



Le 11 JUIN 2024

Palais de la Bourse

**Sophie MORIN Médecin de SOS INSTITUT BERGONIE** 

4<sup>E</sup> POST-ASCO EN NOUVELLE-AQUITAINE



#### Liens d'intérêts

Aucun (de Queen B à Queen T...!)



# La star

Des SOS à l'ASCO





# Comparative Effectiveness Trial of Early Palliative Care Delivered via Telehealth versus In Person among Patients with Advanced Lung Cancer: The REACH PC Trial

Joseph A. Greer PhD & Jennifer S. Temel MD on behalf of:

Chardria Trotter MPH MBA, Vicki A. Jackson MD MPH, Simone Rinaldi APN-BC, Mihir Kamdar MD, Areej El-Jawahri MD, Nora Horick MS, Kedie Pintro MS, Dustin Rabideau PhD, Josephine Feliciano MD, Isaac Chua MD MPH, Konstantinos Leventakos MD, Stacy Fischer MD, Toby C. Campbell MD, Michael W. Rabow MD, Finly Zachariah MD, Laura C. Hanson MD, Sara F. Martin MD, Maria Silveira MD, and the REACH PC Investigators













#### **Enhancing Access to Early Palliative Care**

### ASCO Guidelines for Early Palliative Care

- Recommend integrating palliative care from diagnosis of advanced cancer
- Limited scalability

#### A Promising Solution: Telehealth

- Overcomes access barriers
- Reduces financial toxicity

#### **Study Question:**

Is the effect of delivering early palliative care via secure video equivalent to in-person care for patients with advanced lung cancer?











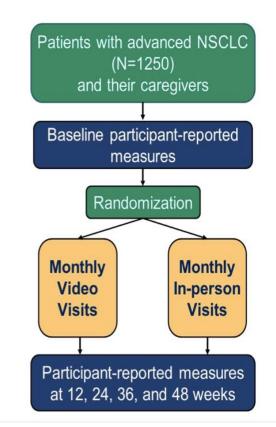
#### **Study Aims and Design**

#### **Primary Aim:**

 To evaluate the equivalence of the effect of delivering early palliative care using video versus in-person visits on patientreported quality of life

#### **Secondary and Exploratory Aims:**

- Satisfaction with care
- Caregiver attendance at study visits
- Mood symptoms



#### **Study Procedures**

- Enrollment: 6/14/2018 to 5/4/2023
- Random assignment (1:1) to groups
- Technology provided if needed
- Intervention:
  - Monthly palliative care visits
  - Initial in-person encounter in video group to establish rapport
  - Clinician documentation of topics discussed during visits







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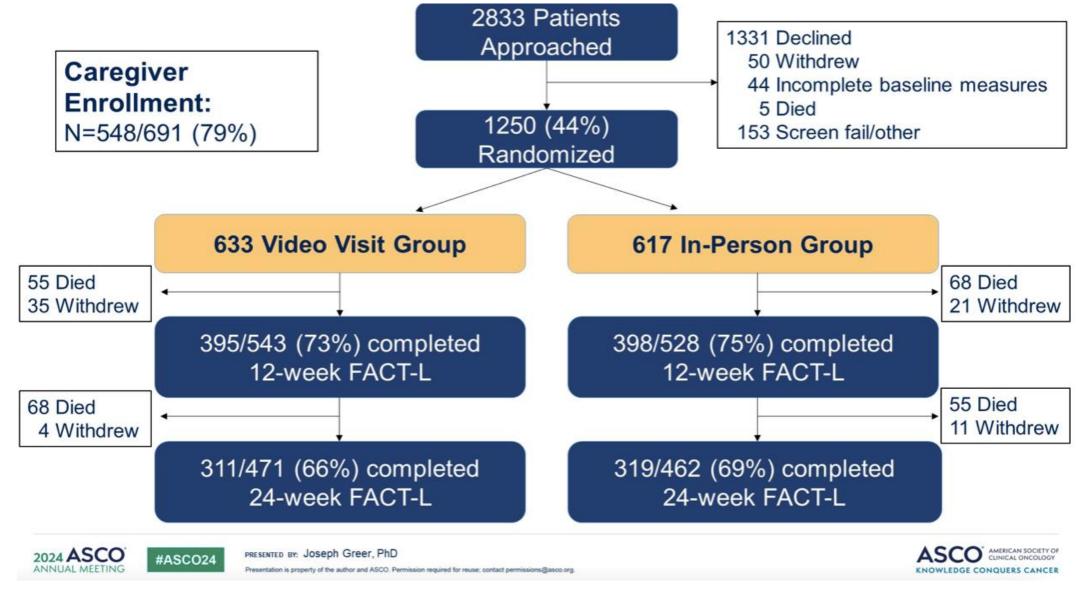




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#### **Patient Characteristics**

Characteristic	Video Visit Group N (%)	In-Person Group N (%)
Age, Mean Years (SD)	65.5 (10.9)	65.5 (10.6)
Woman	356 (56%)	318 (52%)
American Indian/Alaskan Native Asian Black or African American Native Hawaiian/Pacific Islander White Other	4 (<1%) 32 (5%) 57 (9%) 2 (<1%) 524 (83%) 21 (3%)	4 (<1%) 32 (5%) 72 (12%) 4 (<1%) 502 (82%) 10 (2%)
Hispanic or Latino/x	29 (5%)	30 (5%)
Married/Partnered Single/Divorced/Widowed/Other	420 (67%) 210 (33%)	409 (67%) 203 (33%)

#### **Clinical Characteristics**

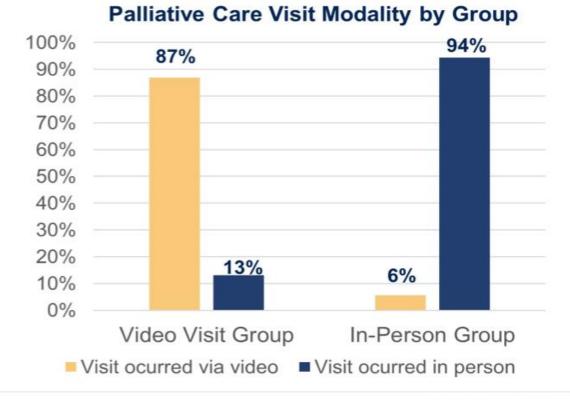
Characteristic	Video Visit Group, N (%)	In-Person Group, N (%)
ALK EGFR ROS RET Other or no mutation	28 (4%) 113 (18%) 6 (<1%) 11 (2%) 475 (75%)	26 (4%) 102 (17%) 0 (0%) 7 (1%) 482 (78%)
Platinum-based doublet chemo (± 3 <sup>rd</sup> agent) Radiation Oral targeted therapy Immunotherapy alone Single agent IV chemotherapy Concurrent chemotherapy and radiation No treatment	257 (41%) 138 (22%) 126 (20%) 93 (15%) 7 (1%) 4 (<1%) 8 (1%)	277 (45%) 123 (20%) 114 (19%) 72 (12%) 8 (1%) 5 (<1%) 18 (3%)



#### Intervention Delivery & Fidelity

Number of Palliative Care Visits by 24 Weeks Mean (SD)

Video Visit	In-Person
4.7 (2.5)	4.9 (2.7)







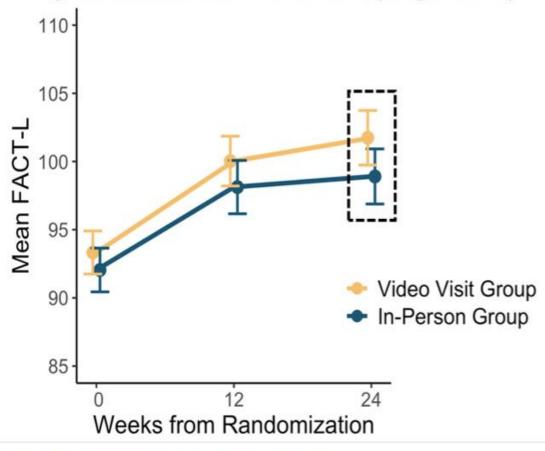
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#### Primary Outcome: Patient Quality of Life (QOL)

Higher scores indicate better QOL (range: 0-136)



## Adjusted Mean FACT-L at 24 Weeks:

Video Visit Group: 99.7

In-Person Group: 97.7

Difference (90% CI): 2.0 (0.1, 3.9)

p=0.04 for equivalence





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#### **Secondary Outcomes at Week 24**

Outcome Measure	Video Visit Group Estimated Mean/Proportion	In-Person Group Estimated Mean/Proportion	Difference 95% (CI)	Р
Satisfaction with Care† Patient report, mean Caregiver report, mean	41.3 37.2	41.0 36.8	0.3 (-1.0, 1.7) 0.4 (-1.5, 2.3)	>0.99 >0.99
Attendance of Caregiver at Visits proportion	36.6%	49.7%	-13.0% (-17.6, -8.6)	<0.001

<sup>†</sup>Higher scores on the Satisfaction and Care Delivery Questionnaire indicate greater satisfaction





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#### CONCLUSION

#### Equivalence de la télé consultation en SOS

- Mise en valeur la TELEMEDECINE
- Etude de SOS
- Changement de pratiques :

Développer la téléconsultation/télémédecine pour une offre de SOS plus écologique, économique et équitable sur la région

Discussion sur l'indication clinique à une téléconsultation







### Multi-Site Randomized Trial of Stepped Palliative Care for Patients with Advanced Lung Cancer

Jennifer S. Temel MD, Vicki A. Jackson MPH, MD, Areej El-Jawahri MD, Simone P. Rinaldi MSN, ANP-BC, ACHPN, Laura A. Petrillo MD, Pallavi Kumar MD, Kathryn A. McGrath MD, Thomas W. LeBlanc MD, Arif H. Kamal MD, Christopher A. Jones MD, Dustin J. Rabideau PhD, Nora Horick MS, Kedie Pintro MS, Emily R. Gallagher Medeiros RN, Kathryn E. Post PhD, RN, ANP-BC, Joseph A. Greer PhD





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#### Early Integrated Palliative Care Models Require Substantial Resources





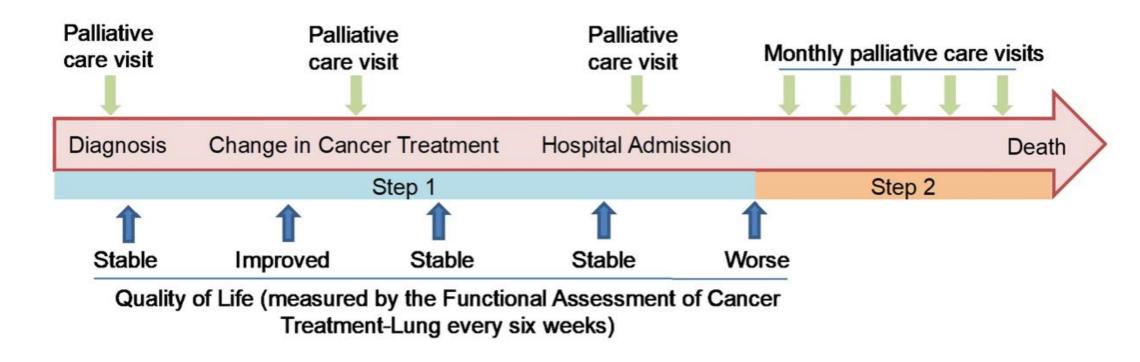
- Interventions delivered by specialty-trained palliative care physicians or advanced practice providers.
- Insufficient number of clinicians for monthly visits with all patients diagnosed with serious cancer.

#### **Study Question**

Can a stepped palliative care intervention tailored to a patient's illness course and care needs achieve the same effect as an early integrated palliative care model while utilizing fewer palliative care resources?



#### **Stepped Care Model of Early Palliative Care**







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#### **Study Design and Aims**

- Multi-site randomized controlled trial of stepped palliative versus early integrated palliative care (monthly palliative care visits).
- Primary Aim: To establish the non-inferiority of stepped palliative care on patient quality of life at 24 weeks, as measured by the Functional Assessment of Cancer Therapy-Lung (FACT-L), with a non-inferiority margin of - 4.5 points.

#### Secondary Outcomes:

- Palliative care utilization as measured by number of visits.
- Patient-reported communication about end-of-life care.
- Hospice utilization as measured by length of stay in hospice.





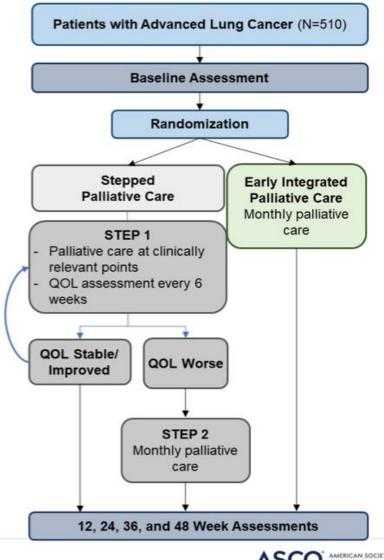
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#### **Study Procedures**

- Research assistants reviewed health records to identify eligible patients.
- After clinician approval, research assistant approached and consented patients.
- Patients were randomized 1:1 to study group.
- Research assistants collected patientreported outcomes throughout the study.
- Enrollment period 2/12/18 12/15/22.



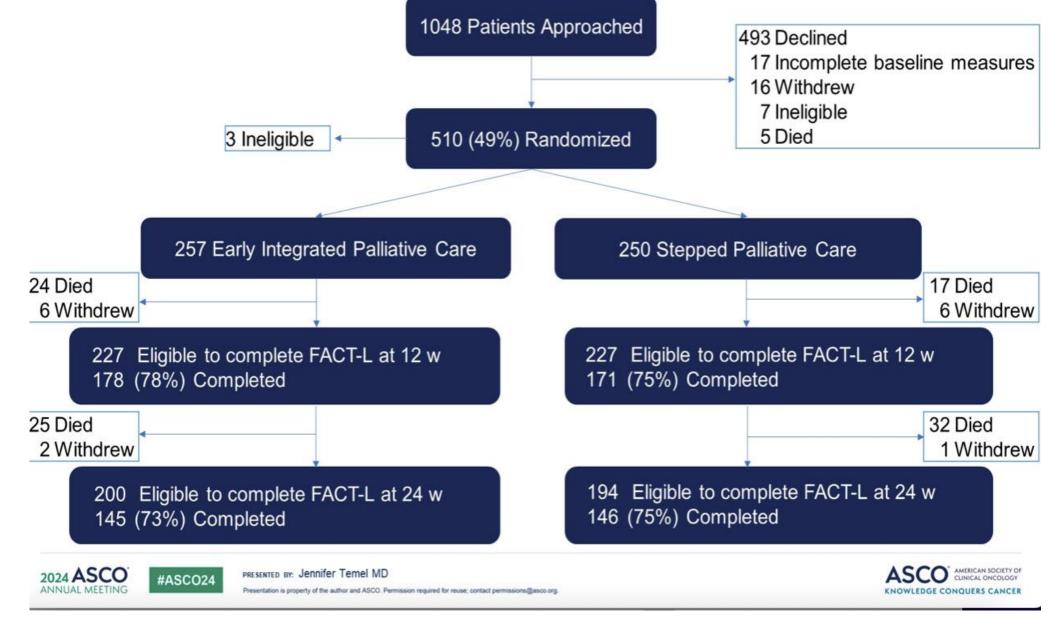




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#### **Participant Characteristics**

Characteristic	Early Integrated PC (N=257)	Stepped PC (N=250)
Age, Mean Years (SD)	66.1 (11.1)	66.8 (9.2)
Woman	130 (51%)	130 (52%)
American Indian or Alaskan Native Asian Black or African American Native Hawaiian or Pacific Islander White	2 (<1%) 11 (4%) 28 (11%) 0 (0%) 212 (83%)	2 (<1%) 3 (1%) 29 (12%) 0 (0%) 215 (86%)
Hispanic or Latino/x	5 (2%)	3 (1%)
ECOG PS 0 ECOG PS 1 ECOC PS 2	64 (25%) 153 (60%) 40 (16%)	61 (24%) 153 (61%) 36 (14%)

PC = palliative care, ECOG PS = Eastern Cooperative Group Performance Status

#### **Disease Characteristics**

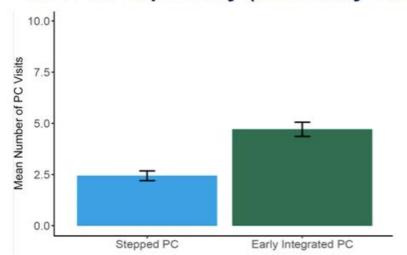
Characteristic	Early Integrated PC (N=257)	Stepped PC (N=250)
Non-Small Cell Lung Cancer Small Cell Lung Cancer Mesothelioma	203 (79%) 47 (18%) 7 (3%)	194 (78%) 53 (21%) 3 (1%)
ALK EGFR ROS RET Other or no mutation	10 (4%) 39 (15%) 2 (<1%) 2 (<1%) 204 (79%)	11 (4%) 37 (15%) 3 (1%) 1 (<1%) 198 (79%)
Platinum-based chemotherapy (+/- 3 <sup>rd</sup> agent) Radiation Oral targeted therapy Immunotherapy alone Single agent intravenous chemotherapy	116 (45%) 38 (15%) 50 (20%) 39 (15%) 9 (4%)	128 (51%) 54 (22%) 40 (16%) 22 (9%) 3 (1%)



#### **Intervention Delivery**

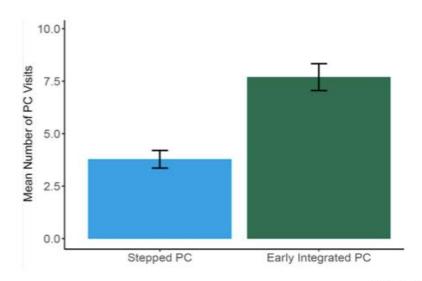
#### Number of PC Visits by Week 24

Adjusted means: 2.4 vs 4.7 visits per patient
Difference (95% CI): -2.3 (-2.7, -1.8)
p < 0.001 for superiority (secondary outcome)



#### Number of PC Visits by Week 48

Adjusted means: 3.8 vs 7.7 visits per patient Difference (95% CI): -3.9 (-4.6, -3.1)



PC = palliative care



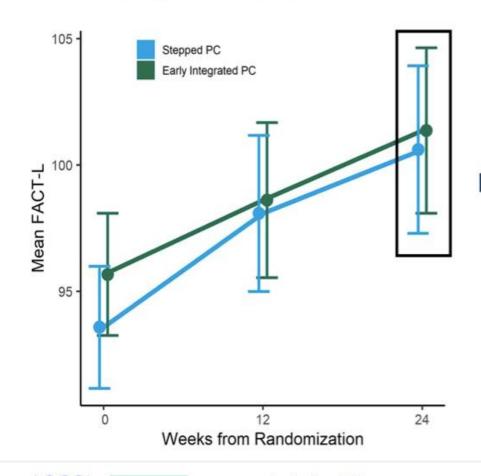
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#### **Quality of Life**



#### **Quality of Life at Week 24**

Adjusted means: 100.6 vs 97.8

Difference (lower one-sided 95% CL): 2.9 (-0.1)

Non-inferiority margin: -4.5

p < 0.001 for non-inferiority (primary outcome)

Higher scores indicate better quality of life

PC = palliative care





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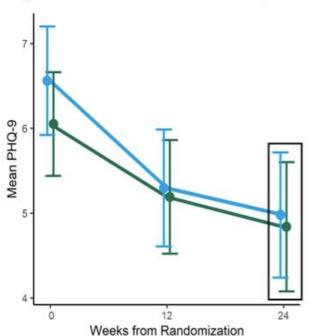




#### **Depression and Coping at Week 24**

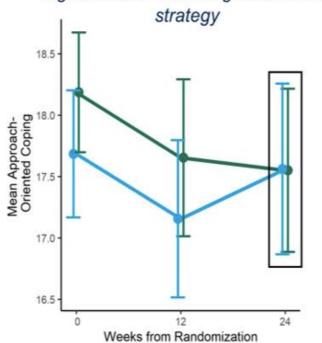
#### Depression

Difference (95% CI): -0.4 (-1.3, 0.5) Higher scores indicate worse depression



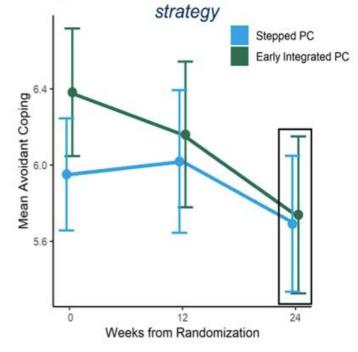
#### **Approach-Oriented Coping**

Difference (95% CI): **0.2 (-0.7, 1.0)**Higher scores indicate greater use of



#### **Avoidant Coping**

Difference (95% CI): **0.0** (-0.5, 0.5) Higher scores indicate greater use of





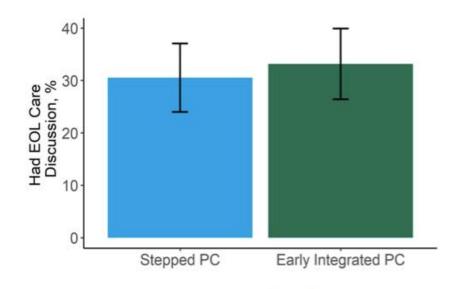


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#### Patient-Reported End-Of-Life Discussion



Bars represent "Yes" responses

#### **End-of-Life Discussion**

Difference (lower one-sided 95% CL): -2.6% (-10.4%)
Non-inferiority margin: -10%

p = 0.09\* for non-inferiority (secondary outcome)

\* significant at pre-specified 0.15 significance level

PC = palliative care





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# Modèle de Soins Palliatifs plus EVOLUTIF et ECONOME

Evolution de pratiques de suivi palliatif et SOS

- Step 1 : Suivi ?
- Step 2 : Accentuation des visites sur les derniers mois de vie avec travail de coordination ville hôpital
- E SANTE et IDEC ?









#### Risk Prediction Model for Taxane-Induced Peripheral Neuropathy (TIPN) in Patients with Early-Stage Cancer Receiving Taxane Therapy: SWOG S1714

Meghna S. Trivedi, Joseph M. Unger, N. Lynn Henry, Amy K. Darke, Daniel L. Hertz, Thomas H. Brannagan, Stephanie J. Smith, Bryan P. Schneider, William J. Irvin Jr, Amanda R. Hathaway, Amy C. VanderWoude, Vinay K. Gudena, Paula Cabrera-Galeana, Mary Orsted, Michael LeBlanc, Michael J. Fisch, Dawn L. Hershman















# S1714: A prospective observational cohort study to develop a predictive model of taxane-induced peripheral neuropathy in cancer patients

Receiving taxane regimen for Stage I-III cancer: Primary NSCLC Primary Breast Primary Ovarian

N = 1,336 enrolled at 105 sites in the NCORP **Baseline Evaluation** 

Study Assessments at Weeks 4, 8, 12, 24, and 52 after registration

Study Assessments at Weeks 104 and 156 after registration

- EORTC QLQ CIPN-20
- CTCAE
- Additional PROs
- Neurosensory testing
- Functional testing
- Biospecimen submission

- EORTC QLQ CIPN-20
- CTCAE
- Additional PROs
- Neurosensory testing
- Functional testing
- Biospecimen submission
- · Treatment adjustments

- EORTC QLQ CIPN-20
- CTCAE



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PRESENTED BY: Meghna S. Trivedi, MD, MS

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NCT# 03939481





#### **RESULTS: Training Set**

- In the training set, adverse risk factors for TIPN were:
  - Receipt of paclitaxel (vs docetaxel)
  - Stage 2/3 (vs 1) disease
  - Planned duration of taxane >12 weeks (vs <12 weeks)</p>
  - Comorbidity: (≥1 vs none)
    - o Diabetes, autoimmune disease, or moderate kidney disease
  - Race/Ethnicity: (vs non-Hispanic White or Asian)
    - Black, Native American, Pacific Islander, multiple race
    - o Hispanic ethnicity

	Overall cohort N=1278	Low risk: 0-1 risk factors N=267 (35%)	High risk: ≥ 2 risk factors N=501 (65%)	p value
% experiencing TIPN	62%	48.7%	70.9%	<.001





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#### Odds of TIPN by model type

Number of Risk **Factors** 

**Odds Ratio** (95% Confidence Interval)

p-value

0.5

1.0

2.5

5.0

#### **Risk Model Ordinal Increase** Per Additional Risk Level Q1 vs. Q2, Q2 vs. Q3, etc. 1.69 (1.49-1.92) <.001 2-level Model High Risk >Median vs. <Median 2.36 (1.86-2.99) <.001 4-Level Model (Quartiles) 0 factors Low Risk [reference] Low-Intermediate Risk 1 factor 1.75 (1.14-2.69) 0.01 2 factors 2.89 (1.91-4.35) <.001 High-Intermediate Risk High Risk 4.93 (3.17-7.68) 3-5 factors <.001





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C-statistic for the 4-level risk prediction model: 0.63 (95% CI, 0.60-0.66)



10.0







# Alliance 222001: A Randomized, Double-Blind, Placebo-Controlled Study of Oxybutynin versus Placebo for the Treatment of Hot Flashes in Men Receiving Androgen Deprivation Therapy

Brad J. Stish, Gina L. Mazza, Jones T. Nauseef, Michael Sandon Humeniuk, Thomas J. Smith, Cindy Tofthagen, Dayssy Alexandra Diaz Pardo, Christopher Chay, Andrew Huang, Kushal Naha, Scott T. Tagawa, Selina Chow, Lucile L. Adams-Campbell, Paul J. Novotny, Charles L. Loprinzi

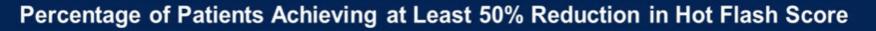




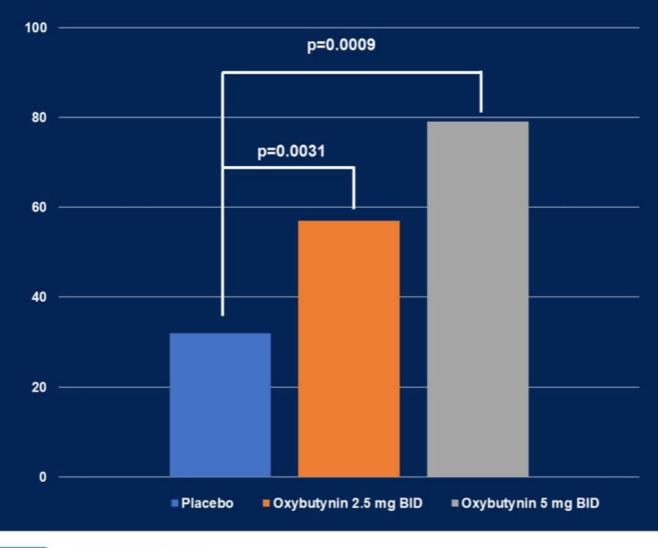
PRESENTED BY: Brad Stish, MD















PRESENTED BY: Brad Stish, MD





