**Question for written answer E-007044/2013**
**to the Commission**
Rule 117
**Anna Rosbach (ECR)**

Subject: Medical devices and translation: further question following incomplete answer

In its answers to my written questions E-004765/2013 and E-004767/2013, the Commission seems to have forgotten to answer some of the questions in E-004765/2013.

Could the Commission therefore please outline:

1. the language requirements applicable to instructions for medical devices for patients in each Member State (listing language requirements for each Member State);

2. the language requirements applicable to instructions for medical devices for healthcare practitioners in each Member State (listing language requirements for each Member State);

3. the language requirements applicable to the labelling of medical devices in each Member State (listing language requirements for each Member State)?