# **Curriculum Vitae**

Name:

## Education

# *Mark one*

MD |\_\_| Nurse |\_\_| Other than MD or Nurse \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ I\_\_I

Year of graduation:

### Positions

Please, contact the HR department.

### Theoretical experience

Have you received supervision related to the SUP-ICU protocol such as regulatory or legislative acts, informed consent, patient safety and GCP?

Yes (describe):

**Describe briefly your experience with clinical trials:**

Keywords: informed consent, applied for approval at scientific/ethic committees, participating in drug related public or private conducted trials, laboratory based projects etc.

Date: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ Signature: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_