

Small Activating RNAs — a Novel Therapeutic Class of Oligonucleotides

Smi reports: Understanding the principles of RNA activation and possibilities of therapeutic benefit at Oligonucleotide Therapeutics and Delivery Conference

LONDON, UNITED KINGDOM, July 6, 2021 /EINPresswire.com/ -- Engage in the latest innovations of oligonucleotide therapeutics with insights into

[immunochemotherapeutics](#), ocular indications and Duchenne Muscular

Dystrophy. Deepen your understanding of crucial delivery methods and available platforms for non-hepatocytic delivery.

Download a copy of your brochure to find out who will be speaking at the upcoming conference in September www.oligonucleotide.co.uk/pr5ein

EMERGING DELIVERY SOLUTIONS

Targeted Delivery of C/EBPa-saRNA by RNA Aptamers

- Understanding the principles of RNA activation and possibilities of therapeutic benefit
- Harnessing small activating RNAs for the treatment of Pancreatic ductal adenocarcinoma
- Evaluating aptamers and appropriate means of targeted delivery
- Looking at lessons learnt from using saRNA for oncological purposes

Nagy Habib, Professor of Surgery, Imperial College, Co-founder MiNA therapeutics, Imperial College London

Small activating RNAs — a novel therapeutic class of [oligonucleotides](#)

- Description of saRNA technology to upregulate transcription and broad applicability to a range of therapeutic targets
- Generation of a lead candidate saRNA to HNF4a for liver disease



- Update on MiNA's lead saRNA clinical agent MTL-CEBPA – Clinical PD and efficacy
Matthew Catley, Research Director, MiNa Therapeutics Ltd

The tool-box approach to improve the performance of siRNA-platform technology

- Interplay of design and function
- Stability and duration of action
- Linker design and valency of ligands
- Is there room for additional improvements?

Adrien Weingartner, Principal Scientist, Group Leader Drug Delivery, Silence Therapeutics

Clinical development of AsiDNA, a first in class decoy agonist oligonucleotide targeting DNA damage response in tumor cells

- Introduction to platON™: proprietary chemistry platform based on a library of decoy agonist oligonucleotides which generates disruptive compounds acting on intracellular DNA-binding targets
- Introduction to AsiDNA the leading decoy agonist generated from platON and targeting DNA damage response function
- Recent preclinical highlights: AsiDNA abrogates resistance to multiple anti-cancer targeted therapies
- Recent clinical development of AsiDNA
- Outlining next steps of clinical development

Wael Jdey, Head of Biology, Onxeo S.A.

Register online: www.oligonucleotide.co.uk/pr5ein

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