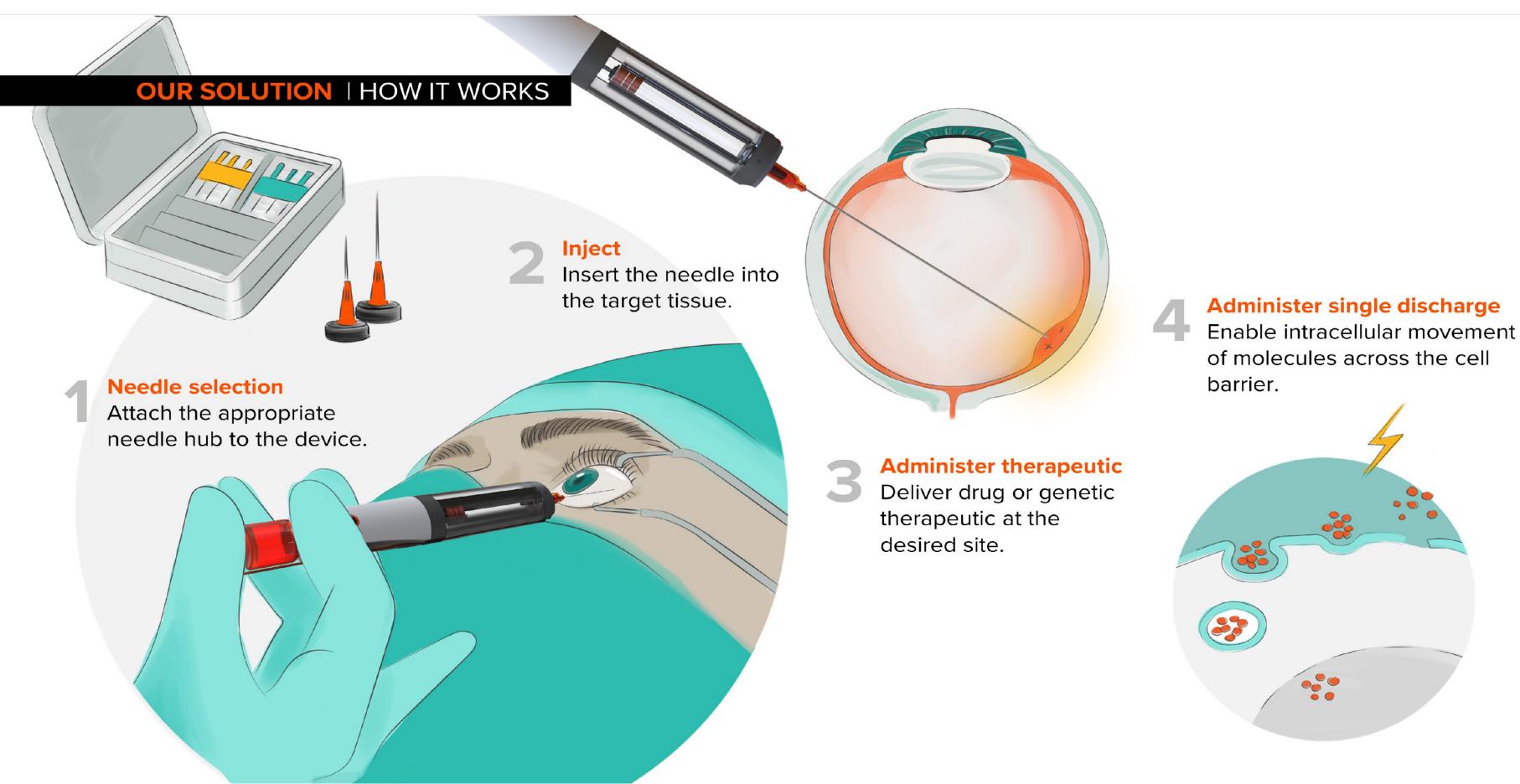
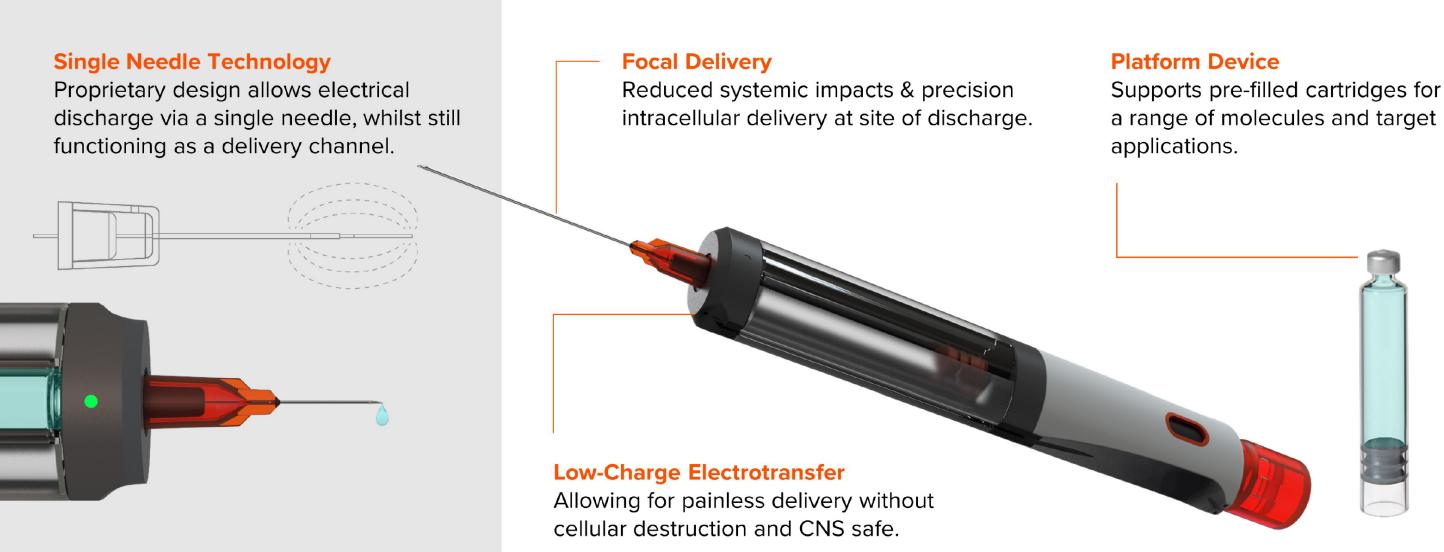
Precision DNA / RNA Electrotherapeutics Platform

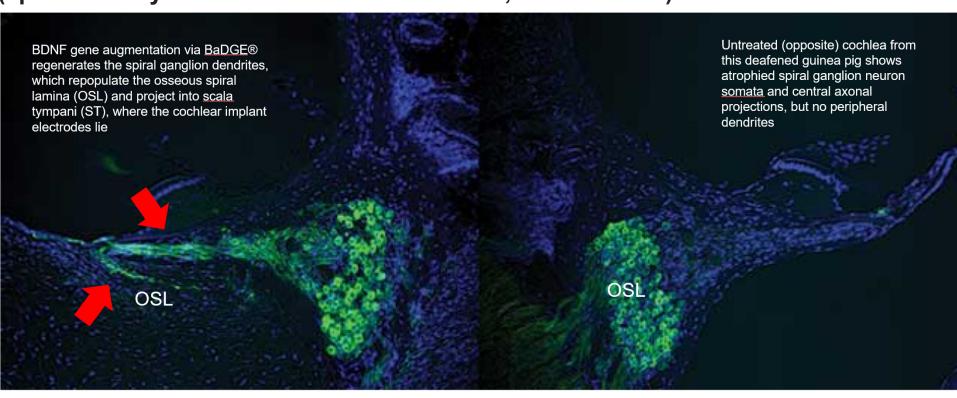






VALIDATION | SCIENTIFIC FEASIBILITY

Hearing-BaDGE® - cochlear nerve regeneration via proprietary neurotrophin - encoding DNA therapeutic (GMP) (sponsored by Passe & Williams Foundation; Cochlear Ltd)



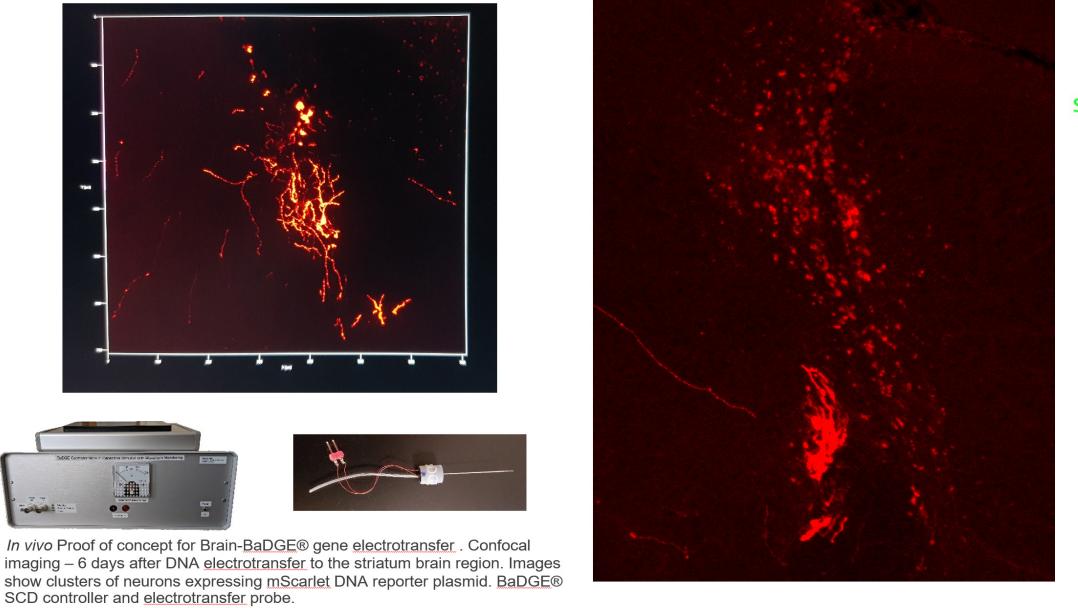
NERVE REGENERATION: Experimentally deafened mammals suffer loss of sensory hair cells followed by atrophy of lear nerve. But a new therapy being tested in guinea pigs may spur the regeneration of the nerve (left, green projections), by placing an implant in the cochlea that can mediate the uptake of a gene therapy construct encoding brain-derived neurotrophic factor (BDNF)

First-in-human clinical trial www.cingt.info; ACTRN12618001556235

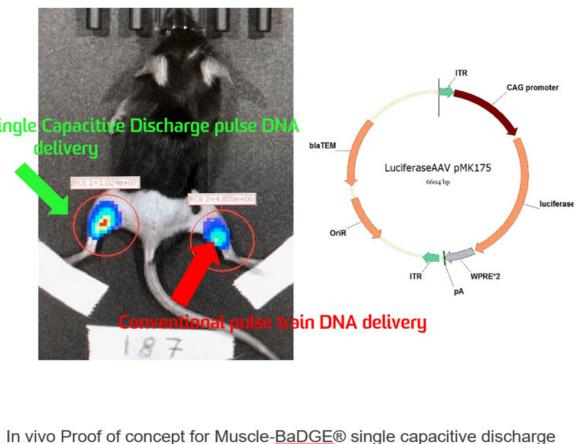
- **Safety:** naked plasmid DNA is not biologically active outside of cells; very limited expression in tissues in the absence of electrotransfer; lowered immunogenicity (no immune responses to viral vector elements). • Target Specificity: BaDGE® provides optimal control of which tissue
- regions achieve take-up of the plasmid DNA into cells, so expression is limited to the region of interest.
- **Versatility:** plasmid DNA does not have the packaging (size) limits inherent in viral vector-based gene therapy; large (complex) gene expression cassettes, including DNA expression regulatory switches, can be used.
- Speed, Efficiency, Duration: Onset of expression within hours, duration of many months
- Lower Regulatory Compliance: transfer of naked DNA to human somatic cells is not subject to regulatory oversight by the office of the Gene Technology Regulator (Australia).

Brain-BaDGE® - neurosuppression to control epilepsy & Parkinsons' Disease (sponsored by NHMRC & Boehringer-Ingelheim)

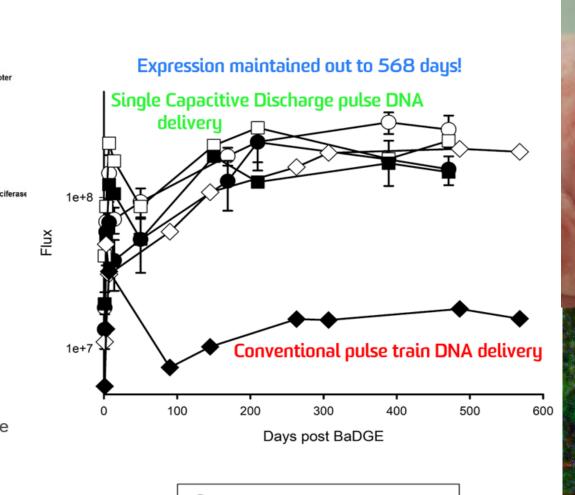




Muscle-Badge - vaccines, immunotherapy & muscle reinnervation (sponsored by Touchlight Genetics UK)

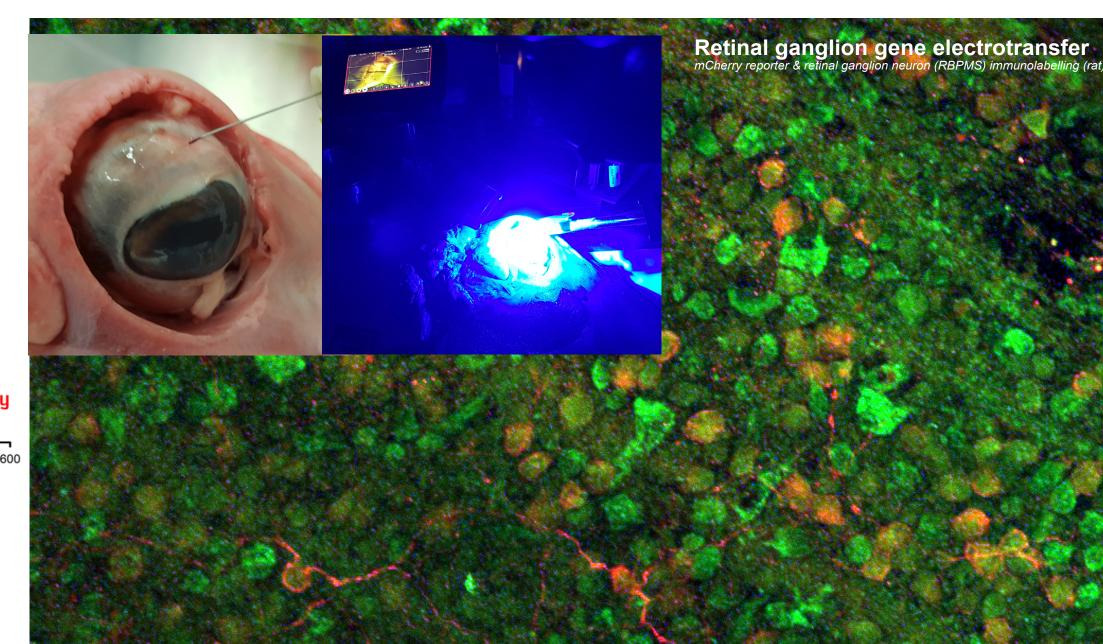


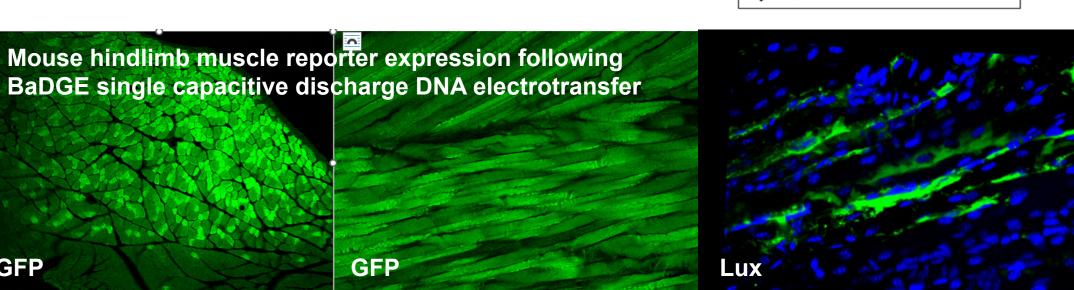
*(SCD) gene electrotransfer . Spectrum CT bioluminescence imaging – mouse (ref. ms187, 250519) at 24 hours post DNA electrotransfer of a luciferase encoding plasmid DNA (LuciferaseAAV pMK175.CAGpluciferase.WPRE), under isoflurane anaesthesia within 30 min of i.p. injection of luciferin substrate to elicit photo emission arising from expression of recombinant luciferase by the hindlimb muscles. SCD left hindlimb, conventional monophasic pulse train (right hindlimb). DNA (1

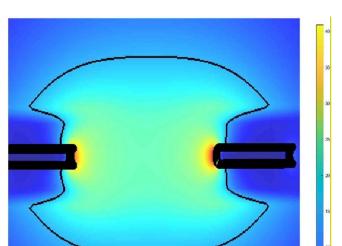


-Col 36 vs 120V - Col 41 vs 250V - Col 46 vs 1 ug/ul Col 48 vs 40ul day vs 187 L = 2.2uF at 120V 1ug/ul 40 ul day vs 187 R = 4 x 5 ms

Vision-Baddel Beautiform for non-viral-based eye therapeutics

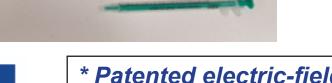






SCD controller and electrotransfer probe





- * Patented electric-field focusing for 'dial-up' control of instantaneous focal DNA/RNA electrotransfer
- * Patented single-pulse delivery
- * > \$5M undiluted research funding & Pharma engagement Peer-reviewed publications
- National and international media coverage
- * Clinical pathway validated





