# MSD Pipeline

4Q2026 Reflecting Pipeline to **02-Nov-2026**

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# Lead-in language

The chart below reflects the company's research pipeline as of **02-Nov-2026**. Candidates shown in Phase 3 include specific products and the date such candidate entered into Phase 3 development. Candidates shown in Phase 2 include the most advanced compound with a specific mechanism or, if listed compounds have the same mechanism, they are each currently intended for commercialization in a given therapeutic area. Small molecules and biologics generally are given MK-number designations and vaccine candidates generally are given V-number designations. Except as otherwise noted, candidates in Phase 1, additional indications in the same therapeutic area (other than with respect to cancer, immunology and certain other indications) and additional claims, line extensions or formulations for in-line products are not shown.

1. Being developed in a collaboration.

Moved forward since last pipeline update.

# MSD pipeline as of 2-Nov-2026

| Phase 2 | Phase 2 | Phase 2 | Phase 2 | Phase 2 | Phase 2 |
|---|---|---|---|---|---|
| Cancer<br>Bladder<br>Cervical<br>Esophageal<br>Gastric<br>Melanoma<br>NSCLC<br>patritumab deruxtecan<br>**MK-1022**[1] | Cancer<br>Heme<br>**MK-1045** | Cancer<br>Solid Tumors<br>calderasib<br>**MK-1084** | Cancer<br>Solid Tumors<br>**MK-2010** | Alzheimer's Disease<br>**MK-2214** | Cancer<br>Bladder<br>Endometrial<br>HCC<br>HNSCC<br>Melanoma<br>NSCLC<br>Ovarian<br>Solid Tumors<br>ifinatamab deruxtecan<br>**MK-2400**[1] |
| Cancer<br>Biliary<br>Esophageal<br>Neoplasm Malignant<br>sacituzumab tirumotecan<br>**MK-2870** | Cancer<br>Prostate<br>**KEYTRUDA®**<br>**MK-3475** | Cancer<br>Heme (US)<br>**KEYTRUDA QLEX™**<br>**MK-3475A** | Cancer<br>Breast<br>Endometrial<br>Ovarian<br>opevesostat<br>**MK-5684** | Cancer<br>Cervical<br>Endometrial<br>Gastric<br>NSCLC<br>SCLC<br>raludotatug deruxtecan<br>**MK-5909**[1] | MASH<br>efinopegdutide<br>**MK-6024** |

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1. Being developed in a collaboration.

Moved forward since last pipeline update.

# MSD pipeline as of 2-Nov-2026

| Phase 2 | Phase 2 | Phase 2 | Phase 2 | Phase 2 | Phase 2 |
|---|---|---|---|---|---|
| Cancer<br>SCLC<br>gocatamig<br>**MK-6070**[1] | Cancer<br>Breast<br>WELIREG™<br>**MK-6482** | Immunology<br>Axial Spondyloarthritis<br>Hidradenitis Suppurativa<br>Psoriatic Arthritis<br>Rheumatoid Arthritis<br>tulisokibart<br>**MK-7240** | Atherosclerosis<br>**MK-7262** | Pulmonary Hypertension due to Left Heart Disease<br>**WINREVAIR™**<br>**MK-7962** | HIV-1 Infection<br>islatravir (+) ulonivirine<br>**MK-8591B** |
| Immunology<br>Ulcerative Colitis<br>**MK-8690** | Cancer<br>Bladder<br>RCC<br>intismeran autogene<br>**V940**[1] | COPD<br>ensifentrine (+)<br>glycopyrrolate<br>**MK-5884A** | | | |

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# MSD pipeline as of 2-Nov-2026

1. Being developed in a collaboration.
2. On FDA partial clinical hold for higher doses of islatravir than those used in current clinical trials.
3. LAGEVRIO™ is available in the U.S under Emergency Use Authorization, which will terminate effective June 29, 2027.
4. Program is in Phase 2/3 studies.

▶ Moved forward since last pipeline update.

| Phase 3 | Phase 3 | Phase 3 | Phase 3 | Phase 3 | Phase 3 |
|---|---|---|---|---|---|
| Cancer<br>Breast<br>patritumab deruxtecan<br>**MK-1022**[1] | Cancer<br>Heme<br>nemtabrutinib<br>**MK-1026** | Cancer<br>CRC<br>NSCLC<br>calderasib<br>**MK-1084** | Influenza<br>efitazanamavir alpha<br>**MK-1406** | Cancer<br>Heme<br>zilovertamab vedotin<br>**MK-2140** | Cancer<br>Esophageal<br>Prostate<br>SCLC<br>ifinatamab deruxtecan<br>**MK-2400**[1] |
| Cancer<br>Bladder<br>Breast<br>Cervical<br>Endometrial (EU)<br>Gastric<br>NSCLC<br>Ovarian<br>sacituzumab tirumotecan<br>**MK-2870** | Diabetic Macular Edema<br>remigromig<br>**MK-3000**[4] | Cancer<br>SCLC<br>**KEYTRUDA®**<br>**MK-3475** | Cancer<br>Myeloproliferative Disorders<br>bomedemstat<br>**MK-3543** | COVID-19<br>**LAGEVRIO®**<br>**MK-4482**[3]<br>**(US)** | Cancer<br>Prostate<br>opevesostat<br>**MK-5684** |
| COPD (EU)<br>**OHTUVAYRE®**<br>**MK-5884** | Cancer<br>Ovarian<br>raludotatug deruxtecan<br>**MK-5909**[1] | Immunology<br>Crohn's Disease<br>Ulcerative Colitis<br>tulisokibart<br>**MK-7240** | Cancer<br>NSCLC<br>SCLC<br>**LYNPARZA®**<br>**MK-7339**[1] | HIV-1 PrEP<br>alimatravir<br>**MK-8527** | HIV-1 Infection<br>islatravir (+) lenacapavir<br>**MK-8591D**[1,2] |
| Neovascular Age-Related Macular Degeneration<br>**MK-8748**[4] | Dengue Fever Virus Vaccine<br>**V181** | Cancer<br>Melanoma<br>NSCLC<br>intismeran autogene<br>**V940**[1] | | | |

# MSD pipeline as of 2-Nov-2026

1. Being developed in a collaboration.
2. Under review for combination use with KEYTRUDA®.
3. Under review in Japan for Lenvima, used in combination with WELIREG®.

▶ Moved forward since last pipeline update.

| **New Molecular Entities** Under Review | **New Molecular Entities** Under Review | **Certain Supplemental Filings** Under Review | **Certain Supplemental Filings** Under Review | **Certain Supplemental Filings** Under Review | **Certain Supplemental Filings** Under Review |
|---|---|---|---|---|---|
| Hypercholesterolemia<br>**LIPFENDRA®**<br>**MK-0616**<br>**(EU, JPN)** | Previously Treated Endometrial Cancer<br>sacituzumab tirumotecan<br>**MK-2870**<br>**(US)** | Respiratory Syncytial Virus (2nd Season)<br>**MK-1654**<br>**ENFLONSIA™**<br>**(US, EU, JPN)** | Cisplatin Eligible Muscle Invasive Bladder Cancer (KNB15)<br>**KEYTRUDA®**<br>**MK-3475**<br>**(JPN)** | Cisplatin Ineligible Muscle Invasive Bladder Cancer (KN905)<br>**KEYTRUDA®**<br>**MK-3475**<br>**(JPN)** | Newly Diagnosed High-Risk Endometrial Cancer (KNB21)<br>**KEYTRUDA®**<br>**MK-3475**<br>**(EU)** |
| HIV-1 Infection<br>**IDVYNSO™**<br>**MK-8591A**<br>**(EU)** | | Clear Cell Renal Cell Carcinoma Following Nephrectomy (LITESPARK-022)<br>**WELIREG®**<br>**MK-6482**[2]<br>**(EU, JPN)** | Previously Treated Advanced Renal Cell Carcinoma (LITESPARK-011)<br>**WELIREG®**<br>**MK-6482**[1,3]<br>**(EU, JPN)** | HIV-1 Infection (Treatment Naive)<br>**IDVYNSO™**<br>**MK-8591A**<br>**(US)** | |

# MSD pipeline as of 2-Nov-2026

1. Approvals obtained within the last 3 months.
2. Being developed in a collaboration.
3. MK-3475A to be marketed under the trade name KEYJECT in Japan.
4. MK-3475A to be marketed under the trade name KEYTRUDA SC in the EU.

▶ Moved forward since last pipeline update.

| New Molecular Entities Approvals[1] | Certain Supplemental Approvals[1] | Certain Supplemental Approvals[1] | Certain Supplemental Approvals[1] | Certain Supplemental Approvals[1] |
|---|---|---|---|---|
| ▶ Cancer Previously Approved Indications (3475A-D77) **KEYTRUDA QLEX™ MK-3475A[3] (JPN, CHN)** | ▶ Cisplatin Eligible Muscle Invasive Bladder Cancer (KNB15) **KEYTRUDA® and KEYTRUDA QLEX™ MK-3475/A[4] (EU)** | ▶ Platinum-Resistant Recurrent Ovarian Cancer (KNB96) **KEYTRUDA® MK-3475 (JPN)** | ▶ Resectable Locally Advanced Head and Neck Squamous Cell Carcinoma (KN689) **KEYTRUDA® MK-3475 (CHN)** | ▶ Advanced, Unresectable, or Metastatic Pheochromocytoma and Paraganglioma (PPGL) (LITESPARK-015) **WELIREG® MK-6482 (CHN)** |
| | ▶ Previously Treated Advanced Renal Cell Carcinoma (LITESPARK-011) **WELIREG® MK-6482[2] (US)** | ▶ Pulmonary Arterial Hypertension (HYPERION) **WINREVAIR™ MK-7962 (US)** | ▶ Children and Adolescents Aged 2 through 17 at Increased Risk for Pneumococcal Disease (STRIDE-13) **CAPVAXIVE™ V116 (JPN)** | |

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# Forward-looking statement

This presentation of Merck & Co., Inc., Rahway, N.J., USA (the "company") includes "forward-looking statements" within the meaning of the safe harbor provisions of the U.S. Private Securities Litigation Reform Act of 1995. These statements are based upon the current beliefs and expectations of the company's management and are subject to significant risks and uncertainties. There can be no guarantees with respect to pipeline candidates that the candidates will receive the necessary regulatory approvals or that they will prove to be commercially successful. If underlying assumptions prove inaccurate or risks or uncertainties materialize, actual results may differ materially from those set forth in the forward-looking statements.

Risks and uncertainties include but are not limited to, general industry conditions and competition; general economic factors, including interest rate and currency exchange rate fluctuations; the impact of pharmaceutical industry regulation and health care legislation in the United States and internationally; global trends toward health care cost containment; technological advances, new products and patents attained by competitors; challenges inherent in new product development, including obtaining regulatory approval; the company's ability to accurately predict future market conditions; manufacturing difficulties or delays; financial instability of international economies and sovereign risk; dependence on the effectiveness of the company's patents and other protections for innovative products; and the exposure to litigation, including patent litigation, and/or regulatory actions.

The company undertakes no obligation to publicly update any forward-looking statement, whether as a result of new information, future events or otherwise. Additional factors that could cause results to differ materially from those described in the forward-looking statements can be found in the company's Annual Report on Form 10-K for the year ended 2025 and the company's other filings with the Securities and Exchange Commission (SEC) available at the SEC's Internet site (www.sec.gov).

# No duty to update

The information contained in the presentation set forth below was current as of **2-Nov-2026**. While this presentation remains on the company's website the company assumes no duty to update the information to reflect subsequent developments. Consequently, the company will not update the information contained in the presentation and investors should not rely upon the information as current or accurate after **2-Nov-2026**.

The chart reflects the MSD research pipeline as of **2-Nov-2026**.

Candidates shown in Phase 3 include specific products. Candidates shown in Phase 2 include the most advanced compound with a specific mechanism in a given therapeutic area. Phase 1 candidates are not shown.