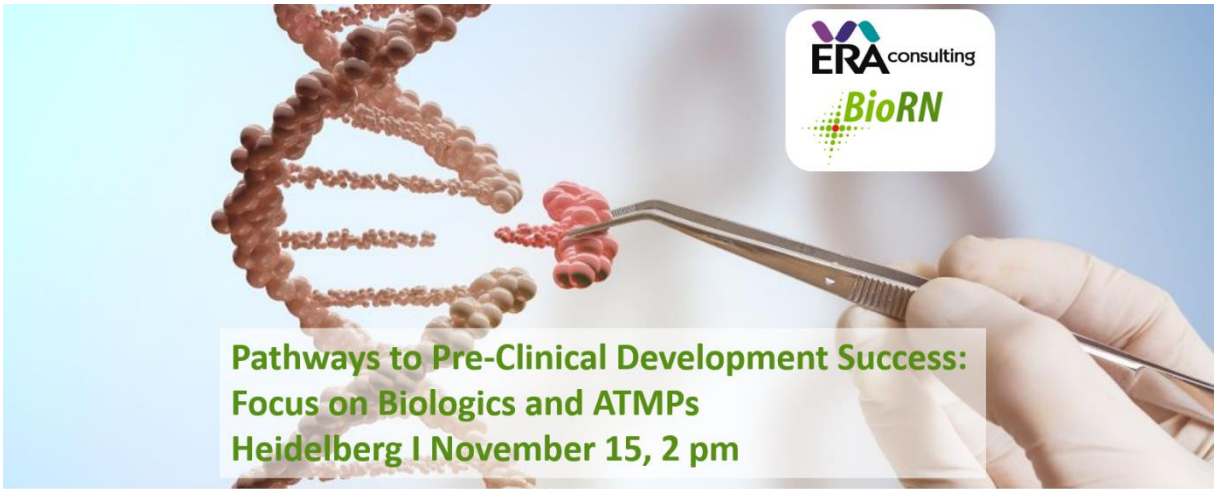

Pathways to Pre-Clinical Development Success: Focus on Biologics and ATMPs >>

When: 15 November 2018, at 2 pm
Where: Technology Park – Conference Center, Heidelberg

This event will introduce technical and regulatory strategies for pre-clinical development, with a focus on biologicals and advanced therapy medicinal products (ATMPs). The sessions are designed to provide attendees with a regulatory scientific approach for successful translation of biologicals and ATMPs into the clinic. We will aim to provide an understanding of pre-clinical development pathways and requirements for biologicals and ATMPs leading to the most expedited regulatory strategy in Europe, the US and beyond. Potential challenges in the development and registration of these products will also be discussed, to provide practical advice as well as highlight key aspects determining successful regulatory submissions.

Who should attend?

- R&D scientists
- Biotech entrepreneurs
- Industry related professionals
- Students and Post-doctoral scientists
- Investors

Agenda (Speakers Profile)

13:45 – 14:00	Registration
14:00 – 14:05	Introduction by BioRN
14:05 – 14:35	**A Regulatory Strategy Adds Value in Pre-clinical Development - Dr Veronika Alt**
14:35 – 15:05	**Quality Counts in Pre-clinical Development for Cell and Gene Therapies – Dr Dianne Jackson-Matthews**
15:05 – 15:30	Afternoon Tea
15:30 – 16:30	**Critical factors in Successful Pre-clinical Testing - Dr Lesley Earl**
16:30 – 17:30	Reception/Networking