

Food and Agriculture Organization of the United Nations



# Residue Monograph prepared by the meeting of the Joint FAO/WHO Expert Committee on Food Additives (JECFA), 87th Meeting 2019

# β-apo-8'-CAROTENAL

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# β -apo-8'-CAROTENAL

|                    | Prepared at the 87 <sup>th</sup> JECFA (2019) and published in FAO<br>Monographs 23 (2019), superseding specifications prepared at the<br>74thJECFA (2011) and published in FAO Monographs 11 (2011). A<br>group ADI of 0-5 mg/kg bw expressed as the sum of carotenoids<br>including $\beta$ -carotene, $\beta$ -apo-8'-carotenal, and the methyl and ethyl<br>esters of $\beta$ -apo-8'-carotenoic acid was established at the 18 <sup>th</sup><br>JECFA (1974).  |
|--------------------|---|
| SYNONYMS           | CI Food Orange 6; CI (1975) No. 40820; INS No. 160e   |
| DEFINITION         | These specifications apply to $\beta$ -apo-8'-carotenal which consists<br>predominantly of all-trans- $\beta$ -apo-8'-carotenal and may also contain<br>minor quantities of other carotenoids such as all-trans-<br>crocetindialdehyde, all-trans- $\beta$ -apo-12'-carotenal and all-trans- $\beta$ -<br>carotene. Commercial preparations of $\beta$ -apo-8'-carotenal intended<br>for use in food are prepared from $\beta$ -apo-8'-carotenal meeting these<br>specifications and are formulated as suspensions in edible oil,<br>emulsions and water dispersible powders. These preparations may<br>also contain cis isomers.   |
| Chemical names     | ß-Apo-8'-carotenal, 8'-apo-ß-carotene-al<br>2E,4E,6E,8E,10E,12E,14E,16E)-2,6,11,15-tetramethyl-17-(2,6,6-<br>trimethyl-1-cyclohexenyl)heptadeca-2,4,6,8,10,12,14,16-octaenal  |
| C.A.S. number      | 1107-26-2   |
| Chemical formula   | C <sub>40</sub> H <sub>40</sub> O   |
| Structural formula | All- <i>trans</i> -β-apo-8'-carotenal (main compound)   |
|                    | $\begin{array}{c} H_3C \\ H_3C \\ H_3 \\ H_3$ |
| Formula weight     | 416.65  |
| Assay              | Not less than 96% total colouring matters   |
| DESCRIPTION        | Deep violet crystals with metallic lustre or crystalline powder;<br>sensitive to oxygen and light and should therefore be kept in a light-<br>resistant container under inert gas.  |
| FUNCTIONAL USES    | Colour  |

## CHARACTERISTICS

**IDENTIFICATION** 

| <u>Solubility</u> (Vol. 4)  | Insoluble in water; slightly soluble in ethanol; sparingly soluble in vegetable oils.  |
|---|--|
| <u>Spectrophotometry</u><br>(Vol. 4)                                  | Determine the absorbance of the diluted sample solution used in the Method of Assay at 460 nm and 488 nm. Determine the absorbance at 332 nm of a solution containing a ten-fold higher concentration as that of the diluted sample solution used in the Method of Assay. The ratio $A_{488}/A_{460}$ is between 0.77 and 0.85. The ratio of $A_{332}/A_{460}$ is between 0.63 and 0.75.   |
| PURITY  |  |
| <u>Sulfated ash</u> (Vol. 4)  | Not more than 0.1%<br>Test 2 g of the sample (Method I)  |
| Carotenoids other<br>than ß-apo-8'-<br>carotenal                      | Not more than 3% of total colouring matters.<br>See description under TESTS  |
| <u>Lead</u> (Vol. 4)  | Not more than 2 mg/kg.<br>Determine using a method appropriate to the specified level. The<br>selection of sample size and method of sample preparation may be<br>based on the principles of the methods described in Volume 4<br>(under "General Methods, Metallic Impurities").  |
| TESTS   |  |
| PURITY TESTS  |  |
| <u>Carotenoids other</u><br><u>than ß-apo-8'-</u><br><u>carotenal</u> | <ul> <li>Chromatographic system</li> <li>HPLC equipped with a UV/Vis detector or a photodiode array detector, refrigerated auto sampler and integrator</li> <li>Detector wavelength: 463 nm</li> <li>Column: Reverse phase C18; Suplex pkb-100 (250 x 4.6 mm, 5 μm) from Supelco or equivalent</li> <li>Mobile phase: In a 1000 ml volumetric flask, dissolve 50 mg BHT in 20 ml 2-propanol and add 0.2 ml N-ethyldiisopropylamine, 25 ml 0.2% aqueous ammonium acetate solution, 455 ml acetonitrile, and 3pprox 450 ml methanol. Mixture cools and contracts. Allow to reach room temperature and dilute to volume with methanol. Discard after 2 days.</li> <li>Isocratic elution</li> <li>Column temperature: 30°</li> <li>Flow rate: 0.6 ml/min</li> <li>Injection volume: 10 μl</li> <li>Temperature of the autosampler: (3pprox 15°)</li> </ul> |

Run time: 4pprox.. 35 min

#### **Reagents**

- Butylated hydroxytoluene (BHT), reagent grade
- 2-Propanol, HPLC grade
- N-ethyldiisopropyl-amine, reagent grade
- Ammonium acetate, reagent grade
- Acetonitrile, HPLC grade
- Methanol, HPLC grade
- Ethanol, HPLC grade
- Tetrahydrofuran, HPLC grade

### Sample solution

Weigh accurately (to ±0.1 mg) 0.010 g of the sample and dissolve in tetrahydrofuran (stabilized with 0.025% BHT). Transfer to a 100 ml volumetric flask and bring to volume with tetrahydrofuran. Dilute to the ratio of 1:10 with ethanol.

#### **Procedure**

Inject the sample solution using the conditions detailed under *Chromatographic system*. The retention times for all-*trans*- $\beta$ apo-8'-carotenal is in the range of 7-9 min and corresponds to the largest peak in the chromatogram. The relative retention times of minor carotenoids with respect to the retention time of all-*trans*- $\beta$ -apo-8'-carotenal are: all-*trans*crocetindiadebyde (0.54); all-*trans*- $\beta$ -apo-12'-carotenal

crocetindialdehyde (0.54); all-*trans*- $\beta$ -apo-12'-carotenal (0.84); all-*trans*- $\beta$ -carotene (2.55).

Integrate the areas of the peaks in the chromatogram.

#### **Calculation**

Calculate the percentage of carotenoids other than  $\beta$ -apo-8'carotenal (%, w/w) using the following formula:

Carotenoids other than 
$$\beta$$
 – apo – 8' – carotenal  $\left(\%, \frac{w}{w}\right)$   
=  $\frac{A_{\text{total}} - A_{\beta-\text{apo}-8'-\text{carotenal}}}{A_{\text{total}}}$ 

where

A<sub>total</sub> is the sum of the area of all the peaks in the chromatogram, excluding the solvent peak (area units); and

 $A_{\beta-apo-8'-carotenal}$  is the area of the peak of  $\beta$ -apo-8'-carotenal in the chromatogram (area units).

## METHOD OF ASSAY <u>Total colouring matters content by spectrophotometry</u>

Proceed as directed under Total Colouring Matters Content – Colouring Matters Content by Spectrophotometry, Procedure 2, using the following conditions: Sample weight (W): 0.08 g ( $\pm$ 0.01 g); Volume of the three volumetric flasks: V<sub>1</sub> = V<sub>2</sub> = V<sub>3</sub> = 100 ml; Volume of the two pipets:  $v_1 = v_2 = 5$  ml; Specific absorbance of the standard:  $A_{1cm}^{1\%} = 2640$ ; Wavelength of maximum absorption:  $\lambda_{max}$  about 461 nm. <u>Calculation</u> Calculate the percentage of total colouring matters using the following formula:

Total colouring matters (%, w/w) = 
$$\frac{A \times V_1 \times D}{A_{1cm}^{1\%} \times W}$$

where

A is the absorbance of the twice-diluted sample solution at 461  $\,\mathrm{nm};\,\mathrm{and}$ 

D is the dilution factor  $(V_2 x V_3)/(v_1/v_2)$ .