
**Sterilization of medical devices —
Guidance on the requirements for the
validation and routine processing of
ethylene oxide sterilization processes
using parametric release**

*Stérilisation des dispositifs médicaux — Lignes directrices concernant
les exigences de validation et de traitement de routine des procédés de
stérilisation à l'oxyde d'éthylène par libération paramétrique*





COPYRIGHT PROTECTED DOCUMENT

© ISO 2020

All rights reserved. Unless otherwise specified, or required in the context of its implementation, 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
CP 401 • Ch. de Blandonnet 8
CH-1214 Vernier, Geneva
Phone: +41 22 749 01 11
Email: copyright@iso.org
Website: www.iso.org
Published in Switzerland