



 **iPSC**

Comprehensive Solution

Provided by uBriGene's iPSC Production Platform



Your bridge from ATMPs concept to commercialization



Autonomous IP of
GC-DNA Production Process



Grade A Production
Environment
Single-use Enclosed
Production



Full Process Compliant with
GMP Standards

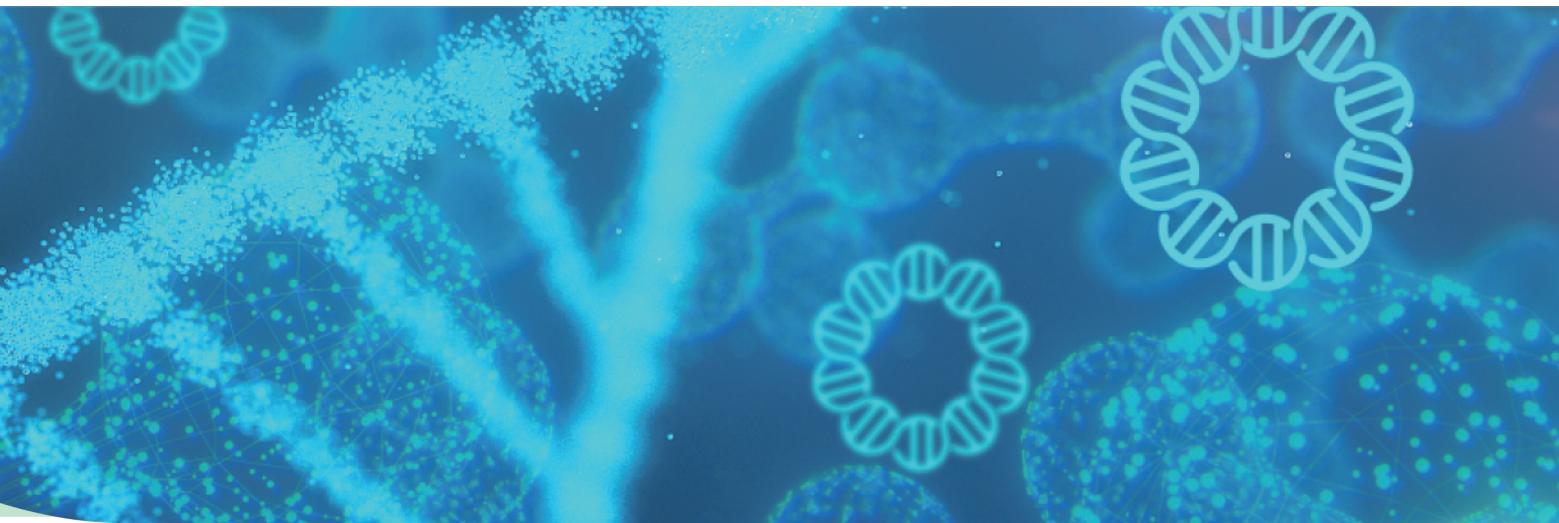
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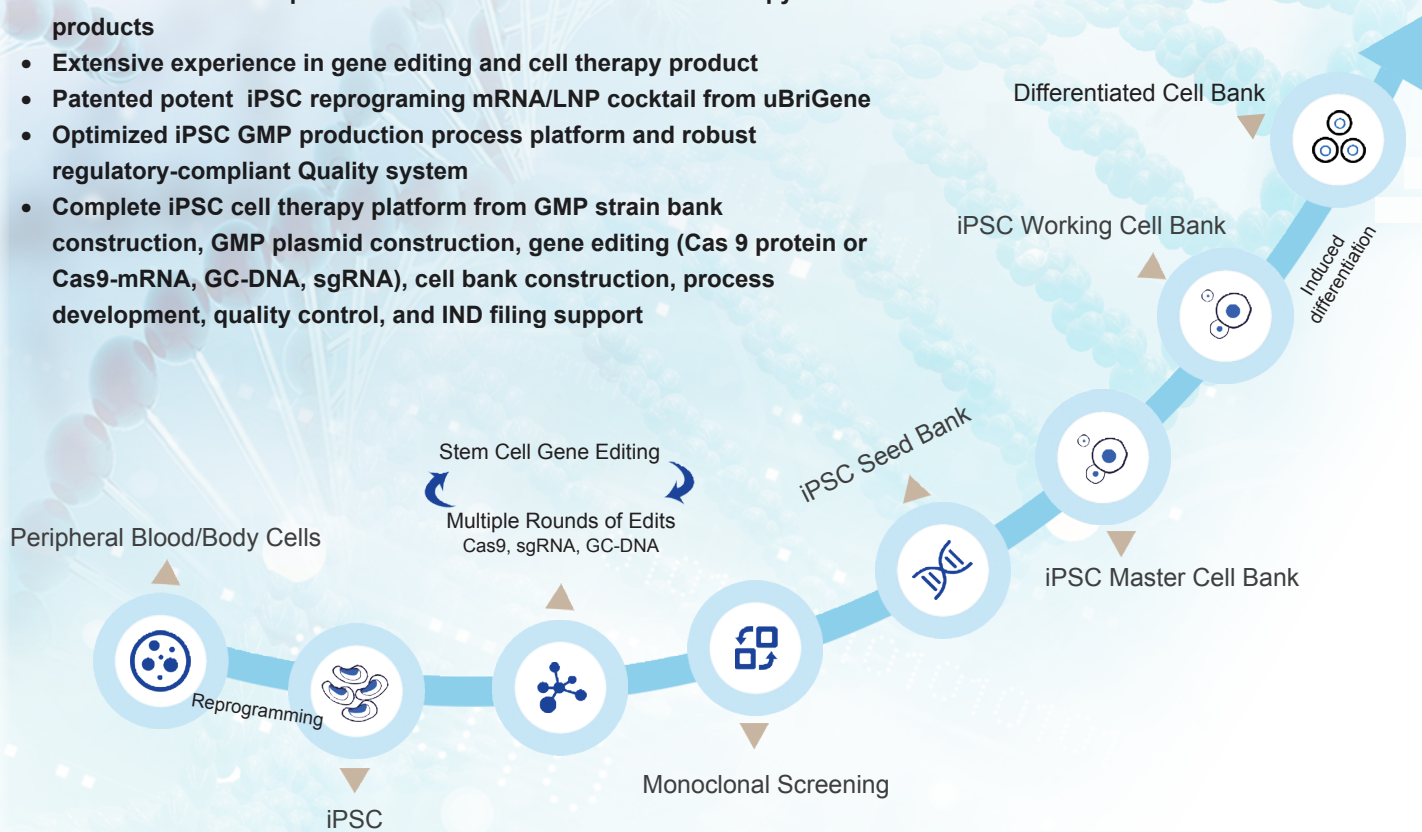
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uBriGene's Robust iPSC Platform

- uBriGene offers comprehensive CDMO services for iPSC therapy products
- Extensive experience in gene editing and cell therapy product
- Patented potent iPSC reprogramming mRNA/LNP cocktail from uBriGene
- Optimized iPSC GMP production process platform and robust regulatory-compliant Quality system
- Complete iPSC cell therapy platform from GMP strain bank construction, GMP plasmid construction, gene editing (Cas 9 protein or Cas9-mRNA, GC-DNA, sgRNA), cell bank construction, process development, quality control, and IND filing support



IIT

Non-registered research grade plasmid, gene editing components, development and production of iPSC cell bank



IND

iPSC cell therapy new drug clinical filing plasmid, gene editing components and cell bank development and production



Clinial Trial Grade

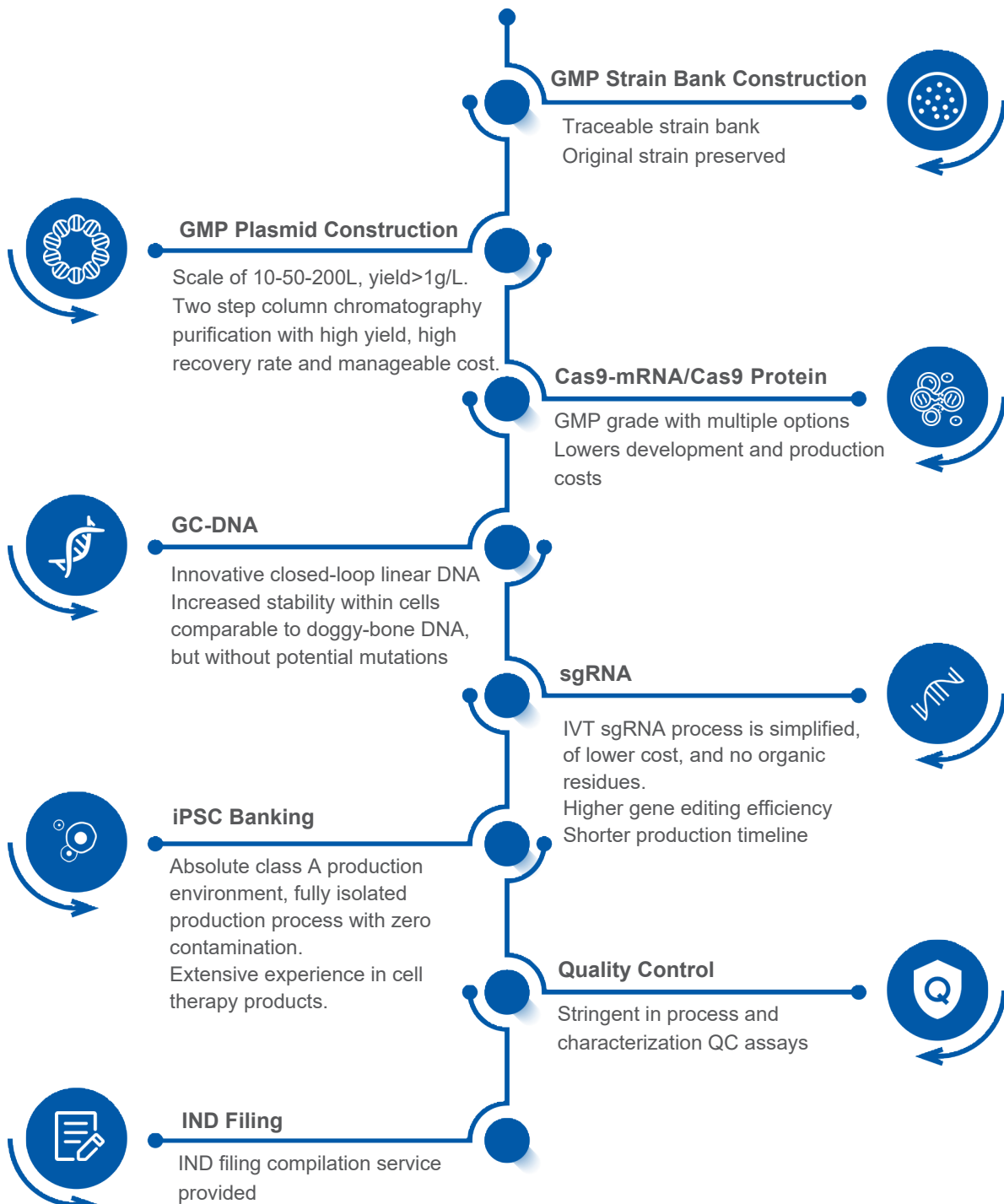
iPSC cell therapy new drug plasmid, gene editing components and clinical stage I/II/III cell bank production



Commercialization

iPSC cell therapy new drug commercialized GMP production

iPSC Cell Therapy Comprehensive Solution

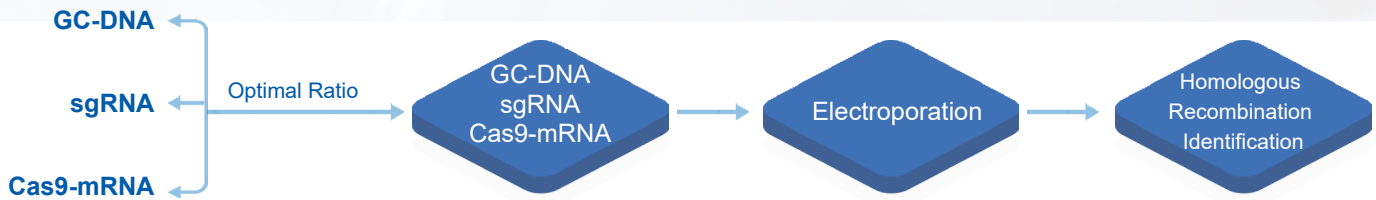


Gene Editing Process Flow

Using GC-DNA, sgRNA, Cas9 as basic components



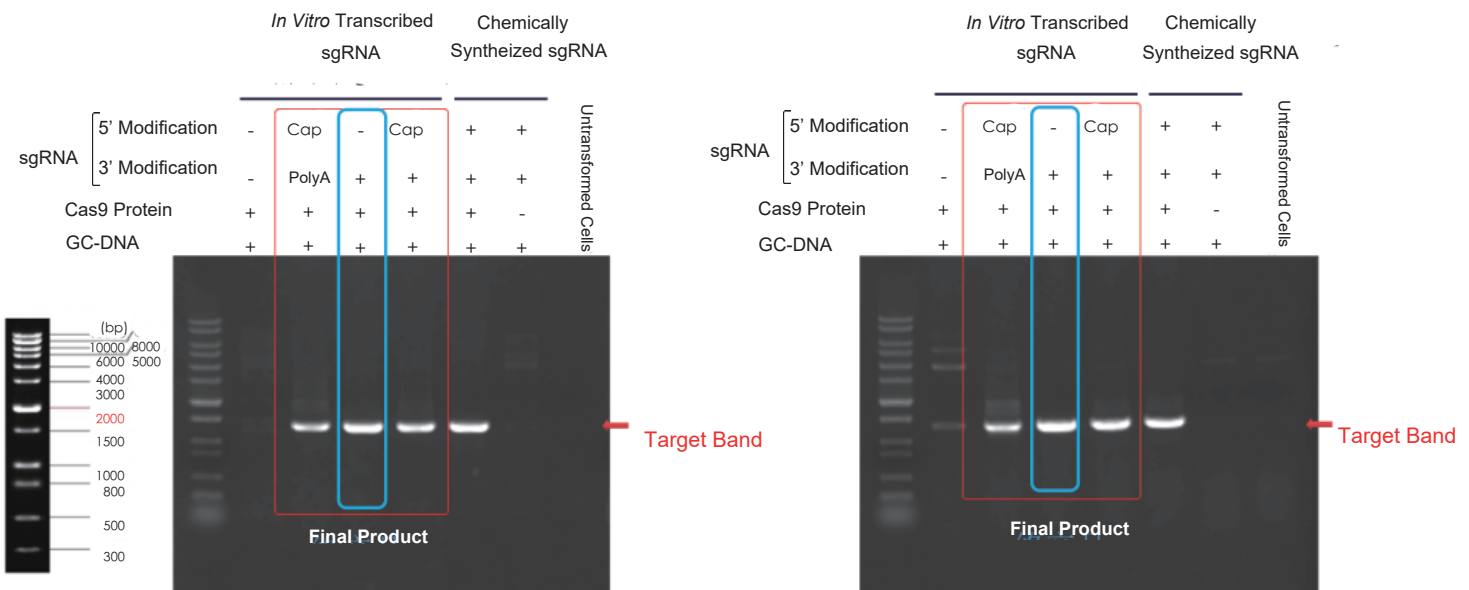
Using GC-DNA, sgRNA, Cas9-mRNA as basic components



Higher Gene Editing Efficiency with sgRNA-IVT

PCR Identification of Left Homologous Arm: Target fragment size after recombination is 1304 bp

PCR Identification of Right Homologous Arm: Target fragment size after recombination is 1321 bp



Results

1. Modified sgRNA has better stability and higher efficiency in guiding homologous recombination compared to unmodified sgRNA which is prone to degradation.
2. Our 3' sgRNA modification is a built-in process in the template, no additional modification step.
3. sgRNA prepared by *in vitro* transcription has high production yield, low cost and no organic residue compared to chemically synthesized sgRNA.