

INNOVATIVE SCIENCE

LEADING MEDICINES

Noteworthy in 2012

For more information on any of these medicines, visit
Pfizer Pharmaceutical Products.

Xeljanz

OWNING POTENTIAL



Xeljanz (tofacitnib) is the first new oral disease-modifying antirheumatic drug approved for rheumatoid arthritis in more than ten years and the first rheumatoid arthritis treatment in a new class of medicines known as Janus kinase (JAK) inhibitors. Unlike biologic therapies, which work outside the cell, Xeljanz targets the inflammation associated with rheumatoid arthritis from inside the cell. Specifically, Xeljanz inhibits the JAK pathways, which are signaling pathways inside cells that are used by pro-inflammatory cytokines (proteins that facilitate communication between cells). Approved in the U.S. for the treatment of adults with moderately to severely active rheumatoid arthritis who have had an inadequate response or intolerance to methotrexate, Xeljanz is currently under review by several regulatory agencies around the world, including in Europe and Japan.

Bosulif

Bosulif (bosutinib) is for the treatment of adult patients with chronic, accelerated, or blast phase Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) with resistance, or intolerance to prior therapy. Once-daily Bosulif was studied in a broad range of patients with CML, including CML patients treated with imatinib followed by a second-generation tyrosine kinase inhibitor. Bosulif addresses an unmet need in the CML treatment landscape.

Eliquis

Eliquis (apixaban) gained approval to reduce the risk of stroke and systemic embolism in patients with nonvalvular atrial fibrillation in the U.S., the 27 countries in the European Union, Canada, and Japan, and is under review in other countries. Eliquis, an oral Factor Xa inhibitor anticoagulant, has demonstrated superior risk reductions versus warfarin in three key outcomes of stroke and systemic embolism, major bleeding and all-cause death in patients with nonvalvular atrial fibrillation. In 2007, Pfizer and Bristol-Myers Squibb entered into a worldwide collaboration to develop and commercialize this oral anticoagulant discovered by Bristol-Myers Squibb.

Inlyta

Inlyta (axitinib) is indicated for the treatment of advanced kidney cancer when one prior drug treatment for this disease has not worked or has stopped working. It is the first treatment to demonstrate superior progression-free survival in a Phase III study compared with sorafenib in second-line advanced kidney cancer. Since its approval, Inlyta has established its utility in this setting, where it is an important treatment option for many patients with advanced renal cell carcinoma after failure of one prior systemic therapy.

Quillivant XR

Quillivant XR (methylphenidate hydrochloride) for extended-release oral suspension, CII, is the first once-daily liquid medication approved in the U.S. for the treatment of attention deficit/ hyperactivity disorder (ADHD). This recently approved medicine is expected to be available in pharmacies in the U.S. in early 2013. With the acquisition of NextWave Pharmaceuticals in 2012, Pfizer gained exclusive North American rights to market Quillivant XR.



Our Top 10 Best Selling Medicines in 2012

For more information on any of these medicines, visit
Pfizer Pharmaceutical Products.

Lyrica

(pregabalin)

\$4,158 million

Lipitor

(atorvastatin)

\$3,948 million

Enbrel

(Outside the U.S. and Canada)

(etanercept)

\$3,737 million

Prevnar 13/Prevenar 13

(pneumococcal polysaccharide conjugate vaccine)

\$3,718 million

Celebrex

(celecoxib)

\$2.719 million

Viagra

(sildenafil)

\$2,051 million

Norvasc

(amlodipine besylate)

\$1,349 million

Zyvox

(linezolid)

\$1,345 million

Sutent

(sunitinib malate)

\$1,236 million

Premarin Family

(conjugated estrogens)

\$1,073 million