

# **Department of Health**Therapeutic Goods Administration

## Compositional guideline for Molybdenum trioxide

## Name of the ingredient

Molybdenum trioxide (AAN)

## Definition of the ingredient

Molecular formula: MoO<sub>3</sub>

Molar mass: 143.94

CAS Number: 1313-27-5

 Table 1.
 Ingredient specific requirements

| Test  | Method reference     | Acceptance criteria  |  |
|---|----------------------|--|--|
| Description   |                      |  |  |
| Appearance  | Visual               | Yellow-green or grey powder  |  |
| Characteristics   |                      |  |  |
| Loss on drying (dried to a constant weight at 100-105 °C) | BP (Appendix IX D)   | Not more than 0.5%   |  |
| Solubility  | BP (General Notices) | Slightly soluble in water. Soluble in concentrated mineral acids, solutions of alkaline hydroxides and ammonia |  |
| Identification  |                      |  |  |
| AAS   | Ph Eur 2.2.23        | Matches spectrum of authenticated reference material   |  |
| Assay   |                      |  |  |
| Molybdenum trioxide                                       | Ph Eur 2.2.23        | 99.5-101.0% (anhydrous basis)  |  |

**Table 2.** Incidental constituents

| Test   | Method reference   | Acceptance criteria     |  |
|--|--------------------|-------------------------|--|
| Incidental metals and non-metals                                       |                    |                         |  |
| Heavy metals (as lead)   | BP (Appendix VII)  | Not more than 10 ppm    |  |
| Arsenic  | Ph Eur 2.4.2       | Not more than 10 ppm    |  |
| Sulfates   | Ph Eur 2.4.13      | Not more than 200 ppm   |  |
| Phosphates   | Ph Eur 2.4.11      | Not more than 5 ppm     |  |
| Ammonium   | BP (Appendix VII)  | Not more than 20 ppm    |  |
| Other organic or inorganic impurities or toxins                        |                    |                         |  |
| Ammonia insoluble matter (10g sample in 120 mL dilute aqueous ammonia) | Literature method¹ | Not more than 0.01% w/w |  |

### **Microbiology**

While substance manufacturers are encouraged to include limits for objectionable microorganisms, it is the product into which those substances are formulated that is subject to a legally binding set of criteria. The Therapeutic Goods Order No. 77 *'Microbiological Standards for Medicines'* mandates that any finished product that contains the ingredient, alone or in combination with other ingredients, must comply with the microbial acceptance criteria set by Clause 9 of the Order.

#### Notes

<sup>1</sup>Reagent Chemicals Ninth Edition, American Chemical Society Specifications

#### **Key to abbreviations:**

AAS = Atomic Absorption Spectrophotometry

BP = British Pharmacopoeia

Ph Eur = European Pharmacopoeia