

DIN EN ISO 15223-1:2022-02 (E)

Medical devices - Symbols to be used with information to be supplied by the manufacturer - Part 1: General requirements (ISO 15223-1:2021)

Contents	Page
European foreword	3
Annex ZA (informative) Relationship between this European standard and the General Safety and Performance Requirements of Regulation (EU) 2017/745 aimed to be covered	5
Annex ZB (informative) Relationship between this European standard and the General Safety and Performance Requirements of Regulation (EU) 2017/746 aimed to be covered	14
Foreword	23
Introduction	24
1 Scope	25
2 Normative references	25
3 Terms and definitions	25
4 General requirements	31
4.1 Future <i>symbols</i>	31
4.2 Requirements for usage	31
4.3 Other <i>symbols</i>	31
5 Symbols	31
Annex A (informative) Guidance and examples of <i>symbol</i> use, including multiple <i>symbols</i>	52
Annex B (informative) Use of general prohibition <i>symbol</i> and negation <i>symbol</i>	58
Bibliography	59