

Regulation (EU) 2017/746 of the European Parliament and of the Council of 5 April 2017 on in vitro diagnostic medical devices and repealing Directive 98/79/EC and Commission Decision 2010/227/EU (Text with EEA relevance)

## CHAPTER VIII

### **COOPERATION BETWEEN MEMBER STATES, MEDICAL DEVICE COORDINATION GROUP, EU REFERENCE LABORATORIES AND DEVICE REGISTERS**

#### *Article 96*

#### **Competent authorities**

The Member States shall designate the competent authority or authorities responsible for the implementation of this Regulation. They shall entrust their authorities with the powers, resources, equipment and knowledge necessary for the proper performance of their tasks pursuant to this Regulation. The Member States shall communicate the names and contact details of the competent authorities to the Commission which shall publish a list of competent authorities.

**Status:**

Point in time view as at 05/04/2017. This version of this provision has been superseded.

**Changes to legislation:**

There are currently no known outstanding effects for the Regulation (EU) 2017/746 of the European Parliament and of the Council, Article 96.