

Sexual functioning and the quality of sexual life in patients with colorectal cancer and their partners.

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Sexual dysfunction is present, to some extent, in most patients after the treatment for colorectal cancer. There is a relationship between both treatment-related (e.g., type of surgery) and psychosocial factors (e.g., depressive symptoms) and...

Ethische beoordeling	Positief advies
Status	Werving nog niet gestart
Type aandoening	-
Onderzoekstype	Observationeel onderzoek, zonder invasieve metingen

Samenvatting

ID

NL-OMON27180

Bron

NTR

Aandoening

Sexual function, Sexual dysfunction,
Quality of (sexual) life, Colorectal cancer

Ondersteuning

Primaire sponsor: Tilburg University

Overige ondersteuning: Dutch Cancer Society

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The objectives of this study will be to examine:

1. The incidence of sexual problems and the extent patients with colorectal cancer are

bothered by these problems across time;

2. The effect of different treatment options on sexual functioning;

3. The association between sexual problems and quality of life;

4. The determinants of sexual problems and quality of sexual life adopting the biopsychosocial approach of patients with colorectal cancer who have been treated with surgery, radiation and/or chemotherapy, and more specifically to the role of personality and partner factors and sexual functioning/quality of sexual life.

Toelichting onderzoek

Achtergrond van het onderzoek

N/A

Doel van het onderzoek

Sexual dysfunction is present, to some extent, in most patients after the treatment for colorectal cancer.

There is a relationship between both treatment-related (e.g., type of surgery) and psychosocial factors (e.g., depressive symptoms) and sexual dysfunction, the quality of sexual life and the partnerrelationship.

Onderzoeksopzet

Questionnaires are completed before treatment (T0), 6 weeks (T1), 3 months (T2), 6 months (T3), and 12 months (T4) after treatment.

Onderzoeksproduct en/of interventie

Patients will receive care as usual. Participants and their partners will complete several questionnaires (e.g., sexual functioning, depressive symptoms, quality of life, personality) before treatment (T0), 6 weeks (T1), 3 months (T2), 6 months (T3), and 12 months (T4) after treatment.

Contactpersonen

Publiek

Postbus 90153
M.J. Traa
Warandelaan 2
Tilburg 5000 LE
The Netherlands
+31 (0)13 4664129

Wetenschappelijk

Postbus 90153
M.J. Traa
Warandelaan 2
Tilburg 5000 LE
The Netherlands
+31 (0)13 4664129

Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

1. Patients who are diagnosed with colorectal cancer in one of the participating centra, and their partners are asked to participate in this study prior to treatment;
2. Patients have to be between 18 and 75 years old.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

1. Disease recurrence at baseline or metastases;
2. Poor expression in the Dutch language;
3. Dementia;
4. History of psychiatric illness.

Onderzoeksopzet

Opzet

Type: Observationeel onderzoek, zonder invasieve metingen
Onderzoeksmodel: Parallel
Toewijzing: N.v.t. / één studie arm
Controle: N.v.t. / onbekend

Deelname

Nederland
Status: Werving nog niet gestart
(Verwachte) startdatum: 15-03-2010
Aantal proefpersonen: 600
Type: Verwachte startdatum

Ethische beoordeling

Positief advies
Datum: 12-10-2010
Soort: Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

ID: 36622
Bron: ToetsingOnline
Titel:

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL2452
NTR-old	NTR2568
CCMO	NL32121.008.10
ISRCTN	ISRCTN wordt niet meer aangevraagd.
OMON	NL-OMON36622

Resultaten

Samenvatting resultaten

N/A