

# SLOVENSKI STANDARD SIST EN 12586:2025

01-julij-2025

Nadomešča: SIST EN 12586:2008+A1:2011

#### Izdelki za otroke - Držalo dude - Varnostne zahteve in preskusne metode

Child care articles - Soother holder - Safety requirements and test methods

Artikel für Säuglinge und Kleinkinder - Schnullerhalter - Sicherheitstechnische Anforderungen und Prüfverfahren

# **Feh Standards**

Articles de puériculture - Attache sucette - Exigences de sécurité et méthodes d'essai

### Ta slovenski standard je istoveten z: EN 12586:2025

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97.190 Otroška oprema

Equipment for children

SIST EN 12586:2025

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#### SIST EN 12586:2025

# EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM

# EN 12586

April 2025

ICS 97.190

Supersedes EN 12586:2007+A1:2011

**English Version** 

## Child care articles - Soother holder - Safety requirements and test methods

Articles de puériculture - Attache-sucette - Exigences de sécurité et méthodes d'essai Artikel für Säuglinge und Kleinkinder - Schnullerhalter - Sicherheitstechnische Anforderungen und Prüfverfahren

This European Standard was approved by CEN on 1 December 2024.

CEN members are bound to comply with the CEN/CENELEC Internal Regulations which stipulate the conditions for giving this European Standard the status of a national standard without any alteration. Up-to-date lists and bibliographical references concerning such national standards may be obtained on application to the CEN-CENELEC Management Centre or to any CEN member.

This European Standard exists in three official versions (English, French, German). A version in any other language made by translation under the responsibility of a CEN member into its own language and notified to the CEN-CENELEC Management Centre has the same status as the official versions.

CEN members are the national standards bodies of Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Republic of North Macedonia, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Türkiye and United Kingdom.

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EUROPEAN COMMITTEE FOR STANDARDIZATION COMITÉ EUROPÉEN DE NORMALISATION EUROPÄISCHES KOMITEE FÜR NORMUNG

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#### EN 12586:2025 (E)

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### **European foreword**

This document (EN 12586:2025) has been prepared by Technical Committee CEN/TC 252 "Child care articles", the secretariat of which is held by AFNOR.

This European Standard shall be given the status of a national standard, either by publication of an identical text or by endorsement, at the latest by October 2025, and conflicting national standards shall be withdrawn at the latest by April 2026.

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

This document supersedes EN 12586:2007+A1:2011.

EN 12586:2024 includes the following significant technical changes with respect to EN 12586:2007+A1:2011:

- The standard has been completely renewed and a new structure was given. A whole series of new aspects had to be considered.
- The list of materials used currently for soother holders is wider than the list in the former edition, therefore additional materials have been included (silicone elastomer, rubber, TPE, leather, resinbonded materials).
- Introduction: Double testing should be avoided for environmental protection reasons. Therefore, substances regulated by REACH like phthalates, nickel are not included any more.
- 7.3: The list of chemical elements to be tested for migration of certain elements has been enlarged to cover 19 elements in accordance with the last amendment of Directive 2009/48/EC.

— 7.4: Colour fastness test was changed and split in first action and final action method.

- 7.7: Limits and test methods for formaldehyde release from different materials have been revised and modified and included in a separate paragraph. The test and the limit for formaldehyde release is set based on the pragmatic approach that independently of the material type the amount of formaldehyde migrating from the entire product into saliva simulant shall not exceed the healthbased limit.
- 7.8: Migration limits for BPA, phenol have been revised and set based on the recent amendments of Appendix C to Annex II of Directive 2009/48/EC.
- 7.9 and 7.10: The list of preservatives was enlarged to cover wood and leather preservatives too.
- Annex B: The specific migration limits have been specified based on the migration scenario usually applied in the childcare article standards drafted by TC 252 WG5, considering the allocated daily intake, the body weight, and the likely exposure.

This document has been prepared under a standardization request addressed to CEN by the European Commission. The Standing Committee of the EFTA States subsequently approves these requests for its Member States.

For the relationship with EU Legislation, see informative Annex ZA, which is an integral part of this document.

NOTE Information about interpretations of CEN TC 252 Standards can be found in the collection of interpretation requests CEN/TR 16411:2022 [31].

Any feedback and questions on this document should be directed to the users' national standards body. A complete listing of these bodies can be found on the CEN website.

According to the CEN-CENELEC Internal Regulations, the national standards organisations of the following countries are bound to implement this European Standard: Austria, Belgium, Bulgaria, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Republic of North Macedonia, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Türkiye and the United Kingdom.

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#### Introduction

Accidents caused by soothers are few and any resulting in the death of a child are unknown. However, a small number of children die as a result of accidents where a soother is involved, but in these cases, death almost always results from strangulation by a cord being used to hang the soother around the child's neck. Such cords are usually either home-made or made for some other purpose, e.g. a coiled key ring holder.

The main aim of this document is to eliminate the risk of strangulation. This has only been made possible by severely restricting the length of the soother holder. The length should be sufficient for its purpose while not being so long that the strap can encircle the neck and strangle even the smallest child. The lack of reported incidents involving accidents or fatalities since its introduction would suggest that this document has achieved this main aim.

A functional soother holder is not to be considered as a toy. However, a soother holder may contain a part or be designed in such a way as to present a dual use. If the soother holder, in addition to its functional use, has a significant play value, it could also be considered as a toy.

Chemical hazards have been considered by the Technical Committee for all construction materials normally encountered in soother holders. Where relevant, requirements have been derived and included in this document. Restrictions for phthalates, for poly aromatic hydrocarbons (PAH) and for the release of nickel have not been included in this document as they are regulated by REACH [1] (See also part 7.1). Flame retardants are not included as they are regulated by REACH and Regulation (EU) 2019/1021 [2] and do not have any relevance to functional soother holders.

Products with claimed biocidal effect or intended biocidal action are falling under the scope of Regulation (EU) No 528/2012 concerning the marketing and use of biocidal products [3].

A normative annex has been included which contains translations of warning phrases into the main European languages.

It is recommended that manufacturers and suppliers operate to EN ISO 9001 for quality management systems [4]. It is also recommended that laboratories operate to ISO/IEC 17025, General requirements for the competence of testing and calibration laboratories [5].

It is also important that the product or its packaging bears a traceability marking (e.g. batch coding) to enable traceability of the product through the supply chain.