

Johnson & Johnson Innovative Medicine Pipeline Key Events in 2024*

POTENTIAL APPROVALS US/EU

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| | <p>US OPSUMIT (macitentan) EU Pediatric Pulmonary Arterial Hypertension (TOMORROW)</p> |
| ✓ | <p>US OPSYNVI (macitentan/tadalafil STCT) EU Pulmonary Arterial Hypertension</p> |
| ✓ | <p>US EDURANT (rilpivirine) EU HIV pediatric 2-12 year old</p> |
| ✓ | <p>US^ BALVERSA (erdafitinib) EU Urothelial Cancer (THOR)</p> |
| | <p>US DARZALEX (daratumumab) Frontline multiple myeloma transplant eligible (PERSEUS)</p> |
| ✓ | <p>US CARVYKTI (cilta cabtagene autoleu cel) ✓ EU Relapsed Refractory multiple myeloma w/1-3 PL (CARTITUDE-4)</p> |
| ✓ | <p>US RYBREVANT (amivantamab) ✓ EU Frontline Non Small Cell Lung Cancer in combination with chemotherapy (PAPILLON)</p> |
| | <p>US RYBREVANT / lazertinib EU Non Small Cell Lung Cancer 2L (MARIPOSA-2)</p> |
| | <p>US RYBREVANT / lazertinib EU Non Small Cell Lung Cancer (MARIPOSA)</p> |

PLANNED SUBMISSIONS US/EU

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| | <p>US OPSUMIT (macitentan) Pediatric Pulmonary Arterial Hypertension (TOMORROW)</p> |
| ✓ | <p>EU UPTRAVI (selexipag) Pediatric Pulmonary Arterial Hypertension (SALTO)</p> |
| ✓ | <p>EU REKAMBYS HIV Adolescents</p> |
| | <p>US nipocalimab EU Generalized Myasthenia Gravis</p> |
| | <p>US SPRAVATO (esketamine) monotherapy Treatment Resistant Depression (TRD4005)</p> |
| ✓ | <p>US RYBREVANT (amivantamab) ✓ EU Subcutaneous (PALOMA-3)</p> |
| ✓ | <p>US DARZALEX (daratumumab) ✓ EU Frontline multiple myeloma transplant eligible (PERSEUS)</p> |

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| | <p>US SIMPONI (golimumab) EU Pediatric Ulcerative Colitis</p> |
| ✓ | <p>EU STELARA (ustekinumab) Pediatric Crohn's Disease</p> |
| | <p>US TREMFYA (guselkumab) Pediatric Psoriasis</p> |
| ✓ | <p>US TREMFYA (guselkumab) ✓ EU Crohn's Disease (GALAXI)</p> |
| | <p>US TREMFYA (guselkumab) Pediatric Juvenile Psoriatic Arthritis</p> |
| ✓ | <p>US TREMFYA (guselkumab) ✓ EU Ulcerative Colitis Monotherapy (QUASAR)</p> |
| | <p>US TREMFYA (guselkumab) Ulcerative Colitis Subcutaneous Induction (ASTRO)</p> |
| ✓ | <p>US TREMFYA (guselkumab) ✓ EU Crohn's Disease Subcutaneous Induction (GRAVITI)</p> |

POTENTIAL CLINICAL DATA

Phase III

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| ✓ | <p>TREMFYA (guselkumab) Crohn's Disease (GALAXI)</p> |
| ✓ | <p>TREMFYA (guselkumab) Ulcerative Colitis Monotherapy (QUASAR)</p> |
| ✓ | <p>RYBREVANT (amivantamab) Subcutaneous (PALOMA-3)</p> |
| | <p>ERLEADA (apalutamide) High Risk Prostate Cancer (PROTEUS)</p> |
| ✓ | <p>seltorexant Adjunctive treatment for major depressive disorder with insomnia symptoms</p> |
| ✓ | <p>nipocalimab Generalized Myasthenia Gravis</p> |
| | <p>TREMFYA (guselkumab) Crohn's Disease Subcutaneous Induction (GRAVITI)</p> |
| | <p>aticaprant Adjunctive Major Depressive Disorder (VENTURA 1)</p> |
| ✓ | <p>SPRAVATO (esketamine) monotherapy Treatment Resistant Depression (TRD4005)</p> |
| | <p>DARZALEX (daratumumab) Amyloidosis (ANDROMEDA)</p> |

Phase II

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| | <p>JNJ-4804 Co-antibody Therapeutic Psoriatic Arthritis (AFFINITY)</p> |
| ✓ | <p>nipocalimab Sjogren's Disease (DAHLIAS)</p> |
| | <p>TAR-200 (RIS/gemcitabine plus cetrelimab) Non Muscle Invasive Bladder Cancer (SunRISe-1)</p> |



*This information is as of July 17, 2024 to the best of the Company's knowledge. Johnson & Johnson assumes no obligation to update this information. ^ BALVERSA US Full Approval

✓ = Achieved