STN

Rádiologická ochrana Kriériá spôsobilosti laboratórií používajúcich pri biologickej dozimetri CBMN skúšku v periférnych krvných lymfocytoch (ISO 17099: 2014)

STN EN ISO 17099

40 1406

Radiological protection - Performance criteria for laboratories using the cytokinesis block micronucleus (CBMN) assay in peripheral blood lymphocytes for biological dosimetry (ISO 17099:2014)

Táto norma obsahuje anglickú verziu európskej normy. This standard includes the English version of the European Standard.

Táto norma bola oznámená vo Vestníku ÚNMS SR č. 02/18

Obsahuje: EN ISO 17099:2017, ISO 17099:2014

STN EN ISO 17099: 2018

EUROPEAN STANDARD NORME EUROPÉENNE EUROPÄISCHE NORM

EN ISO 17099

October 2017

ICS 13.280

English Version

Radiological protection - Performance criteria for laboratories using the cytokinesis block micronucleus (CBMN) assay in peripheral blood lymphocytes for biological dosimetry (ISO 17099:2014)

Radioprotection - Critères de performance pour les laboratoires pratiquant la dosimétrie biologique par le test des micronoyaux avec blocage de la cytodiérèse (CBMN) dans les lymphocytes du sang périphérique (ISO 17099:2014)

This European Standard was approved by CEN on 13 September 2017.

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, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and United Kingdom.



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

CEN-CENELEC Management Centre: Avenue Marnix 17, B-1000 Brussels

© 2017 CEN All rights of exploitation in any form and by any means reserved worldwide for CEN national Members.

Ref. No. EN ISO 17099:2017 E

EN ISO 17099:2017 (E)

Contents	Page
European foreword	3

European foreword

The text of ISO 17099:2014 has been prepared by Technical Committee ISO/TC 85 "Nuclear energy, nuclear technologies, and radiological protection" of the International Organization for Standardization (ISO) and has been taken over as EN ISO 17099:2017 by Technical Committee CEN/TC 430 "Nuclear energy, nuclear technologies, and radiological protection" 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 April 2018, and conflicting national standards shall be withdrawn at the latest by April 2018.

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.

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, Former Yugoslav Republic of Macedonia, France, Germany, Greece, Hungary, Iceland, Ireland, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, Norway, Poland, Portugal, Romania, Serbia, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey and the United Kingdom.

Endorsement notice

The text of ISO 17099:2014 has been approved by CEN as EN ISO 17099:2017 without any modification.

INTERNATIONAL STANDARD

ISO 17099

First edition 2014-11-15

Radiological protection —
Performance criteria for laboratories
using the cytokinesis block
micronucleus (CBMN) assay in
peripheral blood lymphocytes for
biological dosimetry

Radioprotection — Critères de performance pour les laboratoires pratiquant la dosimétrie biologique par analyse des micronoyaux par blocage de la cytokinèse (CBMN) dans les lymphocytes du sang périphérique



ISO 17099:2014(E)



COPYRIGHT PROTECTED DOCUMENT

© ISO 2014

All rights reserved. Unless otherwise specified, no part of this publication may be reproduced or utilized otherwise in any form or by any means, electronic or mechanical, including photocopying, or posting on the internet or an intranet, without prior written permission. Permission can be requested 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

Cor	itent	S	Page		
Foreword					
Intro	ductio	n	v i		
1	Scop	e	1		
2	Tern	ns and definitions	1		
3		onucleus assay methodology used in this standard			
3	3.1	General			
	3.2	General requirement of the laboratory			
	3.3	Requests for analysis and blood sampling			
	3.4	Cell culturing			
	3.5 3.6	Staining Microscopy			
	3.7	Visual scoring of slides			
	3.8	Automated analysis			
4	Conf	identiality of personal information			
	4.1	Overview			
	4.2	Applications of the principle of confidentiality			
5	Laho	ratory safety requirements			
	5.1	Overview			
	5.2	Microbiological safety requirements			
	5.3	Chemical safety requirements			
	5.4	Optical safety requirements			
	5.5	Safety plan	10		
6		oration source(s), calibration curve, and minimum resolvable dose			
	6.1	Calibration source(s)			
	6.2 6.3	Calibration curve Background micronucleus frequency			
	6.4	Minimum resolvable dose measurement			
7	Resn	onsibility of the customer			
	Responsibility of the CBMN laboratory				
8	8.1	Setup and sustainment of the QA program			
	8.2	Responsibility during service			
9	Accia	lental overexposure involving few individuals	14		
9	9.1	Procedure for scoring micronuclei in binucleated cells			
	9.2	Criteria for converting a micronucleus yield into an estimate of absorbed dose			
	9.3	Reporting of Results	15		
10	Popu	ılation triage	17		
	10.1	General	17		
	10.2	Use of a CBMN assay network for large scale exposures	17		
	10.3	Procedure for scoring micronuclei in binucleated cells			
	10.4 10.5	Criteria for converting a micronucleus yield into an estimate of absorbed dose			
11					
11	Quai 11.1	ity assurance and quality control Overview			
	11.2	Quality Assurance			
	11.3	Quality control			
Anne	ex A (in	formative) Sample data sheet for recording micronuclei in binucleated cells	20		
	_	formative) Automation of micronuclei scoring			
		formative) Instructions for customer (sample)			
	(111	·			

STN EN ISO 17099: 2018

ISO 17099:2014(E)

Annex D (informative) Sample questionnaire	24
Annex E (informative) Sample of report for single assessment	26
Annex F (informative) Example group sample report	27
Bibliography	29

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.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see www.iso.org/directives).

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. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see www.iso.org/patents).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation on the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the WTO principles in the Technical Barriers to Trade (TBT), see the following URL: Foreword — Supplementary information.

The committee responsible for this document is ISO/TC 85, *Nuclear energy, nuclear technologies, and radiological protection*, Subcommittee SC 2, *Radiological protection*.

Introduction

The purpose of this International Standard is to define the use of the cytokinesis block micronucleus (CBMN) assay with human peripheral blood lymphocytes for biological dosimetry of exposure to ionizing radiation. This assay is intended to be applied for accidental or malevolent exposures involving a) up to a few casualties to provide individual full dose estimates or b) in a triage mode to populations to provide interim dose estimates for individuals.

The CBMN assay is an alternative cytogenetic technique, which is possibly simpler and faster to perform than the dicentric assay (ISO 19238:2014, ISO 21243:2008). It is also routinely used to demonstrate exposure to genotoxic agents, other than ionizing radiation, which is not covered in this International Standard. Although culture of the blood samples is slightly longer than for dicentrics, the scoring of micronuclei in binucleated lymphocytes is easier.

As was done with the dicentric assay, the CBMN assay has been adapted for the emergency triage of large-scale multi casualty radiation accidents. The blood volume required for sufficient number of scorable binucleated cells is similar than required for the dicentric assay. Again, the faster counting speed for micronuclei compensates for the extended culture time. In addition, the CBMN assay can be performed in an automated mode.

This International Standard provides a guideline on how to perform the CBMN assay for dose assessment using documented and validated procedures. Dose assessment using the CBMN assay has relevance in medical management, radiation-protection management, record keeping, and medical/legal requirements. This International Standard is divided into two parts, according to the use of CBMN assay: radiation exposure of a few individuals or population triage in a large radiological event.

A part of the information in this International Standard is contained in other international guidelines and scientific publications, primarily in the International Atomic Energy Agency's (IAEA) technical reports series on biological dosimetry. However, this International Standard expands and standardizes the quality assurance and quality control, the criteria of accreditation and the evaluation of performance. This International Standard is generally compliant with ISO/IEC 17025 "General requirements for the competence of testing and calibration laboratories" with particular consideration given to the specific needs of biological dosimetry. The expression of uncertainties in dose estimations given in this International Standard complies with the "ISO-guide for the expression of uncertainty in measurement" (former GUM) and the ISO 5725-all parts.

Radiological protection — Performance criteria for laboratories using the cytokinesis block micronucleus (CBMN) assay in peripheral blood lymphocytes for biological dosimetry

1 Scope

This International Standard addresses the following:

- a) confidentiality of personal information for the customer and the laboratory;
- b) laboratory safety requirements;
- c) radiation sources, dose rates, and ranges used for establishing the calibration reference dose-effect curves allowing the dose estimation from CBMN assay yields and the minimum resolvable dose;
- d) performance of blood collection, culturing, harvesting, and sample preparation for CBMN assay scoring;
- e) scoring criteria;
- f) conversion of micronucleus frequency in binucleated cells into an estimate of absorbed dose;
- g) reporting of results;
- h) quality assurance and quality control;
- i) informative annexes containing examples of a questionnaire, instructions for customers, a microscope scoring data sheet, a sample report and advice on strengths and limitations of current automated systems for automated micronucleus scoring.

koniec náhľadu - text ďalej pokračuje v platenej verzii STN

© ISO 2014 – All rights reserved