

# Home parenteral nutrition in advanced cancer patients

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- Increase in use of HPN for cancer patients in the US and parts of Europe
- Unclear definition of palliative care



Aim:

 To investigate the experiences of HPN from the perspective of advanced cancer patients and their family members.

Orrevall et al, Clinical Nutrition, 2004, 2005



### Palliative home care services

- Physician directed
- Multi-professional
- Medical care in the patient's home
- Available 24h/7days per week







#### **Participants**

- 13 patients
  47-79 years old
- 6 patients on chemotherapy treatment
- 11 family members (spouse/daughter)



#### **HPN treatment**

 HPN was delivered in "all-in-one-bags" via subcutaneous venous access port, 2 to 7 nights per week, and administered by a nurse in the palliative home care team.



#### **HPN treatment**

- 2 patients on total HPN
- 9 patients on partial HPN and oral intake
- 2 patients had recently stopped HPN



#### Results

- Nutritional situation prior to HPN
- Decision to start HPN
- Experience of receiving HPN



- Patients described periods of weight loss
- Weight loss was at first seen as something positive
- Later a sign of disease and death



- Meals were a source of worry and often despair
- Meals were no longer pleasurable social events
- Food served only a survival function



#### **Family member**

When there's a person sitting at the kitchen-table with a vomit bag in one hand and a fork in the other, it's not the way things should be.



- Family members tried to help the patients eat more
- Family members felt totally responsible for patients' nutritional care



 During hospital visits attention was not focused on nutritional issues



- All patients had tried oral nutritional supplements with limited success
- Dietitians were used as consultants and did not follow the patient during the illness trajectory



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- Patient and family had come to realise they were unable to solve nutritional problems by themselves
- HPN was seen as necessary for the survival of the patient
- The desperate and chaotic situation in the family led to willingness to accept HPN



#### Patient

I just kept losing weight, and I was so tired that I didn't have enough energy to do anything. And then the nurse visited me, and she said, "You'll have this drip thing because it's going to do you good". And it did, because I stopped losing weight.



- Sense of relief and security
- Effect on quality of life
- Effect on weight and energy, strength and activity
- Effect on food intake and mealtimes



#### Patient

There's just this sense of security that I have the drip, because – if it's hard to eat during the evening and I feel that I don't want to. I don't have the energy. To stuff something in when you have no appetite at all is no fun. But now I can eat a little less, more calmly, and think - yes, in any case I'm getting the nourishment that I know I need from the drip.



 Autonomy and dependency in relation to HPN



#### Patient

This drip means that I'm not getting worse, you know. Because it's very difficult not to be able to do things oneself, to have to call for help every time you have to go to the toilet and get out of bed. Since I've had the drip, I can manage by myself in a whole different Way.



#### Patient

I felt right away that it helped. Maybe not the first day, but the first week, and I got out of bed and cooked food.



 HPN treatment means regular visits from the home care nurses.



#### Patient

I'm more fit, food tastes better, and then they come also. They're so nice, very nice the whole group. We have the doctor, who we can call anytime at all if something happens and I can see that my husband also became much happier because he thought he had the responsibility.



Restrictions in family life



- Restrictions in the family life
- Sleep disturbance



- HPN-free evenings and nights
- Possibility of changing routines



- Patients and family members expressed physical, social and psychological benefits from HPN treatment.
- Benefits of HPN treatment were generally said to outweigh negative aspects.



## **Time from first HPN to death**

No of	
patients	Months
2	1-3
3	4-6
4	7-12
3	13-19
1	20-33



#### Aim:

 To investigate the prevalence and use of oral, enteral and parenteral nutritional support in palliative home care services.

Orrevall et al, Supportive Care in Cancer, 2009



#### **Methods**

Patients recruited from:

- 21 palliative home care services
- telephone interviews with 621 patients



#### **Participants (n=621)**

- Mean age 67 years
- Mean BMI 22
- 68% nutritionally at risk
- 40% died within 4 months post-interview
- >25% lived >1 year post-interview



#### Nutritional support (n=621)

- 55% reported use of nutritional support
- 41% oral nutritional support alone
- 14% artificial nutrition



## Artificial nutrition (n=621)

- 14% Artificial nutrition
  - 3% enteral tube feeding
  - 11% parenteral nutrition



## **Characteristics of HPN-users**

- Gastrointestinal malignancies most common
- >1/3 had lost >20% of pre-cancer weight
- Mean BMI was in lower normal range
- The majority survived more than 4 months after first HPN-treatment at home



## Use of HPN

- Almost 3/4 were able to eat
- More than half used oral nutritional support products
- The number of treatments/week varied greatly
- Six out of 68 patients reported discomfort related to HPN feeding solutions



## Most common reason for use of HPN as reported by patients

- Concerns about body weight
- Concerns about nourishment



## **Use of artificial nutrition**

- Use of enteral tube feeding was limited
- HPN was used to supplement oral intake rather than only for patients with a non functioning GI tract
- HPN was generally not introduced at the end of life



