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Global Duchenne Muscular Dystrophy Market Survey and Trend Research 2018

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Abstracts

Summary

Duchenne muscular dystrophy is an x-linked genetic disorder that affects mostly boys. In Duchenne, boys begin to show signs of muscle weakness as early as two to five years of age. The disease gradually weakens the skeletal or voluntary muscles in the arms, legs and trunk. Due to progressive muscle weakness, Duchenne patients are often wheelchair bound between the ages of seven and 13 years old. At a later stage, the boys' respiratory and cardiac muscles are also affected and for most boys, respiratory and cardiac failure are major causes of death, often prevalent by the age of 20.

This report describes the development of the industry by upstream & downstream, industry overall and development, key companies, as well as type segment & market application and so on, and makes a scientific prediction for the development industry prospects on the basis of analysis, finally, analyzes opportunities for investment in the industry at the end of the report.

Industry Chain

Raw Materials

Cost

Technology

Consumer Preference

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Industry Overall:

- History
- Development & Trend
- Market Competition
- Trade Overview
- Policy

Region (North America, Europe, Asia-Pacific, South America, Middle East, Africa):

- Regional Market
- Production Development
- Sales
- Regional Trade
- Regional Forecast

Company (PTC Therapeutics, Sarepta Therapeutics, Bristol-Myers Squibb, ITALFARMACO, Daiichi Sankyo, Daiichi Sankyo, Solid Biosciences, Summit Therapeutics, FibroGen, NS Pharma, Pfizer, ReveraGen BioPharma, Wave Life, Genethon, Santhera Pharmaceuticals etc.):

- Company Profile
- Product & Service
- Business Operation Data

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