

## Declaration of Conformity

**DYNEX TECHNOLOGIES, spol. s r.o.**  
**Vodičkova 791/41, 110 00 Praha 1**  
**Czech Republic**

We declare under our sole responsibility that the product

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**DYNABLOT Plus (product name)**

**D7144-P7 (order number)**

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meets the demands of Directives

**Directive 98/79/EC**      **In vitro diagnostic medical devices**

**Directive 2014/30/EU**      **Electromagnetic compatibility Directive**

**Directive 2011/65/EU**      **on the restriction of the use of certain hazardous substances in electrical and electronic equipment**

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**The following standards are applied:**

**EN 61326:2013**      Electrical equipment for measurement, control and laboratory use – EMC requirements

**EN 14971:2012**      Medical devices – Application of risk management to medical devices

**EN 61010-1:2011**      Safety requirements for electrical equipment for measurement, control and laboratory use - Part 1: General requirements

**EN 61010-2-101:2003**      Safety requirements for electrical equipment for measurement, control and laboratory use - Part 2-101: Particular requirements for in vitro diagnostics (IVD) medical equipment

**EN ISO 13485:2012**      Medical devices

Praha, 11. 7. 2012  
Date of revision: 19.8.2016  
Place and date of issue

  
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