

Center for Gene & Cell Therapy

遺伝子・細胞治療センター

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IMSUT hospital has been playing a crucial role in clinical gene therapy and stem cell transplantation in Japan. In order to reinforce this clinical development even further, IMSUT established the Center for Gene & Cell Therapy (CGCT) in 2014. CGCT particularly focuses on the development of gene therapy and cell therapy for intractable cancer and chronic diseases such as oncolytic virus therapy, engineered T cell therapy, gene therapy for neurological disorders using AAV vectors, T cell therapy for post-transplant viral infections, and cell therapy using mesenchymal stem cells.

1. Gene therapy for amyotrophic lateral sclerosis and spinocerebellar ataxia type 6

Shin-ichi Muramatsu

In sporadic amyotrophic lateral sclerosis (ALS) patients, down regulation of the RNA-editing enzyme, adenosine deaminase acting on RNA 2 (ADAR2), is death-causing molecular abnormality that occurs in motor neurons. Gene delivery of the ADAR2 using adeno-associated virus (AAV) vectors in conditional ADAR2 knockout mice effec-

tively prevented progressive motor dysfunction without any adverse effects. We have started to produce GMP grade AAV vectors that express ADAR2 for a clinical trial. In collaboration with Chicago university, we have developed miR-based gene therapy for spinocerebellar ataxia type 6 (SCA 6). SCA6 is caused by abnormal expansions of the polyglutamine tract within a second CACNA1A gene product, $\alpha 1$ ACT. Selective translational block of SCA6-associated $\alpha 1$ ACT by delivering miR-3191-5p protected from the Purkinje cell degeneration and ataxia in a mouse model.

Publications

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