

Cannabis Partnerships Congress

Advancing Cannabinoid Science by Driving Effective Research and Overcoming Hurdles in Therapeutic Production and Regulatory Approval

Featured Speakers

CHAIRS



Bruce Bloom, CEO
CURES WITHIN REACH



Dean Petkanas,
Chief Medical Officer,
KANNALIFE SCIENCES



Jeremy Unruh,
General Counsel,
PHARMACANN LLC



Thoma Kikis,
Chief Marketing Officer,
KANNALIFE SCIENCES



Veronica Arroyave,
Sr Director, Global Health Programs,
PARTNERSHIP FOR QUALITY MEDICAL DONATIONS



Lezli Engelking,
President and Founder,
FOCUS



Elias Jackson, Ph.D.,
CEO,
VYRIPHARM



Kimberly Lincoln,
Manufacturing Scientist,
ROCHE



Brian Fisher,
VP of Operations and Corporate Partnerships,
THE STURGE-WEBER FOUNDATION



Jeffrey Chen,
Director,
UCLA CANNABIS RESEARCH INITIATIVE

Featured Topics

Governmental and Regulatory



- ▶ Regulatory hurdles and anticipated changes
- ▶ Patient needs in medicinal cannabis
- ▶ Exploring the collaborative environment in associations or unions
- ▶ Establishing and adopting industry quality standards

Patient Advocacy



- ▶ Driving the research through patient-led requests (PTSD, stroke, neuroprotection)
- ▶ Understanding patient needs in cannabidiol neuroprotection and recovery

Academic/Research



- ▶ Lab-based case for healthcare driven cannabis efficacy
- ▶ Science and economics in quality control phylos science
- ▶ Understanding the role of cannabis in a cross-section of pathologies
- ▶ Cannabis quality controls for consistent and effective ingredients

Medical



- ▶ Medical relevance and proven areas of efficacy
- ▶ Navigating barriers to effective research
- ▶ Cannabis efficacy in Immune Therapy

Analytical



- ▶ Quality Control to decrease time to compound approval
- ▶ Cannabis testing for regulatory/compliance standardization

Dear Colleague,

The Cannabis Partnership Congress brings together all of the key stakeholders in a collaborative environment to face the scientific, medicinal, and governmental challenges of cannabis-based product development. Despite recent changes in state policy and the increasing rate of cannabis use and its implications for health, the federal government has yet to legalize cannabis. When facing the continued restrictions on research into the health benefits of cannabis, what strategies must pharma companies follow to bring these therapeutics to market?

Clinical research with cannabis needs special approval from FDA, while obtaining the plants themselves involves a separate time-consuming process through NIDA. The Cannabis Partnership Congress helps you navigate all of these regulatory and development challenges.

Join us this March in San Francisco and learn how to:

- ▶ Face the regulatory restrictions on cannabis R&D
 - ▶ Develop industrial standards for cannabis
 - ▶ Ensure compliance in cannabis marketing
 - ▶ Weave cannabis into your rare disease pipeline
 - ▶ Analyze the future of cannabis lab research
-and much more! We look forward to seeing you there!

WHO SHOULD ATTEND

This conference is designed for representatives from pharmaceutical, medical device, and biotechnology companies with responsibilities in the following areas:

- ▶ Cannabis
- ▶ Cannabinoid Science
- ▶ Strategic Partnerships
- ▶ Program and Pharmaceutical Portfolio Development
- ▶ Translational Medicine
- ▶ Medical Science Liaisons
- ▶ Asset Management
- ▶ Preclinical Staging Development
- ▶ Source Management
- ▶ Research and Development
- ▶ Regulatory Affairs
- ▶ Patient Advocacy and Networks
- ▶ Legislative Health Aids
- ▶ Strategic Partnerships
- ▶ Cannabis Clinical Trial Management/ Operations
- ▶ Clinical Research
- ▶ Medical/Clinical Affairs
- ▶ Personalized Medicine
- ▶ Patient Advocacy/Access
- ▶ Commercial Operations
- ▶ Lab Management
- ▶ Toxicology

This conference is also of interest to:

- ▶ Clinical Labs
- ▶ Data Analysis Information Technology
- ▶ Clinical Research Organizations
- ▶ Regulatory Consultants
- ▶ Marketing Firms
- ▶ Contract Manufacturing



8:00	Registration and Continental Breakfast	12:30	Luncheon
8:45	<p>Conference Co-Chairpersons' Opening Remarks</p> <p>Dean Petkanas, <i>Chief Medical Officer</i>, KANNALIFE SCIENCES</p> <p>Bruce Bloom, <i>CEO</i>, CURES WITHIN REACH</p>	1:30	<p>Breakout Groups</p> <ul style="list-style-type: none"> > Group 1: Roundtable option on how to attain rapid legalization > Group 2: What activities legislatively support research and increased research locations > Group 3: Consortium topics and successful partnerships to obtain NIH and/or DoD funding <p>Veronica Arroyave, <i>Sr. Director, Global Health Programs</i>, PARTNERSHIP FOR QUALITY MEDICAL DONATIONS</p>
9:00	<p>Gain Insight into a Growing Market Within Existing Regulatory Frameworks</p> <ul style="list-style-type: none"> > Understand the current state of regulation > Recognize the limitations of clinical development > The future of cannabinoid therapeutics > Gain a profound understanding of the cannabis market growth <p>Dean Petkanas, <i>Chief Medical Officer</i>, KANNALIFE SCIENCES</p>	2:30	<p>Ensure Compliance in Cannabis Marketing</p> <ul style="list-style-type: none"> > Cannabis producers and retailers have incorrectly understood the limits of FDA's authority as it applies to the promotion of cannabis products > Ensure that no direct disease claims are made > Overview of FDA cracking down on these claims and issuing warning letters > FDA oversight as it relates to promotional marijuana claims > Impact of FDA regulation and the types of discussions that can be appropriately conducted <p>Darshan Kulkarni, <i>Pharm.D., M.S., Esq., Advisor</i>, AMERICAN SOCIETY OF CANNABIS PHARMACISTS</p>
9:45	<p>Monitor Governmental Awareness and Legislative Research Options</p> <ul style="list-style-type: none"> > Updates in current legislative standards > Proposed legislation, legislative branch overview > Attorney General position and standards enforcement stance <p>Brian Fisher, <i>VP of Operations and Corporate Partnerships</i>, THE STURGE-WEBER FOUNDATION</p>	3:15	Networking Break
10:30	Networking Break	4:00	<p>Accelerate Cannabis Compound Approval for Rare Diseases</p> <ul style="list-style-type: none"> > Steps toward licensing > Partnering with FDA, patient groups and institutions Increasing compliance, faster trial enrollment, minimize patient fallout > Rare Disease Indication which can lead to shorter timeline > Legislative support through multiple patient advocacy groups
11:00	<p>Understand Cannabis Chemical Composition in Order to Develop Industrial Standards</p> <ul style="list-style-type: none"> > Lab information management for cannabis quality control > Understand chemical components and develop standards > Critical tools for lab management with cannabis > Quality control from growth to oil and review synthetic options <p>Kimberly Lincoln, <i>Manufacturing Scientist</i>, ROCHE</p>	4:45	<p>Interactive Partnerships Discussion: Maximize Strategic Partnerships in Order to Ensure Patient Safety and Launch a Successful Product</p> <ul style="list-style-type: none"> > Ensure high-priority patient safety > Review of regulatory landscape > Setting the stage for technology advancements > Impact of all stakeholders to continue cannabis support <p>Dr. Bruno Battistini, <i>President, CEO and Scientific Director</i>, NEW BRUNSWICK HEALTH RESEARCH FOUNDATION</p>
11:45	<p>Review the Barriers and Routes to Success for Efficacy Research</p> <ul style="list-style-type: none"> > PTSD interim research results > Neuro protection and the military > Geriatric population quality of life improvement research > Barrier Busters > FDA, NIH and patentable compounds <p>Thoma Kikis, <i>Chief Marketing Officer</i>, KANNALIFE</p>	5:30	Day One Concludes

- 8:00 Registration and Continental Breakfast
- 8:45 Conference Co-Chairpersons' Recap of Day One
Dean Petkanas, Chief Medical Officer, KANNALIFE SCIENCES
Bruce Bloom, CEO, CURES WITHIN REACH
- 9:00 **Understand Why State and Federal Governments Must Work Together to Align Regulations**
 > Cost implications and effects on Healthcare
 > Overview of safety and quality controls in medicinal cannabinoids
 > Understand current legislations
Elias Jackson, Ph.D., CEO, VYRIPHARM BIOPHARMACEUTICAL
- 9:45 **Forecast the Future of Lab Research in Cannabidiol and Improvements in Analytics and Quality**
 > Lab information management for cannabis quality control
 > Critical tools for lab management with cannabis
 > Public vs. private cannabis testing
 > Quality control from growth to oil and review of the synthetic options
- 10:30 Networking Break
- 11:00 **Update QC and Patient Experience Communications for Medical Cannabis**
 > Enhance cannabidiol patient case study in seizures
 > CBD vs. alcohol case study reviewed
 > Medicinal cannabidiol communications and branding
Mark Reichman, EVP, Client Services Director, GUIDEMARK HEALTH, Cannabis Patient Advocate
- 11:45 **Innovative Treatments for Patients Through Drug Repurposing**
 > Opportunities for partnerships in advancing science
 > Alzheimer's, Anxiety and Answers
 > Partnering results: Israel, Canada and the U.S.
Bruce Bloom, CEO, CURES WITHIN REACH

- 12:30 Luncheon
- 1:30 **Panel Discussion: Regulatory Review: Current Legal Questions and Likely Future Developments**
 > Observe the current state of cannabis within the federal government
 > Identify current and future regulations
 > Provide an in-depth look at future cannabis practices
Lezli Engelking, President and Founder, FOCUS
Douglas Brenneman, Scientific Advisory Board, KANNALIFE SCIENCES
Jeremy Unruh, General Counsel and Chief Compliance Officer, PHARMACANN
- PANEL**
- 2:30 **Integrate Medical Cannabis With Traditional Medicine**
 > Understand the cannabis components that can impact a treatment
 > Explore how doses can be impacted
 > Minimize side effects
Jerry Bryant, Managing Member, VYRIPHARM BIOPHARMACEUTICAL
- 3:15 Networking Break
- 4:00 **Utilize Crowdsourcing in Response to Federal Research Restrictions**
 > Accelerate our understanding of the health impacts of cannabis
 > Utilize mobile health, electronic medical records, and artificial intelligence for crowdsourcing
 > International cannabis research landscape and how to leverage strategic strengths amongst nations
 > Formulations (whole plant vs. pure compounds)
 > IP/commercialization of cannabis-based therapeutics
 > Preferred delivery routes
Jeffrey Chen, Director, UCLA CANNABIS RESEARCH INITIATIVE
- 4:45 Congress Concludes

Media Partners



VENUE INFORMATION

Hilton San Francisco Airport Bayfront

600 Airport Boulevard | Burlingame, CA 94010

To make reservations, please call 1-800-HILTONS and request the negotiated rate for **ExL's March Meetings**. You may also make reservations online using the following weblink: <http://bit.ly/2ysQD5h>. The group rate is available until **February 20, 2018**. Please book your room early, as rooms available at this rate are limited.

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Registration Fees for Attending ExL's Cannabis Partnerships Congress

EARLY BIRD PRICING—Register by January 26, 2018

Life Science	\$1,595
Service Providers	\$1,895
Academic / Non-Profit	\$1,195

STANDARD PRICING

Life Science	\$1,795
Service Providers	\$2,095
Academic / Non-Profit	\$1,395

ONSITE PRICING

Life Science	\$1,995
Service Providers	\$2,295
Academic / Non-Profit	\$1,595

Group Discount Program

Save 25% per person when registering four

For every three simultaneous registrations from your company, you will receive a fourth complimentary registration to the program (must register four at one time). This is a savings of 25% per person.

Save 15% per person when registering three

Can only send three? You can still save 15% off every registration.

Offers may not be combined. Early Bird rates do not apply. To find out more about how you can take advantage of these group discounts, contact our offices at (201) 871-0474.

March 12-13, 2018

Hilton San Francisco Airport Bayfront | San Francisco, CA

Cannabis Congress



YES! Register me for this conference!

Name: _____ Title: _____

Company: _____ Dept.: _____

Address: _____

City: _____ State: ____ Zip: _____

Email: _____

Phone: _____ Fax: _____

Method of Payment: Check Credit Card

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MasterCard Visa Discover AMEX

Card Number: _____

Exp. Date: _____ CVV: _____

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CONFERENCE CODE: C1033

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- Four weeks or more: A full refund (minus a \$295 processing fee) or a voucher to another ExL event valid for 12 months from the voucher issue date.
- Less than four weeks: A voucher to another ExL event valid for 12 months from the voucher issue date.
- Five days or less: A voucher (minus a \$395 processing and documentation fee) to another ExL event valid for 12 months from the voucher issue date.

To receive a refund or voucher, please contact our offices at (201) 871-0474.

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ExL Events' liability is limited to the conference registration fee in the event of a cancellation and does not include changes in program date, content, speakers and/or venue.

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