

# Prevention of radiotherapy side-effects by early hyperbaric oxygen administration

<b>Submission date</b> 04/04/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 04/04/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 18/09/2008	<b>Condition category</b> Cancer	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

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

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

**Protocol serial number**  
MEC 2005-318; NTR617

## Study information

**Scientific Title**

**Study objectives**

To prevent radiotherapy side-effects with early hyperbaric oxygen administration.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Ethics approval received from the local medical ethics committee

**Study design**

Randomised, active controlled, parallel group trial

**Primary study design**

Interventional

**Study type(s)**

Treatment

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

Nasopharyngeal cancer, oropharyngeal cancer

**Interventions**

Randomisation of hyperbaric oxygen (one group with and one group without). Treatment consists of 30 sessions of 2 hours a day.

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

1. Xerostomia
2. Mucositis
3. Trismus

**Key secondary outcome(s))**

1. Tumour control
2. Quality of life

**Completion date**

06/02/2009

## **Eligibility**

**Key inclusion criteria**

1. Curative intent
2. Presence of oropharynx and nasopharynx tumours

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**Key exclusion criteria**

Palliative treatment

**Date of first enrolment**

06/02/2006

**Date of final enrolment**

06/02/2009

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

Netherlands

**Study participating centre**

Erasmus Medical Centre

Rotterdam

Netherlands

3075 EA

**Sponsor information****Organisation**

Erasmus Medical Centre (The Netherlands)

**ROR**

<https://ror.org/018906e22>

**Funder(s)****Funder type**

Hospital/treatment centre

**Funder Name**

Erasmus Medical Centre (The Netherlands)

**Results and Publications**

Individual participant data (IPD) sharing plan

**IPD sharing plan summary**

Not provided at time of registration