Pharmaceutical Business Clinical Development as of April 30, 2020

<In-house development>

| Code (Generic Name) | Potential Indication/Dosage form | Mechanism | | Phase (Region) | Origin | Note |
|--|---|---|---|--|------------|--|
| JTZ-951 (enarodustat) | Anemia associated with chronic kidney disease /Oral | HIF-PH inhibitor | Increases red blood cells by stimulating production of erythropoietin, an erythropoiesis-stimulating hormone, via inhibition of HIF-PH. | NDA filed (Japan) Phase1 (Overseas) | In-house | Co-development with Torii |
| JTE-052 (delgocitinib) | Atopic dermatitis (pediatric) /Topical | JAK inhibitor | Suppresses overactive immune response via inhibition of Janus kinase (JAK) related to immune signal. | Phase 3 (Japan) | - In-house | Co-development with Torii |
| | Autoimmune/allergic diseases /Oral, Topical | | | Phase1 (Japan) | | |
| JTE-051 | Autoimmune/allergic diseases /Oral | Interleukin-2 inducible T cell kinase inhibitor | Suppresses overactive immune response via inhibition of the signal to activate T cells related to immune response. | Phase2 (Overseas) | In-house | |
| JTE-451 | Autoimmune/allergic diseases /Oral | RORy antagonist | Suppresses overactive immune response via inhibition of ROR γ related to Th 17 activation. | Phase2 (Overseas) | In-house | |
| JTT-251 | Type 2 diabetes mellitus /Oral | PDHK inhibitor | Decreases blood glucose by activation of pyruvate dehydrogenase (PDH) related to carbohydrate metabolism. | Phase1 (Overseas) | In-house | |
| JTT-662 | Type 2 diabetes mellitus /Oral | SGLT1 inhibitor | Suppresses postprandial hyperglycemia and normalizes blood glucose level via inhibition of SGLT1. | Phase1 (Overseas) | In-house | |
| JTE-761 | Autoimmune/allergic diseases /Oral | RORy antagonist | Suppresses overactive immune response via inhibition of ROR γ related to Th 17 activation. | Phase1 (Overseas) | In-house | |
| JTT-751 (ferric citrate hydrate) | Iron-deficiency anemia/Oral | Oral iron replacement | Corrects iron-deficiency anemia by using absorbed iron for synthesis of hemoglobin. | Phase3 (Japan) | In-license | Licensed from Keryx Biopharmaceuticals Co-development with Torii Additional indication |

Clinical trial phase presented above is based on the first dose.

<Licensed compounds>

| Compound (JT's code) | Licensee | | Mechanism | Note |
|-------------------------------|------------------------------------|---------------------|---|------|
| trametinib | Novartis | MEK inhibitor | Inhibits cellular growth by specifically inhibiting the activity of MAPK/ERK pathway. | |
| Anti-ICOS monoclonal antibody | AstraZeneca | ICOS antagonist | Suppresses overactive immune response via inhibition of ICOS which regulates activation of T cells. | |
| delgocitinib | LEO Pharma ROHTO Pharmaceutical | JAK inhibitor | Suppresses overactive immune response via inhibition of Janus kinase (JAK) related to immune signal. | |
| enarodustat | JW Pharmaceutical Salubris | HIF-PH inhibitor | Increases red blood cells by stimulating production of erythropoietin, an erythropoiesis-stimulating hormone, via inhibition of HIF-PH. | |

Updates since the previous announcement on February 7, 2020:
• Price Listing on the NHI Reimbursement List and Launch of CORECTIM® Ointment 0.5% in Japan.(April 22, 2020)