



CONFERENCE ON RARE DISEASE AND ORPHAN DRUGS

1st and 2nd October





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EVENT OVERVIEW

The orphan drug market is continuously growing over the years now at twice the growth rate of overall prescription drug market due to the arising interest from pharmaceutical companies for underlying benefits and involvement of many smaller biotechnology start-up companies who are primarily focused on rare diseases. However still there are lot of attention that needs to be given since the ratio of one having rare disease is 1 in 2000 leading to the lack of knowledge about disease mechanisms, along with FDA/Marketing approvals which take years.

The goal of this Conference is uniting all well-known clinical researchers, biotech professionals, Medical Advisories, Academicians and decision makers to discuss possible ways to accelerate orphan drug development and access to rare disease patients, including the introduction of recent technologies and products to help aid the access of orphan drugs.

In the pioneer edition of DBMR Conference on Rare Disease and Orphan Drugs come join us in this enriching knowledge fest where experts will speak on new technologies and market dynamics perceptions. Together let's move ahead in the future of medical science and technology.



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KEY HIGHLIGHTS

- Orphan drug: Development trends and strategies
- Challenges in Access to and Affordability of Medicines for Rare Diseases
- How to Commercialize an Orphan Drug
- Non-profit drug discovery enabling platforms vs advocacy organizations - challenges in pharma perception of foundations
- Defining of 'Orphan Drugs': Beyond National Boundaries
- The "One and Done" Era for Orphan Drugs
- Regulatory Framework for Drugs for Rare Diseases

WHO SHOULD ATTEND?

- Medical Affairs ○
- Medical Advisory ○
- Clinical Operations ○
- Rare Disease Research ○
- Patient Advocacy ○
- Commercial Officer ○
- Orphan Drug Therapeutics ○
- Regulatory Affairs ○
- Rare Disease Charity Officers ○



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KEY HIGHLIGHTS

- Gain Insights on new developments and strategies aimed to improve patient access to orphan drugs
- Know how to overcome challenges being faced by Biotech Companies.
- More than 250+ Listeners
- Exhibit your organizations expertise
- Panel Discussions among Industry Experts and Academics and Scholars
- TMeet the upcoming minds of Rare Disease and Orphan Drugs
- One on One Networking Opportunities



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AGENDA (DAY ONE)

8:30 AM EST	Registration (Telephonic And Virtual)
9:30 AM EST	Introduction To DBMR
10:00 AM EST	Addressing The Greatest Unmet Need Across All Rare Diseases: Information
11:15 AM EST	Orphan drug: Development trends and strategies
12:30 PM EST	Advocacy And The Need To Hear The Rare Disease Patients Voice Within Strategic Efforts
1:45 PM EST	A Case Study (Epidiolex) Demonstrating How To Use Special FDA Designations To Efficiently Reach The US Market
3:00 PM EST	The Future Of Rare Diseases: Developing And Accessing The Next Generation Of Therapies
4:15 PM EST	Patient involvement in drug development for rare diseases
5:15 PM EST	How to Commercialize an Orphan Drug
6:15 PM EST	Rare vs. Common Diseases
7:00 PM EST	DBMR Quantum Leap Awards



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AGENDA (DAY TWO)

8:30 AM EST	Registration
9:30 AM EST	Challenges being faced by Orphan drug development Companies
10:00 AM EST	Non-profit drug discovery enabling platforms vs advocacy organizations - challenges in pharma perception of foundations
11:15 AM EST	Defining of 'Orphan Drugs': Beyond National Boundaries
12:30 PM EST	Treatment and Advanced Therapies for Rare Diseases
1:45 PM EST	Social Entrepreneurship In Rare Diseases: It Takes Concerted Efforts Between Technology And Advocacy To Accelerate Therapies For Rare Diseases
3:00 PM EST	The "One and Done" Era for Orphan Drugs
4:15 PM EST	Examining industry strategies for leveraging patient data and AI to advance rare disease patient identification, R&D, and commercialization
5:15 PM EST	Regulatory Framework for Drugs for Rare Diseases
6:15 PM EST	Rare Mental and Behavioural Disorders
7:00 PM EST	DBMR Closing Remarks



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OUR SPEAKERS



JONATHAN MORTON

RARE DISEASE CONSULTANT
AMICULUM

AGENCY LEAD
COMRADIS LIMITED



RICH HORGAN

FOUNDER AND PRESIDENT
CURE RARE DISEASE



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OUR SPEAKERS



ALICE RATHJEN

CO-FOUNDER AND CEO
DNA COMPASS



HARSHA K RAJASIMHA

FOUNDER AND CEO
JEEVA INFORMATICS SOLUTIONS

FOUNDER AND CHAIRMAN
INDOUSRARE

MEMBER OF THE BOARD OF ADVISORS
SYNERGY



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OUR SPEAKERS



KELLY AUSTIN

FOUNDER, REGULATORY SPECIALIST
(ODD, FAST TRACK, BREAKTHROUGH, RMAT)
KLA BREAKTHROUGH CONSULTING



LYDIA L. SEIDERS

PATIENT
EDUCATION COUNCIL MEMBER
APLASTIC ANEMIA AND
MDS INTERNATIONAL FOUNDATION
ADVISOR
LIV4THECURE
ADVISOR
RARE DISEASE GHANA INITIATIVE



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OUR SPEAKERS



GAVIN JONES

DIRECTOR OF RARE DISEASE
OPEN HEALTH



HECTOR DE LA RIVA

COMMERCIAL LEAD OCULAR
RARE DISEASE – GENE THERAPY
NOVARTIS



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OUR SPEAKERS



DIANA STEFANI-HUNYADY

HEAD OF GLOBAL MEDICAL AFFAIRS
VIELA BIO

CO-FOUNDER
LIFE KEY SCIENCES LLC



CHRISTINE VON RAESFELD

FOUNDER AND CEO
PEOPLE WITH EMPATHY

TECHNICAL EXPERT PANELIST
CENTERS FOR MEDICARE &
MEDICAID SERVICES



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OUR SPEAKERS



SHAHAR KEINAN

CEO AND CO-FOUNDER
POLARISqb



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OUR CLIENTS





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BLOG: <https://www.databridgemarketresearch.com/digital-conference/conference-on-rare-disease-and-orphan-drugs/group-delegate-registration>