

2
3 **Method Name: Determination of Lycopene in Infant and Adult/ Pediatric**
4 **Nutritional Formula**

5
6 **Approved by:** Stakeholder Panel for Infant Formula and Adult Nutritionals

7 **Final version date:**

8 **Effective date:**

9
10 **Intended Use:** Reference method for dispute resolution.

11
12 **1. Applicability:**

13 Determination of total¹ Lycopene (CAS 502-65-8) in all forms of infant, adult, and/or
14 pediatric formula (powders, ready-to-feed liquids, and liquid concentrates).

15
16 **2. Analytical Technique:**

17 Any analytical technique that meets the following method performance
18 requirements is acceptable.

19
20 **3. Definitions:**

21 Accuracy (Corresponds to the VIM definition for “trueness”).

22 The closeness of agreement between the average of an infinite number of replicate
23 measured quantity values and a reference quantity value.

24
25 **Adult/Pediatric Formula**

26 Nutritionally complete, specially formulated food, consumed in liquid form, which may
27 constitute the sole source of nourishment [AOAC Stakeholder Panel on Infant
28 Formula and Adult Nutritionals (SPIFAN); 2010], made from any combination of milk,
29 soy, rice, whey, hydrolyzed protein, starch, and amino acids, with and without intact
30 protein.

31
32 **Infant formula**

33 Breast-milk substitute specially manufactured to satisfy, by itself, the nutritional
34 requirements of infants during the first months of life up to the introduction of
35 appropriate complementary feeding (Codex Standard 72 – 1981), made from any
36 combination of milk, soy, rice, whey, hydrolyzed protein, starch, and amino acids,
37 with and without intact protein.

38
39 **Limit of Detection (LOD)**

40 The minimum concentration or mass of analyte that can be detected in a given matrix
41 with no greater than 5% false positive risk and 5% false negative risk.

42
43 **Limit of Quantitation (LOQ)**

44 The minimum concentration or mass of analyte in a given matrix that can be reported
45 as a quantitative result

46
47 **Lycopene**

48 IUPAC name: (6E,8E,10E,12E,14E,16E,18E,20E,22E,24E,26E)-
49 2,6,10,14,19,23,27,31-octamethyldotriaconta-2,6,8,10,12,14,16,18,20,22,24,26,30-
50 tridecaene, CAS number: 502-65-8. Figure 1.

51
52

¹ Include *cis* and *trans* isomers if they are separated

53 Repeatability
54 Variation arising when all efforts are made to keep conditions constant by using
55 the same instrument and operator, and repeating during a short time period.
56 Expressed as the repeatability standard deviation (SD_r); or % repeatability
57 relative standard deviation (%RSD_r).

58
59 Reproducibility
60 The standard deviation or relative standard deviation calculated from among-
61 laboratory data. Expressed as the reproducibility relative standard deviation
62 (SD_R); or % reproducibility relative standard deviation (% RSD_R).

63
64
65
66
67
68

4. **Method Performance Requirements:**
See Table 1.

Table 1. Method Performance requirements^a

Analytical range	1–50 ^b
Limit of Quantitation (LOQ)	≤ 1 ^b
Recovery	90-110%
Repeatability (RSD _r)	8%
Reproducibility (RSD _R)	15%
^a Concentrations apply to: a) 'ready-to-feed' liquids "as is"; b) re-constituted powders (25 g into 200 g of water); and c) liquid concentrates diluted 1:1 by weight.	
^b μg /100 g reconstituted final product	

69
70
71
72
73
74
75
76
77
78
79
80
81
82
83
84
85

5. **System suitability tests and/or analytical quality control:**
Suitable methods will include blank check samples, and check standards at the lowest point and midrange point of the analytical range. Methods must be capable of resolving lycopene from α-carotene and β-carotene.

6. **Reference Material(s):**
SRM 1869. Please contact Dr. Melissa Phillips, Research Chemist, NIST for materials at melissa.phillips@nist.gov or (301) 975-4134.

7. **Validation Guidance:**
Recommended level of validation: *Official Methods of Analysis*SM.

8. **Maximum Time-To-Result:** No maximum time.

86 Figures:

87

88

89

90

91

92

93

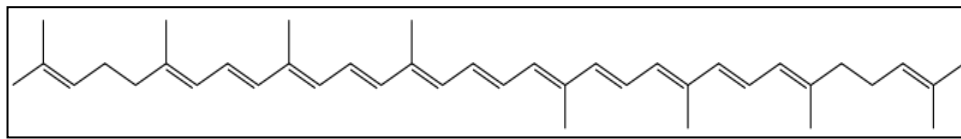


Figure 1: Molecular structure of lycopene.

DRAFT