

ISO 17351:2013-01 (E)

Packaging - Braille on packaging for medicinal products

Contents		Page
Foreword		iv
Introduction		v
1	Scope	1
2	Terms and definitions	1
3	General requirements for medicinal product packaging	1
3.1	Product identification	1
3.2	Braille spacing convention	2
3.3	Braille character sets	2
4	Determination of Braille legibility	2
4.1	Principles of Braille legibility compliance	2
4.2	Braille cell dot height	2
4.3	Altered Braille labelling	3
Annex A (normative) Methods of verification		4
Annex B (informative) Braille characteristics and recommendations		5
Annex C (informative) Technology for the application of Braille to packaging for medicinal products		7
Annex D (informative) Guidance on Braille specifications and artwork generation		10
Annex E (informative) Braille character sets		12
Bibliography		13