Ripening and Various Injuries: the Crosslink between Genetics and Phytohormones- A review

Salman AloufiFruit Ripening and Various InjurFACTORand ethylene biosynthetic genes to
promote ethylene biosynthesis during apple fruit
ripening. The Plant Cell, 29(6), 1316-1334.

- Li, X., Wang, Y., Chen, S., Tian, H., Fu, D., Zhu, B., et al. (2018). Lycopene is enriched in tomato fruit by CRISPR/Cas9-mediated multiplex genome editing. Front. Plant Sci. 9:559. doi: 10.3389/fpls.2018.00559
- Li, Z., Jiang, G., Liu, X., Ding, X., Zhang, D., Wang, X., ... & Duan, X. (2020). Histone demethylase SIJMJ6 promotes fruit ripening by removing H3K27 methylation of ripening-related genes in tomato. *New Phytologist*, 227(4), 1138-1156.
- Liang, Z., Chen, K., Li, T., Zhang, Y., Wang, Y., Zhao, Q., et al. (2017). Efficient DNA-free genome editing of bread wheat using CRISPR/Cas9 ribonucleoprotein complexes. Nat. Commun. 8:14261. doi: 10.1038/ncomms14261
- Liu, K., Yuan, C., Feng, S., Zhong, S., Li, H., Zhong, J., ... & Liu, J. (2017). Genome-wide analysis and characterization of Aux/IAA family genes related to fruit ripening in papaya (Carica papaya L.). *BMC* genomics, 18(1), 1-11.
- Liu, R. et al. A DEMETER-like DNA demethylase governs tomato fruit ripening. Proc. Natl Acad. Sci. USA 112, 10804–10809 (2015).
- Lor, V. S., Starker, C. G., Voytas, D. F., Weiss, D., and Olszewski, N. E. (2014). Targeted mutagenesis of the tomato procera gene using transcription activatorlike effector nucleases. Plant Physiol. 166, 1288–1291. doi: 10.1104/pp.114. 247593
- Lu, P. T. et al. Genome encode analyses reveal the basis of convergent evolution of fleshy fruit ripening. Nat. Plants 4, 784–791 (2018).
- Lü, P., Yu, S., Zhu, N., Chen, Y. R., Zhou, B., Pan, Y., ... & Zhong, S. (2018). Genome encode analyses reveal the basis of convergent evolution of fleshy fruit ripening. *Nature Plants*, *4*(10), 784-791.
- Malnoy, M., Viola, R., Jung, M.-H., Koo, O.-J., Kim, S., Kim, J.-S., et al. (2016). DNA-free genetically edited grapevine and apple protoplast using CRISPR/Cas9 ribonucleoproteins. Front. Plant Sci. 7:1904. doi: 10.3389/fpls.2016.01904
- Malzahn, A., Lowder, L., and Qi, Y. (2017). Plant genome editing with TALEN and CRISPR. Cell Biosci. 7:21. doi: 10.1186/s13578-017-0148-4
- Martín-Pizarro, C., Triviño, J. C., and Posé, D. (2018). Functional Analysis of TM6 MADS Box Gene in the Octoploid Strawberry by CRISPR/Cas9 Directed Mutagenesis. bioRxiv [Preprint]. doi: 10.1101/351296
- McAtee PA, Richardson AC, Nieuwenhuizen NJ, Gunaseelan K, Hoong L, Chen X, et al. The hybrid non-ethylene and ethylene ripening response in kiwifruit (Actinidia chinensis) is associated with differential regulation of MADS-box transcription factors. BMC Plant Biology. (2015); 15:304. https://doi.org/10. 1186/s12870-015-0697-9 PMID: 26714876

- Medina-Puche, L. *et al.* MYB10 plays a major role in the regulation of flavonoid/phenylpropanoid metabolism during ripening of Fragaria x ananassa fruits. *J Exp Bot* **65**, 401–417 (2014).
- Merchante, C. *et al.* Ethylene is involved in strawberry fruit ripening in an organ-specific manner. *J Exp Bot* **64**, 4421–4439 (2013).
- Meyer, R. S., and Purugganan, M. D. (2013). Evolution of crop species: genetics of domestication and diversification. Nat. Rev. Genet. 14, 840–852. doi: 10.1038/ nrg3605
- Minas IS, Tanou G, Karagiannis E, Belghazi M, Molassiotis A. Coupling of Physiological and Proteomic Analysis to Understand the Ethylene- and Chilling-Induced Kiwifruit Ripening Syndrome. Frontiers in Plant Science. (2016); 7(120). https://doi.org/10.3389/fpls.2016.00120 PMID: 26913040
- Mingying, L. *et al.* Transcriptome sequencing and de novo analysis for bamboo using the illumina platform. *Plos one* **7**, 1–11 (2012).
- Molina-Hidalgo, F. J. *et al.* The strawberry (Fragaria x ananassa) fruit-specific rhamnogalacturonate lyase 1 (FaRGLyase1) gene encodes an enzyme involved in the degradation of cell-wall middle lamellae. *J Exp Bot* **64**, 1471–1483 (2013).
- Moyano-Cañete, E. *et al.* FaGAST2, a strawberry ripening-related gene, acts together with FaGAST1 to determine cell size of the fruit receptacle. *Plant Cell Physiol* **54**, 218–236 (2013).
- Mworia EG, Yoshikawa T, Salikon N, Oda C, Asiche WO, Yokotani N, et al. Low-temperature-modulated fruit ripening is independent of ethylene in 'Sanuki Gold' kiwifruit. Journal of Experimental Botany. (2012); 63(2):963–71. https://doi.org/10.1093/jxb/err324 PMID: 22058408
- Nakajima, I., Ban, Y., Azuma, A., Onoue, N., Moriguchi, T., Yamamoto, T., et al. (2017). CRISPR/Cas9mediated targeted mutagenesis in grape. PLoS One 12:e0177966. doi: 10.1371/journal.pone.0177966
- Nishitani, C., Hirai, N., Komori, S., Wada, M., Okada, K., Osakabe, K., et al. (2016). Efficient genome editing in apple using a CRISPR/Cas9 system. Sci. Rep. 6:31481. doi: 10.1038/srep31481
- Olubero Asiche W, Witere Mitalo O, Kasahara Y, Tosa Y, Gituma Mworia E, Ushijima K, et al. Effect of Storage Temperature on Fruit Ripening in Three Kiwifruit Cultivars. The Horticulture Journal. (2017); 86(3):403–10. <u>https://doi.org/10.2503/hortj.OKD-028</u>
- Ozden, O. *et al.* SIRT3 deacetylates and increases pyruvate dehydrogenase activity in cancer cells. *Free Radic Biol Med* **76**, 163–172 (2014).
- Paniagua, C. *et al.* Fruit softening and pectin disassembly: an overview of nanostructural pectin modifications assessed by atomic force microscopy. *Ann Bot* **114**, 1375–1383 (2014).
- Peng, A., Chen, S., Lei, T., Xu, L., He, Y., Wu, L., et al.

Fruit Ripening and Various Injuries: the Crosslink between Genetics and Phytohormones- A review

Salman Aloufi Fruit Ripening and Various Injuri (2017). Engineering cankerresistant plants through CRISPR/Cas9-targeted editing of the susceptibility gene CsLOB1promoter in citrus. Plant Biotechnol. J. 15, 1509–1519.

- Posé, S. *et al.* Insights into the effects of polygalacturonase FaPG1 gene silencing on pectin matrix disassembly, enhanced tissue integrity, and firmness in ripe strawberry fruits. *J Exp Bot* **64**, 3803–3815 (2013).
- Ren, C., Liu, X., Zhang, Z., Wang, Y., Duan, W., Li, S., et al. (2016). CRISPR/Cas9- mediated efficient targeted mutagenesis in Chardonnay (Vitis vinifera L.). Sci. Rep. 6:32289. doi: 10.1038/srep32289
- Ring, L. *et al.* Metabolic interaction between anthocyanin and lignin biosynthesis is associated with peroxidase FaPRX27 in strawberry fruit. *Plant Physiol* **163**, 43– 60 (2013).
- Rodríguez-Leal, D., Lemmon, Z. H., Man, J., Bartlett, M. E., and Lippman, Z. B. (2017). Engineering quantitative trait variation for crop improvement by genome editing. Cell 171, 470–480.e8. doi: 10.1016/j.cell.2017.08.030
- Ron, M., Kajala, K., Pauluzzi, G., Wang, D., Reynoso, M. A., Zumstein, K., et al. (2014). Hairy root transformation using Agrobacterium rhizogenes as a tool for exploring cell type-specific gene expression and function using tomato as a model. Plant Physiol. 166, 455–469. doi: 10.1104/pp.114.239392
- Salvatierra, A., Pimentel, P., Moya-León, M. A. & Herrera, R. Increased accumulation of anthocyanins in Fragaria chiloensis fruits by transient suppression of FcMYB1 gene. *Phytochemistry* **90**, 25–36 (2013).
- Sánchez-Sevilla, J. F., Vallarino, J. G., Osorio, S., Bombarely, A., Posé, D., Merchante, C., ... & Valpuesta, V. (2017). Gene expression atlas of fruit ripening and transcriptome assembly from RNA-seq data in octoploid strawberry (Fragariax ananassa). *Scientific reports*, 7(1), 1-13.
- Sato, S. et al. The tomato genome sequence provides insights into fleshy fruit evolution. Nature 485, 635–641 (2012).
- Schaart, J. G. *et al.* Identification and characterization of MYB-bHLH-WD40 regulatory complexes controlling proanthocyanidin biosynthesis in strawberry (Fragaria × ananassa) fruits. *New Phyto* **197**, 454– 467 (2013).
- Shen, Y. Y. & Rose, J. K. C. ABA Metabolism and Signaling in Fleshy Fruits. In: Abscisic Acid: Metabolism, Transport and Signaling (Zhang D. P., ed.), pp. 271–286, Springer Science + Business Media Dordrecht (2015).
- Sheng, X. & Liu, Y. Theoretical study of the catalytic mechanism of E1 subunit of pyruvate dehydrogenase multienzyme complex from Bacillus stearothermophilus. *Biochemistry* **52**, 8079–8093 (2013).
- Soyk, S., Müller, N. A., Park, S. J., Schmalenbach, I.,

Jiang, K., Hayama, R., et al. (2016). Variation in the flowering gene SELF PRUNING 5G promotes dayneutrality and early yield in tomato. Nat. Genet. 49, 162–168.

- Sun, J. H. *et al.* New evidence for the role of ethylene in strawberry fruit ripening. *Journal of Plant Growth Regulation* **32**, 461–470 (2012).
- Svitashev, S., Schwartz, C., Lenderts, B., Young, J. K., and Cigan, A. M. (2016). Genome editing in maize directed by CRISPR–Cas9 ribonucleoprotein complexes. Nat. Commun. 7, 1–7. doi: 10.1038/ncomms13274
- Symons, G. M., Chua, Y.-J., Ross, J. J., Quittenden, L. J., Davies, N. W., and Reid, J. B. (2012). Hormonal changes during non-climacteric ripening in strawberry. J. Exp. Bot. 63, 4741–4750. doi: 10.1093/jxb/ers147
- Thurman, R. É. et al. The accessible chromatin landscape of the human genome. Nature 489, 75–82 (2012).
- Tian, S., Jiang, L., Gao, Q., Zhang, J., Zong, M., Zhang, H., et al. (2016). Efficient CRISPR/Cas9-based gene knockout in watermelon. Plant Cell Rep. 36, 399– 406. doi: 10.1007/s00299-016-2089-5
- Tian, Y., Bai, S., Dang, Z., Hao, J., Zhang, J., & Hasi, A. (2019). Genome-wide identification and characterization of long non-coding RNAs involved in fruit ripening and the climacteric in Cucumis melo. *BMC plant biology*, *19*(1), 1-15.
- Wang, W. Q., Wang, J., Wu, Y. Y., Li, D. W., Allan, A. C., & Yin, X. R. (2020). Genome-wide analysis of coding and non-coding RNA reveals a conserved miR164-NAC regulatory pathway for fruit ripening. *New Phytologist*, 225(4), 1618-1634.
- Wang, Y., Xu, F., Feng, X. & MacArthur, R. L. Modulation of Actinidia arguta fruit ripening by three ethylene biosynthesis inhibitors. *Food Chem* **173**, 405–4013 (2015).
- Xiao, K., Chen, J., He, Q., Wang, Y., Shen, H., & Sun, L. (2020). DNA methylation is involved in the regulation of pepper fruit ripening and interacts with phytohormones. *Journal of experimental botany*, *71*(6), 1928-1942.
- Xiao-Jiao, H., Yang-Dong, W., Yi-Cun, C., Li-Yuan, L. & Qing-Ke, W. Transcriptome sequencing and expression analysis of terpenoid biosynthesis genes in Litsea cubeba. *Plos one* **8**, 1–14 (2013).
- Xu, X. *et al.* High-throughput sequencing and degradome analysis identify miRNAs and their targets involved in fruit senescence of Fragaria ananassa. *PLoS One* **8**, e70959 (2013).
- Yan, J. et al. Transcription factor binding in human cells occurs in dense clusters formed around cohesin anchor sites. Cell 154, 801–813 (2013).
- Yang, Y., Zhu, G., Li, R., Yan, S., Fu, D., Zhu, B., ... & Zhu, H. (2017). The RNA editing factor SIORRM4 is required for normal fruit ripening in tomato. *Plant Physiology*, *175*(4), 1690-1702.

- Yue, P., Lu, Q., Liu, Z., Lv, T., Li, X., Bu, H., ... & Wang, A. (2020). Auxin-activated MdARF5 induces the expression of ethylene biosynthetic genes to initiate apple fruit ripening. *New phytologist*, 226(6), 1781-1795.
- Zhang, Z., & Li, X. (2018). Genome-wide identification of AP2/ERF superfamily genes and their expression during fruit ripening of Chinese jujube. *Scientific reports*, *8*(1), 1-16.
- Zhong, S. et al. Single-base resolution methylomes of tomato fruit development reveal epigenome modifications associated with ripening. Nat. Biotechnol. 31, 154–159 (2013).
- Zhou, D., Shen, Y., Zhou, P., Fatima, M., Lin, J., Yue, J., ... & Ming, R. (2019). Papaya CpbHLH1/2 regulate carotenoid biosynthesis-related genes during papaya fruit ripening. *Horticulture research*, *6*(1), 1-13.
- Zorrilla-Fontanesi, Y. *et al.* Genetic analysis of strawberry fruit aroma and identification of O-methyltransferase FaOMT as the locus controlling natural variation in mesifurane content. *Plant Physiol* **159**, 851–870 (2012).