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Indoor Decorating and Refurbishing Materials – Limit of Harmful Substance of Wood Based Furniture

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Foreword

The Chapter 4 in this Standard is mandatory, while the rest are recommended.

Since January 1, 2002, the products produced by the manufacturer shall implement this Standard; the transition period is 6 months; Since July 1, 2002, the products that don't conform to this Standard shall stop selling on the market.

This Standard was proposed by China Light Industry Council.

This Standard shall be under the jurisdiction of National Standardization Center of Furniture.

Drafting organizations of this Standard: National Standardization Center of Furniture, School of Wood Industry of Nanjing Forestry University, and Beijing Building Materials Academy of Sciences Research.

Participating drafting organizations of this Standard: Tiantan Furniture Branch of Beijing Tiantan Co., Ltd., Beijing Easyhome Decoration Development Co., Ltd., Shenzhen Dafuhao Industrial Development Co., Ltd., Shunde Meihua Home Manufacturing Co., Ltd., and Shunde Hualong Coating Industrial Co., Ltd.

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Indoor Decorating and Refurbishing Materials – Limit of Harmful Substance of Wood Based Furniture

1 Scope

This Standard specifies the limit requirements, test methods and inspection rules for the harmful substance in the indoor wood based furniture.

This Standard is applicable to the various indoor wood based furniture products.

2 Normative Reference

The following documents contain the provisions which, through reference in this Standard, become the provisions of this Standard. For dated references, their subsequent amendments (excluding corrigendum) or revisions do not apply to this Standard. However, the parties who enter into agreement based on this Standard are encouraged to investigate whether the latest versions of these documents are applicable. For undated reference documents, the latest versions apply to this Standard.

GB/T 6682-1992 Water for Analytical Laboratory Use - Specification and Test Methods (neq ISO 3696:1987)

GB/T 9758.1-1988 Paints and Varnishes - Determination of "Soluble" Metal Content -Part 1: Determination of Lead Content - Flame Atomic Absorption Spectrometric Method and Dithizone Spectrophotometric Method (idt ISO 3856-1:1984)

GB/T 9758.4-1988 Paints and Varnishes - Determination of "Soluble" Metal Content - Part 4: Determination of Cadmium Content - Flame Atomic Absorption Spectrometric Method and Polarographic Method (idt ISO 3856-4:1984)

GB/T 9758.6-1988 Paints and Varnishes - Determination of "Soluble" Metal Content -Part 6: Determination of Total Chromium Content of the Liquid Portion of the Paint - Flame Atomic Absorption Spectrometric Method (idt ISO 3856-6:1984)

GB/T 9758.7-1988 Paints and Varnishes - Determination of "Soluble" Metal Content - Part 7: Determination of Mercury Content of the Pigment Portion of the Paint and of the Liquid Portion of Water-Dilutable Paints - Flameless Atomic

- --- Glass mortar: diameter (100~120) mm;
- --- Crystalizing dish: diameter 120mm, height 60mm;
- **5.1.2.6** Small-mouth plastic bottle: 500mL, 1000mL.

5.1.3 Reagent

- --- Potassium iodide (KI): analytically pure;
- --- Potassium dichromate (K₂Cr₂O₇): guaranteed reagent;
- --- Sodium thiosulfate (Na₂S₂O₃ 5H₂O): analytically pure;
- --- Mercury iodide (Hgl₂): analytically pure;
- --- Anhydrous sodium carbonate (Na₂CO₃): analytically pure;
- --- Sulfuric acid (H₂SO₄): ρ=1.84g/mL, analytically pure;
- --- Hydrochloric acid (HCL): ρ=1.19g/mL, analytically pure;
- --- Sodium hydroxide (NaOH): analytically pure;
- --- lodine (l₂): analytically pure;
- --- Soluble starch: analytically pure;
- --- Acetylacetone (CH₃COCH₂COCH₃): guaranteed reagent;
- --- Ammonium acetate (CH₃COONH₄): guaranteed reagent;
- --- Formaldehyde (CH₂O) solution: concentration 35%~40%.

5.1.4 Specimen preparation

5.1.4.1 Specimen sampling

The specimen shall take sampling from the exit-factory qualified products that conform to the test provisions. If the product is made of several kinds of wood materials, then sampling from the component of each material.

- **5.1.4.2** The specimen shall be prepared within the distance about 50mm away from the furniture component edge.
- **5.1.4.3** Specimen specification: length (150±1)mm, width (50±1)mm.

5.1.4.4 Specimen quantity

The quantity of the specimen is 10 blocks. When preparing the specimen, consider the

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- **5.2.2.3** Membrane filter: pore size 0.45µm.
- **5.2.2.4** Magnetic stirrer: the outer layer of stirrer is plastic or glass.
- **5.2.2.5** Single-scale pipette: 25mL.
- **5.2.2.6** White volumetric flask: 50mL.
- **5.2.2.7** Scraper: tool with sharp blade.

5.2.3 Reagent

The used reagent is analytically pure, the used water meet the requirement for Grade-III water stipulated in GB/T 6682-1992.

5.2.3.1 Hydrochloric acid solution

0.07 mol/L, 1 mol/L, 2 mol/L.

5.2.3.2 Nitric acid solution

Mass fraction is 65%~68%.

5.2.4 Preparation of coating powder

Use the scraper to scratch proper amount of coating from the coasting surface of furniture product, crush the scratched coating by magnetic stirrer under the room temperature, then make it penetrate through the 0.5mm metal sieve, and wait for treatment.

5.2.5 Test procedures

5.2.5.1 Sample treatment

Weigh 0.5g (accurate to 0.0001g) of sieved powder, put it into white volumetric flask; add 25mL of 0.07 mol/L hydrochloric acid solution, mix 1min, determine its acidity; if pH is greater than 1.5, shake the solution as drop the hydrochloric acid solution with concentration of 2 mol/L till pH decreases to 1.0~1.5. Stir such mixture continuously under room temperature for 1h, after that stand for 1h, then use membrane filter to filter and keep in dark place.

The sample treatment shall be finished within 4h. If it can't finish within 4h, add 25mL of 1 mol/L hydrochloric acid solution to treat the sample, the treatment method can see above.

5.2.5.2 Determination of soluble metal content

5.2.5.2.1 Determination of soluble lead content shall be performed as per the Chapter

3 in GB/T 9758.1-1988.

- **5.2.5.2.2** Determination of soluble cadmium content shall be performed as per the Chapter 3 in GB/T 9758.4-1988.
- **5.2.5.2.3** Determination of soluble chromium content shall be performed as per GB/T 9758.6-1988.
- **5.2.5.2.4** Determination of soluble mercury content shall be performed as per GB/T 9758.7-1988.

5.2.5.3 Result calculation

Calculate the soluble metal content as per Formula (2), accurate to 0.1mg/kg.

$$C = \frac{(a_1 - a_0) \times 25 \times F}{m} \qquad \dots (2)$$

Where:

- C (Lead, cadmium, chromium, mercury) soluble content, in mg/kg;
- a_0 Blank concentration of 0.07mol or 1mol of hydrochloric acid solution, in $\mu g/mL$;
- a_1 Concentration of test solution (lead, cadmium, chromium, mercury) determined from the standard curve, in μ g/mL;
- F Dilution factor;
- 25 Extracted hydrochloric acid solution, in mL;
- m Weighed sample amount, in g.

6 Inspection Rules

- **6.1** All required contents listed in Chapter 4 of this Standard are the type test items.
- **6.1.1** In case the normal production, after certain period or accumulating certain output, the periodic type test shall be performed once. The cycle is generally 1 year.
- **6.1.2** In case one of the following situations, the type test shall be performed:
 - --- When the trial stereotype of new product;
 - --- When production process and raw materials have great changes;
 - --- When product restores production after long-term shutdown;

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