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# Effect of drying methods on chemical constituents and flour of coriander (*Corianderum sativum*) leaves

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#### Abstract

The present study was undertaken to study the effects of blanching and drying methods on two different coriander varieties *viz* local variety (V1) and hybrid variety (V2). Sun and cabinet drying were employed post blanching which was carried out at  $85^{\circ}$  C for 3.5 min. Blanching decreased the total chlorophyll, chlorophyll b, total phenol, total carotenoid, ascorbic acid, essential oil content and luminosity of dried leaves while increased chlorophyll a content, redness, yellowness and total solids in dried samples. Cabinet drying recorded 36.34% and 32.80% decrease in total chlorophyll content of V1 and V2 respectively against 44.16% decrease in V1 and 38.58% in V2 for total chlorophyll content during sun drying. However, a higher percentage of total phenol content was lost in cabinet dried samples as compared to sun dried samples for both coriander varieties. The study revealed minimum loss of nutrients in cabinet dried leaves and thus cabinet drying at 60° C is better technique for drying of coriander leaves. Storage studies of the dried leaves revealed that they can be safely stored for 3 months in LDPE pouches under ambient conditions.

Keywords: Blanching, cabinet dryer, coriander, LDPE, sun drying

#### Introduction

Coriander (Corianderum sativum) is an annual herbaceous plant that belongs to the family Umbelliferae. Coriander or "dhanaya" is a culinary and medicinal crop which finds extensive use in food and pharmaceutical industry worldwide for its rich nutritional composition <sup>[1]</sup>. The leaves of plant are rich in volatile oils, flavonoid glycosides (quercetin, iso-quercetin, and rutin), caffenic acid, minerals (calcium, phosphorus and iron), carotene, fibre, and carbohydrates. It also contains essential vitamins and minerals in profuse amounts <sup>[2]</sup>. Coriander is also known for its therapeutic effects. Various researchers have reported antioxidant, anti-bacterial, anti-fungal, anti-cholinesterase activities as well as memory improving power and cholesterol lowering action of coriander extracts <sup>[3]</sup>. Fresh leaves are used to garnish various cuisines and mask odour of various foods. Moreover, they act as appetite stimulant and flavouring agent. However, the high perishability and huge post-harvest losses of coriander limits its availability to every corner of world round the year and thus lowers its commercial value. One of the ways to increase the shelf-life of coriander and make it available throughout the year is by drying the leaves as the demand for high quality, minimally processed, shelfstable dried vegetables are on rise. Drying is the most common and widely practised method for shelf-life extension of coriander leaves. Drying inhibits the growth and proliferation of micro-organisms<sup>4</sup>. In addition, leaves undergo blanching before drying which improves their sensorial characteristics. Several studies have been conducted in past on drying behaviour of coriander leaves. Hihat et al.<sup>[5]</sup>. Studied the effect of oven and microwave drying on phenolic content and anti-oxidant activity of coriander leaves. Khanum et al. [6] investigated the antioxidant activity and mineral content of dried and fresh coriander leaves. Effect of drying on essential oil content of coriander leaves was studied by Pirbaloutia et al. [7]. Yet none of the research work provides a suitable drying method for coriander leaves. Thus, present work was undertaken to study the effect of sun and cabinet drying as well as blanching on physical characteristics and biochemical constituents of fresh coriander leaves. Sun drying and cabinet drying were employed because of low application cost and easy handling.

#### **Materials and Methods**

Two coriander varieties were used in this research *viz* local variety (V1) & hybrid variety (SH-DH-1) (V2). Local variety was procured from the markets of Shalimar, Kashmir & the hybrid variety was harvested from the fields of SKUAST -K, Shalimar. The leaves were de-stemmed, washed, and drained. Afterwards, they were blanched at 85° C for 3.5 min, cooled immediately, and dried using sun (3 days) and cabinet dryer (60° C for 6 hours). The dried leaves were then packed in LDPE pouches and stored under ambient conditions (25° C  $\pm$  3) for a period of 60 days. *Total chlorophyll, chlorophyll a and chlorophyll b* Estimation of chlorophyll was done by following the procedure of Jones *et al.* <sup>[8]</sup> using 80% acetone. The absorbance of chlorophyll extracts was measured at 650nm using UV–Vis spectrophotometer (UV 2401 PC, Shimadzu Co., Singapore).

mg Chlorophyll a/g tissue = 12.7 (A<sub>663</sub>) – 2.69 (A<sub>645</sub>) ×  $\frac{V}{1000 \times W}$ 

mg Chlorophyll a/b tissue = 22.9 (A<sub>645</sub>) – 4.68 (A<sub>663</sub>) ×  $\frac{V}{1000 \times W}$ 

mg total chlorophyll = 20.2 (A<sub>645</sub>) – 8.02 (A<sub>663</sub>) ×  $\frac{V}{1000 \times W}$ 

Where

A = absorbance at specific wave length V = Final volume of chlorophyll extract in 80% acetone W = Fresh weight of tissue artesated

W = Fresh weight of tissue extracted

## Total Phenols, total carotenoids and ascorbic acid

The total phenols were estimated according to the Folin-Ciocalteu method<sup>[9]</sup>. For ascorbic acid content determination. 2, 6-dichlorophenol indophenol (DCPIP) titration method described by Rao and Deshpande <sup>[10]</sup> was followed. Carotenoid content of coriander leaves were quantified by following the spectrophotometric procedure given by Zakaria and Simpson <sup>[11]</sup> and absorbance was taken at 450nm. Essential Oil estimation Essential oil content of coriander leaves was estimated by steam distillation method outlined by CHEM 333L Organic Chemistry Laboratory (Revision 2.1). Microbial examination Standard serial dilution and pour plate technique was used for microbial analysis of the samples <sup>[12]</sup>. Instrumental colour Colour measurement of dried coriander leaves was carried out by the following the method of Hunt<sup>13</sup> using a hunter colorimeter model "Lab scan XE" (Hunter associates laboratory, USA) using universal software, based on three colours co-ordinates namely L\*, a\* and b\* value. L\* represents the lightness index, a\* represents red-green, while b\* represents yellow-blue colour components. The instrument was calibrated using a standard white and black reference tile. Sensory evaluation A group of 10 semi-trained judges carried out the sensory evaluation of dried coriander leaves using 5point scale. The dried leaves were dissolved in water and the solution was subjected to organoleptic testing. Before judges could commence the test, they were acquainted with the use of rating method, terminology for each attribute and sensory characteristics. Samples were randomly presented to judges and were asked to rate them on the basis of flavour, colour and taste. Overall acceptability was calculated as average of the other parameters. Moisture content, water activity and rehydration ratio Moisture content as determined by laboratory oven method <sup>[14]</sup>. Water activity of the sample was analyzed using a water activity analyzer (Pre-Aqua Lab, Water Activity Analyser). Rehydration ratio was calculated by the below given formula:

Rehydration ratio = 
$$\frac{Wr}{Wd}$$

Where

 $W_r$  = weight of rehydrated sample (g)  $W_d$  = weight of dried sample (g)

## **Total solids**

Total solid of dehydrated coriander leaves were determined by drying the samples in oven and following the method given by Ranganna<sup>[15]</sup>. Total solids percentage was calculated as per the below given equation:

Total solids (%) = 100- moisture content (%).

## **Statistical Analysis**

All the analyses were carried out in triplicate and the results were provided as mean value. Statistical analysis was analysed using SPSS statistics for Windows version 20.0 Armonk, New York. A factorial CRD was employed to test the significance of data at 5% level.

## **Results and Discussion**

Total chlorophyll, chlorophyll a and chlorophyll b Drying method as well as blanching and variety had significant (p < 0.05) effects on total chlorophyll content of coriander leaves (Table 1a). Blanching decreased the total chlorophyll content in case of both the varieties irrespective of drying method. Total chlorophyll content in case of V1 before drying was 3.99mg/100g which decreased significantly to 3.55mg/100g after sun drying and 3.97mg/100g after cabinet drying. Similarly, in case of V2, total chlorophyll content before drying was 3.81mg/100g which decreased to 3.45mg/100g after sun drying and 3.76mg/100g after cabinet drying which signifies higher loss of total chlorophyll content in sun dried leaves of both varieties (Table 1a). This may be adduced to the fact that blanching causes conversion of chlorophyll to stable chlorophyll ides due to action of chlorophyllase enzyme. Similar results have been reported by Ahmed et al.<sup>[4]</sup> for drying characteristics of coriander leaves. In case of unblanched samples, sun as well as cabinet drying caused a significant (p < 0.05) decrease in total chlorophyll content of V1 from an initial value of 3.99mg/100g to 2.21mg/100g and 2.54mg/100g respectively. Likewise, a decrease from 3.81mg/100g to 2.34mg/100g and 2.56mg/100g was recorded for V2 during sun and cabinet drying respectively. This might be due to exposure of coriander leaves to less drying time and controlled temperatures in cabinet dryer as compared to sun. Higher drying time causes faster conversion of chlorophyll to brown pigment pheophytins [4]. A similar result has been observed by Rocha et al. [16] on the drying of basil. Chlorophyll a as well as chlorophyll b content of both varieties were significantly affected by all the three parameters. Blanching significantly (p < 0.05) increased chlorophyll a in V1 from 2.34mg/100g to 2.36mg/100g in sun dried samples and 2.63mg/100g in cabinet dried leaves while decreased chlorophyll b content from 1.65mg/100g to 1.34mg/100g and 1.19mg/100g when both the drying techniques were employed. Likewise for V2, blanching enhanced chlorophyll a content from 2.12mg/100g to 2.25mg/100g in case of sun drying and 2.56mg/100g for cabinet dried leaves while reduced chlorophyll b content from 1.69mg/100g to 1.20mg/100g when both drying techniques were employed which signifies higher retention of chlorophyll a in cabinet dried samples as compared to sun dried leaves. This might be due to exposure of coriander

leaves to less drying time and controlled temperatures in cabinet dryer as compared to sun. Higher drying time causes faster conversion of chlorophyll to brown pigment pheophytin <sup>[4]</sup>. A similar result has been observed by Rocha *et al.* <sup>[16]</sup> on the drying of basil.

For unblanched samples, sun drying resulted in 41% decrease in V1 and 46% loss in V2 while cabinet drying caused only 34% and 36% loss in V2 which implies higher retention of chlorophyll a in cabinet dried samples. A similar reduction of 63.03% and 39.39% and 28.40% and 27.81% in chlorophyll b content of unbleached sun and cabinet dried samples of V1 and V2 respectively was recorded. A rapid degradation in chlorophyll a content of sun-dried coriander leaves was observed as compared to chlorophyll b content of sun-dried coriander leaves (Table 1). This is due to higher sensitivity of Chlorophyll a to pheophytinization <sup>[17]</sup>.

## **Total phenols**

Total phenolic content of coriander leaves was significantly (p < 0.05) influenced by drying method and blanching treatment. The phenolic content of fresh coriander leaves of V1 was recorded as 62.41mg/100g which decreased to 43.39mg/100g and 16.42mg/100g after sun and cabinet drying of blanched samples respectively. Similar decrease from 60.07mg/100g to 45.02 and 16.12mg/100g after sun and cabinet drying treatments was recorded in case of blanched hybrid (V2) coriander leaves (Table 1a). Some phenolic compounds are bound to cells of the plant which limit their solubility in water. Blanching promotes leaching of phenolics in water <sup>[18]</sup> by disrupting the cell wall material of plant cells, thereby releasing the phenolic compounds into water <sup>[19]</sup>. The other possible reason for decrease in phenolic content upon blanching may be due to the formation of phenolic complexes with other leaf constituents (carbohydrates, proteins, antinutritional factors) which limit their extractability <sup>[20]</sup>. Similar decrease in phenolic compounds have been also reported by Ironic et al.<sup>[21]</sup> in blanched Adansonia digestate. In case of unbranched samples, sun drying lead to 30.12% and 24.55% reduction whereas cabinet drying showed 73.14% and 72.78% loss in total phenol content of V1 and V2 respectively. Thus, a predominant decrease in total phenolic content was recorded in case of cabinet drying as compared to sun dying for both the varieties (Table 1a). This decline in phenolics is adduced to intense and prolonged heating during cabinet drying which leads to thermal degradation of phenolic compounds during <sup>[22]</sup> as phenolics decompose readily at elevated temperatures <sup>[23]</sup>. In addition, during drying, phenolic compounds undergo oxidation in presence of polyphenol oxidase (PPO), which results in intermolecular condensation reactions thus, reducing their content <sup>[24]</sup>. Katsube *et al.* <sup>[25]</sup> and Chan *et al.* <sup>[26]</sup> have reported a similar decline in poly phenolic content of mulberry leaves and ginger leaves after drying respectively. Francisco et al. (2010)<sup>[19]</sup> reported that some phenolic compounds are known to be insoluble form in combination with plant cell wall components. During blanching (high temperature 90° C and time (10-15 min),

disruption of the cell wall of the plant may occur leading to leaching out of the soluble phenolic Francisco *et al.* (2010)<sup>[19]</sup> reported that some phenolic compounds are known to be insoluble form in combination with plant cell wall components. During blanching (high temperature 90° C and time (10-15 min), disruption of the cell wall of the plant may occur leading to leaching out of the soluble phenolic Francisco *et al.* (2010)<sup>[19]</sup> reported that some phenolic compounds are known to be insoluble form in combination

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oil All the three factors viz variety, drying method and blanching had a significant (p < 0.05) effect on essential oil content of coriander leaves. The essential oil content of fresh coriander leaves of V1 was found to be 0.16% and 0.12% for V2. Blanching decreased the essential oil content in V1 from 0.16% to 0.021% and 0.026% in sun and cabinet drying respectively. A similar decrease in essential oil content of V2 coriander leaves was observed from 0.12% to 0.018% in sun dried samples and 0.020% in cabinet dried leaves (Table 1a). This decrease in essential oil content of coriander leaves during blanching may be attributed to high temperatures<sup>33</sup> as high temperature causes breakage of oil cells, subsequently leading to loss of volatile oil <sup>[34]</sup>. In unblanched leaves of V1, a reduction of 85% and 82.5% and for V2, a loss of 81.66% and 75.83% of essential oil was recorded during sun and cabinet drying respectively. During drying process, essential oil along with water is dragged to leaf surface mainly by diffusion which explains the loss of essential oil content during drying <sup>[35]</sup>. Saied et al. <sup>[36]</sup> also reported higher loss of essential oil content when Mentha long folia leaves were dried under sun. Colour the colour parameters for fresh leaves of both the coriander varieties varied significantly (p < 0.05) (Table 1b). Blanching had a significant (p < 0.05) decreasing effect on L\* value of coriander leaves from 37.35 for fresh V1 coriander leaves to 25.36 and 34.20 in case of sun and cabinet dried coriander leaves respectively. Similarly, L\* value of V2 coriander leaves also decreased with blanching from 39.36 to 25.92 and 34.36 during sun and cabinet drying respectively. Excessive loss of natural colour pigments due to leaching during blanching increased lightness of the leaves <sup>[27]</sup>. The other possible reason for decrease in luminosity of dried coriander leaves might be formation of chlorophyll ides which increase the colour saturation of coriander leaves <sup>[37]</sup>. In unbleached leaves of V1 and V2, sun drying reduced L\* value by 29.26% and 31.60% while cabinet drying caused 1.7% and 7.26% reduction in luminosity of coriander leaves. At elevated drying temperatures coriander leaves undergo nonenzymatic browning which decreased the lightness in leaves <sup>[38]</sup>. Drying as well as blanching significantly effected a\* value of coriander leaves. Blanching enhanced the redness of V1 coriander leaves from -6.34 to -4.26 and -5.77 in case of sun and cabinet dried leaves respectively. For V2, redness of sun and cabinet dried leaves increased from -7.67 to -5.24 and -6.74 respectively. High temperature promotes non-enzymatic browning reaction, and turn the samples less greenish<sup>38</sup>. In unblanched samples, a 73.34% and 12.46% increase in redness and 13.21% and 2.04% decrease in b\* value of V1 while 68.05% and 27.11% increase in redness and 7.18% and 2.86% decrease in greenness of V2 variety was recorded during sun and cabinet drying respectively. Blanching causes degradation of chlorophyll to pheophytin which increases the yellowness of dried leaves <sup>[17]</sup>. Dwivedy *et al.* <sup>[39]</sup> found that after microwave drying of Indian Borage leaves all the three chromatic coordinates values decreased, but the reduction in L\* values was not significant.

## **Total solids**

Drying, blanching as well as variety had a significant (p<0.05) effect on total solid content of dried coriander leaves (Table 1b). The total solid content of fresh coriander leaves of V1 was 14.91% which increased to 86.63% and 86.90% in

case of sun and cabinet dried blanched leaves respectively. In V2 blanched leaves, total solid content increased from 14.67% to 86.71% in case of sun drying and 86.87% in case of cabinet dried leaves. Moreover, for unblanched leaves, an increase in 82.81% and 82.83% total solid content of V1 samples was observed while V2 leaves recorded 83.06% and 83.07% increase in total solid content during sun and cabinet drying which implied a predominant increase in total solid content of dried leaves during cabinet drying. The pronounced increase in total solids of coriander leaves during cabinet drying may be due to greater loss of moisture. An efficient and accelerated heat transfer and mass transfer rates are encountered during cabinet drying, which removes moisture even from the centre of material <sup>[40]</sup>. Storage Studies Biochemical evaluation of different coriander varieties after sun and cabinet drying revealed that blanched cabinet dried samples proved superior in terms of quality. In order to check the shelf stability of dried coriander leaves, the sample was packed in LDPE and stored under ambient conditions ( $25^{\circ}C \pm$ 3) for a period of 90 days. The stored samples were analysed for moisture gain, water activity, microbial analysis (bacterial count and fungi count), rehydration ratio and sensory evaluation (on 5-point scale) at an interval of 30 days. The change in selected quality attributes during storage are depicted in Table 2. Storage period had a significant (p < .05) effect on all the quality parameters. Moisture content of the dried coriander leaves increased from 13.10% at the beginning to 16.51% at the end of storage. This gain in moisture content of samples over storage period is because of the hygroscopic nature of the dried leaves [41]. In addition, ingress of water vapour through micro cracks which develops in packaging material also contributes to moisture gain <sup>[42]</sup>. Similar result has been observed by Razak et al. <sup>[43]</sup> during storage of Orthosiphon stamineus dried leaf. A significant (p < 0.05) increase in water activity of dried coriander leaves was reported over storage time of 90 days. At the beginning of storage, water activity was 0.44 which increased up to 0.61 with the advancement of storage. This increase in water activity is due to increase in moisture content of dried leaves <sup>[44]</sup>. The bacterial and fungi count of dried samples were recorded as 3 cfu/g and 0.31cfu/g respectively at 0<sup>th</sup> day of storage which. Increased to 250cfu/g and 3.10cfu/g respectively at the completion of storage period of 90 days. This increase in microbial flora of dried leaves is due to increase in water activity and moisture content of samples with storage45. Rehydration ratio of the dried leaves was significantly (p < 0.05) affected by storage time. At the beginning of storage, rehydration ratio of 1.60 was recorded which increased to 2.21 at the end of 90th day of storage. A gradual increase in rehydration ratio over time can be adduced to increased porosity of the cellular structure of dried leaves46. Similar results have been reported by Khedkar and Roy47 in mango slices. Storage period had a significant effect on overall acceptability of dried coriander leaves. Overall acceptability of dried samples decreased over storage period of 90 days from 3.76 to 2.32 which may be ascribed to loss of crisp texture due to ingress of moisture content by dried leaves. In addition, the stored leaves lost colour and flavour upon reconstitution with the storage time which further decreased the acceptability of dried coriander leaves.

Drying Method (D)	Total Chl	orophy	vll (mg	/100g)	Chlor	ophyll	a (mg	/100g)	Chlore	ophyll I	b (mg/	100g)	Total	phenol	l (mg/	100g)	Total carot	tenoid	s (mg/1	100g)	Ascorbi	c acid (	(mg/10	0g)	F	Essential	oil (%	,)
	1		V	/2	V	/1		2	1			2	1			V2	V1		V	/2	V1			V2	V	/1	,	V2
Fresh coriander leaves	3.99	)	3.	81	2.	34	2.	12	1.6	65	1.	69	62.4	41	60	0.07	31.89		32	.20	113.3	0	10	07.0	0.	16	0	0.12
	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2
Sun (D1)	3.55	2.21	3.45	2.34	2.36	1.60	2.25	1.13	1.34	0.61	1.20	1.21	43.39	43.60	45.02	45.32	12.24	12.54	12.26	13.20	23.42	24.01	22.71	23.07	0.021	0.024	0.018	0.022
% increase or decrease	11.02	44.61	9.44	38.58	0.85	31.62	6.13	46.69	18.78	63.03	29	28.40	30.47	30.13	25.05	24.55	61.61	60.67	61.92	59	79.32	78.80	78.77	78.43	86.87	85	85	81.66
Cabinet (D2)	3.97	2.54	3.76	2.56	2.63	1.54	2.56	1.34	1.19	1.00	1.20	1.22	16.42	16.76	16.12	16.35	22.87	23.21	22.18	22.92	46.40	51.39	46.76	51.98	0.026	0.028	0.020	0.029
% increase or decrease	0.50	36.34	1.31	32.80	12.39	57.26	20	36.80	27.87	39.39	29	27.81	73.69	73.14	73.16	72.78	28.28	27.21	31.11	28.81	59.04	54.64	56.30	51.42	83.75	82.5	83.33	75.83
C.D. (p<0.05)	D: 0.032 0.045; T 0.045; T >	V: 0.03 < V: 0.0	32; D × 045; D	<b>V</b> :	0.048 0.048	3; V: 0. ; T × V	.034; E 7: 0.04	0 × V: 8; D ×		; V: 0.0 Γ × V: 0	032; D 0.045;	$\times$ V:	0.092;	; V: 0.0 × V: N	)65; D J.S; D	$\times$ V:	D: 0.025; 0.035; V: 0.0	25; D	$\times$ V: 0.	.035; T	D: 0.025 0.035; V: 0 T × V: 0.03	.025; E	$\mathbf{V} \times \mathbf{V}$ :	0.035;	V: 0.00	2; T: 0.00 )2; D × V S; D × T	': N.S;	$T \times V$ :
		V: 0.0	63			$T \times V$	: 0.068			$\times$ V: 0	.063			V: N	N.S													

## Table 1a: Effect of drying method and blanching on various biochemical constituents of coriander leaves.

V1: Local variety; V2: T1: blanched; T2: unblanched; C.D: Critical Difference

## Table 1b: Effect of drying method and blanching on physical properties of coriander leaves

Drying Method (D)	L*				a*				b*				Total Solids			
	V1			V2	V1		V	/2	V1		V	2	V1		V	2
Fresh coriander leaves	37.35		39.36		-6.34		-7.67		28.38		30.75		14.91		14.67	
	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2	T1	T2
Sun (D1)	25.36	26.42	25.92	26.92	-4.26	-1.69	-5.24	-2.45	28.45	24.62	32.41	28.54	86.63	86.77	86.71	86.60
% increase or decrease	32.10	29.26	34.19	31.60	32.80	73.34	31.68	68.05	2.4	13.21	5.39	7.18	82.78	82.81	83.08	83.06
Cabinet (D2)	34.20	36.70	34.36	36.50	-5.77	-5.55	-6.74	-5.59	30.33	27.80	32.72	29.87	86.90	86.85	86.87	86.67
% increase or decrease	8.43	1.7	12.70	7.26	9	12.46	12.12	27.11	6.87	2.04	6.40	2.86	82.84	82.83	83.11	83.07
	D: 0.034; T	: 0.034;	$D \times T$ : 0.	048; V: 0.034;	D: 0.030; T: 0.03	$0; D \times $	T: 0.043	3; V:	D: 0.032; T: 0.03	$32; D \times 32$	T: 0.045	5; V:	D: 0.015; T	: 0.015;	$D \times T$ : 0.0	022; V:
C.D. (p<0.05)	$D \times V$ : 0.048; $T \times V$ : 0.048; $D \times T \times V$ :				$0.030; D \times V: 0.043; T \times V: 0.043; D \times T$			; $\mathbf{D} \times \mathbf{T}$	$0.032; D \times V: 0.045; T \times V:$			; $\mathbf{D} \times \mathbf{T}$	$0.015; D \times V: 0.022; T$		T × V: 0.022; D ×	
		(	0.068		× V: 0.061				× V: 0.063					031		

V1: Local variety; V2: T1: blanched; T2: unblanched; C.D: Critical Difference

#### Conclusion

The results of the study revealed that blanching treatment retained various nutrients in the leaves and enhanced the colour of dried leaves. Moreover, cabinet drying was found superior to sun drying in maintaining biochemical and physical properties of leaves. Storage studies of the dried leaves revealed that they can be safely stored for 3 months in LDPE pouches under ambient conditions. The study can help in drying of various other leaves of underutilized plants with several medicinal benefits.

Table 2: Effect of storage time on quality parameters of dried coriander leaves

Storage Time	Moisture content (%)	aw	Bacterial count(cfu/g × 10 <sup>2</sup> )	Fungal count(cfu/g)	<b>Rehydration ratio</b>	<b>Overall acceptability</b>
0 <sup>th</sup> day	13.10	0.44	0.03	0.31	1.60	3.76
30 DAS	14.22	0.49	0.06	1.54	1.80	3.40
60 DAS	15.30	0.54	1.45	2.28	2.05	2.70
90 DAS	16.51	0.61	2.50	3.10	2.21	2.32
C. D (p<0.05)	0.601	0.066	0.081	0.085	0.096	0.086

DAS: Days after storage; cfu: colony forming unit; C.D: Critical Difference

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