

Quality attributes of fermented beverages from rose (*Rosa species*) and cinnamon (*Cinnamomum verum* J. Presl)

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ABSTRACT

An investigation was carried out to examine the quality attributes of fermented beverages obtained from the combination of rose petals and cinnamon powder. The experiment consisted of 12 treatments and 3 replications in CRD design. The treatment T₁ and T₂ were made of 100% rose and 100% cinnamon each. The other treatments consists of; T₃ (99% rose + 1% cinnamon), T₄ (98% rose + 2% cinnamon), T₅ (97% rose + 3% cinnamon), T₆ (96% rose + 4% cinnamon), T₇ (95% rose + 5% cinnamon), T₈ (94% rose + 6% cinnamon), T₉ (93% rose + 7% cinnamon), T₁₀ (92% rose + 8% cinnamon), T₁₁ (91% rose + 9% cinnamon), T₁₂ (90% rose + 10% cinnamon) respectively. The yeast *Saccharomyces cerevisiae* was used for fermentation. The maximum nutritional values in freshly fermented beverages were; phenol (199.76 GAE/100 ml), anthocyanin (7.00 mg/100 ml), alcohol (12.41%), ascorbic acid (5.46 mg/100 ml), reducing sugar (1.58%), non-reducing sugar (8.00%), total sugar (1.99%) and TSS (19.67 °B). The treatment with 100% cinnamon showed strong antimicrobial properties throughout experimentation. However, the proportion of cinnamon added to the beverages significantly improved the nutritional value of the beverages. With the storage period, phenol, ascorbic acid, and alcohol content in the beverages increased whereas other parameters were found to be fluctuating during storage. The beverages obtained from the combination of rose petal and cinnamon powders contained a high amount of vitamins and antioxidants and were highly acceptable.

Key words : Alcoholic beverages, Cinnamon, Fermented beverages, Flower fermentation, Wine

Introduction

Fermentation, a metabolic process in which carbohydrates rich substrates are converted into alcohol, carbonic acid and carbon dioxide with the help of microorganisms such as yeast and bacteria anaerobically is one of the oldest, simplest and cheapest techniques adopted by man for the preservation of food since ancient times, which is still prevalent in different cultures and regions. The yeast *Saccharomyces*

cerevisiae is mostly used which predominantly controls the flavor of the final products of the fermented products (Joshi *et al.*, 2009). Fermentation process enhances the nutritional and functional properties of substrate and formation of bioactive and bioavailable end products due to transformation. Fermented food provides many health benefits such as antioxidants, antimicrobial, antifungal, anti-inflammatory, antidiabetics and atherosclerotic activity (Sanlier *et al.* 2017), anticancer, anti-HIV

(Almeida *et al.* 2011) reduce anti-nutritional components (Onkuwa *et al.* 2007). Besides, fermentation is used for bio-economy, green chemistry, which gives us important products like medicines, bioethanol, antibiotics, plant growth regulators etc. According to archaeological evidences, fermentation method is as old as 7000-6000 B.C in Jiahu China, 6000 in Georgia and 5000 in India.

Fermentation is commonly done with carbohydrate-rich organic substrates mostly cereals or fruits. However, there are about 30,000 flower species in the world that are reported to be edible and many are rich in fermentable sugar, phytochemicals, and bioactive compounds (Shaheen *et al.*, 2017). Flowers are particularly rich in antioxidants; phenol, flavonol, flavonoids, carotenoids, anthocyanin, etc. which mitigates the excess oxidative stress. The consumption of flowers has been in practice since ancient times in Rome, Greece, and China for thousands of years. They used flowers to add aesthetic looks to various dishes and to enhance the organoleptic synergy between traditional foods and flower aromas (Benvenuti and Mazzoncini, 2020). Rose flower is one such, which has been used since time immemorial for various purposes. A woody perennial plants native to Asia, Europe and Africa is valued for its effective medicinal properties which were first documented in Sumerians and Chinese. The rose petals have been consumed in many cultures for many years especially as jam (Jat *et al.*, 2018, Butcaru *et al.*, 2017) teas, cakes, flavor extracts and used in medicinal practices for remedy of various illness (Friedman *et al.*, 2007). Rose petal extracts act as antioxidants (Kart and Cagindi, 2017), anticancer (Tatke *et al.*, 2015), neuroprotective (Esfandiary *et al.*, 2015), antidepressant (Tirupati and Golla, 2016), anti-HIV (Gao *et al.*, 2013), anticonvulsant (Homayoun *et al.*, 2015), antidiabetic (Ju *et al.*, 2014), antimicrobial (Swati *et al.*, 2016; Laxmi *et al.*, 2017), and hepatoprotective (Achutan *et al.* 2003; Akbari *et al.*, 2013).

Whereas, cinnamon (*Cinnamomum verum* J. Presl) is one of the oldest and dearest spices known to mankind since time immemorial. The word cinnamon is derived from Greek word 'Kinnamon' which means 'sweet wood'. The cinnamaldehyde and eugenol are two principle constituents of cinnamon oil which attributed to its aroma and flavor. Almost every parts of cinnamon tree; bark, leaves, flowers, fruits and roots are of medicinal or culinary uses. Cinnamon is used for essential oil extraction, per-

fumes, pharmaceutical products, condiments and flavoring additives (Kumar, 2014) processed foods. It acts as antidiabetic (Sartorius *et al.*, 2014; Song *et al.*, 2013), anti-cardiovascular activities (Rahman *et al.*, 2013), anticancer (Xie *et al.*, 2018; Yang *et al.*, 2015), antioxidants (Ervina *et al.*, 2016; Abeysekara *et al.*, 2013), neuroprotective (Bae *et al.*, 2018; Peterson *et al.*, 2009), hepatoprotective (El-Kholy *et al.*, 2019), antimicrobial (Utcharikiat *et al.*, 2016), anti-HIV (Cornell *et al.*, 2016), anti-convulsant (Cuan *et al.*, 2018), anti-inflammatory (Han and Parker, 2007; Hong *et al.*, 2012), and antidepressant (Sohrabi *et al.*, 2017) etc. The main chemical constituents of cinnamon are mostly cinnamyl alcohol, coumarin, cinnamic acid, cinnamaldehyde, anthocyanin, eugenol and linalool.

People's awareness about such nutraceutical values of flowers and other plant products, and changes in modern food habits have them started to think about rediscovering the ancient ethnobotanical traditions. There has been a report of a rapid increase in demand for fresh edible flowers and its consumptions worldwide in recent times and studies are going on to explore the edible flower having potential for human diet as food, supplements, or additives (Loizzo *et al.*, 2016). However, there is a research gap in processing these nutritious flowers into drinkable beverages through the fermentation process and their nutritional assessment. This may be because the plant contains numerous anti-nutritional components, which may be poisonous sometimes, or maybe the scepticism about the undesirable end product after fermentation. Therefore, considering the importance of both rose and cinnamon as ethnomedicinal plants the present investigation was carried out to analyze the combined effects of rose and cinnamon on the nutraceutical quality of beverages obtained through the fermentation process. The rose was used as a major component in the study, while the cinnamon was added as a small supplement.

Materials and Method

The experiment was carried out in the laboratory of the Department of Horticulture, North Eastern Hill University, Tura Campus, Meghalaya, India. The average temperature during the study period was 16.71 to 30.62 °C, whereas humidity was 64.13 to 82.37%. The experiment consisted of 12 treatment combinations comprised of grounded rose petals

and cinnamon; T₁= (100% rose); T₂= (100% cinnamon); T₃= (99% rose + 1% cinnamon); T₄= (98% rose + 2% cinnamon); T₅= (97% rose + 3% cinnamon); T₆= (96% rose + 4% cinnamon); T₇= (95% rose + 5% cinnamon); T₈= (94% rose + 6% cinnamon); T₉= (93% rose + 7% cinnamon); T₁₀= (92% rose + 8% cinnamon); T₁₁= (91% rose + 9% cinnamon) and T₁₂= (90% rose + 10% cinnamon) respectively.

The TSS of the mixed solution was maintained at 20 °Brix by the addition of sugar, while pH was maintained at 3.5 by adding citric acid. Each treatment was prepared separately and poured into a sterilized conical flask of 500 ml in 3/4th of the total volume. The conical flasks were sealed airtight with polythene and nylon thread and then pasteurized for 15 minutes. After cooling down the samples, yeast (20ml/180 ml) was inoculated and then kept in a dry, dark, and ambient environment for fermentation. Racking, fining and filtration were done after completion of fermentation of samples (18.27 days). The TSS (Refractometer), reducing sugar (Lane and Enyon, 1923), non-reducing sugar (Lane and Enyon, 1923), acidity (AOAC, 1990), ascorbic acid (Ranganna, 2004), total sugar (Lane and Enyon, 1923), phenolic content (Ranganna, 2004) alcohol (digital alcoholometer Alex-500 in % by v/v), anthocyanin (Ranganna, 2004) were analyzed in the freshly fermented product as well as during its storage periods. The acceptability of the beverages was measured by score by panelists on 5.00 points Hedonic Scale. The statistical analysis of the significance of data was analyzed with the method described by Panse and Sukhatme (1967) and the analysis of variance (ANOVA) of data was carried out by techniques suggested by Raghuramula *et al.* (1983).

Results and Discussion

Sensory evaluation of beverages

As shown in Figure 1 the acceptability of the beverages was significantly different among the treatments in fresh stage as well as during storage. The highest acceptability score was found in treatment T₁₀ (3.79) which was at par with T₉ (3.51), and the lowest was in T₄ (2.35) in freshly fermented beverages. The acceptability score changed over the storage period which was in line with Mukisa *et al.* (2012a). In the subsequent months T₉ had the highest score of acceptability. Treatment T₄ had the low-

est acceptability throughout study period. The freshly fermented beverages had a clear reddish-pink color with a sweet rosy aroma and slight astringency which could be due to the presence of phenolic compounds reported by Lopez *et al.* (1983a). The color of beverages changed from clear reddish-pink to dark red during storage, which might be due to continuous oxidation or an increase in phenolic content during storage. This was in accordance to

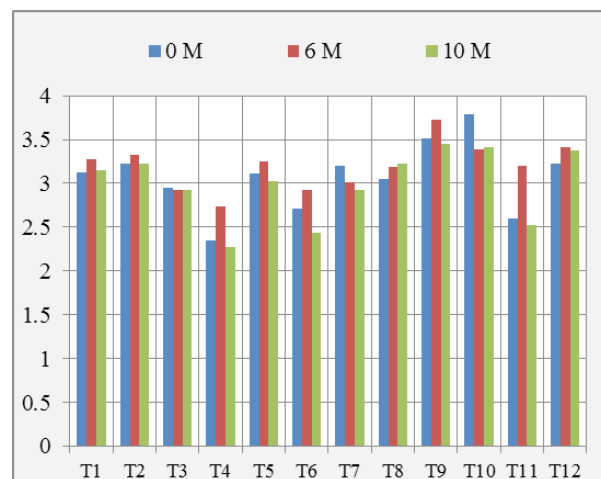


Fig. 1. Changes in the organoleptic score (5.00 points Hedonic Scale) of fermented beverages during storage

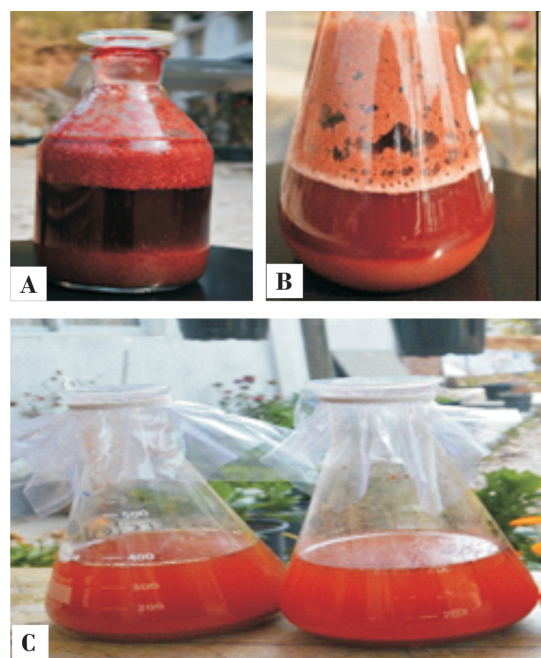


Fig. 2A. Mother Culture for inoculation; B. Treatment with combination of rose and cinnamon; C. Pure treatment of cinnamon

Lopez *et al.* (1983b) that the phenolic component contributes greatly to the color of the fermented products.

Nutritional assessment of fermented beverages from rose and cinnamon

TSS

As depicted in Table 1 the treatment T₂ had the highest TSS (19.67 °Brix) which was significantly different from rest of the treatments. Treatment T₁₂ (8.73 °Brix) had second highest TSS which was at par with T₁ (8.20), T₄ (8.47), T₉ (8.27) and T₁₁ (8.20) in freshly fermented beverages. The TSS content in beverages fluctuates during the storage. After 10th months of storage, the highest TSS was observed in T₂ (19.64 °Brix, with just 0.16% reduction) which was significantly different from other treatments. With 8.34 °Brix, T₄ had second highest TSS content which was at par with rest of the treatments. A minor and irregular change in TSS content was noticed during storage. A similar of fluctuation in TSS of fermented beverages from Mahua flower was reported by Singh *et al.* (2013a).

Total sugar

As shown in Table 2 the treatment T₁₂ had the highest (1.99%) total sugar content followed by T₅ (1.91%) which was at par with T₃ (1.88%), and T₆ (1.89%) in freshly fermented beverages. While the lowest (1.67%) was in T₁. The total sugar content in the beverages decreased over the storage period in all treatments. After 10 months of storage, T₁ had the

highest total sugar (1.57%) content which was at par with T₂, T₃ (1.54%), T₆ (1.55%), and T₁₂ (1.53%). These were in line with Okemini *et al.* (2017) reported 1.25% total sugar in fermented Tiger Nut beverages. Over the storage period, the lowest reduction of total sugar content was recorded in T₁ (5.98%) followed by T₂ (12.92%), whereas, maximum in T₁₁ (30.89%), followed by T₁₀ (29.03%). The reduction percent of total sugar in the treatments with higher concentration of cinnamon was relatively higher than the pure treatments of rose and cinnamon alone. The regular decreased in total sugar content might be due to the continuous conversion of sugar into alcohol by viable yeast as reported by Mukisa *et al.* (2012b) in fermented Obushera beverages; Singh *et al.* (2013b) in fermented Mahua flower.

Reducing sugar

As depicted in Table 3 the reducing sugar content in beverages changed throughout the storage period irregularly. In freshly fermented beverages, the highest reducing sugar was found in T₁₀ (1.58%) which was at par with T₃ (1.56%) whereas the lowest was in T₇ (1.15%) which was at par with T₈ (1.19%). This was in agreement with Yadav *et al.* (2009a) who reported 0.08-2.21% reducing sugar in the fermented beverage from Mahua flower. After 10 months of storage, treatment T₂ and T₆ had the highest (1.46%) reducing sugar content which was at par with T₃ (1.45%), T₁, T₅, and T₈ (1.42%) whereas the minimum was in T₁₂ (1.15%) which was at par with T₁₁ (1.20%). The highest increment of reducing sugar after 10 months of storage was found in T₂

Table 1. Changes in TSS content in beverages during beverages during storage (%)

Treatments	0 M	6 M	10 M
T ₁	8.20	7.80	8.00
T ₂	19.67	19.60	19.64
T ₃	8.13	7.93	8.03
T ₄	8.47	8.20	8.34
T ₅	8.00	8.07	8.04
T ₆	7.80	8.20	8.00
T ₇	8.00	7.60	7.80
T ₈	7.80	8.00	7.90
T ₉	8.27	8.20	8.24
T ₁₀	8.47	7.20	7.84
T ₁₁	8.20	7.87	8.04
T ₁₂	8.73	7.73	8.23
C.D (5%)	0.53	0.59	0.56
S.E(m)	0.18	0.20	0.19

Table 2. Changes in Total Sugar content in storage (p Brix)

Treatments	0 M	6 M	10 M
T ₁	1.67	1.64	1.57
T ₂	1.78	1.75	1.55
T ₃	1.88	1.84	1.54
T ₄	1.79	1.72	1.40
T ₅	1.91	1.88	1.46
T ₆	1.89	1.85	1.55
T ₇	1.79	1.77	1.39
T ₈	1.84	1.79	1.45
T ₉	1.74	1.71	1.34
T ₁₀	1.86	1.82	1.32
T ₁₁	1.78	1.73	1.23
T ₁₂	1.99	1.96	1.53
C.D (5%)	0.04	0.02	0.06
S.E(m)	0.01	0.01	0.02

and T₆ (20.66%) whilst it decreased by 18.35% (highest) in treatment T₁₀ followed by 14.17% in T₁₂. This was in accordance to Singh *et al.* (2013b) who reported decreasing reducing sugar with a storage period from 5.76 to 0.48% in fermented beverages from Mahua flower during storage.

Non-reducing sugar

From the perusal of data shown in Table 4 the non-reducing sugar content in the fermented beverages were significantly different among the treatments and decreased continuously with the storage period. The T₅ (0.80%) had the highest non-reducing sugar which was at par with T₈ (0.74%), T₁₂ (0.73%), and T₆ (0.72%) in fresh conditions. While the least was recorded in T₁₀ (0.25%) followed by T₃ (0.38%). Similar was resulted by Thakur and Sharma (2017a) with 0.82% non-reducing sugar in fermented pomegranate beverages. A slight increase in non-reducing sugar was noticed in treatment T₃ (0.55%), and T₁₀ (0.48%) while in other treatments it was reduced after 6 months of storage. After 10 months of storage, the maximum non-reducing sugar content was found in T₁₂ (0.43%) and the minimum was recorded in T₉ and T₁₁ (0.09%) which was at par with T₅, T₆ (0.11%) and T₇, T₈ and T₁₀ (1.10%) respectively. The minimum reduction of non-reducing sugar (41.09%) with storage period was also recorded in T₁₂, while maximum reduction (86.48%) was recorded in T₈.

Acidity

From the data shown in Table 5 the acidity of fermented beverages was recorded to be reducing over

the time of storage except for the treatments T₃ and T₁₀ where acidity remained unchanged till the tenth months of storage. The treatment T₅ (0.32%) had the highest acidity and minimum in T₂ (0.22%) which was at par with T₆ and T₁₀ (0.24%) in freshly fermented beverages which were in line with Yadav *et al.* (2009b) reported 0.63% acidity in fermented beverages from Mahua flower. The acidity of the beverage remained unchanged in treatment T₃ (0.29%), T₁₀ (0.24%), T₁₁ (0.28%), and T₁₂ (0.25%) after 6 months of storage, while in rest, it was reduced. After 10 months of the storage period, the acidity of beverages was found maximum in T₅ (0.28%) which was at par with T₃ (0.27%). The maximum reduction in acidity was recorded in pure treatments; T₂ (22.72%) and T₁ (13.79%) respectively after 10 months. It was observed that reduction (%) in acidity in combined treatment of rose and cinnamon was comparatively lower than pure treatments (T₁ and T₂). The least reduction of acidity was found in T₇ and T₁₁ with 3.57% each. A similar finding of decreasing acidity was reported by Iheke *et al.* (2017) in fermented locust bean flour.

Ascorbic acid

As depicted in Table 6 the treatment T₄ contained a maximum amount of ascorbic acid (5.51 mg/100ml) which was at par with T₅ (4.80 mg/100 ml) and T₉ (5.46 mg/100 ml) and the least was in T₁ and T₁₂ (4.32 mg/100 ml). A similar was reported by Yadav *et al.* (2009c) 4.27 mg/100 ml ascorbic acid in fermented beverages from Mahua flower. In the sixth month of storage, an increment of ascorbic acid was

Table 3. Changes in reducing sugar in beverages during storage (%).

Treatments	0 M	6 M	10 M
T ₁	1.26	1.49	1.42
T ₂	1.21	1.29	1.46
T ₃	1.56	1.36	1.45
T ₄	1.29	1.30	1.33
T ₅	1.21	1.25	1.42
T ₆	1.21	1.32	1.46
T ₇	1.15	1.33	1.35
T ₈	1.19	1.38	1.42
T ₉	1.40	1.44	1.32
T ₁₀	1.58	1.41	1.29
T ₁₁	1.30	1.44	1.20
T ₁₂	1.34	1.44	1.15
C.D (5%)	0.14	0.08	0.07
S.E(m)	0.04	0.03	0.02

Table 4. Changes in non-reducing sugar in beverages during storage (%).

Treatments	0 M	6 M	10 M
T ₁	0.46	0.22	0.14
T ₂	0.63	0.53	0.12
T ₃	0.38	0.55	0.13
T ₄	0.55	0.49	0.14
T ₅	0.80	0.69	0.11
T ₆	0.72	0.59	0.11
T ₇	0.69	0.51	0.10
T ₈	0.74	0.48	0.10
T ₉	0.45	0.35	0.09
T ₁₀	0.25	0.48	0.10
T ₁₁	0.54	0.36	0.09
T ₁₂	0.73	0.60	0.43
C.D (5%)	0.08	0.08	0.01
S.E(m)	0.02	0.03	0.01

recorded in all treatments. However, it was noticed that it started to decline at tenth month. The maximum ascorbic acid content after 10 months of storage was found in T₇ (4.75 mg/100 ml) which was at par with T₈ (4.56 mg/100 ml), T₁₂ (4.48 mg/100 ml), and T₉ (4.42 mg/100 ml). About 80.95% to 98.06% of ascorbic acid was retained in different treatments till the tenth month of storage. This reduction in ascorbic acid content might be due to the oxidation and reduction reaction in beverages, as reported by Wahab *et al.* (2005) that growing yeast metabolizes the ascorbic acid and reduce it. Furthermore, Fleet and Heard (1993) also resulted that yeast uses ascorbic acid as a source of carbon. On the contrary, a small decrease in ascorbic acid content at the tenth month may be due to the presence or continuous

increase of phenol which prevents the oxidation reactions reported by Mancini *et al.* (1998a) that phenolic content inhibits the oxidative process in fermented beverages.

Anthocyanin

As shown in Table 7 the treatment T₁₂ contained maximum amounts of anthocyanin (7.00 mg/100ml), which was at par with T₁ (6.71 mg/100 ml), T₃ (6.77 mg/100 ml), T₄ (6.72 mg/100 ml), T₅ (6.59 mg/100 ml), and T₁₁ (6.67 mg/100 ml) whereas the least was in T₂ (0.50 mg/100 ml) in the freshly fermented beverage. At sixth months of storage, there was huge reduction in anthocyanin content in the beverages with maximum reduction in T₁₂ (80%) to lowest in T₂ (38%) respectively. Except T₂, the antho-

Table 5. Changes in acidity content in beverages during storage (%)

Treatments	0 M	6 M	10 M
T ₁	0.29	0.28	0.25
T ₂	0.22	0.21	0.17
T ₃	0.29	0.29	0.29
T ₄	0.26	0.25	0.25
T ₅	0.32	0.31	0.30
T ₆	0.24	0.22	0.22
T ₇	0.28	0.27	0.27
T ₈	0.27	0.26	0.26
T ₉	0.27	0.26	0.26
T ₁₀	0.24	0.24	0.24
T ₁₁	0.28	0.28	0.27
T ₁₂	0.25	0.25	0.24
C.D (5%)	0.03	0.04	0.05
S.E(m)	0.01	0.01	0.01

Table 6. Changes in the ascorbic acid content in beverages during storage (mg/100ml)

Treatments	0 M	6 M	10 M
T ₁	4.32	6.08	3.70
T ₂	4.37	4.87	3.34
T ₃	4.40	5.85	3.96
T ₄	5.51	6.47	4.39
T ₅	4.80	6.08	4.36
T ₆	4.46	6.98	4.32
T ₇	4.51	6.97	4.75
T ₈	4.65	4.82	4.56
T ₉	5.46	6.92	4.42
T ₁₀	4.40	7.34	4.05
T ₁₁	4.43	6.22	4.08
T ₁₂	4.32	6.10	4.48
C.D (5%)	0.75	0.66	0.68
S.E(m)	0.26	0.22	0.23

Table 7. Changes in anthocyanin content in beverages during storage (mg/100ml)

Treatments	0 M	6 M	10 M
T ₁	6.71	1.71	1.19
T ₂	0.50	0.31	0.25
T ₃	6.77	1.62	1.30
T ₄	6.72	1.78	1.33
T ₅	6.59	1.74	1.25
T ₆	5.68	1.69	1.25
T ₇	6.17	1.63	1.21
T ₈	6.18	1.56	1.30
T ₉	5.80	1.52	1.19
T ₁₀	6.20	1.50	1.18
T ₁₁	6.67	1.43	1.16
T ₁₂	7.00	1.40	1.21
C.D (5%)	0.64	0.2	0.13
S.E(m)	0.21	0.07	0.04

Table 8. Changes in phenolic content in beverages during storage (mg/100ml/GAE)

Treatments	0 M	6 M	10 M
T ₁	157.96	172.45	215.25
T ₂	29.23	78.69	72.87
T ₃	151.97	164.63	205.59
T ₄	157.13	171.62	218.91
T ₅	116.33	148.14	192.1
T ₆	164.63	173.12	212.42
T ₇	169.29	171.79	217.58
T ₈	165.12	177.95	208.59
T ₉	199.76	202.59	221.58
T ₁₀	182.11	192.94	215.58
T ₁₁	169.62	174.78	198.6
T ₁₂	166.62	174.78	202.59
C.D (5%)	30.52	29.93	14.47
S.E(m)	10.39	10.19	4.92

cyanin content in all treatments was reduced to more than 70% at 6th month.

After 10 months, T₄ (1.33 mg/100 ml) had the highest anthocyanin content which was at par with all the treatments except T₂ (0.25 mg/100 ml). The maximum retention of anthocyanin was recorded in T₄ (19.89%), whereas maximum reduction was observed in T₁₂ (82.71%) followed by T₁ (82.26%) during storage. This reduction in anthocyanin content in beverages could be due to changes in pH, temperature, exposure to light, and oxidation during handling of the beverages which causes degradation or maybe continuously metabolized by the viable yeast. Mousavi *et al.* (2011a) reported a similar result of decreasing anthocyanin content with storage period in fermented pomegranate juice.

Phenol

From the perusal data in Table 8 it can be seen that the phenolic content in fermented beverages increased with the storage period, except in treatment T₂ which declined at tenth month of storage. In freshly fermented beverages, T₉ had highest amount of phenol (199.76 GAE/100 ml) which was at par with T₇ (169.29 GAE/100 ml), T₁₀ (182.11 GAE/100ml), and T₁₁ (169 GAE/100 ml). At sixth months of storage, maximum increment in phenolic content was recorded in cinnamon pure treatment (T₂) with 49.46 GAE/100ml) whereas least increment was recorded in T₇ (2.5 GAE/100ml).

After 10 months, T₉ (221.58 GAE/100 ml) had maximum phenol content which was on par with T₄ (218.91 GAE/100 ml) and T₇ (217.58 GAE/100 ml), whereas the lowest was in T₂ (72.87 GAE/100 ml). The phenolic content in T₉ remained consistently highest throughout the study period. The maximum increase in phenolic content was recorded in T₅ (from 116.33 to 192.10 GAE/100 ml) by 65.13% after 10 months. This was in agreement with Ifie *et al.* (2016) reported a continuous increase in phenolic content in fermented wine from *Hibiscus sabdariffa* by 41.26% to 45.70% after 40 days of storage; Thakur and Sharma (2017c) also resulted 206.64 mg/100 ml phenolic content in fermented pomegranate beverage. This is also in line with Mousavi *et al.* (2011b) that enzymes such as α -glycosidase derived from fermentative microorganisms hydrolyze the complex phenolic compounds to simpler types and increased the quantitative amount of total phenolic compounds during fermentation.

Alcohol

The alcohol content changed concomitantly with the storage period except in treatment T₁ (100% rose) where it declined as depicted in Table 9. The maximum alcohol content was recorded in T₃ (12.41%) which was at par with T₁ (12.32%), T₅ (12.24%), and T₆ (12.27%) in freshly fermented beverages while the least was in T₂ (0.10%). The alcohol content in treatment T₂ was incomparable with other treatments. It might be due to the antimicrobial (anti-yeast) properties of cinnamon which inhibited the fermentation process, and consequently slowed down the conversion of sugar into an alcohol. A similar study was reported by Fraser *et al.* (2007); Moritz *et al.* (2012) that cinnamon inhibits the fermentation action and microbial counts in the culture. The antimicrobial effect of cinnamaldehyde is also reported by Utcharykiat *et al.* (2016).

Table 9. Changes in alcohol content in beverages during storage (%)

Treatments	0 M	6 M	10 M
T ₁	12.32	12.24	12.04
T ₂	0.10	0.64	0.71
T ₃	12.41	12.49	12.93
T ₄	12.03	12.51	12.96
T ₅	12.24	12.52	13.18
T ₆	12.27	12.95	13.17
T ₇	12.17	12.68	13.27
T ₈	12.13	12.39	13.17
T ₉	12.19	12.33	13.22
T ₁₀	11.89	12.27	13.09
T ₁₁	12.19	12.43	13.51
T ₁₂	12.07	12.15	13.41
C.D (5%)	0.17	0.45	0.25
S.E(m)	0.05	0.15	0.08

After 6 months, maximum alcohol content was found in T₆ (12.95%) which was at par with T₇ (12.68%) and T₅ (12.65%) and the minimum was in T₂ (0.64%). After 10 months, the maximum alcohol content was recorded in T₁₁ (13.51%) which was at par with T₁₂ (13.41%) and T₇ (13.27%) whereas the least was in T₂ (0.71%). The increasing rate of alcohol content was noticed from 9.77-9.99% in various treatments which was following Okemini *et al.* (2017) reported that alcohol content in fermented Tiger nut increased with storage period. While decreasing in alcohol content was noticed in T₁ up to 2.27%. The small increments of alcohol content in T₂

and decreasing trend in T_1 could be attributed to slower conversion of total sugar, in which reduction of total sugar was recorded comparatively lower than others during storage period.

Summary and Conclusion

The fermented beverages from rose and cinnamon had an attractive reddish-pink in color with rosy aroma. In freshly fermented condition, the highest acceptability was recorded in T_{10} (3.79) which was at par with T_9 (3.52) and least in T_4 (2.35). In the subsequent storage period, the acceptability of the beverages changed. The T_9 (3.73) had the highest acceptability at tenth month. While T_4 remained the least acceptable treatment throughout the study period.

The TSS content in the freshly fermented beverages was recorded highest in T_2 (19.67pB) which was significantly different from all the treatments. It remained highest till 10th month of the storage study. The total sugar content was recorded highest in T_{12} (1.99%) while lowest in T_1 (1.67%). After 10 months of storage, the total sugar content declined with highest recorded in T_1 (1.57%) which was at par with T_2 , T_3 (1.54%), T_6 (1.55%) and T_{12} (1.53%). The reduction in total sugar content in the beverages with storage period was recorded lowest in pure treatments of rose and cinnamon compare to the treatments with combination of two.

The reducing sugar content of the freshly fermented beverages was found highest in T_{10} (1.58%) which was at par with T_3 (1.56%) while lowest in T_7 (1.15%) which was at par with T_8 (1.19%). After 10 months of storage, T_2 and T_6 had the highest reducing sugar (1.46%) which was at par with T_3 (1.45%), T_1 , T_5 and T_8 (1.42%). Meanwhile the highest non-reducing sugar was recorded in T_5 (0.80%) which was at par with T_8 (0.74%), T_{12} (0.73%) and T_6 (0.72%) respectively in fresh condition. Whereas least was found in T_{10} (0.25%).

The acidity of the treatment T_3 and T_{10} remained unchanged throughout the study period. The highest acidity was recorded in T_5 (0.32%) while minimum in T_2 (0.22%) which was at par with T_6 and T_{10} (0.24%). After 10 months, the highest acidity was recorded in T_5 (0.28%) which was at par with T_3 (0.27%). The maximum reduction in acidity was observed in pure treatments T_2 (22.72%) and T_1 (13.79%) respectively. The initial highest ascorbic acid content was found in T_4 (5.51 mg/100ml) which was at par with T_5 (4.80 mg/100ml) while least in T_1

and T_{12} (4.32 mg/100 ml). The highest increment in ascorbic acid content in fermented beverages was recorded in sixth month. However, it started to decline in tenth month. The highest ascorbic acid content was found in T_7 (4.75 mg/100 ml) which was at par with T_8 (4.56 mg/100 ml), T_{12} (4.48 mg/100 ml) and T_9 (4.42 mg/100 ml). About 80.95% to 98.06% ascorbic acid was retained at tenth month of storage.

The highest anthocyanin content was recorded in T_{12} (7.00 mg/100 ml) which was at par with T_1 (6.71 mg/100 ml), T_3 (6.77 mg/100 ml), T_4 (6.72 mg/100 ml), T_5 (6.59 mg/100 ml) and T_{11} (6.67 mg/100 ml) respectively while least was in T_2 (0.50 mg/100 ml). A huge reduction in anthocyanin content was recorded during sixth month of storage (38-80%). After 10 months of storage, highest anthocyanin content was recorded in T_4 (1.33 mg/100 ml) which was at par with all the treatments except T_2 (0.25 mg/100 ml). At tenth month, highest retention of anthocyanin was recorded in T_4 (19.89%) whereas maximum reduction was in T_{12} (82.71%).

The phenolic content in the beverages was found highest in T_9 (1.99.76 GAE/100 ml) which was at par with T_7 (169.29 GAE/100 ml), T_{10} (182.11 GAE/100 ml) and T_{11} (169.00 GAE/100ml). The highest increment in phenolic content was recorded in T_2 while lowest in T_7 during sixth month of storage. After 10 months of storage, T_9 (221.58 GAE/100 ml) had highest phenolic content which was at par with T_4 (218.91 GAE/100 ml) and T_7 (217.58 GAE/100 ml) while lowest in T_7 (72.87 GAE/100 ml). The highest increment in phenolic content was recorded in T_5 (65.13%) with storage period.

The alcohol content in the beverages changed concomitantly with the storage period except in T_1 (100% rose) where it declined continuously. In fresh condition, the maximum alcohol content in the beverages was recorded highest in T_3 (12.41%) which was at par with T_1 (12.32%), T_5 (12.24%) and T_6 (12.27%) while least was in T_2 (0.10%). After 10 months of storage, T_{11} (13.51%) had the highest alcohol content which was at par with T_{12} (13.41%) and T_7 (13.27%) while least in T_2 (0.71%). The alcohol content increased from 9.77% to 9.99% in various treatments.

Based on the analyses of certain characteristics and quality of fermented beverages, the treatment T_9 (93% rose + 7% cinnamon) was the best treatment in terms of organoleptic score, vitamin C content (ascorbic acid) and anti-oxidants counts (anthocyanin, phenol). However, in terms of alcohol content in

the beverages, the best treatment was T₁₁ (91% rose + 9% cinnamon). The pure treatment of cinnamon (100% cinnamon) in T₂ showed strong antimicrobial property by inhibiting the fermentation process throughout the study period. However, it was found that the proportion of cinnamon added in the beverages in combination with rose improved the nutritional quality of the beverages as well as prevent the oxidation of important nutrients such as ascorbic acid.

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Conflicts of interest

The authors declare no conflict of interest.

References

- Abeyssekera, W.P.K.M., Premakumara, G.A.S. and Ratnasooriya, W.D. 2013. In vitro antioxidant properties of leaf and bark extracts of Ceylon cinnamon (*Cinnamomum zeylanicum* Blume). *Tropical Agricultural Research Journal*. 24(2): 128-138.
- Achuthan, C.R., Babu, B.H. and Padikkala, J. 2003. Antioxidant and hepatoprotective effects of *Rosa damascena*. *Pharmaceutical Biology*. 41(5): 357-361.
- Akbari, M., Kazerani, H.R., Kamrani, A. and Mohri, M. 2013. A preliminary study on some potential toxic effects of *Rosa damascena* Mill. *Iranian Journal of Veterinary Research*. 14(3): 232-236.
- Al-Dhubiab, B.E. 2012. Pharmaceutical applications and phytochemical profile of *Cinnamomum burmanii*. *Pharmacognosy Reviews*. 6(12): 125-131.
- Almeida, H., Amaral, M.H. and Lobão, P. 2011. Drugs obtained by biotechnology processing. *Brazilian Journal of Pharmaceutical Sciences*. 47(2): 199-207.
- Anonymous, 2018. *United States Department of Agriculture, National Nutrient Database for Standard Reference (USDA)*.
- AOAC, 1990. *Official Methods of Analysis*. 15th edition. Association of Official Analytical Chemists, Arlington, VA.
- Bae, W.Y., Choi, J. and Jeong, J. 2018. The neuroprotective effects of cinnamic aldehyde in an MPTP mouse model of Parkinson's disease. *International Journal of Molecular Sciences*. 19(2): 551.
- Benvenuti, S. and Mazzoncini, M. 2021. The Biodiversity of Edible Flowers: Discovering New Tastes and New Health Benefits. *Frontiers Plant Sciences*. 11: 1-14.
- Butcaru, A.C., Stănică, F., Nicolae, and Velcea, M. 2017. Preliminary studies regarding the production of Jam from organic rose petal. *Food Science and Technology*. 74(2): 1-8.
- Chericoni, S., Prieto, J.M., Iacopini, P., Cioni, P. and Morelli, I. 2005. In vitro activity of the essential oil of *Cinnamomum zeylanicum* and eugenol in peroxynitrite-induced oxidative processes. *Journal of Agricultural and Food Chemistry*. 53(12): 4762-4765.
- Cuan, Y., He, X., Zhao, Y., Yang, J., Bai, Y., Sun, Y., Zhang, Q., Zhao, Z., Wei, X. and Zheng, X. 2018. Anticonvulsant activity of halogen-substituted cinnamic acid derivatives and their effects on glycosylation of PTZ induced chronic epilepsy in mice. *Molecules*. 23(1): 1-31.
- El-Kholy, W.M., Hemieda, F.A.E. and Elabani, G.M. 2019. Role of cinnamon extract in the protection against amoxicillin/clavulanate-induced liver damage in rats. *Journal of Pharmacy and Biological Sciences*. 14(1): 14- 21.
- Ervina, M., Nawu, Y.E. and Esar, S.Y. 2016. Comparison of in vitro antioxidants activity of infusion, extracts and fractions of Indonesian cinnamon (*Cinnamomum burmanii*) bark. *International Food Research Journal*. 23(3) : 1346-1350.
- Esfandiary, E., Karimipour, M., Mardani, M., Ghanadian, M., Alaei, H.A., Mohammadnejad, D. and Esmaeili, A. 2015. Neuroprotective effects of *Rosa damascena* extract on learning and memory in a rat model of amyloid- β -induced Alzheimer's disease. *Advance Biomedical Research*. 4(131): 1-18.
- Fleet, G.H. and Heard, G.M. 1993. Yeasts- Growth during fermentation. *Wine Microbiology and Biotechnology. Harwood Academic Publications, Chur, Switzerland*. 8: 27-54.
- Fraser, G.R., Chaves, A.V., Wang, Y., McAllister, T.A., Beauchemin, K.A. and Benchaar, C. 2007. Assessment of the effects of cinnamon leaf oil on rumen microbial fermentation using two continuous culture systems. *Journal of Dairy Science*. 90(5): 2315-2328.
- Friedman, H., Rot, I., Agami, O., Vinokur, Y., Rodov, V., Rajnick, N., Umiel, N., Ganot, L., Shmuel, D. and Matan, E. 2007. Edible flower new crop with potential health benefits. *Acta Horticulturae*. 755(36): 283-290.
- Gao, X.M., Shu, L.D., Yang, L.Y. and Shen, Y.Q. 2013. Phenylethanoids from the flowers of *Rosa rugosa* and their biological activities. *Bulletin-Korean Chemical Society*. 34(1): 246-248.
- Guimaraes, R., Barros, L., Carvalho, A. M., Isabel, C. and Ferreira, F. R. 2010. Studies on chemical constituents and bioactivity of *Rosa micrantha*: An alternative

- antioxidants source for food, pharmaceutical, or cosmetic applications. *Journal of Agricultural and Food Chemistry*. 58(10): 6277-6284.
- Han, X. and Parker, T.L. 2007. Anti-inflammatory activity of cinnamon (*C. zeylanicum*) barks essential oil in human skin disease model. *Pythotherapy Research*. 31(7): 1034-1038.
- Homayoun, M., Seghatoleslam, M., Pourzaki, M., Shafieian, R., Hosseini, M. and Bideskan, A.E. 2015. Anticonvulsant and neuroprotective effects of *Rosa damascena* hydro-alcoholic extract on rat hippocampus. *Avicenna Journal of Phytomedicines*. 5(3): 260-270.
- Hong, J., Yang, G., Kim, Y.B., Eom, S.H., Lew, J. and Kang, H. 2012. Anti-inflammatory activity of cinnamon water extracts *in vivo* and *in vitro* LPS induced models. *Complementary and Alternative Medicine*. 12(237): 1-8
- Ifie, I., Marshall, L.J., Ho, P. and Williamson, G. 2016. *Hibiscus sabdariffa* (Roselle) extracts and wine: Phytochemical profile, physico-chemical properties, and carbohydrate inhibition. *Journal of Agricultural Food and Chemistry*. 64(24): 4921-4931.
- Iheke, E., Oshodi, A., Omoboye, A. and Ogunlalu, O. 2017. Effect of fermentation on the physicochemical properties and nutritionally valuable minerals of Locust Bean (*Parkia biglobosa*). *American Journal of Food Technology*. 12(6): 379-384.
- Jat, R., Mahawer, L.N., Bairwa, H.L., Meena, R.H., Pilania, S. and Singh, M. 2018. Sensory evaluation and microbial analysis of rose petal jam. *Journal of Pharmacognosy and Phytochemistry*. 7(5): 617-620.
- Joshi, V.K., Sharma, S. and Devi, M.P. 2009. Influence of different yeasts strains on fermentation behavior, physico-chemical, and sensory qualities of Plum wine. *Natural Production Radiance*. 8(4): 445-451.
- Ju, J. E., Joo, Y.H., Chung, N., Chung, S.Y., Han, S.H. and Lee, Y.W. 2014. Antidiabetic effects of red rose flowers in streptozotocin-induced diabetic mouse. *Journal of Korean Society for Applied Biological Chemistry*. 57(4): 445-448.
- Kart, D. and Çađindi, O. 2017. Determination of antioxidant properties of dry rose tea. *International Journal of Secondary Metabolite*. 4(3): 384-390.
- Kumar, N. 2014. *Introduction to Spices, Plantation Crops, Medicinal and Aromatic Plants*. Oxford and IBH Publishing. Pp. 296-336.
- Lane, J.H. and Eynon, L. 1923. Determination of reducing sugar by means of Fehling's solution with methylene blue indicator as internal indicator. *Journal of the Society of Chemical Industry*. 43: 32-36.
- Laxmi, S.K., Sandhya D.D., Geetha S. and Lakshmi S.M. 2017. Biochemical and antimicrobial analysis of rose petals (*Rosa indica*). *European Journal of Pharmaceutical and Medical Research*. 4(7): 637-640.
- Loizzo, M. R., Pugliese, A., Bonesi, M., Tenuta, M. C., Menichini, F., Xiao, J. and Tundis, R. 2016. Edible flowers: A rich source of phytochemicals with antioxidant and hypoglycemic properties. *Journal of Agricultural and Food Chemistry*. 64(12): 2467-2474.
- Lopez, Y., Gordon, D.T. and Fields, M.L. 1983. Release of Phosphorus from phytate by natural lactic fermentation. *Journal of Food Science*. 48(3): 935-954.
- Mancini-Filho, J.A., Van-Koijij, D.A., Mancini, P., Cozzolino, F.F. and Torres, R.P. 1998. Antioxidant activity of cinnamon (*Cinnamomum zeylanicum*, Breyne) extracts. *Bollettino Chimico Farmaceutico*. 137(11): 443-447.
- Moritz, C.M.F., Rall, V.L.M., Saeki, M.J. and Júnior, F.J. 2012. Inhibitory effect of essential oils against *Lactobacillus rhamnosus* and starter culture in fermented milk during its shelf-life period. *Brazilian Journal of Microbiology*. 43(2): 1147-1156.
- Mousavi, Z.E., Mousavi, S.M., Razavi, S.H., Emmam-Djomeh, Z. and Kiani, H. (2011). Effect of fermentation on pomegranate juice by *Lactobacillus planatarum* and *Lactobacillus acidophilus* on the antioxidants activity and metabolism of sugars, organic acids, and phenolic compounds. *World Journal of Microbiology and Biotechnology*. 27(2): 123-128.
- Mukisa, I., Cmbk, M., Byaruhanga, Y.B. Langsrud, T. and Narvhus, J. 2012. Changes in physico-chemical properties and flavour compounds during fermentation of different obushera (sorghum and millet) beverages. *African Journal of Food, Agriculture, Nutrition and Development*. 12(6): 6665-6684.
- Okemini, O.F. and Dilim, I.C. 2017. Physico-chemical properties and sensory evaluation of wine produced from Tiger nut (*Cyperus esculentus*). *International Journal of Chemical Technology Research*. 10(12): 155-164.
- Onkuwa, G.I. and Ogbogu, N.I. 2007. Studied on the effect of fermentation on the quality and physico-chemical properties of cassava based Fufu products made from two cassava varieties NR8212 and Nwangbisi. *Journal of Food Technology*. 5(3): 261-264.
- Pansee, V. G. and Sukhatme, P. V. 1967. *Statistical Methods for Agricultural Workers*. 2nd edition., Indian Council of Agricultural Research, New Delhi.
- Peterson, D.W., George, R.C. and Scaramozzino, F. 2009. Cinnamon extract inhibits tau aggregation associated with Alzheimer's disease *In vitro*. *Journal of Alzheimer's Disease*. 17(3): 585-597.
- Raghuramula, H., Madhavan, N. K. and Sundaram, K. 1983. A manual of laboratory technology. *National Institute of Nutrition, Indian Council of Medical Research, Hyderabad*.
- Rahman, S., Begum, H., Rahman, Z., Ara, F., Iqbal, M.J. and Yousuf, A.K.M. 2013. Effect of cinnamon (*Cinnamomum cassia*) as a lipid lowering agent on hypercholesterolemic rats. *Journal of Enam Medical College*. 3(2): 94-98.
- Ranganna, S. 2004. *Handbook of Analysis and Quality Con-*

- trol for Fruit and Vegetable Products*. 3rd edition., Tata McGraw Hill Publication Corporation Limited, New Delhi, India.
- Sanlier, N., Gockcen, B.B. and Sezgin, A.C. 2017. Health benefits of fermented foods. *Critical Reviews in Food science and Nutrition*. 0(0): 1-22.
- Sartorius, T., Peter, A., Schulz, N., Drescher, A., Bergheim, I., Machann, J., Schick, F., Siegel-Axel, D., Schurmann, A., Weigert, C., Haring, H. and Hennige, A.M. 2014. Cinnamon extract improves insulin sensitivity in the brain and lowers liver fat in mouse models of obesity. *Public Library of Science*. 9(3): 1-12.
- Shaheen, S., Ahmad, M. and Haroon, N. 2017. Edible wild plants: A solution to overcome food insecurity. In *Edible Wild Plants: An Alternative Approach to Food Security*, Springer International Publishing, Cham, Switzerland. pp. 41–57.
- Singh, Kushal, Kumar, Gunjeet, Saha, T. N. and Kumar, R. 2013. *Post-Harvest Technology of Cut Flowers*. Venus Printers and Publishers, New Delhi. Pp. 26-28.
- Singh, Rupesh., Mishra, B.K., Shukla, K.B., Jain, N.K., Sharma, K.C., Kumar, Sunil., Kant, Krishna. and Ranjan, J.K. 2013. Fermentation process for alcoholic beverage production from mahua (*Madhuca indica* J. F. Mel.) flowers. *African Journal of Biotechnology*. 12(39): 5771-5777.
- Sohrabi, R., Pazgoohan, N., Seresht, H.R. and Amin, B. 2017. Repeated systemic administration of the cinnamon essential oil possesses anti-anxiety and antidepressant activities in mice. *Iranian Journal of Basic Sciences*. 20(6): 708-714.
- Song, F., Li, H., Sun, J. and Wang, S. 2013. Protective effects of cinnamic acid and cinnamic aldehyde on isoproterenol-induced acute myocardial ischemia in rats. *Journal of Ethnopharmacology*. 150(1) : 125-130.
- Swati, D.B., Chakravarthy, A., Mutalik, S. and Devkar, R. 2016. Determination of antibacterial and antifungal properties of rose extract- An *in vitro* study. *International Journal of Pharmacognosy and Phytochemical Research*. 8(10) : 1695-1697.
- Tatke, P.A., Patil, P.S. and Gabhe, S.Y. 2015. *In vitro* antioxidants and free radical scavenging activity of extracts of *Rosa damascena* flower petals. *American Journal of Phytomedicine and Clinical Therapeutics*. 3(9): 589-601
- Thakur, M. and Sharma, R.K. 2017. Development of probiotic pomegranate beverage and its physico-chemical and microbial characterization. *International Journal of Pure and Applied Bioscience*. 5(1): 35-41.
- Tirupathi, H. and Golla, P. 2016. To evaluate and compare antidepressant activity of *Rosa damascena* in mice by using forced swimming test. *International Journal of Basic and Clinical Pharmacology*. 5(5): 1949-1952.
- Utcharyiakiat, I., Surassmo, S., Jaturanpinyo, M., Khuntayaporn, P., Traidej, M. and Chomnawang. 2016. Efficacy of cinnamon bark oil and cinnamaldehyde on anti-multidrug resistant *Pseudomonas aeruginosa* and the synergistic effects in combination with other antimicrobial agents. *Bio-Medicine Central Complementary and Alternative medicine*. 16(158): 1-7.
- Wahab, O. Okunowo., Rufus, O. Okotore and Akinniyi, A.O. 2005. The alcoholic fermentative efficiency of indigenous yeast strains of different origin on orange juice. *African Journal of Biotechnology*. 4(11): 1290-1296.
- Xie, G.Y., Ma, J., Guan, L., Liu, X-M., Wang, A. and Hu, C. 2018. Proliferation effects of cinnamon extract on human HeLa and HL-60 tumor cell lines. *European Review for Medical and Pharmacological Science*. 22(16): 5347-5354.
- Yadav, Preeti., Garg, Neelima. and Diwedi, D.H. 2009. Effect of location of cultivar, fermentation temperature and additives on the physico-chemical and sensory qualities on mahua (*Madhuca indica* J. F. Gmel.) wine preparation. *Natural Product Radianc*. 8(4): 406-418.
- Yang, X.Q., Zheng, H., Ye, Q., Li, R.Y. and Chen, Y. 2015. Essential oil of cinnamon exerts anti-cancer activity against head and neck squamous cell carcinoma via attenuating epidermal growth factor receptor - tyrosine kinase. *Journal of Balkan Union of Oncology*. 20(6): 1518-1525.
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