## A study on the leading optimization for cancer immunotherapy of regulatory T cell suppression/depletion mechanism using CD25-ApDC (CD25 aptamer-drug-conjugate)

## **APTAMER SCIENCES, Inc.**



| ONCOLOGY                 | Lead  |
|--------------------------|---|
| Product Type             | Aptamer-Drug conjugate (ApDC)   |
| Indication               | Solid tumor   |
| Target                   | CD25 (Regulatory T cell)  |
| MoA(Mechanism of Action) | <ul> <li>CD25-targeted aptamer blocks IL-2 signaling in regulatory T cells (Tregs) while it is not blocking the IL-2/IL-2R interaction in effector T cells (Teffs).</li> <li>CD25-targeted aptamer-drug conjugate (CD25-ApDC) leads to increase Teffs and APC activity in the tumor by selectively depleting Tregs.</li> <li>Selective down-regulation and depletion of Treg which is a key immunosuppressive agent can improve overall immune response in the tumor.</li> </ul>  |
|                          | 1 Selective Treg depletion  |
| Competitiveness          | <ul> <li>CD25-targeted aptamer blocks IL-2 signaling in regulatory T cells (Tregs) while it is not blocking the IL-2/IL-2R interaction in effector T cells (Teffs)</li> <li>CD25-ApDC is a selective CD25 Treg depleter resulting in the increase of Teff/Treg ratio in the tumor.</li> <li>CD25-specific aptamer inhibits the immunosuppressive activity of Treg by blocking IL-2 binding and downstream signaling, resulting in the decrease of TGF-beta secretion.</li> <li>Relatively shorter half-life, rapid clearance, superior tissue permeability of an aptamer compared to an antibody enable the effective removal of Tregs in the tumor with minimizing systemic target-related adverse effects such as autoimmunity due to long-term depletion of Treg.</li> </ul> |
| Development Stage        | Lead  |
| Route of Administration  | intravenous injection (IV)  |

