

# Maillard Browning: Pros and Cons in Dairy and Food Industries

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## **Abstract**

*The Maillard reaction, also known as non-enzymatic browning reaction between reducing sugars and amino groups. This reaction is responsible for the formation of several compounds, called MRP (Maillard Reaction Products) for food products. The Maillard reaction produces undesirable effects during the processing and storage of different liquid foods such as milk or fruit juices whereas for other solid foods the changes are favorable (in the case of bread, breakfast cereals, candies, coffee, chocolate, etc.). The Maillard reaction is a complex reaction, since it is influenced by many factors such as temperature, pH, time, water activity, type and concentration of reactant source and sugar involved. In heated food products, two carcinogenic compounds are produced during maillard reactions, acrylamide and imidazoquinoline. Furosine and HMF are two compounds that indicate the extent of the maillard reaction related to the type and intensity of the food processing conditions.*

**Keywords:** Maillard reaction, melanoidin, furfural, amadori rearrangement

**Abbreviations:** HMF- Hydroxy- methylfurfural; MRP –MaillardReacton Products; AGEs- Advanced glycation end products

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## **INTRODUCTION**

The Maillard reaction is of most important for quality of foods, in particular for heated foods. It induces browning of foods, has an effect on nutritive value, can have toxicological implications (such as the formation of acrylamide), can produce antioxidant components and it has also a large effect on flavor. Browning of food and food products is one of the major problems in food industry. Browning in a food occurs by oxidation and non oxidation reaction. Oxidative reactions involve enzymes (polyphenols oxidases) and oxygen; it is also called enzymatic browning, while non-oxidative reactions are called maillard reaction [1]. The Maillard reaction is named for L.C. Maillard, who conducted pioneering work on sugar-amino acid condensations from 1912 to 1917. He reported that upon gently heating sugars and amino acids in water, a yellow-brown color developed, called it a maillard reaction. The Maillard reaction (MR) or non-enzymatic glycation is there action of reducing sugars

with amino acids and amino groups of peptides or proteins. This reaction produces a variety of early, intermediate and advanced compounds. The advanced compounds of protein-sugar reactions are referred to as advanced glycation end products or AGEs and melanoidins (more concrete, melanoproteins) [2]. This reaction has become important in the field of food science and medicine [3]. The consumption of maillard Reaction Products (MRPs) has increased in recent decades and there are evidences that these substances are absorbed and may participate in pathological processes such as, cataract, diabetes, degenerative diseases, atherosclerosis and chronic renal failure [4]. Dairy powders are sensitive to the maillard reactions they contain high concentration of lactose and proteins with high lysine level [5]. On the other hand, these compounds are responsible for essential sensory attributes of thermally processed food products, contributing to their appearance, flavor, aroma and texture [6].

### Chemistry of the reaction

The Maillard reaction has been named after the French chemist Louis Maillard (1912) who first described it but it was only in 1953 that the first coherent scheme was put forward by Hodge. In essence, it states that in an early stage a reducing sugar, like glucose, condenses with a compound possessing a free amino group (of an amino acid or in proteins mainly the  $\epsilon$ - amino group of lysine, but also the  $\alpha$ -amino groups of terminal amino acids) to give a condensation product N-substituted glycosilamine, which rearranges to form the Amadori rearrangement product (ARP). The subsequent degradation of the Amadori product is dependent on the pH of the system. At pH 7 or below, it undergoes mainly 1, 2-enolisation with the formation of furfural (when pentoses are involved) or hydroxymethylfurfural (HMF) (when hexoses are involved). At pH >7 the degradation of the Amadori compound is thought to involve mainly 2, 3 enolisation, where reductones, such as 4-hydroxy-5-methyl-2,3-dihydrofuran-3-one (HMFone), and a variety of fission products, including acetol, pyruvaldehyde and diacetyl are formed. All these compounds are highly reactive and take part in further reactions. Carbonyl groups can condense with free amino groups, which results in the incorporation of nitrogen into the reaction products. Dicarbonyl compounds will react with amino acids with the formation of aldehydes and  $\alpha$ -aminoketones. This reaction is known as the Strecker degradation. Subsequently, in an advanced stage, a range of reactions takes place, including cyclisations, dehydrations, retroaldolisations, rearrangements, isomerisations and further condensations, which ultimately, in a final stage, lead to the formation of brown nitrogenous polymers and copolymers, known as melanoidins. The complexity and the variety of the maillard reaction products have, throughout the years, raised the interest of scientists in different fields of research [7]. New important pathways, not accounted for by Hodge, have been established. In maillard browning most important intermediates in color formation are 3-deoxyosuloses and 3,4-dideoxyosulos-3-enes, which in the case of glucose is 3-deoxyhexosulose (DH) and 3,4-dideoxyhexosuloses-3-ene (DDH) [8]. 3-deoxy-2-hexosuloses, 1- deoxy-2, 3-hexodiuloses and other -dicarbonyl

intermediates can undergo nucleophilic addition reactions with amino acids with subsequent decarboxylation to produce the so called Strecker aldehyde (RHC=O). In [9] Huber and Ledl isolated and characterized 1-deoxy- and 3-deoxyglucosones from heated Amadori products. More recently, in agreement with the previous reports, Tressl, Nittka and Kersten [10] using  $^{13}\text{C}$ -labeled sugars, have given a new perspective to the reaction mechanism. It involves different reaction pathways, in which the key intermediates are the 1-, 3- and 4-deoxyhexosuloses. Moreover, a major influence of the pH is expressed. Along with enolization reactions, the Amadori product and its dicarbonyl derivatives can undergo concurrently retro-aldol reactions producing more reactive C2, C3, C4 and C5 sugar fragments, such as hydroxyacetone derivatives, glyceraldehyde and diketones. Retroaldol reactions become more important at higher pH values. Also, Yaylayan and Huyghues [11] stated that under basic conditions, ARP could generate acetic acid and pyruvaldehyde and other lower sugars in addition to free amino acid. As a result, high pH is suggested to be the main pathway to flavor formation. In addition to retroaldol reactions, three redox mechanisms have been identified, in which hydroxycarbonyls; dicarbonyls and formic acid are involved. Berg and Van Boekel [12] reported formic acid as a main degradation reaction product for the maillard reaction of lactose. Also, Boekel [13] reported formic acid and acetic acid as two main degradation products for the maillard reaction of glucose and fructose. Berg [14] concluded that isomerization and degradation reactions of the sugar are, from a quantitative point of view, more important than the maillard reaction, for conditions as in heated milk. Finally, the central importance of the Amadori product, formerly supposed to be the main intermediate of the reaction, has been questioned in both food [15] and medical fields [16]. The Maillard reaction is influenced by many factors, including temperature, time, initial pH, water activity ( $a_w$ ), physical state of the matrix, reactant concentration and type of carbohydrate involved [17]. In spite of all the work that has been done, the mechanism of the Maillard reaction is still a controversial issue. Mechanism of maillard reaction is shown in Figure 1.

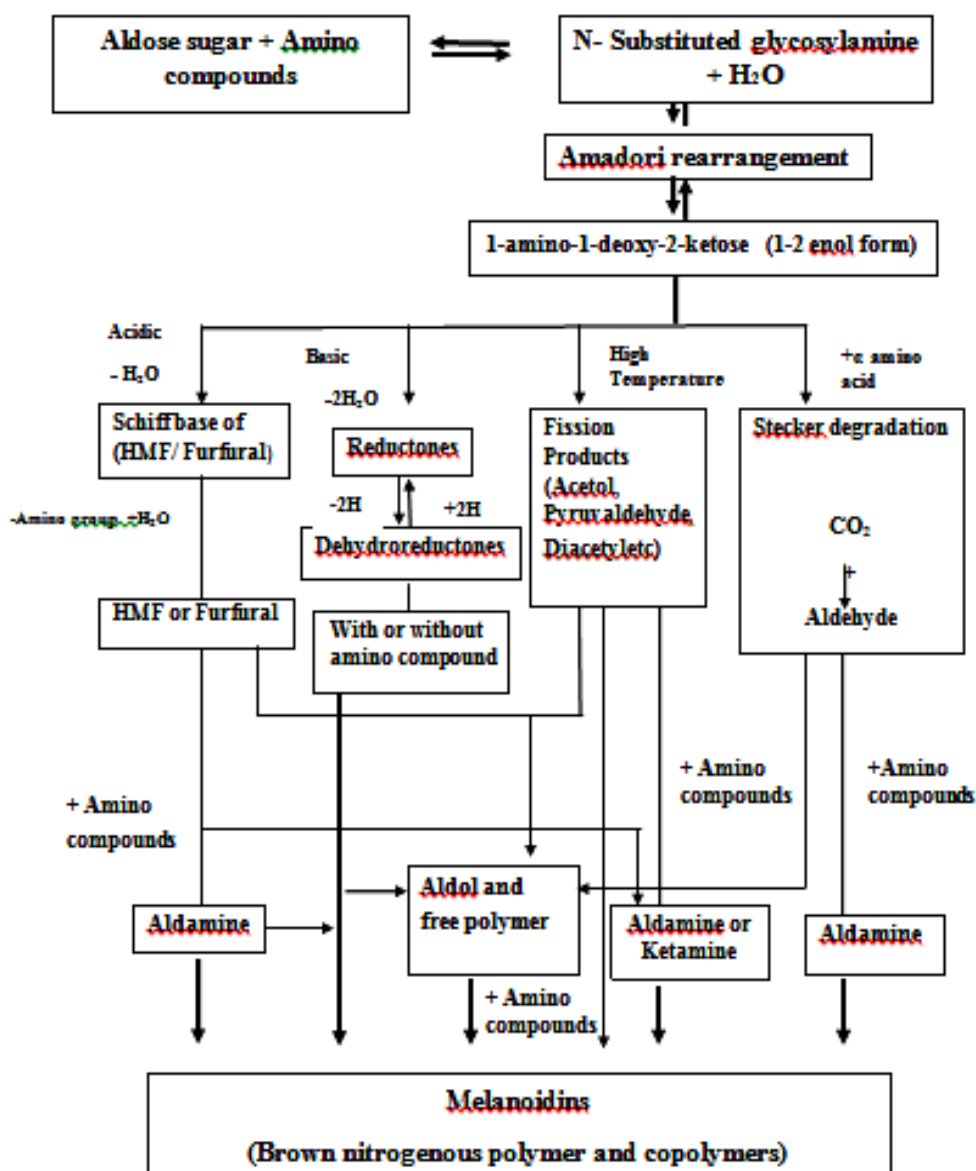


Fig. 1: Mechanism of Maillard Reaction Adapted from Hodge [1].

### Pros of Maillard Reaction in Foods

Maillard reaction occurs during many industrial and domestic thermal treatments of foods. It is widely used because of its role in creating colors, flavors, textures and other functional properties in foodstuffs. Proteins glycosylated without the use of conventional chemical reagents have demonstrated to improve techno-functional properties as heat stability, emulsifying and foaming properties. It has been a central and major challenge in food industry, since the maillard reaction is related to aroma, taste and color, in particularly in traditional processes such as the roasting of coffee and cocoa beans, the baking

of bread and cakes, the toasting of cereals and the cooking of meat [18].

Now-a-days maillard reactions products are proved to have antioxidant properties as well. These products also use to improve the properties of protein as polysaccharide - protein conjugate formed during the maillard reaction have better property than the native one [19].

### Color Formation

Color formation is the primary characteristic of the Maillard reaction. Brown color development during processing and storage is desirable for many products such as baked

foods, coffee, cookies while undesirable in some kinds of food products orange juice, white chocolate, milk and powder egg [20]. Color occurs due to the formation of high molecular weight (>12,000 Daltons) polymeric compounds also known as melanoidins. These are generally formed by the reaction of the Amadori product or/and with other dicarbonyls, i.e., the deoxyosuloses, with amino acids [21]. Predicting and controlling food color development are particularly important for companies to satisfy consumer preference, since a complex array of melanoidins produced by the maillard reaction is strongly dependent on the food matrix composition as well as the technological conditions of the reaction [22]. Melanoidin compound formed by the reaction of amadori product and with other dicarbonyl that is deoxyosuloses with amino acid is a reason for characteristic color of roasted foods such as coffee, cocoa, bread and malt. Color development depends on the reactant, structure of melanoidin, structure of chromatographic moiety and method of isolation of melanoidins [23]. It is very hard to separate the individual compound and structure of melanoidin, due to its complex nature. The composition of melanoidin depends on the binding of amino groups of lysine that bound to proteins. In the food products there are heterogeneous mixtures of food products, which have a different maximum absorbance due to which color formation takes place. In general sugar to amino acid is 1:1 for pentoses and 1.5:1 for glucose [24]. Although the chemical structures and health effects of these compounds produced both in food and model systems have been investigated.

### Flavor and Aroma

Flavor and aroma development due to the maillard reaction depends on the reaction temperature, time, pH, water content and on the type of sugars and amino acids involved [25]. Mostly reaction temperature influences the kinetics parameters, and time determines the type of flavor compounds formed. The intermediate and final stages of the maillard reaction are the most important in flavor development [26]. Maillard reaction between different carbohydrate and proteins gives different end products that produce versatile flavors. The volatile products of the maillard reaction depend on kind of sugar that

undergoes dehydration/fragmentation and produces furans, pyrones, cyclopentenones, carbonyls and acids. The amino acid degradation during maillard reaction produces aldehydes and sulfur compounds; and upon further interactions they create pyrroles, pyridines, imidazoles, pyrazines, oxazoles, thiazoles, and others. All of these compounds are volatiles and associated with flavor production. The amino acid proline gives rise to typical bread, rice and popcorn flavors while other sulfur containing amino acids (methionine), generate a highly intense smell of potatoes (Gu). Cysteine and glucose generally produce sulfur compounds, but in oxidized condition they synthesize pyrazine and furan, while cysteine when reacts with ribose gives 2-methyl-3-furanthiol, an important flavor contributor in meat. Glutathione and glucose produce thiophene, thiazole and cyclic polysulfites at pH 6 and 8 while furan at acidic pH. Pyrazines and alkyl pyrazines are associated with the flavor and aroma of cooked (roasted) and nutty, respectively. Alkyl pyridines add green, bitter, astringent and burnt flavor, while furans, furanones and pyranones impart sweet, burnt, pungent and caramel-like flavors [27]. Maillard reaction mechanism is to develop a sweet, caramel-type flavor in the final whipped cream product [28]. The flavor compounds developed in the advanced stage of the maillard reaction likely contribute to the characteristic flavor of UHT processed milk [29]. Sulfur containing maillard odorants constitute the most powerful aroma compounds and often play, although at trace levels, a dominant role in the flavor of cooked meats. These volatile compounds are responsible for the flavor and aroma of stewed beef juice, boiled trout, French fries, bread crust, cooked chicken, roasted chicken, boiled beef, cocoa powder, peanuts, pilsner, roasted beef, popcorn and coffee [30]. The intermediate and final stages of the maillard reaction are the most important to flavor development, especially the so-called Strecker degradation step, in which amino acids are degraded by dicarbonyls formed previously in the reaction, leading to the amino acids deamination and decarboxylation [31]. The oxygen-containing aroma compounds 2,3-butanedione, 2,3-pentanedione, methylpropanal, 3-methylbutanal, phenylacetaldehyde, 3-hydroxy-4,5-dimethyl-

2(3H) furanone and 2,5-dimethyl- 4-hydroxy-3(2h)-furanone occur in concentration ranging from 1µg/kg up to 100 mg/kg. The nitrogen-containing aroma compounds 2-ethyl-3,5dimethylpyrazine, 2,3-diethyl-5-methylpyrazine and 2-acetyl-1- pyrroline are present in food in an order of magnitude of 0.001–10 mg/kg. On the whole, sulfur containing maillard odorants constitute the most powerful aroma compounds and often play, although at trace levels, a dominant role in the flavor of cooked meats. These volatile compounds are responsible for the flavor and aroma to stewed beef juice, boiled trout, French fries, bread crust, cooked chicken, roasted chicken, boiled beef, cocoa powder, peanuts, pilsner, roasted beef, popcorn and coffee. Mixtures containing amino acids other than cysteine or methionine in combination with reducing sugars are characterized mostly by caramel and jammy smell [32]. The food industry invests great effort trying to create synthetic flavors and aromas by reconstituting combinations of these compounds. The process of creating synthetic flavors is limited since the subtleties of flavor perception are many and varied, and although electronic noses may detect these compounds, human sensory perception is considered essential to validate instrumental data [33].

### Texture

Texture is complex sensory attribute of structure, mechanical and surface properties of food that is judge through the senses of vision, hearing, and touch, acceptability of food mainly bakery products depends on texture. According to Szczesniak [34] texture is the sensory and functional manifestation of the structural, mechanical and surface properties of foods detected through the senses of vision, hearing, touch and kinesthetic. Maillard reaction influences the texture of food via protein cross-linking and its extent and nature. Functional properties of foods can be manipulated by altering the extent and nature of these cross linking. This protein cross linking by maillard reaction also improves their digestibility [35]. Maillard reaction influences the texture of food via protein cross-linking. Manipulation of the extent and nature of such protein cross-linking during food processing offers a means by which the

food industry can modify the functional properties of food [36].

But the extent of how much protein cross-linking affects food texture in processed foods and how to control this parameter to maximize food quality is not yet known [37]. Protein cross-linking by the maillard reaction will affect not only texture, but the protein digestibility as well.

Although maillard reaction effects on food color, flavor and aroma are well understood and used by the food industry, its effects on food texture has attracted less attention from the scientific community. However, this is a promising tool for texture development.

### Antioxidant Property

In the literature antioxidant activity of maillard reaction products have been noticed. Maillard reactions products like amino reductone, heterocyclic compounds, or high molecular melanoid have antioxidant properties. MRPs prepared from aqueous ribose-lysine and fructose lysine show antioxidant activity, they have radical chain breaking activity, metal chelation, and decomposition of hydrogen peroxide and scavenging of oxygen species [38]. Due to different reaction conditions and reactant reactivity MRPs exhibit varying antioxidant activity [39]. Whey protein isolates and sugar react to form maillard reaction products have good antioxidant activity. This antioxidant property can be used to protect the lipid oxidation in food products. Maillard reaction products prepared by heating mixture with higher initial pH showed higher antioxidant activity [40].

### Improve Functional Properties of Protein

Proteins have wide application in food industry because of their functional properties. Their functional properties depend on physicochemical and intrinsic structure. Maillard reaction was used to improve the functional properties like heat stability, emulsifying foaming stability of  $\beta$ -lactoglobulinglycated with several sugars (arabinose, galactose, glucose, lactose). The functional properties of proteins can be modified by chemical, physical and enzymatic treatments. But these treatments also modify the ability of proteins to form gel, water

retention, solubility, foaming and emulsifying capacity [41]. Application of maillard reaction improve the functional properties of protein is the new era of research. In maillard reaction, protein interacts with polysaccharide to form protein - polysaccharide conjugates which have different properties. These properties depend on both protein conformation and polysaccharide e characteristic such as hydrophobicity, viscosity, chain length and number of joints [42].

This protein polysaccharide conjugates have better emulsifying, foaming property, protein solubility and heat stability then their native proteins. Various studies also revealed that maillard reaction improve the gelation properties of protein-polysaccharide conjugates system, especially gel mechanical property [43].

#### Cons of Maillard Browning Reaction

Maillard reaction products not only have beneficial effect but some of maillard reaction products also have harmful effect on human health. The most common harmful effect of maillard reaction products are:

1. Protein nutritional impairment [44].
2. Stale or “cardboard” flavors have also been reported in different types of dried dairy ingredients, including milk powders due to the maillard reaction [45].  
Nutritional damage in lactose-hydrolysed milk by maillard reaction was reported in liquid milk [46].
3. Loss of nutritive value due to blockage of lysine residues which are no longer available for digestion (early, advanced and final maillard reaction reduced digestibility and inhibition of enzymes [47].
4. Destruction of essential amino acids and reduction in bioavailability: - Amino acid lysine is more prone to destruction during maillard reaction especially in breakfast cereals near about 20–54% lysine undergoes destruction during processing [48].
5. Decrease in bioavailability of minerals: - MRPs chelates the minerals, interfere with their solubility and absorption in guts, result in reducing the bioavailability of minerals (calcium, iron, magnesium and dietary phosphorus) by chelating them and interfere with their solubility [49].
6. Exhibit mutagenic activity: - Maillard reaction product exhibit mutagenic activity. Cysteamine a decarboxylated compound of cysteine that is formed during maillard reaction exhibit mutagenic activity, also sometime it is further degrades to thiazolidine which also have mutagenic activity. Various products form during maillard reactions like 1,2 carbonly compound (diacetyl, maltol), dihydroxyacetone, glyceraldehydes, glyoxal, methyl glyoxal and glyoxylic acid as proved to have mutagenic activity [50] caramelized sugar can induce high frequency chromosomal breakage. Maillard products from ketose sugar exhibit higher mutagenic activity as compared to aldose isomer due to difference in reaction mechanism [51].
7. Produces off flavor: - Maillard browning reaction products like furfural, furane, pyrole, pyrazine produces off flavor. These compounds also responsible for altering the milk flavor [52].
8. Other harmful effects: - Maillard reaction product, 5- hydroxymethylfurfural (HMF) is considered a potentially carcinogenic to human [53]. The formation of advanced glycation end products (AGEs) is observed during normal ageing and occurs inside as well as outside of cells [54]. These compounds, when cross linking with proteins profoundly affect protein functionality and irreversibly modify chemical properties and functions of diverse biological structures [55], which seems to be implicated in inflammatory processes and diabetic complications, such as nephropathy and vascular disease [56, 57]. AGEs accumulate in various tissues during aging, including skin, neural, vascular, renal and cardiac tissues, collagens and crystalline lens. In the skin, glycation is involved in many metabolic processes and, along with aging, affects the functionality of certain cells, such as the synthesis of fibroblasts, enzyme activation of matrix degradation (metalloproteinases) and the organization of the matrix [58]. It is proposed that the accumulation of the advanced glycation end products (AGEs). The receptor for AGEs in the retina could play a significant role in the initiation and progression of age-related macular degeneration and

cataracts [59]. Mostafa et al. [60] showed that AGEs level in plasma proteins are elevated in patients with diabetes. The high blood glucose levels favor the occurrence of spontaneous reactions (glycation) between glucose and proteins, resulting in the formation and excessive deposition of AGEs [61]. In patients with renal failure AGEs accumulation occurs due to the decrease in the extent of degradation and elimination from the body and, also, to increased exposure to oxidative stress. On the other hand, the AGEs and products derived from the process of oxidation promote damage in the renal tissue, leading to greater accumulation of AGEs, creating a vicious cycle.

## CONCLUSION

Coping with the maillard reaction food and the effects of the reaction products on health is important to the improvement and development of food products. The maillard reaction has positive as well as negative aspects in food industry. The positive contributions of the maillard reaction are sensory attributes generation, such as color, flavor, aroma and texture. The negative aspects are off-flavor development, flavor loss, discoloration, and loss of protein nutritional value. In the food industry, the role of flavor and color either desirable or undesirable is the key in the manufacture of products of consistent sensory quality. Contradictory knowledge about the effects of maillard reaction products on health indicates that studies are needed to further expand the AGEs and MRPs database as well as development of methods for reducing MRPs generation during home cooking and food processing. Understanding the chemical, nutritional and toxicological consequences of browning reactions and related transformations, *in vitro* and *in vivo*, can lead to better and safer foods and improved human health.

## REFERENCES

1. Hodge J. Chemistry of browning reactions in model systems. *Journal of Agriculture and Food Chemistry* 1953; 1(15):928–943p.
2. Ames J. M. Applications of the Maillard reaction in the food industry. *Journal of Food Chemistry* 1998; 62:431–439p.
3. Finot P. Historical Perspective of the Maillard Reaction in Food Science. *Ann. N.Y. Acad. Science* 2005; 1043(1): 1–8p.
4. Wang H. Y., Qian H., Yao W. R. Melanoidins Produced by the Maillard Reaction: Structure and Biological Activity. *Journal of Food Chemistry* 2011; 128(3): 573–584p.
5. Palombo R., Gertler A., Saguy I. A simplified method for determination of browning in dairy powders. *Journal of Food Science* 1984; 49: 1609–1613p.
6. Gerrard J. New Aspects of Ageing Chemistry. Recent Developments Concerning the Maillard Reaction. *Australian Journal of Chemistry* 2002; 55(5):299–310p.
7. Ericksson C. *Maillard Reaction in Food: Chemical, Physiological and Technological Aspects in Progress in Food and Nutrition Science* 5, Pergamon Press: 1981.
8. D. J. McWeeny, M. E. Knowles, J. F. Hearne. The Chemistry of Non-enzymic Browning in Foods and its Control by Sulphites. *Journal of the Science of Food and Agriculture* 1974; 25 (6):735–746p.
9. Huber B., Ledl F. Formation of 1-amino-1,4-dideoxy-2,3-hexodiulose and 2-aminoacetylfurans in the Maillard Reaction. *Journal of Carbohydrate Research* 1981; 215–220p.
10. Tressl R., Nittka C., Kersten E. Formation of Isoleucine-specific Maillard Products from [D-glucose and [1-13C]-D-fructose]. *Journal of Agriculture and Food Chemistry* 1995; 43:1163–1169p.
11. Yaylayan V.A., Huyghues, D. Retro-aldol and Redox Reactions of Amadori Compounds: Mechanistic Studies with Various Labeled D-[13C] Glucose. *Journal of Agriculture and Food Chemistry* 1996; 44:672–681p.
12. Berg, H.E. Reactions of Lactose during Heat Treatment of Milk: A Quantitative Study, PhD thesis, Wageningen University, NL 1993.
13. Van Boekel, M. A. Formation of Flavor Compounds in the Maillard Reaction. *Biotechnology Advances* 2006; 24(2):230–233p.

14. Berg H.E., Van Boekel, M.A.J.S. Degradation of Lactose during Heating of Milk. Reactions Pathway Netherlands. *Milk and Dairy Journal* 1994; 48:157–175p.
15. Molero-Vilchez, M.D., Wedzicha B.L. A New Approach to Study the Significance of Amadori Compounds in the Maillard Reaction. *Journal of Food Chemistry* 1997; 58:249–254p.
16. Fu M. et al. Glycation, Glycooxidation, and Cross-linking of Collagen by Glucose: Kinetics, Mechanisms, and Inhibition of Late Stages of the Maillard reaction<sup>1</sup> Diabetes 1994; 43:676–683p.
17. Buza T. P., Baisier, W. M. In H. G. Schwartzberg & R. W. Hartel (Eds.), *The Kinetics of Nonenzymatic Browning Physical Chemistry of Foods* New York, USA: Marcel Dekker Inc. 1992; 595–649p.
18. Cristina Delgado-Andrade, Isabel Seiquer, Ana Haro et al. Development of the Maillard Reaction in Foods Cooked by Different Techniques. Intake of Maillard-derived Compounds. *Journal of Food Chemistry* 2010; 122:145–153p.
19. Foerster A., Henle, T. Glycation in Food and Metabolic Transit of Dietary AGEs (advanced glycation end-products): Studies on the Urinary Excretion Pyrraline. *Biochemical Society Transactions* 2003; 31:1383–1385p.
20. Ramírez-Jiménez, A., García-Villanova, B., Guerra-Hernández, E. Hydroxymethylfurfural and Methylfurfural Content of Selected Bakery Products. *Food Research International* 2000; 33:833–838p.
21. Wang He-Ya., Qian H., Yao, Wei-Rong. Melanoidins Produced by the Maillard Reaction: Structure and Biological Activity. *Journal of Food Chemistry* 2011; 128(3):573–584p.
22. Lievonen S. M., Roos Y. H. Nonenzymatic Browning in Amorphous Food Models: Effects of Glass Transition and Water. *Journal of Food Science* 2002; 67: 2100–2106p.
23. Del Castillo, M. D., Ames, J. M., et al. Effects of Roasting on the Antioxidant Activity of Coffee Brews. *Journal of Agriculture and Food Chemistry* 2002; 50: 3698–3703p.
24. Yu Ai-Nong & Zhang, Ai-Dong. The Effect of pH on the Formation of Aroma Compounds Produced by Heating a Model System Containing L-ascorbic Acid with L-threonine/L-serine. *Food Chemistry* 2010; 119(1):214–219p, ISSN 0308- 8146.
25. Van Boekel, M. A. Formation of Flavor Compounds in the Maillard reaction. *Biotechnology Advances* 2006; 24(2): 230–233p. ISSN 0734-9750.
26. Rizzi G. The Strecker Degradation of Amino Acids: Newer Avenues for Flavor Formation. *Food Reviews International* 2008; 24: 416–435p.
27. Cerny C. The Aroma Side of the Maillard Reaction. *Annals of the New York Academy of Sciences* 2008; 1126(1):66–71p.
28. Davies C., Labuza, T. *The Maillard Reaction: Application to Confectionery Products*. In: Zeigler G, editor. Confectionary science. Pennsylvania: Penn State Univ. Press. 1997; 35–66p.
29. Basta G. Schmidt, A.M., Caterina R., Advanced Glycation End Products and Vascular Inflammation: Implications for Accelerated Atherosclerosis in Diabetes. *Cardiovascular Research* 2004; 63(4): 582– 592p.
30. Wagner K., Reichhold S., Koschutnig K., et al. The Potential Antimutagenic and Antioxidant Effects of Maillard Reaction Products used as "Natural Antibrowning" Agents. *Molecular Nutrition & Food Research* 2007; 51(4): 495 – 504p.
31. Somoza V. Five Years of Research on Health Risks and Benefits of Maillard Reaction Products. *Molecular Nutrition & Food Research* 2005; 49(7):663– 672p.
32. Szczesniak A. S. Texture is a Sensory Property. *Food Quality and Preference* 2002; 13(4): 215–225p.
33. Rufián-Henares, J., Delgado-Andrade, C. Effect of Digestive Process on Maillard Reaction Indexes and Antioxidante Properties of Breakfast Cereals. *Food Research International* 2009; 42 (3): 394–400p.
34. Antonio D., Troisea, S., Vincenzo F. Encapsulation and Maillard Reaction. *Trends in Food Science & Technology* 2013; 30: 127–132p.
35. Gerrard J. A. Protein-Protein Crosslinking in Food: Methods, Consequences,



- Applications. *Trends Food Science and Technology* 2002; 13(12):391– 399p.
36. Gu F.L. *et al.* Structure and Antioxidant Activity of High Molecular Weight Maillard Reaction Product from Casein-Glucose. *Journal of Food Chemistry* 2010; 120:505–511p.
37. Vhangani L.N., Wyk J.V. Antioxidant Activity of Maillard Reaction Products Derived from Fructose-lysine and Ribose-lysine Model System. *Food Chemistry* 2013; 137:92–98p.
38. Sun Q., Luo Y. Effect of Maillard Reaction Conditions on Radical Scavenging Activity of Porcine Haemoglobin Hydrolysate-Sugar Model System. *International Journal of Food Science and Technology* 2011; 46(2):358–364p.
39. Morales F. J., Van Boekel M. A. J. S. A Study on Advanced Maillard Reaction in Heated Casein/Sugar Solutions: Fluorescence Accumulation. *International Dairy Journal* 1997; 7:675–683p.
40. Delgado-Andrade C., Seiquer I., García M.M., *et al.* Increased Intake of Maillard Reaction Products Reduces Phosphorous Digestibility in Male Adolescents. *Journal of Nutrition and Dietetics* 2011; 27(1):86–91p.
41. Rutherford S.M., Torbatinejad N.M., Moughan P.J. Available (ileal digestible reactive) Lysine in Selected Cereal-based Food Products. *Journal of Agriculture and Food Chemistry* 2007; 54(25): 9453–7p.
42. Rerat A. *et al.* Nutritional and Metabolic Consequences of the Early Maillard Reaction of Heat Treated Milk in the Pig. Significance for man. *European Journal of Nutrition* 2002; 4(1): 1–11p.
43. Palombo R., Gertler A., Saguy I. A Simplified Method for Determination of Browning in Dairy Powders. *Journal of Food Science*.1984; 49:1609–1613p.
44. Evangelisti F., *et al.* Deterioration of Protein Fraction by Maillard Reaction in Dietetic Milks. *Journal of Dairy Research* 1999; 66: 237–243p.
45. Booth K., *et al.* Kinetic Studies of Formation of Antigenic Advanced Glycation End Products (AGEs). *Journal of Biological Chemistry* 1997; 272: 5430–5437p.
46. Torbatinejad N.M., Rutherford S.M., Moughan P.J. Total and Reactive Lysine Contents in Selected Cereal-Based Food Products. *Journal of Agricultural and Food Chemistry* 2005; 5(11): 4454–4458p.
47. Delgado-Andrade C., Seiquer I., García M.M., *et al.* Increased Intake of Maillard Reaction Products Reduces Phosphorous Digestibility in Male Adolescents. *Journal of Nutrition* 2011; 27(1): 86–91p.
48. Bjeldanes L. F., Chew H. Mutagenicity of 1,2-dicarbonyl Compounds: Maltol, Kojic Acid, Diacetyl, and Related Substances. *J.Mutat. Res.*1979; 67: 367–371p.
49. Purlis E. Browning Development in Bakery Products A Review. *Journal of Food Engineering* 2010; 99(3): 239–249p.
50. Fox & McSweeney. *Advanced Dairy Chemistry: Lactose, Water, Salts and Minor Constituents.*1998; 3: 13–14p.
51. Capuano E., Fogliano V. Acrylamide and 5-hydroxymethylfurfural (HMF): A Review on Metabolism, Toxicity, Occurrence in Food and Mitigation Strategies. *Journal of Food Science and Technology* 2011; 44(4): 793 –810p, ISSN 1082-0132.
52. García M.M., Seiquer I., Navarro M.P. Influence of Diets Rich in Maillard Reaction Products on Calcium Bioavailability. Assays in Male Adolescents and in Caco-2 Cells. *Journal of Agricultural and Food Chemistry* 2009; 57:9532–9538p.
53. Barbosa J.H.P., Oliveira S.L., Seara L.T. The Role of Advanced Glycation End-Products (AGEs) in the Development of Vascular Diabetic Complications. *Brazilian Archives of Endocrinology and Metabolism* 2008; 52(6): 940–950p.
54. Jakus V., Rietbrock N. Advanced Glycation End-Products and the Progress of Diabetic Vascular Complications. *J. Physiol. Res.* 2004; 53(2):131–142p.
55. Linden E. *et al.* Endothelial Dysfunction in Patients with Chronic Kidney Disease Results from Advanced Glycation End Products (AGE)-Mediated Inhibition of Endothelial Nitric Oxide Synthase through RAGE Activation. *Clinical Journal of American Society Nephrol* 2008; 3(3): 691–698p.
56. Hartog J.H.W., Voors A.A., Bakker S.J.L., *et al.* Advanced Glycation End-products

- (AGEs) and Heart Failure: Pathophysiology and Clinical Implications. *European Journal of Heart Failure* 2007; 9(12):1146–1155p.
57. Nguyen C. Toxicity of the AGEs Generated from the Maillard Reaction: On the Relationship of Food-AGEs and Biological-AGEs. *Molecular Nutrition & Food Research* 2006; 50(12):1140–1149p.
58. Mostafa A.A., et al. Plasma Protein Advanced Glycation End Products, Carboxymethyl Cysteine, and Carboxyethyl Cysteine, are Elevated and Related to Nephropathy in Patients with Diabetes. *Molecular Cell Biochemistry* 2007; 302(1):35–42p.
59. Mahmoud E., et al. Comparison between the Impact of Two Different Exercise Protocols on Advanced Glycation End Products in Type 2 Diabetic Rats. *Journal of Life Science* 2013; 10(3).
60. Hartog H., et al. Advanced Glycation End-products (AGEs) and Heart Failure and Pathophysiology and Clinical Implications. *European Journal of Heart Failure* 2007; 9(12):1146–1155p.
61. Monti S. M., et al. LC/MS Analysis and Antioxidative Efficiency of Maillard Reaction Products from a Lactose–lysine Model System. *Journal of Agricultural and Food Chemistry* 1999; 47:1506–1513p.