

Irish Standard I.S. EN ISO 3107:2011

Dentistry - Zinc oxide/eugenol cements and zinc oxide/non-eugenol cements (ISO 3107:2011)

© NSAI 2011

No copying without NSAI permission except as permitted by copyright law.

Incorporating amendments/corrigenda/National Annexes issued since publication:					
incorporating amenuments/	compenua/NauUIIal AIIIIE	nes issueu siiile publil	-auon.		
The National Standards Authority of Ireland (NSAI) produces the following categories of formal documents:					
I.S. xxx: Irish Standard – national specification based on the consensus of an expert panel and subject to public consultation.					
S.R. xxx: Standard Recomn panel and subject to public cons	nendation - recommendat ultation.	ion based on the cons	ensus of an expert		
SWiFT xxx: A rapidly developed recommendatory document based on the consensus of the participants of an NSAI workshop.					
This document replaces: EN ISO 3107:2004					
This document is based on: EN ISO 3107:2011	Published: 11 March, 2011				
This document was published under the authority of the NSAI and comes into effect on:			ICS number: 11.060.10		
11 March, 2011					
NSAI	T +353 1 807 3800	Sales:			
1 Swift Square, Northwood, Santry Dublin 9	F +353 1 807 3838 E standards@nsai.ie	T +353 1 857 6730 F +353 1 857 6729 W standards.ie			
	W NSAI.ie	Standards.ic			
Údarás um Chaighdeáin Náisiúnta na hÉireann					

# EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM

**EN ISO 3107** 

March 2011

ICS 11.060.10

Supersedes EN ISO 3107:2004

### **English Version**

# Dentistry - Zinc oxide/eugenol cements and zinc oxide/noneugenol cements (ISO 3107:2011)

Médecine bucco-dentaire - Ciments dentaires à base d'oxyde de zinc-eugénol et à base d'oxyde de zinc sans eugénol (ISO 3107:2011)

Zahnheilkunde - Zinkoxid-Eugenolzemente und eugenolfreie Zinkoxidzemente (ISO 3107:2011)

This European Standard was approved by CEN on 28 February 2011.

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, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland and United Kingdom.



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

Management Centre: Avenue Marnix 17, B-1000 Brussels

# EN ISO 3107:2011 (E)

Contents	Page
Foreword	3

**EN ISO 3107:2011 (E)** 

### **Foreword**

The text of ISO 3107:2011 has been prepared by Technical Committee ISO/TC 106 "Dentistry" of the International Organization for Standardization (ISO) and has been taken over as EN ISO 3107:2011 by Technical Committee CEN/TC 55 "Dentistry" the secretariat of which is held by DIN.

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 September 2011, and conflicting national standards shall be withdrawn at the latest by September 2011.

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

This document supersedes EN ISO 3107:2004.

According to the CEN/CENELEC Internal Regulations, the national standards organizations 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, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland and the United Kingdom.

### **Endorsement notice**

The text of ISO 3107:2011 has been approved by CEN as a EN ISO 3107:2011 without any modification.

This is a free page sample. Access the full version online.

I.S. EN ISO 3107:2011

This page is intentionally left BLANK.

This is a free page sample. Access the full version online.

# I.S. EN ISO 3107:2011 INTERNATIONAL STANDARD

**ISO** 3107

Fourth edition 2011-03-01

# Dentistry — Zinc oxide/eugenol cements and zinc oxide/non-eugenol cements

Médecine bucco-dentaire — Ciments dentaires à base d'oxyde de zinceugénol et à base d'oxyde de zinc sans eugénol



ISO 3107:2011(E)

### PDF disclaimer

This PDF file may contain embedded typefaces. In accordance with Adobe's licensing policy, this file may be printed or viewed but shall not be edited unless the typefaces which are embedded are licensed to and installed on the computer performing the editing. In downloading this file, parties accept therein the responsibility of not infringing Adobe's licensing policy. The ISO Central Secretariat accepts no liability in this area.

Adobe is a trademark of Adobe Systems Incorporated.

Details of the software products used to create this PDF file can be found in the General Info relative to the file; the PDF-creation parameters were optimized for printing. Every care has been taken to ensure that the file is suitable for use by ISO member bodies. In the unlikely event that a problem relating to it is found, please inform the Central Secretariat at the address given below.



### COPYRIGHT PROTECTED DOCUMENT

### © ISO 2011

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized in any form or by any means, electronic or mechanical, including photocopying and microfilm, without permission in writing from either ISO at the address below or ISO's member body in the country of the requester.

ISO copyright office
Case postale 56 • CH-1211 Geneva 20
Tel. + 41 22 749 01 11
Fax + 41 22 749 09 47
E-mail copyright@iso.org
Web www.iso.org

Published in Switzerland

# ISO 3107:2011(E)

Cont	<b>ents</b>	Page
Forewo	ord	iv
Introdu	ıction	
1	Scope	1
2	Normative references	1
3	Terms and definitions	1
4	Classification	1
5 5.1 5.2	Requirements Performance requirements Biocompatibility	1
6	Sampling	2
7 7.1 7.2 7.3 7.4 7.5	Test methods	2 3 6
8 8.1 8.2	Marking, labelling and packagingPackaging Marking and instructions for use	8 8
Bibliog	raphy	10

### **Foreword**

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

International Standards are drafted in accordance with the rules given in the ISO/IEC Directives, Part 2.

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 3107 was prepared by Technical Committee ISO/TC 106, *Dentistry*, Subcommittee SC 1, *Filling and restorative materials*.

This fourth edition cancels and replaces the third edition (ISO 3107:2004), which has been technically revised. It also incorporates the Technical Corrigendum ISO 3107:2004/Cor.1:2006.

The main changes are that the

- a) classification types have been consolidated into two,
- b) compressive strength limit has been reduced to reflect materials in current use,
- c) text on interpretation of compressive test results has been modified, and
- d) lower setting time limit has been lowered to reflect materials in current use.

ISO 3107:2011(E)

# Introduction

Specific qualitative and quantitative requirements for freedom from biological hazard are not included in this International Standard, but it is intended that in assessing possible biological or toxicological hazards, reference be made to ISO 10993-1 and ISO 7405.

This is a free page sample. Access the full version online.

I.S. EN ISO 3107:2011



**Product Page** 

- Dooking for additional Standards? Visit Intertek Inform Infostore
- Dearn about LexConnect, All Jurisdictions, Standards referenced in Australian legislation