ISO 21976:2018 (E)

Packaging — Tamper verification features for medicinal product packaging

Contents

Foreword

Introduction

1 Scope

2 Normative references

3 Terms and definitions

4 General requirements

4.1 Tamper verification features
4.2 Purpose of tamper verification features
4.3 Application and use of tamper verification features
4.4 Check of tamper verification features

5 Categories of tamper verification features

5.1 General
5.2 Folding boxes closed with glue
5.2.1 Description
5.2.2 Criteria of tamper verification
5.2.3 Verification
5.3 Specially constructed folding boxes
5.3.1 Description
5.3.2 Criteria of tamper verification
5.3.3 Verification
5.4 Sealing labels and tapes
5.4.1 Description
5.4.2 Criteria of tamper verification
5.4.3 Verification
5.5 Film wrappers
5.5.1 Description
5.5.2 Criteria of tamper verification
5.5.3 Verification
5.6 Sleeves
5.6.1 Description
5.6.2 Criteria of tamper verification
5.6.3 Verification
5.7 Breakable or tear-away closure
5.7.1 Description
5.7.2 Criteria of tamper verification
5.7.3 Verification
5.8 Container inner seal or membrane
5.8.1 Description
5.8.2 Criteria of tamper verification
5.8.3 Verification
5.9 Display blister pack
5.9.1 Description
5.9.2 Criteria of tamper verification
5.9.3 Verification
5.10 Flexible packaging
5.10.1 Description
5.10.2 Criteria of tamper verification
5.10.3 Verification
5.11 Blow-fill-and-seal-container (BFS)

5.11.1 Description
5.11.2 Criteria of tamper verification
5.11.3 Verification
5.12 New and emerging technologies

Annex A (informative) Additional information regarding tamper verification features

Page count: 16