

# RISE UP: <u>Revolutionizing Investigations to StEp Up Prevention for Breast Cancers:</u> November 1-3, 2024 San Francisco, California

This new, interdisciplinary conference focuses on a bold reimagining of breast cancer prevention and treatment through cross-disciplinary exchange of ideas and out-of-the-box thinking. We will focus on minimizing the long-term side effects associated with therapeutic interventions using early endpoints, recognizing that a key to better uptake in preventive therapies is better tolerability. An important goal of the meeting is to foster interdisciplinary approaches across the gynecology and oncology and primary care fields and build collaborations focused on reducing breast cancer incidence.

A primary goal of RISE UP is to consider breast cancer prevention across a woman's life course, and translate scientific findings to actionable, accessible medications and strategies to reduce breast cancer risk among individuals and communities. Currently, there is strong evidence that those who live in more impoverished areas often have more aggressive forms of breast cancer. We hope this meeting will spur new treatment and prevention strategies that will bring important options to those at most risk for breast cancer and who are diagnosed with the most aggressive disease. Through this multi-disciplinary conference, bringing together epidemiologists, policy makers, and more, we hope to present research that specifically examines the impact of deprivation, racism, environmental exposure burden, and place of residence on the etiology and incidence of breast cancer to better understand these disparities.

This conference is designed for women's health providers of all specialties, scientists, advocates, policymakers and innovators to RISE UP to the challenge of reducing breast cancer incidence and mortality in our lifetime

At the completion of the conference, participants should be able to:

- Identify new classifications and treatments for breast cancer.
- Discuss new ways to assess risk for breast cancer and what is available for risk reduction.
- Survey the potential for hormonal products as risk-reducing interventions for breast cancer.
- Analyze the impact of restricted access to reproductive services on the health and well-being of women.
- Create opportunities to lower the barriers to participation in trials through patient-centered design and strategies to make trials more accessible regardless of insurance.
- Identify the differences in biologic effects on breast tissue and other organs between selective estrogen receptor modulators and tissue specific estrogen complexes being studied for breast cancer risk reduction.
- Describe the amount of sustained weight/fat loss likely needed for significant breast cancer risk reduction and metabolic pathways impacted for breast cancer risk reduction.
- Evaluate the challenges and potential benefits in studying the GLP-1 and GIP receptor agonists for breast cancer risk reduction.
- Describe the impact of new restrictions on abortion since the Supreme Court's Dobbs decision, including the effects on people seeking services, on general obstetrical care, and on medical education.
- Examine potential common denominators for increased risk in maternal mortality and breast cancer in African American women.

# Program Schedule/Agenda

# Friday, November 1, 2024

12:00 PM- 12:15PM	Welcome: A Paradigm Shift in Women's Health: A Holistic View of hormone cycle control and breast cancer risk reduction (Laura Esserman, MD, MBA (UC San Francisco))			
12:15 PM- 12:30PM	Keynote Address: The Launch of the Birth Control Pill and How It Should Inform Decisions Today (Liz Watkins, PhD (UC Riverside))			
12:30 PM- 12:30PM	Session 1: Tumor Classification in Optimizing Therapy and Outcomes Introduction: Laura van't Veer, PhD, UC San Francisco			
12:30 PM- 12:45PM	Response Predictive Subtypes (RPS): origins, evolution, and future directions (Denise Wolf, PhD (UC San Francisco))			
12:45 PM- 12:55PM	Analysis of I-SPY2-990 Data to Reveal New Pathways of Resistance in Response Predictive Subtype-Negative and Response Predictive Subtype-Discordant Breast Cancers (Tam Binh V. Bui, MSc, MD (UC San Francisco))			
12:55 PM- 1:05PM	Improving risk stratification and prediction of response to neoadjuvant therapy (NAT) by serial monitoring of circulating tumor DNA (ctDNA) in high-risk early-stage breast cancer (Mark Magbanua, PhD (UC San Francisco))			
1:05 PM-1:15PM	The steroid hormone receptor signature (SRS) in stage II/III triple negative breast cancer correlates with highly actionable functional protein drug target activation and			
1:15 PM-1:25PM	is associated with early recurrence (Julie Wulfkuhle (George Mason University)) Examining APOBEC3B in TCGA and ICGC Breast Cancer Datasets Reveals Altered Drug Metabolism Pathways (Joel Pardo (University of Minnesota))			
1:25 PM-1:40PM	Diagnosis 2030: What Al Can (and Can't) do for you (Sandy Borowsky, MD (UC Davis))			
1:40 PM-2:00PM	Summary Panel (moderated by Lajos Pusztai, MD, DPhil (Yale), and Amy Delson, Patient Advocate)			
2:15 PM-2:20PM	Session 2: Imaging as a Predictive Biomarker in Clinical Trials Early Endpoint Introduction (moderated by Nola Hylton, PhD (UCSF), Elissa Price, MD (UCSF), Diane Heditsian, Patient Advocate, Jan Tomlinson, Patient Advocate)			
2:20 PM-2:45PM 2:45 PM-2:55PM	Ultrafast MRI in Post Treatment Assessment (Yiming Gao, MD (NYU)) Patient Perspectives (Jan Tomlinson, Patient Advocate, Diane Heditsian, Patient Advocate)			
2:55 PM-3:02PM	Temporal Changes in Mammographic Breast Density as a Predictive Biomarker across the Breast Cancer Continuum (Gretchen Gierarch, PhD, MPH (NIH))			
3:02 PM-3:09PM	Real World Study of the Effect of T+Ai on Background Parenchymal Enhancement (BPE) (Wayne Tilley, PhD (University of Adelaide))			
3:09 PM-3:16PM	Interim Results from a Multimodality Screening Program Reveal Low Recall Rates and Breast Cancer Detection in BRCA Mutation Carriers Undergoing Biannual DCE-MRI (Hatice Basdag (University of Chicago))			
3:16 PM-3:23PM	Short-term declines in breast density by treatment-associated endocrine symptoms after tamoxifen therapy: Examining the role of CYP2D6 phenotype and tamoxifen metabolites (Cody Ramin, PhD (Cedars Sinai Medical Center)			
3:23 PM-3:30PM	Validation of an MRI-Based Predictive Model for Treatment Tailoring in the I-SPY 2 Trial (Wen Li, PhD (UC San Francisco))			
3:30 PM-3:45PM	Panel Discussion of Speakers			
3:45 PM-4:15PM	Break and Exhibitors Light Refreshments			
4:15-6:30pm	Session 3: Increasing Efficiency While Decreasing Toxicity of Local and Systemic Therapy (NOT FOR CME CREDIT)			
6:30 PM-7:30PM	Poster Session #1 (NOT FOR CME CREDIT): Tumor classifiers, imaging endpoints, toxicity reduction			

# Saturday, November 2, 2024

	Session 4: Risk Prediction and Biomarkers for Prevention Trials
8:00 AM-8:15AM	Introduction (Adetunji Toriola, MD, PhD, MPH (WUSTL))
8:15 AM-8:30AM	Senescence-Based Deep Learning Predicts Breast Cancer Risk Using H&E Core Biopsy Images From Healthy Women (Mark Powel, MD, MPH (Buck Institute for
	Research on Aging))
8:30 AM-8:45AM	Predicting Risk of Future Breast Cancer based on Screening MRI Features
0.50 AW 0.45AW	(Suleeporn Sujichantararat, PhD (University of Washington))
8:45 AM-9:00AM	Stromal Inflammation as a Driver of Parity-related Breast Cancer Etiologic
	Heterogeneity: Implications for Precision Prevention in a Sub-Saharan African
	Population (Mustapha Abubakar, MD, PhD (National Cancer Institute, NIH))
9:00 AM-9:15AM	Characterizing the Immune and Non-immune Landscape of Premalignant and
	Preinvasive Breast Lesions (Vagmi Luhar (National Cancer Institute))
9:15 AM-9:20AM	Advocate Presentation: Carmelita Austin-Schreher (Patient Advocate)
9:20 AM-9:30AM	Session Summary (Laura Fejerman, PhD (UC Davis))
	Session 5: Applying the Lessons of Breast Cancer Biology to Prevention
9:35 AM-9:40AM	Introduction (Carol Fabian, MD (University of Kansas))
9:40 AM-	PrevenNng Breast Cancer with Molecules TargeNng the Estrogen Receptor:
10:00AM	SERMs,
	SERDs, and TSECs (Barry Komm, PhD)
10:00 AM-	DCIS: RECAST - An Adaptive Platform Trial to Identify Women With Hormone
10:08AM	Receptor Positive DCIS who are Candidates for Active Surveillance (Kelly Hewitt,
40.00.414	MD (University of Utah))
10:08 AM-	Refining tamoxifen dose for premenopausal breast cancer risk reduction
10:16AM	(RENAISSANCE). A Phase II single-arm trial (Seema Khan, MD (Northwestern University)
10:16 AM-10:24	Beneficial effects of bazedoxifene plus conjugated estrogens (Duavee) on breast
10.10 AW 10.24	cancer risk and metabolic biomarkers in a rat model of obesity, menopause, and
	breast cancer (Erin Giles, PhD (University of Michigan))
10:24 AM-	Breaking the Obesity-Breast Cancer Link: Diet, Drug and Surgical Strategies
10:45AM	(Stephen Hursting, PhD (University of North Carolina))
10:45 AM-	Resistance and aerobic Exercise for Prevention in women with Dense breasts (REP-
10:53AM	D): A Trial in Progress (Jennifer Ligibel, MD (Dana Farber Cancer Institute))
10:53 AM-	Time restricted Eating And Metformin (TEAM) in invasive breast cancer or DCIS: a
11:01AM	randomized window of opportunity trial. Preliminary safety analysis (Irene Maria
11:01 AM-	Briata, PhD (E.O. Ospedali Galliera))
11:09AM	Low dose TamOxifen and LifestylE changes for bReast cANcer prevention (TOLERANT Study): Study Protocol of a Randomized Phase II Biomarker Trial in
TT.USAW	Women at Increased Risk for Breast Cancer (Matteo Lazzeroni, MD (European
	Institute of Oncology IRCCS))
11:09 AM-	Q&A, Summary, Persistence in Prevention Award
11:30AM	
11:30 AM-	Poster Session #2 (NOT FOR CME CREDIT): Prevention studies in progress,
1:15PM	Turnaway study
	Session 6: The Consequences of the Dobbs Decision: A Look at the Impact in
	Women's Health
2:15 PM-2:45PM	The Effect of the Dobbs Decision on Patients and Providers Nationally and in the
	Southeast (Jamila Perritt, MD (Physicians for Reproductive Health))
2:45 PM-3:00PM	Care Post-Roe Study: Documenting Poor Quality Care Since Dobbs (Daniel
0.00 DM 0 45DM	Grossman, MD (UC San Francisco))
3:00 PM-3:15PM	The Turnaway Study and End of Roe Study (Diane Greene Foster, PhD (UC San
3:15 PM-3:30PM	Francisco))  Effects of Dobbs on Medical Education (Josie Urbina, MD (UC San Francisco))
3:30 PM-3:45PM	Q&A and Discussion

	Session 7 Panel: Breast Cancer and Maternal Mortality Disparities in African American Women: Connection or Coincidence?
4:00 PM-4:05PM	Introduction (Andrea Jackson, MD, MAS (UC San Francisco))
4:05 PM-4:15PM	Infertility and Pelvic Floor Disorders in Black Women (Oluwateniola Brown, MD (Northwestern University))
4:15 PM-4:30PM	Obesity, Inflammation and Uterine Fibroids (Obianuju Sandra Madueke-Laveaux, MD, MPH (University of Chicago))
4:30 PM-4:35PM	Evolutionary trajectory of breast cancer in diverse populations (Olufunmilayo Olopade, MD (University of Chicago))
4:35 PM-4:45PM	Targeting inflammatory pathways in breast cancer (examples from I-SPY) (Frederick Howard, MD (University of Chicago))
4:45 PM-4:50PM	Advocate Perspective (Ricki Fairley, Patient Advocate)
4:50 PM-5:00PM	Panel Discussion (moderated by Olufunmilayo Olopade, MD (University of Chicago), Andrea Jackson, MD, MAS (UCSF), Ricki Fairley (Patient Advocate))
	Session 8: Improving Diversity in Clinical Trials
5:00 PM-5:05PM	Introduction (Sara Horton, MD (Howard University), Kim Rhoads, MD, MS, MPH (UC San Francisco))
5:05 PM-5:30PM	Case Study: ISPY 2 Trial and ACCESS Initiative (Priya Jayachandran, MD (University of Southern California), Dominic Zavala (University of Southern California))
5:30 PM-5:45PM	Financial Barriers and Insurance Coverage in Clinical Trials (Laura Esserman, MD, MBA (UC San Francisco), Dana Dornsife (Lazarex Cancer Foundation), Maimah Karmo (Patient Advocate)
5:45 PM-6:00PM	Poster Session #3 (NOT FOR CME CREDIT): Environmental exposures, novel prevention strategies, risk prediction, risk factors

# Sunday, November 3, 2024

7:00 AM-8:00AM	Poster Session #4 (NOT FOR CME CREDIT): Cycle Control, Post-partum weaning, menopause control, modifiable risk factors
8:00 AM-8:30AM	<b>Session 9:</b> Lecture: The Science of the Sex Hormone Ecosystem Across a Women's Lifetime (Carol Lange, PhD (University of Minnesota))
8:30 AM-8:35AM	Session 10: Postpartum Breast Cancer: Opportunities for Improving Treatment and Prevention? (Virginia Borges, MD (University of Colorado))
8:35 AM-8:40AM	Lynda Weatherby's Story (Lynda Weatherby, Patient Advocate)
8:40 AM-9:00AM	Studies of postpartum breast cancer inform novel treatment options (Traci Lyons,
9:00 AM-9:10AM	PhD ())
	Breastfeeding Attributable Risk of Triple Negative Breast Cancer in the US (Lajos
9:10 AM-9:20AM	Pusztai, MD, Dphil (Yale))
	PPBC Vit D Liver Metabolism (Pepper Schedin, PhD (Oregon Health Sciences
9:20 AM-9:30AM	University (OHSU))
	Identifying Gaps in Care Among Patients with Pregnancy and Post-Partum
9:30 AM-9:40AM	Associated
9:40 AM-9:50AM	Breast Cancer (Malcom Su, MD (University of Texas Southwestern))
	Lynda Weatherby's Story cont. (Lynda Weatherby, Patient Advocate)
9:50AM-10:15AM	Panel Discussion and Q&A
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10:15AM-	Session 11: Making Preven9on and Holis9c Care a Priority in Women's Health
11:15AM	(NOT FOR CME CREDIT)
11:15AM-	Session 12: Ideas to Implementa9on: Incorpora9ng Breast Cancer Preven9on
12:45PM	into a Holistic Approach to Women's Health(NOT FOR CME CREDIT)

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Dana Dornsife	Speaker	Honorarium - Bristol Meyer Squibb
Mikael Eriksson	Speaker	iCAD Medical Patent License
Laura Esserman	Organizing Committee Member	Investigator initiated trial for high risk DCIS funded by Moderna for Dcis phase 1 study
Silvia Formenti	Session Chair	A consultant for AstraZeneca, Bayer, Bristol Myers Squibb, Eisai, Elekta, EMD Serono/Merck, Genentech/Roche, GlaxoSmithKline, Janssen, Medimmune, Merck US, Nanobiotix, Regeneron, Varian, ViewRay and receives research support from Arcus, Bristol Myers Squibb, Celldex, Merck, Regeneron, and Varian.
Judy Garber	Organizing Committee Member	Consulting or Advisory Role: Novartis, Kronos Bio, GV20 Therapeutics, Belharra Therapeutics, Inc, Earli, Inc Research Funding: Novartis, Ambry Genetics, InVitae, Amgen Other Relationship: AACR, Diana Helis Henry Medical Foundation, James P. Wilmot Foundation, Adrienne Helis Malvin Medical Research Foundation, Breast Cancer Research Foundation, Facing our Risk of Cancer Empowered
Frederick Howard	Speaker	Consultant for Novartis and Leica Biosystems
Barry Komm	Presenter	Consultant for Sermonix Pharmaceuticals
Obianuju Sandra Madueke-	Presenter	Honoraria - Intuitive Surgical, PI Investigator - Sumitomo
Laveaux		Pharmaceutical

Olufunmilayo Olopade	Session Chair	Financial relationships with CancerIQ, 54gene, HealthWell Solutions, and Tempus, as well as research funding from Ayala Pharmaceuticals, Cepheid, Color Genomics, Novartis, and Genentech
Lajos Pusztai	Session Chair	L.P. has received consulting fees and honoraria for advisory board participation from Pfizer, Astra Zeneca, Merck, Novartis, Bristol-Myers Squibb, Stemline-Menarini, GlaxoSmithKline, Genentech/Roche, Personalis, Daiichi, Natera, Agendia, Exact Sciences, Radionetics, and institutional research funding from Seagen, GlaxoSmithKline, AstraZeneca, Merck, Pfizer and Bristol Myers Squibb
Wayne Tilley	Speaker	Scientific Advisor for Ellipses Pharma Limited and Havah Therapeutics
Hope Rugo	Session Chair	Consulting roles with Chugai, Puma, Sanofi, Napo, and Mylan; Reports institutional research support from AstraZeneca, Daiichi Sankyo, F. Hofmann-La Roche AG/Genentech, Gilead Sciences, Lilly, Merck & Co., Novartis Pharmaceuticals Corporation, Pfizer, Stemline Therapeutics, OBI Pharma, Ambrx and Greenwich Pharma
Julie Wulfkuhle	Presenter	Reports other from Theralink Technologies outside the submitted work; in addition, J. Wulfkuhle has a patent to 10,823,738-B2 issued, licensed, and with royalties paid.
Douglas Yee	Organizing Committee	Consulting fees from Martell Diagnostics; and honoraria and travel for speaking at the 'International Breast Cancer Conference'

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# **Exhibitor Acknowledgement**

We would like to acknowledge the following companies and organizations for exhibiting at this conference.

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# **Additional Support Acknowledgement**

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Breast Cancer Research Foundation