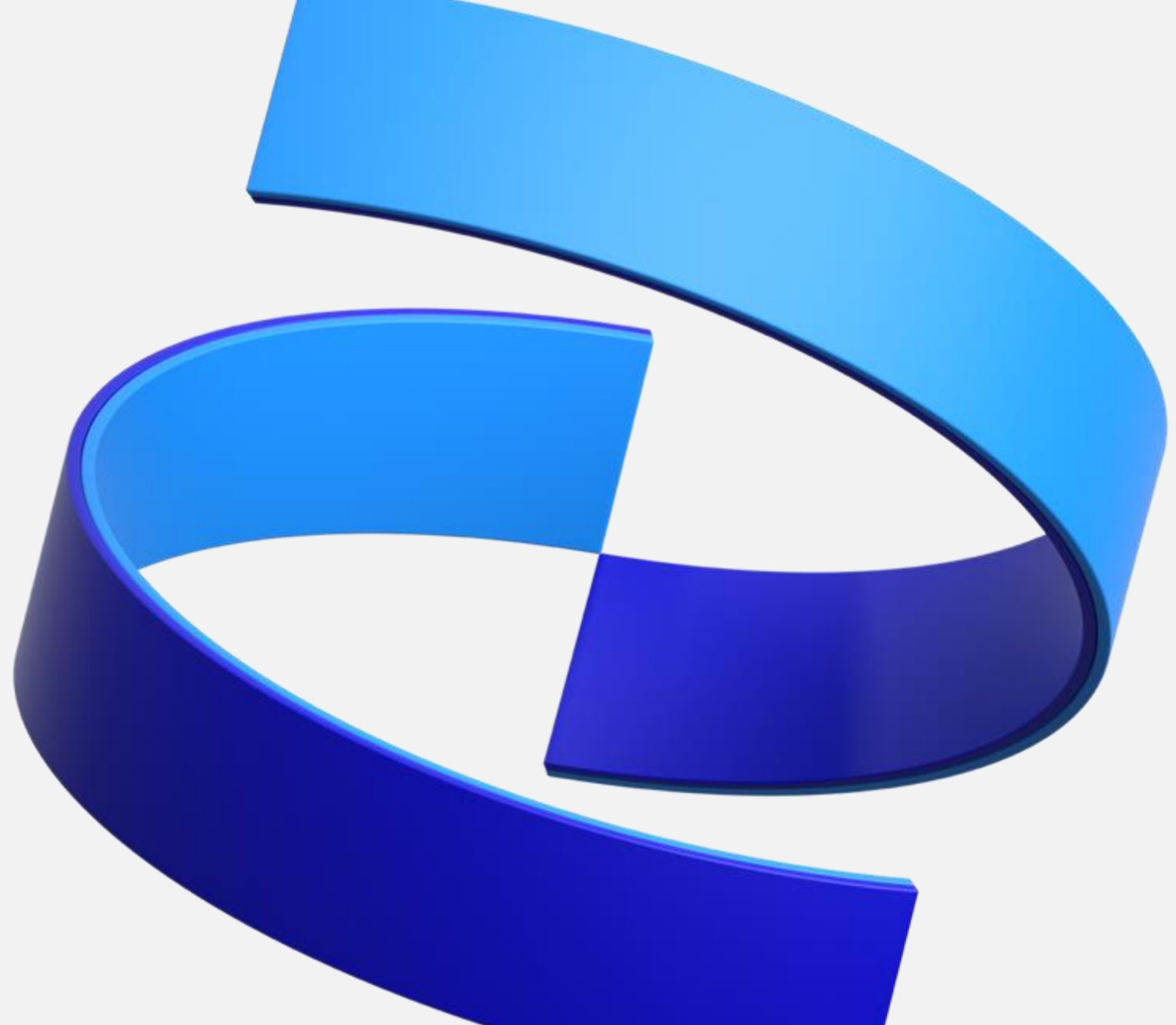




Pfizer Pipeline

Nov 2, 2021



Disclaimer

- As some programs are still confidential, some candidates may not be identified in this list. In these materials, Pfizer discloses Mechanism of Action (MOA) information for some candidates in Phase 1 and for all candidates from Phase 2 through regulatory approval. With a view to expanding the transparency of our pipeline, Pfizer is including new indications or enhancements, which target unmet medical need or represent significant commercial opportunities. The information contained on these pages is correct as of Nov 2, 2021.
- Visit www.pfizer.com/pipeline, Pfizer's online database where you can learn more about our portfolio of new medicines and find out more about our Research and Development efforts around the world.



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Pfizer Pipeline Snapshot



Pfizer Pipeline Snapshot as of November 2, 2021

Pipeline represents progress of R&D programs as of November 2, 2021

- 13 programs advanced or are new
- 16 programs discontinued since last update
- Included are 59 NMEs, 35 additional indications

Recent Approvals

- TICOVAC™ (tick-borne encephalitis (TBE) vaccine) for active immunization to prevent TBE in individuals 1 year of age and older (U.S.)
- COMIRNATY® (COVID-19 Vaccine, mRNA) to prevent COVID-19 in individuals 16 years of age and older. COMIRNATY is the first COVID-19 vaccine to be granted approval by the FDA (U.S.)
- * COMIRNATY® Booster (Pfizer/BioNTech COVID-19 vaccine) received Emergency Use Authorization (EUA) from FDA on Sep 22, 2021 for individuals 65 years of age and older, and individuals ages 18 through 64 within certain high-risk groups; received conditional marketing authorization from the EMA on Oct 5, 2021 for 18 years of age and older. COMIRNATY® received EUA from FDA (U.S.) for 5 to 11 years old age group on Oct 29, 2021



Pfizer Pipeline Snapshot as of July 28, 2021

Pipeline represents progress of R&D programs as of July 28, 2021

- 8 programs advanced or are new
- 5 programs discontinued since last update
- Included are 66 NMEs, 34 additional indications

Recent Approvals

- XTANDI™ (enzalutamide) for adult men with metastatic hormone-sensitive prostate cancer (mHSPC, also known as metastatic castration-sensitive prostate cancer or mCSPC) (E.U.)
- MYFEMBREE® (relugolix 40 mg, estradiol 1 mg, and norethindrone acetate 0.5 mg), for the management of heavy menstrual bleeding associated with uterine fibroids in premenopausal women (U.S.)
- PREVNAR 20™ (Pneumococcal 20-valent Conjugate Vaccine) for the prevention of invasive disease and pneumonia caused by the 20 Streptococcus pneumoniae (pneumococcus) serotypes in the vaccine in adults ages 18 years and older (U.S.)
- * COMIRNATY® (Pfizer/BioNTech COVID-19 vaccine) received Emergency Use Authorization from FDA (U.S.) and Conditional Marketing Authorization from the EMA (E.U.) for 12-15 years old age group

Inflammation and Immunology



Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
abrocitinib (PF-04965842)	JAK1 Inhibitor	Atopic Dermatitis (PRIORITY REVIEW, BREAKTHROUGH – U.S., E.U.)	Registration	New Molecular Entity
Xeljanz (tofacitinib)	JAK Inhibitor	Ankylosing Spondylitis (U.S., E.U.)	Registration	Product Enhancement
ritlecitinib (PF-06651600)	JAK3/TEC Inhibitor	Alopecia Areata (BREAKTHROUGH)	Phase 3	New Molecular Entity
Dekavil*	IL-10	Rheumatoid Arthritis (Biologic)	Phase 2	New Molecular Entity
PF-06480605	TNFSF15 Blocker	Ulcerative Colitis (Biologic)	Phase 2	New Molecular Entity
ritlecitinib +/- PF-06650833; ritlecitinib + PF-06650833 + tofacitinib	JAK3/TEC Inhibitor IRAK4 Inhibitor JAK Inhibitor	Rheumatoid Arthritis	Phase 2	New Molecular Entity
PF-06650833	IRAK4 Inhibitor	Hidradenitis Suppurativa	Phase 2	Product Enhancement
ritlecitinib (PF-06651600)	JAK3/TEC Inhibitor	Ulcerative Colitis	Phase 2	Product Enhancement
ritlecitinib (PF-06651600)	JAK3/TEC Inhibitor	Crohn's Disease	Phase 2	Product Enhancement
ritlecitinib (PF-06651600)	JAK3/TEC Inhibitor	Vitiligo	Phase 2	Product Enhancement
Eucrisa (crisaborole)	PDE4 Inhibitor	Stasis Dermatitis	Phase 2	Product Enhancement
PF-06823859	interferon, beta 1, fibroblast (IFNB1) Blocker	Dermatomyositis (Biologic) (ORPHAN - U.S., E.U., PRIME - E.U.)	Phase 2	New Molecular Entity
PF-07038124	Topical PDE4 Inhibitor	Atopic Dermatitis	Phase 2	New Molecular Entity
PF-06835375	Chemokine Inhibitor	Lupus (Biologic)	Phase 1	New Molecular Entity
PF-07054894	CCR6 Antagonist	Inflammatory Bowel Disease	Phase 1	New Molecular Entity
PF-07242813	CD1a inhibitor	Atopic Dermatitis (Biologic)	Phase 1	New Molecular Entity

► Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com

• Regulatory Designations – See Definitions in Backup

*Clinical trial to be conducted by Philogen S.p.A

Internal Medicine (1 of 2)



Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
► relugolix fixed dose combination	Oral GnRH receptor antagonist	Combination with estradiol and norethindrone acetate for Endometriosis (U.S.)	Registration	Product Enhancement
relugolix fixed dose combination	Oral GnRH receptor antagonist	Combination with estradiol and norethindrone acetate for contraceptive efficacy	Phase 3	Product Enhancement
ervogastat (PF-06865571)	Diacylglycerol O-Acyltransferase 2 (DGAT2) Inhibitor	Non-alcoholic Steatohepatitis (NASH) with Liver Fibrosis	Phase 2	New Molecular Entity
ervogastat (PF-06865571) + clesacostat (PF-05221304)	Diacylglycerol O-Acyltransferase 2 (DGAT2) Inhibitor; Acetyl CoA-Carboxylase (ACC) Inhibitor	Non-alcoholic Steatohepatitis (NASH) with Liver Fibrosis	Phase 2	New Molecular Entity
vupanorsen (PF-07285557)	Angiopoietin Like 3 (ANGPTL3) Antisense Oligonucleotide	Severe Hypertriglyceridemia, Cardiovascular Risk Reduction	Phase 2	New Molecular Entity
danuglipron (PF-06882961)	Glucagon-like peptide 1 receptor (GLP-1R) Agonist	Diabetes Mellitus-Type 2	Phase 2	New Molecular Entity
danuglipron (PF-06882961)	Glucagon-like peptide 1 receptor (GLP-1R) Agonist	Obesity	Phase 2	Product Enhancement

► Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com

• Regulatory Designations – See Definitions in Backup

Internal Medicine (2 of 2)

Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
PF-06946860	Growth Differentiation Factor 15 (GDF15) Monoclonal Antibody	Cachexia (Biologic)	Phase 1	New Molecular Entity
PF-07081532	Glucagon-like peptide 1 receptor (GLP-1R) Agonist	Diabetes Mellitus-Type 2 and Obesity	Phase 1	New Molecular Entity
danuglipron (PF-06882961) + PF-06865571	Glucagon-like peptide 1 receptor (GLP-1R) Agonist; Diacylglycerol O-Acyltransferase 2 (DGAT2) Inhibitor	Non-alcoholic Steatohepatitis (NASH) with Liver Fibrosis	Phase 1	New Molecular Entity
PF-07258669	Melanocortin-4 receptor (MC4R) Antagonist	Geriatric Anorexia	Phase 1	New Molecular Entity
PF-07202954	Diacylglycerol O-Acyltransferase 2 (DGAT2) Inhibitor	Non-alcoholic Steatohepatitis (NASH) with Liver Fibrosis	Phase 1	New Molecular Entity

► Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com

• Regulatory Designations – See Definitions in Backup

Oncology (1 of 3)



Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
Lorbrena (lorlatinib)	ALK inhibitor	1 st Line ALK Non-Small Cell Lung Cancer (E.U.)	Registration	Product Enhancement
Bavencio (avelumab)	Anti PD-L1	1 st Line Non-Small Cell Lung Cancer (Biologic)	Phase 3	Product Enhancement
Ibrance (palbociclib)	CDK 4,6 kinase inhibitor	ER+/HER2+ Metastatic Breast Cancer (PATINA)	Phase 3	Product Enhancement
sasanlimab (PF-06801591) + Bacillus Calmette-Guerin (BCG)	Anti-PD-1	Non-Muscle-Invasive Bladder Cancer (Biologic)	Phase 3	New Molecular Entity
Talzenna (talazoparib)	PARP inhibitor	Combo w/ Xtandi (enzalutamide) for 1 st Line Metastatic Castration Resistant Prostate Cancer	Phase 3	Product Enhancement
Talzenna (talazoparib)	PARP inhibitor	Combo w/ Xtandi (enzalutamide) for DDR-deficient Metastatic Castration Sensitive Prostate Cancer	Phase 3	Product Enhancement
Xtandi (enzalutamide)	Androgen receptor inhibitor	Non-metastatic High-Risk Castration Sensitive Prostate Cancer	Phase 3	Product Enhancement
Braftovi (encorafenib) + Erbitux® (cetuximab)	<i>BRAF</i> kinase inhibitor and anti EGFR	1 st line <i>BRAF</i> -mutant Metastatic Colorectal Cancer	Phase 3	Product Enhancement
Braftovi (encorafenib) + Mektovi (binimetinib) + Keytruda® (pembrolizumab)	<i>BRAF</i> kinase inhibitor and MEK inhibitor and anti PD-1	<i>BRAF</i> -mutant Metastatic or Unresectable Locally Advanced Melanoma	Phase 3	Product Enhancement
► Elranatamab	BCMA-CD3 Bispecific Antibody	Multiple Myeloma Double-Class Exposed (Biologic)	Phase 3	New Molecular Entity

► Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com

- Regulatory Designations – See Definitions in Backup
- Erbitux® is a registered trademark of ImClone LLC
- Keytruda® is a registered trademark of Merck Sharp & Dohme Corp.

Oncology (2 of 3)

Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
Bavencio (avelumab)	Anti PD-L1	Combo w/Talzenna (talazoparib) for Locally Advanced (Primary or Recurrent) or Metastatic Solid Tumors (Biologic)	Phase 2	Product Enhancement
Bavencio (avelumab)	Anti PD-L1	Combo w/Talzenna (talazoparib) for Solid Tumors with a BRCA or ATM defect (Biologic)	Phase 2	Product Enhancement
Braftovi (encorafenib) + Mektovi (binimetinib)	<i>BRAF</i> kinase inhibitor and MEK inhibitor	1 st line and 2 nd line <i>BRAF</i> -mutant Metastatic Non-Small Cell Lung Cancer	Phase 2	Product Enhancement
Talzenna (talazoparib)	PARP inhibitor	2 nd Line Metastatic Castration-Resistant Prostate Cancer	Phase 2	Product Enhancement
ARV-471	ER-targeting PROTAC® protein degrader	ER+/HER2- Metastatic Breast Cancer	Phase 2	New Molecular Entity
Elranatamab	BCMA-CD3 Bispecific Antibody	Multiple Myeloma Triple-Class Refractory (Biologic)	Phase 2	Product Enhancement

► Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com

- PROTAC® is a registered U.S. trademark of Arvinas.



Oncology (3 of 3)

Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
PF-06647020	protein tyrosine kinase 7 (PTK7) Targeted Cytotoxicity	Cancer (Biologic)	Phase 1	New Molecular Entity
PF-06821497	EZH2 Inhibitor	Prostate Cancer	Phase 1	New Molecular Entity
PF-06873600	CDK 2,4,6 Inhibitor	Breast Cancer Metastatic	Phase 1	New Molecular Entity
PF-06939999	protein arginine methyltransferase 5 (PRMT5) Inhibitor	Solid Tumors	Phase 1	New Molecular Entity
PF-07062119	GUCY2c CD3 Bispecific Antibody	Solid Tumors (Biologic)	Phase 1	New Molecular Entity
PF-06940434	Integrin alpha-V/beta-8 Antagonist	Solid Tumors (Biologic)	Phase 1	New Molecular Entity
PF-07209960	interleukin 15 (IL15) Activator	Solid Tumors (Biologic)	Phase 1	New Molecular Entity
PF-07220060	CDK4 Inhibitor	Breast Cancer Metastatic	Phase 1	New Molecular Entity
PF-07265807	AXL/MERTK Inhibitor	Solid Tumors	Phase 1	New Molecular Entity
PF-07104091	CDK2 Inhibitor	Breast Cancer Metastatic	Phase 1	New Molecular Entity
PF-07248144	KAT6A Epigenetic modifier	Breast Cancer Metastatic	Phase 1	New Molecular Entity
PF-07284890	<i>BRAF</i> kinase Inhibitor	Melanoma	Phase 1	New Molecular Entity
PF-07284892	SHP2 tyrosine phosphatase Inhibitor	Cancer	Phase 1	New Molecular Entity
lbrance + ARV-471	CDK 4,6 kinase inhibitor ER-targeting PROTAC® protein degrader	ER+/HER2- Metastatic Breast Cancer	Phase 1	Product Enhancement
▶ PF-07257876	CD47xPDL1 Bispecific	NSCLC (Biologic)	Phase 1	New Molecular Entity

▶ Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com

- PROTAC® is a registered U.S. trademark of Arvinas.



Rare Disease



Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
somatrogon (PF-06836922)	Human Growth Hormone Agonist	Pediatric Growth Hormone Deficiency (Biologic) (ORPHAN - U.S., E.U.)	Registration	New Molecular Entity
PF-07265803	p38 Mitogen-Activated Protein Kinase Antagonist	Dilated Cardiomyopathy due To Lamin A/C Gene Mutation (ORPHAN - U.S.)	Phase 3	New Molecular Entity
fidanacogene elaparovec (PF-06838435)	Gene Therapy, coagulation factor IX (F9)	Hemophilia (Biologic) (BREAKTHROUGH, ORPHAN - U.S., E.U., PRIME - E.U.)	Phase 3	New Molecular Entity
giroctocogene fitelparovec (PF-07055480)	Gene Therapy, coagulation factor VIII (F8)	Hemophilia (Biologic) (RMAT, FAST TRACK, ORPHAN - U.S., E.U.)	Phase 3	New Molecular Entity
somatrogon (PF-06836922)	Human Growth Hormone Agonist	Adult Growth Hormone Deficiency (Biologic) (ORPHAN - U.S., E.U.)	Phase 3	Product Enhancement
fordadistrogene movaparovec (PF-06939926)	Gene Therapy, minidystrophin	Duchenne Muscular Dystrophy Ambulatory (Biologic) (FAST TRACK – U.S.; ORPHAN - U.S., E.U.)	Phase 3	New Molecular Entity
marstacimab (PF-06741086)	Tissue Factor Pathway Inhibitor (TFPI)	Hemophilia (Biologic) (FAST TRACK – U.S.; ORPHAN - U.S., E.U.)	Phase 3	New Molecular Entity
PF-06730512	SLIT2 antagonist	Focal Segmental Glomerulosclerosis (FSGS); ROBO2-Fc (Biologic)	Phase 2	New Molecular Entity
recifercept	Soluble recombinant human fibroblast growth factor receptor 3 (FGFR3) decoy	Achondroplasia (Biologic) (ORPHAN - U.S., EU)	Phase 2	New Molecular Entity
PF-06755347	Immunomodulation	Idiopathic thrombocytopenic purpura/Chronic Inflammatory Demyelination Polyneuropathy (Biologic) (ORPHAN - U.S.)	Phase 1	New Molecular Entity
PF-07209326	E-Selectin antagonist	Sickle Cell Disease (Biologic) (ORPHAN - U.S.)	Phase 1	New Molecular Entity
PF-07059013	Hemoglobin, Beta (HBB) Modulator	Sickle Cell Disease (ORPHAN - U.S.)	Phase 1	New Molecular Entity

► Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com
 • Regulatory Designations – See Definitions in Backup

Vaccines (1 of 2)



Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
Comirnaty (Covid-19 Vx)	Prophylactic mRNA Vaccine	COVID-19 Infection (in collaboration with BioNTech) (FAST TRACK, U.S. – 12 to 15 years of age) (E.U. – 12 years of age and older)*	Registration	Product Enhancement
▶ Comirnaty (Covid-19 Vx)	Prophylactic mRNA Vaccine	COVID-19 Infection Booster (in collaboration with BioNTech) (FAST TRACK, U.S.; EU)**	Registration	Product Enhancement
▶ Comirnaty (Covid-19 Vx)	Prophylactic mRNA Vaccine	COVID-19 Infection (in collaboration with BioNTech) (FAST TRACK, U.S.; EU – 5 to 11 years of age)***	Registration	Product Enhancement
PF-06482077	Prophylactic Vaccine	Invasive and Non-Invasive Pneumococcal infections (adult) (E.U.)	Registration	New Molecular Entity
PF-06425090	Prophylactic Vaccine	Primary <i>Clostridioides difficile</i> infection (FAST TRACK)	Phase 3	New Molecular Entity
PF-06482077	Prophylactic Vaccine	Invasive and Non-Invasive Pneumococcal infections (pediatric) (BREAKTHROUGH, FAST TRACK)	Phase 3	Product Enhancement
PF-06928316	Prophylactic Vaccine	Respiratory Syncytial Virus Infection (maternal) (FAST TRACK)	Phase 3	New Molecular Entity
PF-06886992	Prophylactic Vaccine	Serogroups ABCWY Meningococcal Infections (adolescent and young adults)	Phase 3	New Molecular Entity
▶ PF-06928316	Prophylactic Vaccine	Respiratory Syncytial Virus Infection (older adult)	Phase 3	Product Enhancement

* Comirnaty (Pfizer/BioNTech COVID-19 vaccine) received Emergency Use Authorization (EUA) from FDA on Dec 11, 2020 and FDA BLA on Aug 23, 2021 for 16 years of age and older; received conditional marketing authorization (CMA) from the EMA on Dec 21, 2020 for 16 years of age and older. 12-15 years old age group received EUA from FDA on May 10, 2021 and conditional marketing authorization from the EMA on May 28, 2021.

** Comirnaty Booster (Pfizer/BioNTech COVID-19 vaccine) received EUA from FDA on Sep 22, 2021 for individuals 65 years of age and older, and individuals ages 18 through 64 within certain high-risk groups; received CMA from the EMA on Oct 5, 2021 for 18 years of age and older.

*** Comirnaty (Pfizer/BioNTech COVID-19 vaccine) received EUA from FDA on Oct 29, 2021 and submitted to EU for CMA on Oct 15, 2021 for 5 to 11 years of age

▶ Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com

• Regulatory Designations – See Definitions in Backup

Vaccines (2 of 2)

Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
▶ Comirnaty (Covid-19 Vx)	Prophylactic mRNA Vaccine	COVID-19 Infection (in collaboration with BioNTech) (maternal)	Phase 3	Product Enhancement
▶ Comirnaty (Covid-19 Vx)	Prophylactic mRNA Vaccine	COVID-19 Infection (in collaboration with BioNTech) (children 2 to 4 years of age)	Phase 3	Product Enhancement
▶ Comirnaty (Covid-19 Vx)	Prophylactic mRNA Vaccine	COVID-19 Infection (in collaboration with BioNTech) (infants 6 months to <24 months)	Phase 3	Product Enhancement
PF-06842433	Prophylactic Vaccine	Invasive and Non-Invasive Pneumococcal infections (infants and children)	Phase 2	New Molecular Entity
PF-06760805	Prophylactic Vaccine	Invasive Group B Streptococcus Infection (maternal) (FAST TRACK)	Phase 2	New Molecular Entity
PF-07307405	Prophylactic Vaccine	Lyme disease (FAST TRACK)	Phase 2	New Molecular Entity
PF-06886992	Prophylactic Vaccine	Serogroups ABCWY Meningococcal Infections (infants)	Phase 2	Product Enhancement
▶ PF-07252220	Prophylactic mRNA Vaccine	Influenza (adults)	Phase 1	New Molecular Entity

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• Regulatory Designations – See Definitions in Backup

Hospital (Anti-Infectives)



Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
aztreonam-avibactam (PF-06947387)	Beta Lactam/Beta Lactamase Inhibitor	Treatment of infections caused by Gram-negative bacteria	Phase 3	New Molecular Entity
▶ PF-07321332	SARS-CoV-2 3CL protease inhibitor (oral anti-viral)	COVID-19 Infection (high risk population)	Phase 3	New Molecular Entity
▶ PF-07321332	SARS-CoV-2 3CL protease inhibitor (oral anti-viral)	COVID-19 Infection (low risk population)	Phase 3	Product Enhancement
▶ PF-07321332	SARS-CoV-2 3CL protease inhibitor (oral anti-viral)	COVID-19 Infection (post exposure prophylaxis)	Phase 3	Product Enhancement
Fosmanogepix (APX001)	Gwt1 inhibitor	Treatment of invasive fungal infections	Phase 2	New Molecular Entity
PF-07304814	SARS-CoV-2 3CL protease inhibitor (IV anti-viral)	COVID-19 Infection	Phase 2	New Molecular Entity

▶ Indicates that the project is either new or has progressed in phase since the previous portfolio update of Pfizer.com

Programs Discontinued from Development since July 28, 2021

Compound Name	Mechanism of Action	Indication	Phase of Development	Submission Type
tanezumab	Nerve Growth Factor Inhibitor	Chronic Pain due to Moderate-to-Severe Osteoarthritis (OA) (Biologic) (U.S., E.U.)	Registration	New Molecular Entity
tanezumab	Nerve Growth Factor Inhibitor	Cancer Pain (Biologic)	Phase 3	Product Enhancement
brepocitinib (PF-06700841)* PF-06826647*	TYK2/JAK1 Inhibitor TYK2 Inhibitor	Hidradenitis Suppurativa	Phase 2	Product Enhancement
brepocitinib (PF-06700841)*	TYK2/JAK1 Inhibitor	Ulcerative Colitis	Phase 2	New Molecular Entity
brepocitinib (PF-06700841)*	TYK2/JAK1 Inhibitor	Crohn's Disease	Phase 2	Product Enhancement
brepocitinib (PF-06700841)*	TYK2/JAK1 Inhibitor	Vitiligo	Phase 2	Product Enhancement
brepocitinib (PF-06700841)*	TYK2/JAK1 Inhibitor	Psoriatic Arthritis	Phase 2	Product Enhancement
brepocitinib (PF-06700841)*	TYK2/JAK1 Inhibitor	Alopecia Areata	Phase 2	Product Enhancement
brepocitinib (PF-06700841)*	TYK2/JAK1 Inhibitor	Lupus	Phase 2	Product Enhancement
brepocitinib (PF-06700841)*	Topical TYK2/JAK1 Inhibitor	Atopic Dermatitis	Phase 2	Product Enhancement
brepocitinib (PF-06700841)*	Topical TYK2/JAK1 Inhibitor	Psoriasis	Phase 2	Product Enhancement
PF-06826647*	TYK2 Inhibitor	Psoriasis	Phase 2	New Molecular Entity
Bavencio (avelumab)	Anti PD-L1	Combo w/CMP-001 for Head and Neck Cancer (Biologic)	Phase 2	Product Enhancement
PF-06952229	transforming growth factor, beta receptor 1 (TGFB1) Inhibitor	Cancer	Phase 1	New Molecular Entity
ARV-471**	ER-targeting PROTAC® protein degrader	ER+/HER2- Metastatic Breast Cancer	Phase 1	Product Enhancement
PF-06842874	CDK 4,6 Inhibitor	Pulmonary Arterial Hypertension (ORPHAN - U.S.)	Phase 1	New Molecular Entity

*brepocitinib (PF-06700841) and PF-06826647 have been out-licensed

** Change in reporting convention: ARV-471 Ph1 is a part of the same study as on-going ARV-471 Ph2

An abstract 3D graphic composed of several overlapping, curved, blue and purple planes that create a sense of depth and movement, resembling a stylized wave or a series of connected segments. The colors transition from a light blue on the left to a deep purple on the right.

Back-up

Regulatory Designations

- **Fast Track (U.S.)** is a designation available to a product if it is intended, whether alone or in combination with one or more other drugs, for the treatment of a serious or life-threatening disease or condition, and it demonstrates the potential to address unmet medical needs for such a disease or condition. This designation is intended to facilitate development and expedite review of drugs to treat serious and life-threatening conditions so that an approved product can reach the market expeditiously. More information about the qualifying criteria and features of the Fast Track program can be found on the FDA's website.
- **Breakthrough Designation (U.S.)** may be granted to a drug (alone or in combination with 1 or more other drugs) intended to treat a serious or life-threatening disease or condition, and preliminary clinical evidence indicates that the drug may demonstrate substantial improvement over existing therapies on one or more clinically significant endpoints, such as substantial treatment effects observed early in clinical development. A drug that receives breakthrough designation is eligible for all fast track designation features and an FDA commitment to work closely with the sponsor to ensure an efficient drug development program. More information about the qualifying criteria and features of the Breakthrough program can be found on the FDA's website.
- **Orphan Drug (U.S.)** - Orphan drug status may be granted to drugs and biologics that are intended for the diagnosis, prevention, or treatment of rare diseases/disorders that affect fewer than 200,000 people in the U.S., or that affect more than 200,000 persons but where it is unlikely that expected sales of the product would cover the sponsor's investment in its development. More information about the qualifying criteria, features, and incentives involved in an orphan drug designation can be found on the FDA's website.
- **Orphan Drug (E.U.)** - Orphan drug status may be granted to drugs and biologics that are intended for the diagnosis, prevention or treatment of a life-threatening or chronically debilitating condition affecting no more than 5 in 10,000 persons in the European Union at the time of submission of the designation application, or that affect more than 5 in 10,000 persons but where it is unlikely that expected sales of the product would cover the investment in its development. More information about the qualifying criteria, features, and incentives involved in an orphan drug designation can be found on the EMA's website.
- A U.S. drug application will receive a **priority review designation** if it is for a drug that treats a serious condition and, if approved, would provide a significant improvement in safety or effectiveness. A priority designation is intended to direct overall attention and resources to the evaluation of such applications. A priority review designation means that FDA's goal is to take action on the marketing application within 6 months of receipt (compared with 10 months under standard review). More information about the qualifying criteria and features of a priority review designation can be found on the FDA's website.
- **PRIME (E.U.)** - The PRIME scheme is applicable to products under development which are innovative and yet to be placed on the EU market. The scheme aims to support medicinal products of major public health interest and in particular from the viewpoint of therapeutic innovation. Medicines eligible for PRIME must address an unmet medical need, i.e. for which there exists no satisfactory method of diagnosis, prevention or treatment in the Community or, if such a method exists, in relation to which the medicinal product concerned will be of major therapeutic advantage to those affected. A product eligible for PRIME should demonstrate the potential to address, to a significant extent, the unmet medical need, for example by introducing new methods of therapy or improving existing ones. Data available to support the request for eligibility should support the claim to address the unmet medical need through a clinically meaningful improvement of efficacy, such as having an impact on the prevention, onset or duration of the condition, or improving the morbidity or mortality of the disease. EMA will provide early and enhanced support to optimize the development of eligible medicines. Products granted PRIME support are anticipated to benefit from the Accelerated Assessment procedure. More information about the qualifying criteria and features of PRIME and Accelerated Assessment can be found on the EMA's website.
- **Regenerative Medicine Advanced Therapy (RMAT) (U.S.)** is a designation that is granted to regenerative medicine therapies intended to treat, modify, reverse, or cure a serious condition, for which preliminary clinical evidence indicates that the medicine has the potential to address an unmet medical need. The RMAT designation includes all the benefits of the fast track and breakthrough therapy designation programs, including early interactions with FDA.