Japan Tobacco Inc. Clinical Development as of May 10, 2017

<In-house development>

| Code (Generic Name) | Potential Indication/Dosage form | Mechanism | | Phase | Note |
|-----------------------------|---|---|--|-----------------------------------|--|
| JTZ-951 | Anemia associated with chronic kidney disease /Oral | HIF-PHD inhibitor | Increases red blood cells by stimulating production of erythropoietin, an erythropoiesis-stimulating hormone, via inhibition of HIF-PHD. | Phase2(Japan) Phase1(Overseas) | In-house |
| JTE-052 | Autoimmune/allergic diseases /Oral, Topical | JAK inhibitor | Suppresses overactive immune response via inhibition of Janus kinase (JAK) related to immune signal. | Phase2(Japan) | In-house Co-development with Torii |
| JTE-051 | | 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 |
| 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 |
| JTK-351 | HIV infection /Oral | HIV integrase inhibitor | Suppresses blood HIV levels by inhibiting the activity of integrase, an enzyme involved in the replication of HIV. | Phase1(Japan) | In-house |
| JTE-451 | 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) | Iron-deficiency anemia/Oral | Oral iron replacement | Corrects iron-deficiency anemia by using absorbed Iron for synthesis of hemoglobin. | Phase2(Japan) | In-license (Keryx Biopharmaceuticals) Co-development with Torii *additional indication |

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

<Licensed compounds>

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|-------------------------------|-------------------|-------------------|--|---|--|--|--|
| Compound (JT's code) | Licensee | Mechanism | | Note | | | |
| trametinib | Novartis | MEK inhibitor | Inhibits cellular growth by specifically inhibiting the activity of MAPK/ERK Kinase (MEK1/2). | NSCLC, trametinib+dabrafenib U.S., Japan marketing approvals submitted | | | |
| Anti-ICOS monoclonal antibody | MedImmune | ICOS antagonist | Suppresses overactive immune response via inhibition of ICOS which regulates activation of T cells. | | | | |
| JTE-052 | LEO Pharma | JAK inhibitor | Suppresses overactive immune response via inhibition of Janus kinase (JAK) related to immune signal. | | | | |
| JTZ-951 | JW Pharmaceutical | HIF-PHD inhibitor | Increases red blood cells by stimulating production of erythropoietin, an erythropoiesis-stimulating hormone, via inhibition of HIF-PHD. | | | | |

Updates since the previous announcement on February 6, 2017:

<Licensed compounds> Novartis announced on April 3, 2017 that Mekinist® (trametinib) has been approved in EU, in combination with Tafinlar® (dabrafenib), for the treatment of BRAF V600 mutant non-small cell lung cancer (NSCLC). *additional indication