

Improving care for hospitalized elderly patients: The effects of an e-learning tool.

Gepubliceerd: 19-04-2011 Laatst bijgewerkt: 13-12-2022

The implementation of the delirium part of the Vulnerable elderly project will improve when nurses have completed an e-learning tool regarding delirium care for elderly patients.

| | |
|-----------------------------|--------------------------|
| Ethische beoordeling | Positief advies |
| Status | Werving nog niet gestart |
| Type aandoening | - |
| Onderzoekstype | Interventie onderzoek |

Samenvatting

ID

NL-OMON19991

Bron

NTR

Aandoening

delirium
vulnerability
frailty

Ondersteuning

Primaire sponsor: Nederlands instituut voor onderzoek naar de gezondheidszorg (NIVEL), VU Medical Center, EMGO+ Institute

Overige ondersteuning: Dutch Ministry of Health, Welfare and Sport

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

1. % of patients screened for delirium risk;

2. % of patients screened for vulnerability/frailty.

Toelichting onderzoek

Achtergrond van het onderzoek

Research question trial:

Does the use of a complementary e-learning tool improve the implementation of the delirium part of the Vulnerable elderly project?

Design:

A stepped wedge cluster randomized controlled trial.

The Vulnerable elderly project is a nationwide quality improvement project aimed at care for hospitalized elderly (70 years and older). Four healthcare problems are addressed within this project: delirium, falls, malnutrition and functional decline. For our research questions we focus on the delirium aspect of the project.

The Vulnerable elderly project is a part of a larger patient safety program in the Netherlands, similar to the 100.000 Lives campaign in the United States and the Safer Healthcare Now! campaign in Canada.

Methods for data gathering:

Data will be gathered in 20 Dutch hospitals, all of which are still at the beginning stages of the implementation of the Vulnerable elderly project. One internal medicine ward and one surgical ward of each hospital will participate in the study. Patient record reviews of patients 70 years or older will be the main source of data. During 11 months research-nurses will review 10 records per ward per month. All reviews will take place within the same week of the month, using records of patients still admitted or released within the precious week. From these records data concerning the implementation of the Vulnerable elderly project will be gathered.

Intervention:

Every month two randomly selected hospitals will receive access to an e-learning tool

regarding delirium (one aspect of the Vulnerable elderly project). This stepped wedge design will result in data from all hospitals for the control and for the intervention group.

Doel van het onderzoek

The implementation of the delirium part of the Vulnerable elderly project will improve when nurses have completed an e-learning tool regarding delirium care for elderly patients.

Onderzoeksopzet

Monthly record review of still admitted patients or recently released patients in all participating wards. No follow-up per patient.

Onderzoeksproduct en/of interventie

E-learning tool for nurses, aimed at improving knowledge, skills and attitude regarding delirium care for elderly patients. This tool was developed by Noordhoff, in cooperation with a hospital.

Nurses working in the participating wards will be given access to the e-learning tool for a period of 3 months. The estimated time needed to complete the tool and the knowledge test before and after the tool is 4 hours. The period in which each ward is given access is assigned randomly.

Contactpersonen

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

1. 70 years or older;
2. Length of stay 24 hours or longer;
3. Admitted to surgical ward or internal medicine ward.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

N/A

Onderzoeksopzet

Opzet

| | |
|------------------|-----------------------|
| Type: | Interventie onderzoek |
| Onderzoeksmodel: | Cross-over |
| Toewijzing: | Gerandomiseerd |
| Blinding: | Enkelblind |
| Controle: | Geneesmiddel |

Deelname

| | |
|-------------------------|--------------------------|
| Nederland | |
| Status: | Werving nog niet gestart |
| (Verwachte) startdatum: | 20-05-2011 |
| Aantal proefpersonen: | 4000 |
| Type: | Verwachte startdatum |

Ethische beoordeling

Positief advies

Datum: 19-04-2011

Soort: Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register ID

NTR-new NL2747

NTR-old NTR2885

Ander register METC VUmc / wetenschapscommissie VU : 2011/053 / WC2010-112;

ISRCTN ISRCTN wordt niet meer aangevraagd.

Resultaten

Samenvatting resultaten

N/A