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Evaluatepharma orphan drug report 2018

For a long time, pharmaceutical companies have viewed large populations of diseases as the largest possible revenue streams. But those days are long gone. This perception has changed, especially with the stagnation of the prescription drug market in the U.S. and Europe. The pharmaceutical treatment of rare diseases or disorders of orphan drugs has proven to be a viable money creator, and the industry has taken note. As many drug manufacturers have realized, a small group of patients can prove extremely profitable - at the right price. Rare drugs- sometimes godsend for those who suffer from life-threatening conditions that previously did not have treatment-are some of the highest prices for drugs out there. And while it's well documented that these drugs can bring serious income, EvaluatePharma in 2013. The Orphan Drugs Report reveals that orphan drugs now show a higher return on investment than products for larger patient pools. It is not difficult to see the development of orphaned drugs in the ureitu. Companies found in rare disease solutions face less competition, if any, from branded offers and generic competitors. In addition, the designation of orphan medicinal products offers an accelerated regulatory review process and a smaller late development. And with the global orphan drug market set to reach \$127 billion by 2018, accounting for nearly 16% of all prescription drug sales, it becomes another arena for some of pharmaceuticals' biggest breakthroughs in the fiercest patent wars. The list of the best sellers in 2018 includes treatments ranging from Roche (SRHHBY) Rituxan, a megablockbuster haunted by biosimilar manufacturers who have caught on the market appeal, with therapies such as Soliris of Alexion (\$ALXN) and Kalydeco of Vertex (\$VRTX), whose developers are trying to expand the populations of small patients that keep them afloat. Read on to learn how these 20 drugs and their makers will tell the story of an adaptive market over the next 5 years and feel free to share with us your thoughts on the next pharmaceutical wall. -- Carly Helfand (e-mail | Twitter) Lorem ipsum dolor sit amet, consectetur adipiscing elit. Suspendisse varius enim era elementum tristique. Duis cursus, mi quis viverra ornare, era dolor interdum nulla, ut commodo diam libero vitae erat. Aenean faucibus nibh et justo cursus id rutrum lorem imperdiet. Nunc ut sem vitae risus tristique posuere. EvaluatePharma recently released its orphan drug market current situation in its orphan drug report in 2018. The Orphan Medicinal Products Act promotes the development of medicines used for the treatment of rare diseases, as such medicines have not been considered financially sustainable low number of patients with sick patients. In the United States, rare orphan medicines for rare diseases are manufactured to treat less than 200,000 patients. According to this year's report, sales of orphan medicines are expected to rise by 11% per year by 2024, compared with 6.4% of the overall growth in the pharmaceutical market in the same period, the report says. Celgene has won the distinction of the pharmaceutical company, which has the highest sales of orphaned drugs due to lenalidomide (Revlimid) due to its multiple myeloma indication, earning the company \$8.1 million in 2017 alone. By 2024, the drug is projected to reach \$12 billion in sales worldwide. Although Celgene produced the most sales from its orphan drugs, Alexion Pharmaceuticals retained the name of the most expensive orphan drug, eculizumab (Soliris), for paroxysmal nocturnal hemoglobinuria and myasthenia gravis treatments. Estimatepharma's project to reach \$5.3 billion in sales by 2024. Amgen is currently developing a biosimilar candidate for eculizumab. In addition, the report found that the average price of the top 100 orphans in the United States (as ranked by sales) grew by 5.2% year-on-year between 2013 and 2017, compared with 9.2% for the top 100 non-orphanage drugs. Finally, EvaluatePharma estimates that oncology will be the leading therapeutic area for orphan drugs in 2024, accounting for about 50% of the total market in 2024 worldwide. The second leading area is blood disorders, which are believed to account for about 12% of the market. By 2024, rare drugs are projected to account for one-fifth of the world's prescription drug sales, reaching nearly \$262 billion. The FDA's orphan drug modernization plan has increased the number of orphan names in the United States. Overall, the FDA has given more orphan drug names than any other regulatory body, read the report. LONDON & BOSTON & TOKYO--(BUSINESS WIRE)--May 29, 2018--Continuing unmet medical demand is set to fuel the growth of the global orphan drug market, with sales projected to grow more than 11% year-on-year to \$262 billion by 2024. This year's report shows that every year, the increase in rare drugs can be ordered. However, the average price per patient per year in 2017 still managed to reach \$147,308, more than four times the average cost of non-orphanage drugs of \$30,708. While the overall 2024 revenue forecast for orphan drugs shows that growth is twice as high as that of non-orphaned drugs, if you look at the current R&D forecasts for orphaned medicines, they show similar cagrs of about +132%, said Dr. Karen Pomeranz, lead analyst and report author. It is possible that increased price pressures contribute to this balance. However, other factors are likely to as well as, for example, the expected level of penetration and allocation and the differences in the number of products expected to be released in each segment. Rate Orphan Drugs Report 2018 Highlights: Evaluate the Orphanage Drug Report 2018, based on commercial exploration and consensus predictions from evaluatepharma®. The report can be downloaded from www.evaluategroup.com/OrphanDrug2018### About Evaluate Ltd. Evaluate is a reliable provider of commercial intelligence, including product sales and consensus forecasts for commercial teams and their advisors in the global life sciences industry by 2024. We help our customers make high-value decisions through the highest quality, timely, must-have data and insights, along with personalized, expert customer service. Our online subscription services include the pharmaceutical, biotechnology and medtech sectors. Our Custom Services Group provides design analytical and data services. EP Vantage, our independent, award-winning editorial team, offers data-driven, forward-looking news, comments and analysis on a daily basis. For more information, visit: www.evaluategroup.com. Twitter: [@evaluatepharma](https://twitter.com/evaluatepharma), [@evaluatemedtech](https://twitter.com/evaluatemedtech), [@evaluateJP](https://twitter.com/evaluateJP)@epclinicaltrial. @epvantage. View the source version of [businesswire.com](https://www.businesswire.com): CONTACT: Evaluate and EP Vantage (General Questions and U.S. Media)Jennifer Dinkel+1 617-936-7783jennifer.dinkel@evaluategroup.comorInstinctif Partners(non-US. Media)Gemma Harris+44 (0) 20 7457 2020 evaluate@instinctif.comKEYWORD: UNITED KINGDOM UNITED STATES EUROPE ASIA PACIFIC NORTH AMERICA MASSACHUSETTS JAPAN INDUSTRIAL KEYWORD: HEALTH BIOTECHNOLOGY PHARMACEUTICAL PROFESSIONALS SERVICES OTHER PROFESSIONAL SERVICE COMMUNICATION RESEARCH OTHER COMMUNICATION SCIENCE SOURCE: RateCopyright Business Wire 2018. BAR: 05/29/2018 04:47 AM/DISC: 05/29/2018 04:47 AM May 29, 2018 04:47 AM Eastern Daylight Time LONDON & BOSTON & TOKYO--(BUSINESS WIRE)--Continuing unmet medical demand is set to fuel the growth of the global orphan drug market, with sales projected to grow by more than 11% per year to \$262 billion by 2024. However, while there is a strong demand for new therapies, the price that orphan drug makers can lead their products can be squeezed. This year's report shows that every year, the increase in rare drugs can be ordered. However, the average price per patient per year in 2017 still managed to reach \$147,308, more than four times the average price of \$30,708 for non-orphans. While the overall 2024 income forecast for orphan medicines shows growth twice as much as non-orphans, if you look at the forecasts for orphans and non-orphans currently under H&D, they show approximately +132% of CAGRs, said Karen Pomeranz, PhD student and lead analyst and author of the report. It is possible that an increase in pressure contributes to this balance. However, there are other factors, such as the expected level of penetration and prescribing, as well as differences in the number of products expected to be released in each segment. Evaluate the Orphan Drugs Report 2018 Highlights: The average price of the top 100 orphan drugs in the U.S. (ranked by sales) grew by 5.2% year-on-year between 2013 and 2017, compared with 9.2% for the top 100 non-orphaned drugs. Revlimid (lenalidomide) is projected to be the US's best-selling orphan drug in 2024. Soliris (eculizumab) generated the highest U.S. income for orphan drugs per patient in 2017. Expected strong sales of Revlimid and Pomalyst (pomalidomide) will ensure Celgene's position as a global leader in orphan drug sales in 2024. Alexion Pharmaceuticals leads the global market for orphan drugs for blood indications. Oncology is a leading area of orphan drug therapy in 2024, accounting for about 50% of the global market. Blood is the second leading area of therapy, accounting for about 12% of the market in 2024. Lanadelumab (Shire) is the most valuable rare drug R&D, with a net present value of \$7.48bn. The FDA's orphan drug modernization plan has increased the number of orphan drug prescribes granted in the US. Overall, the FDA has given more orphan drug names than any other regulatory body. Evaluate the 2018 Orphan Medicine Report based on EvaluatePharma and®. The report can be downloaded from www.evaluategroup.com/OrphanDrug2018### About Estimate Ltd. Estimate is a reliable provider of commercial intelligence, including product sales and consensus forecasts for commercial teams and their advisors in the global life sciences industry by 2024. 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