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

No registrations found.

Ethical review	Positive opinion
Status	Recruiting
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON22762

Source
NTR

Brief title
N/A

Health condition

Oropharyngeal and nasopharyngeal cancer

Intervention

Outcome measures

Primary outcome

1. Xerostomia;
2. Mucositis;
3. Trismus.

Secondary outcome

1. Tumor control;

2. Quality of Life.

Study description

Background summary

After regular treatment of oropharyngeal and nasopharyngeal tumours. Patients are randomized to treatment with or without Hyperbaric Oxygen.

Study objective

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

Intervention

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

Contacts

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Eligibility criteria

Inclusion criteria

1. Curative intent;

2. Presence of oropharynx en nasopharynx tumours.

Exclusion criteria

Palliative treatment.

Study design

Design

Study type:	Interventional
Intervention model:	Parallel
Masking:	Open (masking not used)
Control:	Active

Recruitment

NL

Recruitment status:	Recruiting
Start date (anticipated):	06-02-2006
Enrollment:	180
Type:	Anticipated

Ethics review

Positive opinion

Date:	03-03-2006
Application type:	First submission

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
NTR-new	NL560
NTR-old	NTR617
Other	: MEC 2005-318
ISRCTN	ISRCTN25123615

Study results

Summary results
N/A