

MSD Pipeline

November 1, 2016

MSD Pipeline as of November 1, 2016

Phase 2	Phase 2	Phase 3	Phase 3	Phase 3	Phase 3
Asthma MK-1029	Diabetes Mellitus MK-8521	Alzheimer's Disease verubecestat MK-8931	Cancer Bladder Breast Colorectal Esophageal Gastric Hepatocellular Head and Neck (EU) Hodgkin Lymphoma Multiple Myeloma Renal KEYTRUDA® MK-3475	Diabetes Mellitus ertugliflozin MK-8835 ¹ ertugliflozin+sitagliptin MK-8835A ¹ ertugliflozin+metformin MK-8835B ¹	HIV doravirine MK-1439
Cancer PMBCL ² Advanced solid tumors Nasopharyngeal Ovarian Prostate KEYTRUDA® MK-3475	Hepatitis C MK-3682/ MK-5172(grazoprevir) /MK-8408(ruzasvir) MK-3682B	Atherosclerosis anacetrapib MK-0859	CMV Prophylaxis in Transplant Patients letermovir MK-8228	Ebola Vaccine V920	HABP/VABP ³ bacterial pneumonia ZERBAXA™ MK-7625A
Cancer MK-2206 MK-8628	Pneumoconjugate Vaccine V114	Bacterial Infection relebactam+imipenem/ cilastatin MK-7655A	Diabetes Mellitus sitagliptin+ipragliflozin MK-0431J ¹ (Japan)	Herpes Zoster inactivated VZV vaccine V212	HABP/VABP ³ bacterial pneumonia SIVEXTRO® MK-1986
Cough, including cough w/ IPF ⁴ MK-7264		Heart Failure Vericiguat MK-1242 ¹			



Moved forward since last pipeline update.

1. Being developed in a collaboration.
2. Primary Mediastinal Large B-Cell Lymphoma
3. HABP - Hospital-acquired bacterial pneumonia/
VABP - ventilator-associated bacterial pneumonia
4. Idiopathic Pulmonary Fibrosis

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New Molecular Entities Under Review	New Molecular Entities Under Review	New Molecular Entities Approvals ¹	New Molecular Entities Approvals ¹	New Molecular Entities Approvals ¹	New Molecular Entities Approvals ¹
MK-8237 Allergy, House Dust Mite (US) ²	V419 Pediatric Hexavalent Combination Vaccine (US) ³	BRIDION® MK-8616 Neuromuscular Blockade Reversal (US)	ZERBAXA® MK-7625A Complicated intra-abdominal infections (cIAI) & complicated urinary tract infections (cUTI) (US/EU)	GARDASIL®9 V503 HPV Vaccine for Cancer Prevention (US/EU)	KEYTRUDA® MK-3475 Melanoma (EU)
ZINPLAVA™ MK-6072 <i>Clostridium difficile</i> Infection (EU)	MK-1293 Diabetes Mellitus (US/EU) ⁴	MARIZEV® MK-3102 Diabetes Mellitus (Japan)	ZEPATIER® MK-5172A Hepatitis C (US/EU)	VAXELIS™ V419 Pediatric Hexavalent Combination Vaccine (EU) ³	ZINPLAVA™ MK-6072 <i>Clostridium difficile</i> Infection (US)

1. Approvals obtained within the last 24 months.
2. MK-8237 was being developed as part of a North America partnership with ALK-Abelló (ALK). Merck has given ALK six months' notice that it is terminating the agreement and therefore this compound will be returned to ALK.
3. V419, the investigational pediatric hexavalent combination vaccine, DTaP5-IPV-Hib-HepB, is being developed in partnership with Sanofi Pasteur and if approved in the US, will be commercialized through that partnership. On November 2, 2015, the FDA issued a CRL with respect to V419. The companies are reviewing the CRL and plan to have further discussions with the FDA.
4. Being developed in a collaboration

▶ Moved forward since last pipeline update.

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This presentation includes “forward-looking statements” within the meaning of the safe harbor provisions of the United States Private Securities Litigation Reform Act of 1995. These statements are based upon the current beliefs and expectations of MSD’s management and are subject to significant risks and uncertainties. There can be no guarantees with respect to pipeline products that the products 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; MSD’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 MSD’s patents and other protections for innovative products; and the exposure to litigation, including patent litigation, and/or regulatory actions.

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The chart reflects the MSD's research pipeline as of November 1, 2016.

Candidates shown in Phase III include specific products. Candidates shown in Phase II include the most advanced compound with a specific mechanism in a given therapeutic area. Phase I candidates are not shown.