

## Pharmaceutical Business

### Clinical Development as of August 2, 2024

<In-house development>

| Code (Generic Name)    | Potential Indication/Dosage form            | Mechanism                                       |  | Phase (Region)    | Origin     | Note   |
|------------------------|---|---|--|-------------------|------------|--|
| JTE-052 (delgocitinib) | Autoimmune/allergic diseases /Oral, Topical | JAK inhibitor                                   | Suppresses overactive immune response via inhibition of Janus kinase (JAK) related to immune signal.               | Phase1 (Japan)    | In-house   |  |
| 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(Japan)     | In-house   |  |
|                        |   |   |  | Phase2 (Overseas) |            |  |
| 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   |  |
| JTT-861                | Chronic heart failure /Oral                 | PDHK inhibitor                                  | Improves cardiac function by activation of pyruvate dehydrogenase (PDH) related to carbohydrate metabolism.        | Phase2 (Overseas) | In-house   |  |
| JTE-061 (tapinarof)    | Atopic dermatitis (pediatric) /Topical      | AhR modulator                                   | Suppresses skin inflammation via activation of the aryl hydrocarbon receptor (AhR)                                 | Phase3 (Japan)    | In-license | <ul style="list-style-type: none"> <li>• In-license from Dermavant Sciences GmbH</li> <li>• Co-development with Torii</li> </ul> |
| JTC-064                | Neurodegenerative disease /Oral             | PDHK inhibitor                                  | Improves metabolic abnormalities by activation of pyruvate dehydrogenase (PDH)                                     | Phase1 (Overseas) | In-house   |  |
| JTV-161                | Pulmonary arterial hypertension /Oral       | Pim-1 inhibitor                                 | Suppresses pulmonary vascular cell proliferation by inhibiting Pim-1   | Phase1 (Overseas) | In-house   |  |
| JTE-162                | Autoinflammatory/ Autoimmune diseases /Oral | NLRP3 inhibitor                                 | Suppresses immune response by inhibition of NLRP3 inflammasome   | Phase1 (Overseas) | In-house   |  |

Clinical trial phase presented above is based on the first dose.  
We are also conducting additional studies to examine the potential for use in additional dosage forms.

<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.   |      |
| delgocitinib         | LEO Pharma<br>ROHTO Pharmaceutical | JAK inhibitor    | Suppresses overactive immune response via inhibition of Janus kinase (JAK) related to immune signal.                                    |      |
| enarodustat          | JW Pharmaceutical<br>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 May 9, 2024

•JTE-061: Manufacturing and Marketing Approval of VTAMA® Cream 1% for the Treatment of Atopic Dermatitis and Plaque Psoriasis in Japan (June 24, 2024)