

# Psychometrics of Halpin Nausea and Vomiting (HNV) Scales for Use in Clinical Practice

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Acknowledge:

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# Disclosures

- ▣ The authors have no financial gains
- ▣ The authors are publishing the results
- ▣ Contributors to study
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# Translational Research

*“Knowing is not enough; we must apply.  
Willing is not enough; we must do.”*

-Goethe

Reference: (Olsen, Saunders & McGinnis, 2011)

# What we will discuss & learn?

- ▣ Define translational research
- ▣ State key points of translational research at the bedside in a methodology study.
- ▣ Describe enhancement of RN knowledge with involvement in research application.
- ▣ Identify how “*knowledge*” is enhanced when RNs at the bedside participate in EBP and research projects.
- ▣ Describe the 0-5 Halpin Nausea Vomiting Scale (HNV) methodology research as an example of translational research

# Translational Research/EBP

- ▣ Isn't it one and the same?
- ▣ The history – more prevalent in past decade...the terms popularized in 2002 AMA
- ▣ Found to be essential in moving science forward for better outcomes (Fontanarosa & De Angelis, 2002)

# Translational Research

- ▣ *“Nursing research—and its translation into evidence-based practice and policy—stands as a keystone for improving the health and welfare of people around the world, and at NINR we see CTS and implementation research as essential components of our core mission.”*  
(Grady, 2013)

# Translational Research

- ▣ “Translational research transforms scientific findings or discoveries from basic laboratory, clinical, or population studies into new clinical tools, processes, or applications.

## **Improves:**

patient care

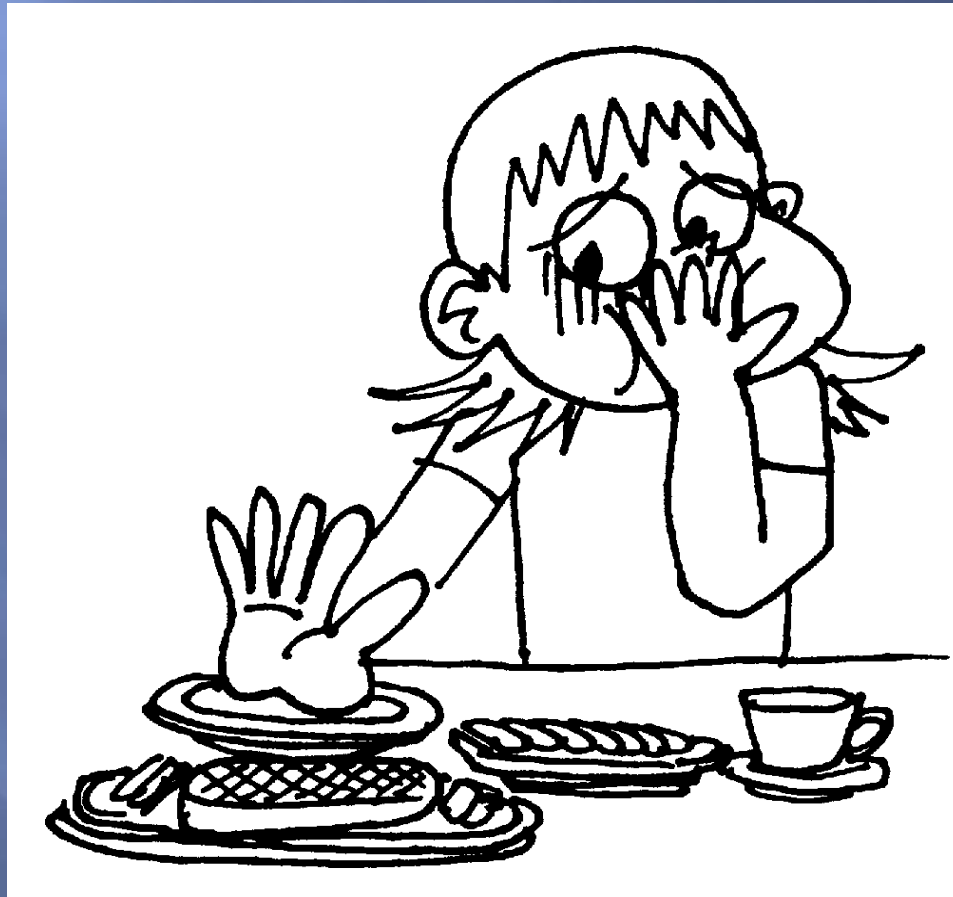
promotes public health.

Intention-build “ bench to bedside.”

Application of scientific findings to clinical practice is the function of translational research.

(Grady, 2010)

# Symptom of Nausea & Vomiting



Symptoms of NV most uncomfortable according to patient reports (Rhodes, 2005; NCCN, 2013)



# Pathophysiology of Nausea and Vomiting

- ▣ CTZ –A zone in the cerebral area of the hypothalamus
- ▣ Sensors
  - Nervous system connection
    - ▣ In gut ( stomach)
    - ▣ Brain pathway
    - ▣ Sympathetic & parasympathetic systems
- ▣ Experience –psychosocial influences

# Problem

- ▣ Nurses in oncology were unable to measure intensity and severity of the NV symptoms with current scales
- ▣ Therefore nurses wanted a 0-5 scale with descriptors to:
  - better understand the patients perceptions of the NV symptoms
  - better treat the symptoms
  - include the patient at the center of the decisions for both

# Telling the Story

- ▣ Review of the literature
  - Multiple scales were available
  - Most were used in outpatient settings
  - Ease of use for inpatient settings remained questionable
- ▣ Pilot test was the first step in 2008
  - Evaluated nurses
    - ▣ satisfaction with the HNV scales
    - ▣ application feasibility
- ▣ Methodology study

# Why nursing needs a NV scale?

- ▣ Inconsistencies existed in communications between patients and health care providers on:
  - Severity
  - Degree
  - Intensity

# PLANNING FOR IMPLEMENTATION

*Physician-nurse champions with  
nursing collaboration*

*Department approvals  
(Organization, Nursing Administration,  
Quality, Research and Information Technology)*

*Study design and IRB approvals*

*Hospitals – two hospitals*

*Data collectors identified and trained*

*Education of staff on rationale across departments*

*Screen for eligibility and if eligible consent participants*

# Methodology Study Design

- ▣ Methodological
  - Psychometrics of HNV tool compared the use of HNV w/ the existing Morrow Scale for NV
  - Sample N=153
- ▣ Consent vs. No consent
- ▣ Institutional Review Board (IRB) approved:
  - Waiver of consent
  - *IRB Decision Rationale*: distress of symptoms & treatment must not be interrupted

# Experimental Groups

- Patients at risk were placed in 3 groups selected in the study N=153

Groups	Admission Baseline	Pre-Rx NV	Rx-anti NV	Post-test NV
Gp I: With NV n=51	↑	↑	Yes	↓
GpII: Chemo Pts n=50	0	↑ chemo	Yes	↓
GpIII: Control n=52	0	0	0	0

**Figure 1: Design of the study**

↑ =Nausea /vomiting expected to be elevated

↓=Nausea /vomiting expected to decrease    Pts= Patients

0 =Nausea /vomiting expected to be low

# Findings

- ▣ HNV tool had high inter-rater accuracy of responses (Kappa test =.851,  $p < .001$ ).
- ▣ Concurrent validity between the HNV tool & Morrow's worst nausea ratings were significant at Time 1 ( $r=.318$ ,  $p=.03$ ).
- ▣ HNV was found to measure fine differences between and within groups, establishing sensitivity.



# Comparability of the groups

- ▣ Demographic and medical background data were cross tabulated with the groups' numbers to assess the comparability of the three groups.
- ▣ Chi square tests were non-significant for ethnicity, gender, coronary artery disease, hypertension, diabetes, congestive heart failure, and other medical surgical diagnosis and risk for PONV.
- ▣ On admission, the groups differed in terms of nausea, as expected!

# Concurrent Validity

- ▣ Calculated between Morrow's drug efficacy rating and changes in Halpin ratings.
- ▣ Gain scores were correlated with Morrow ratings of drug usefulness, where 1 meant "very useful" and 4 meant "doesn't seem to help".
  - The gains in Halpin Nausea ratings had a significant correlation ( $r = -.281$ ,  $p = .019$ ,  $n = 69$ ) with Morrow ratings of drug effectiveness.
  - The Halpin vomiting change scores varied in the expected direction, but the correlation was not significant ( $r = -.201$ ,  $p = .097$ ,  $n = 69$ ).

# Concurrent Validity

- ▣ The highly significant correlations were between the Halpin NV scales and the respective Morrow scales that indicated whether or not the subject was presently experiencing nausea or vomiting.
- ▣ These six correlations ranged from  $-.852$  to  $-.619$ , and were significant at the  $.001$  level.

# HNausea 0-5

	Measure	Descriptions
0	None	No nausea
1	Anticipated	Nausea is <i>anticipated</i> and prophylaxis medications may be given.
2	Mild	Nausea reported. Able to tolerate food or medications by mouth.
3	Moderate	Nausea persisting. Lacks appetite. Able to eat small meals occasionally.
4	Great	Nausea ongoing. No appetite. Unable to tolerate food/medications by mouth.
5	Severe	Nausea with Dry Heaves reported

# HVomiting 0-5

	Measure	Descriptions
<b>0</b>	<b>None</b>	No vomiting
<b>1</b>	<b>Anticipated</b>	Vomiting is <i>anticipated</i> and prophylaxis medications may be given.
<b>2</b>	<b>Mild</b>	1 -2 episodes in 12 hours, small amount of emesis.
<b>3</b>	<b>Moderate</b>	3-5 episodes in 12 hours. Vomiting persist.
<b>4</b>	<b>Great</b>	6 episodes in 12 hours.
<b>5</b>	<b>Severe</b>	≥ 7 episodes in 12 hours, intractable, incessant, retching with emesis.

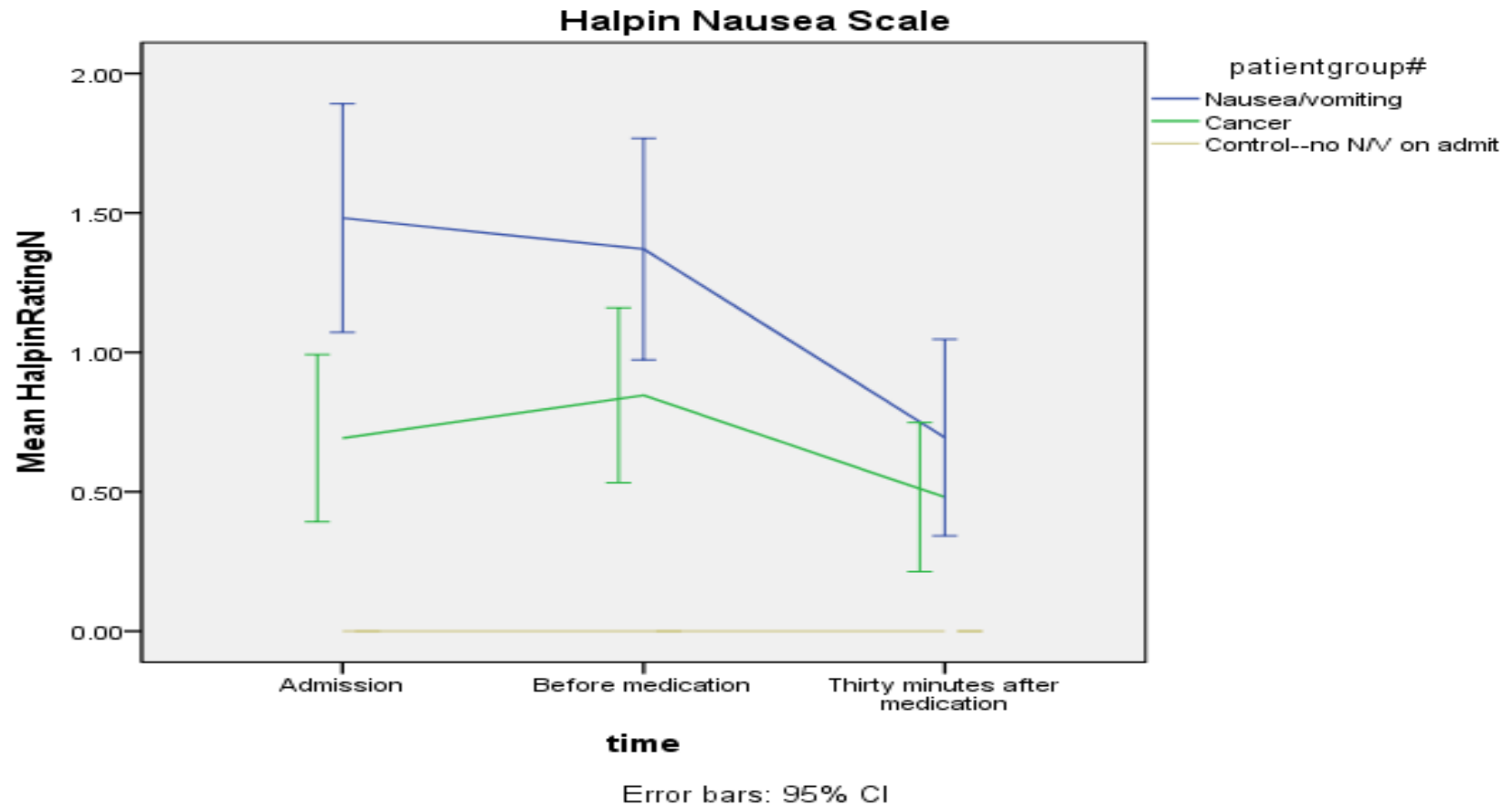
# Sensitivity

- ▣ A two-way analysis of variance with repeated measures on **nausea scores** indicates that there was a significant group effect ( $F(2,160)=29.131$ ,  $p=.000$ ), meaning that the groups differed in their feelings of nausea with a:
  - significant time effect ( $F(1,160)=14.465$ ,  $p=.000$ ) meaning that there were differences between time 1, 2, and 3
  - significant time by group interaction effect ( $F(2,160)=7.306$ ,  $p=.001$ ) meaning that feelings of nausea is jointly determined by both belonging to a specific group and the time period when nausea was measured

# Sensitivity HVS

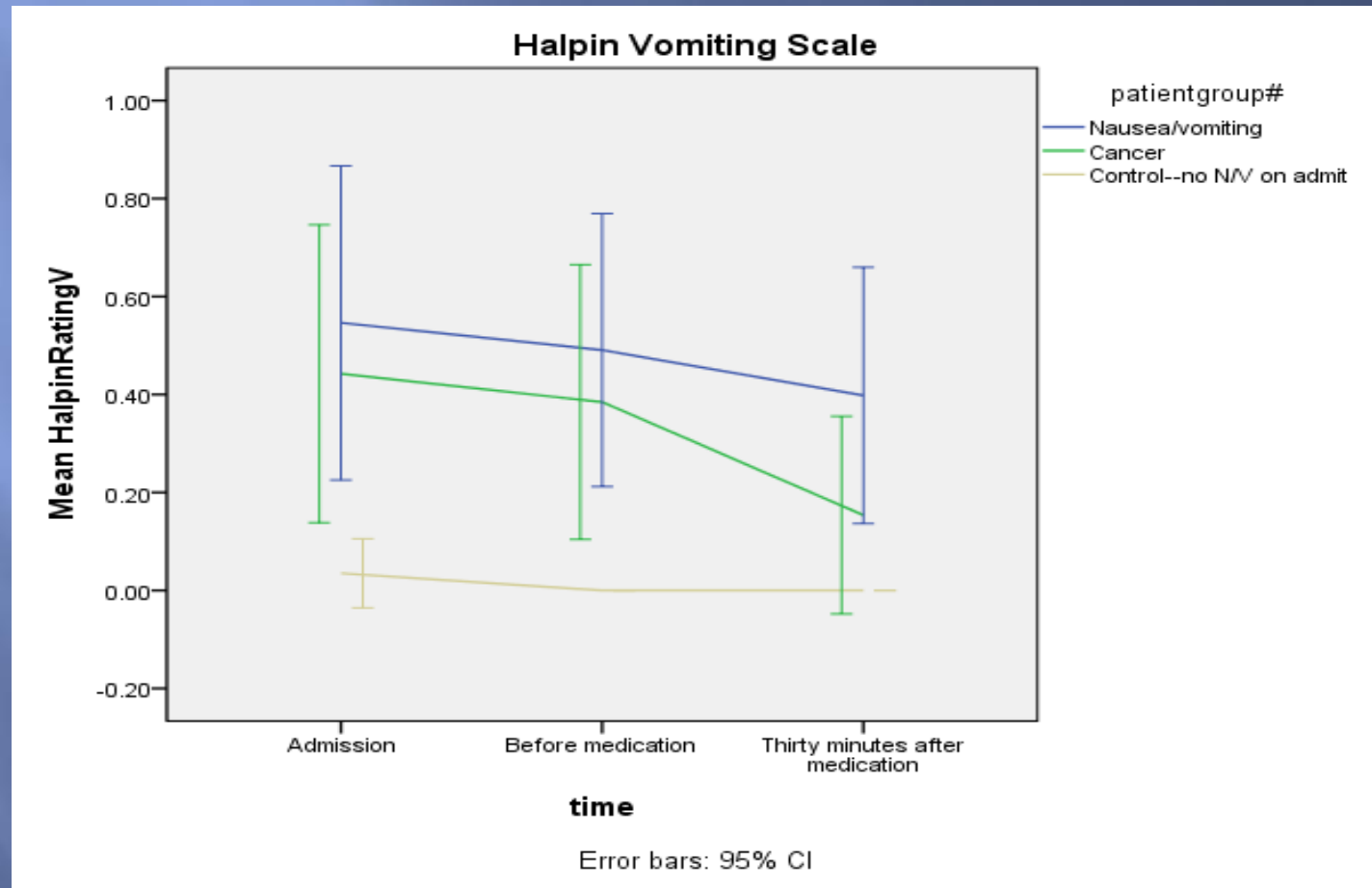
- ▣ For the **vomiting scale**,
  - The group factor was significant
    - ▣ ( $F(2,160) = 5.933, p = .001$ ), as was the time factor
    - ▣ ( $F(1,160) = 6.509, p = .012$ )
  - The group by time interaction was not significant
    - ▣ ( $F(2,160) = 1.414, p = .246$ )
- ▣ As was the case with the nausea scale, belonging to a specific group was a determinant on the feelings of vomiting.

# Figure 2: Group means for nausea ratings at three time intervals





# Figure 3: Group means for vomiting ratings at three time intervals



# Predictability of scales re: PONV

- ▣ Prior to surgery assessment of nausea and vomiting history assist in management of symptoms
  - One of the questions that we asked: *“Is there a relationship between gender and risk for postoperative nausea and vomiting (PONV)?”*
- ▣ A multiple regression was conducted to determine if gender is a predictor variables of PONV:
  - History of smoking, motion sickness, nausea on admission
- ▣ Results were positively related to the gender of the patients ( $F=8.307$ ,  $df=2$ ,  $p=.000$ ).

# Discussion

- ▣ The findings support the HNV scale with descriptors
  - The scale can be translated into practice
  - Used by bedside registered nurses during the care of patients at the time of assessment
  - Nausea and vomiting symptoms are managed when assessed using scales to measure the symptom intensity, severity & duration.
  - HNV scale is a valid and a reliable tool that:
    - Benefits operative patients in conjunction with PONV risk
    - Cancer treatment patients
    - All patients experiencing or expected to experience NV

# Limitations of Study

- ▣ The HNV scales were tested with adult patients.
- ▣ Sensitive for use with children, or patients from different cultures was not tested.
- ▣ The practicality of the tool requires a wider audience of nurses and patients use & qualitative researchers documentation
- ▣ There was a Low *n* for the Morrow ratings

# Future Research

## ▣ Recommend

- Further study on application of the HNV in different age groups responses
- Further study applications of the HNV in specific cultural groups
- Test the tool's practicality and useful in various populations within clinical applications (e.g. Women's Health, Emergency Departments)

## ▣ Thank you! Citosl. Děkuji!

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