INTERNATIONAL STANDARD

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Plastics — Determination of xylenesoluble matter in polypropylene

Plastiques — Détermination des matières présentes dans le polypropylène solubles dans le xylène



Reference number ISO 16152:2005(E)

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Foreword

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The main task of technical committees is to prepare International Standards. Draft International Standards adopted by the technical committees are circulated to the member bodies for voting. Publication as an International Standard requires approval by at least 75 % of the member bodies casting a vote.

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights.

ISO 16152 was prepared by Technical Complete ISO/TC 61, *Plastics*, Subcommittee SC 9, *Thermoplastic materials*.

It replaces ISO 6427:1992, Annex B, which has been technically revised. The revised method tightens the physical parameters of the test to provide improved repeatability and reproducibility.



Introduction

This International Standard specifies a method for the quantitative determination of those components of polypropylene that are soluble in xylene. This new method defines more precisely the factors that have the greatest influence on the repeatability and reproducibility of the determination. The polypropylene is dissolved in hot xylene, the cooled under controlled conditions down to 25 °C, which results in the precipitation of the insoluble fraction. The soluble matter remains in the xylene. The xylene is then evaporated and the residue weighed. The solubles content of polypropylene is important as it has a major influence on the properties of the polypropylene.

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Plastics — Determination of xylene-soluble matter in polypropylene

1 Scope

This International Standard specifies a method for determining the mass fraction of a polypropylene 1.1 homopolymer or copolymer which is soluble in xylene at 25 °C.

A weighed amount of dried sample is dissolved in xylene under reflux conditions, then cooled under 1.2 controlled conditions and maintained at 25 °C to ensure controlled crystallization of the insoluble fraction.

The xylene-soluble fraction then recovered by evaporation of the xylene and determined by weighing 1.3 the residue.

similar to that of the xylene-soluble fraction, such as additives, may 1.4 Other materials with solubilities interfere with the determination.

2 Terms and definitions

me and definitions apply. For the purposes of this document, the following to

2.1

xylene-soluble fraction

 S_{s}

that percentage, by mass, of the polymer that does not recipitate out when a solution of the polymer in xylene is cooled from reflux temperature to +25 °C and held at that temperature for a specified period of time

3 Apparatus

3.1 Reflux condenser, length 400 mm.

Flat-bottomed flask, capacity 400 ml, with one or two necks, or conical flask or cylindrical bottle of 3.2 similar capacity.

3.3 Insulating disc, made of fibreglass or mineral wool.

Magnetic stirrer, with temperature-controlled hotplate, thermostatted oil bath or heating block capable 3.4 of maintaining 140 °C to 150 °C.

- 3.5 Stirrer bar.
- 3.6 Pipette, class A, 200 ml or equivalent.
- 3.7 Pipette, class A, 100 ml or equivalent.
- Glass-stoppered flask, 250 ml. 3.8