

# Combination therapy versus monotherapy: a randomised study on the evolution of inflammatory parameters after ventilator associated pneumonia

<b>Submission date</b> 22/12/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 11/01/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 25/09/2009	<b>Condition category</b> Respiratory	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

## Study information

Scientific Title

**Study objectives**

Does a combination antibiotic therapy of ventilator associated pneumonia improve the inflammatory parameters faster than a monotherapy?

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Yes; 27/02/02; 2002/32

**Primary study design**

Interventional

**Study design**

Randomised unblinded comparative study

**Study type(s)**

Treatment

**Health condition(s) or problem(s) studied**

Ventilator associated pneumonia

**Interventions**

Comparison between treatment with cefepime alone and cefepime associated with amikacin or levofloxacin

**Intervention Type**

Drug

**Phase**

Not Specified

**Drug/device/biological/vaccine name(s)**

Cefepime, amikacin, levofloxacin

**Primary outcome(s)**

Time course evolution of C-reactive protein (CRP) levels, temperature and leucocytosis

**Key secondary outcome(s)**

1. Length of ventilatory support
2. Evolution of PaO<sub>2</sub>/FiO<sub>2</sub>
3. Mortality

**Completion date**

31/12/2003

**Eligibility****Key inclusion criteria**

Adult patients with ventilator associated pneumonia

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**Key exclusion criteria**

1. Patients treated for other infection
2. Immunocompromised patients
3. Patients with life expectancy less than 72 hours

**Date of first enrolment**

01/04/2002

**Date of final enrolment**

31/12/2003

**Locations****Countries of recruitment**

Belgium

**Study participating centre**

General Intensive Care Department

Liege

Belgium

4000

**Sponsor information****Organisation**

Domaine Universitaire du Sart-Tilman (Belgium)

**ROR**

<https://ror.org/00afp2z80>

# Funder(s)

## Funder type

University/education

## Funder Name

Domaine Universitaire du Sart-Tilman (Belgium)

# Results and Publications

## Individual participant data (IPD) sharing plan

### IPD sharing plan summary

Not provided at time of registration

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	01/01/2006		Yes	No