CALCIUM CHLORIDE

Prepared at the 19th JECFA (1975), published in NMRS 55B (1976) and in FNP 52 (1992). Metals and arsenic specifications revised at the 63rd JECFA (2004). An ADI 'not limited' was established at the 17th JECFA (1973)

SYNONYMS INS No. 509

DEFINITION

- Chemical names Calcium chloride
- C.A.S. number 10043-52-4
- Chemical formula Anhydrous: $CaCl_2$ Dihydrate: $CaCl_2 \cdot 2H_2O$ Hexahydrate: $CaCl_2 \cdot 6H_2O$
- Formula weight Anhydrous: 110.99 Dihydrate: 147.02 Hexahydrate: 219.08
- Assay Anhydrous: Not less than 93% Dihydrate: Not less than 99.0% and not more than the equivalent of 107.0% of $CaCl_2 \cdot 2H_2O$ Hexahydrate: Not less than 98.0% and not more than the equivalent of 110% of $CaCl_2 \cdot 6H_2O$
- **DESCRIPTION** Anhydrous: White, deliquescent lumps or porous masses Dihydrate: White, hard, deliquescent fragments or granules Hexahydrate: Colourless, very deliquescent crystals

FUNCTIONAL USES Firming agent

CHARACTERISTICS

IDENTIFICATION

- <u>Solubility</u> (Vol. 4) Anhydrous: Freely soluble in water and ethanol Dihydrate: Freely soluble in water; soluble in ethanol Hexahydrate: Very soluble in water and ethanol
- Test for chloride (Vol. 4) Passes test
- Test for calcium (Vol. 4) Passes test

PURITY

Free alkaliNot more than 0.15% as Ca(OH)2Dissolve 1 g of the sample in 20 ml of freshly boiled and cooled water, and
add 2 drops of phenolphthalein TS. If the solution is pink, the pink colour is
discharged by adding 2 ml of 0.02 N hydrochloric acid.

<u>Magnesium and alkali</u> <u>salts</u>	Not more than 5% Dissolve 1 g of anhydrous calcium chloride, or the corresponding weight of a hydrate, in about 50 ml of water, add 500 mg of ammonium chloride, mix and boil for about 1 min. Quickly add 40 ml of oxalic acid TS, and stir vigorously until precipitation is well established. Immediately add 2 drops of methyl red TS, then add ammonia TS dropwise until the mixture is just alkaline, and cool. Transfer the mixture into a 100-ml cylinder, dilute with water to 100 ml, let stand for 4 h or overnight, and then decant the clear, supernatant liquid through a dry filter paper. To 50 ml of the clear filtrate in a platinum dish add 0.5 ml of sulfuric acid and evaporate the mixture on a steam bath to a small volume. Carefully evaporate the remaining liquid to dryness over a free flame, and continue heating until the ammonium salts have been completely decomposed and volatilized. Finally, ignite the residue to constant weight. The weight of the residue does not exceed 25 mg.
Fluoride (Vol. 4)	Not more than 40 mg/kg (Method III)
<u>Lead</u> (Vol. 4)	Not more than 2 mg/kg Determine using an atomic absorption technique appropriate to the specified level. The selection of sample size and method of sample preparation may be based on the principles of the method described in Volume 4, "Instrumental Methods."
METHOD OF ASSAY	Weigh accurately about 1 g of anhydrous calcium chloride, or the corresponding weight of a hydrate, transfer to a 250-ml beaker, and dissolve in a mixture of 100 ml of water and 5 ml of dilute hydrochloric acid TS. Transfer the solution to a 250-ml volumetric flask, dilute with water to volume and mix. Pipet 50 ml of the solution into a suitable container, add 100 ml of water, 15 ml of sodium hydroxide TS, 40 mg of murexide indicator (amm. purpurate) and 3 ml of naphthol green TS, and titrate with 0.05 M disodium ethylenediaminetetra-acetate until the solution is deep blue in colour. Each ml of 0.05 M disodium ethylenediaminetetraacetate is equivalent to 5.55 mg of CaCl ₂ ; 7.35 mg of CaCl ₂ \cdot 2H ₂ O; or 10.95 mg of CaCl ₂ \cdot 6H ₂ O.