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Committee collaboration Non ctive medical devices, European i A The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, PartIn particular, the different approval criteria needed for the different types of ISO documents should be noted ILNAS-EN ISO Every interested party, which is member of an organization based in Luxembourg, can participate for FREE in the development of Luxembourgish (ILNAS), European (CEN, CENELEC) and International (ISO, IEC) standards: Participate in the design of standards Foresee future developments Participate in technical committee ny or all such patent was prepared by Technical Committee ISO/TC, Transfusion, infusion and injection equipment for m. ISO ISO consists of the following parts, under the general title Plastics collapsible containers for human blood and blood components Any feedback or questions on this document should be directed to the user?s national standards body. A list of all parts in the ISO series can be found on the ISO site. The procedures used to develop this This document supersedes EN ISO According to the CEN-CENELEC Internal Regulations, the national standards organizations of the following countries are ISO (E) Air content The total volume of air contained in the plastics container system divided by the number of containers shall not exceedml. aration, cance. A This document specifies requirements, including performance requirements, for plastics collapsible, non-vented, sterile containers (known as plastics containers) complete with For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization. NOTE Typical plastics container systems are described in ISOWhen used in accordance with the manufacturer's instructions, the plastics container shall be and Committee blood document processing for Standardization was prepared equipment by for Technical Committee Transfusion, infusion and injection, medical accordance with the Agreeme and pharmaceutical t on technical Technical ooperation between ISO and CEN (Vienna Agreement).

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