

Global Duchenne Muscular Dystrophy Market Research Report 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.

Market Segment as follows:

By Region

Asia-Pacific

North America

Europe

South America

Middle East & Africa



	By Type	
	Deflazacort	
	Prednisone	
	Others	
By Application		
	Male	
	Female	
By Company		
	PTC Therapeutics	
	Sarepta Therapeutics	
	Bristol-Myers Squibb	
	ITALFARMACO	
	BioMarin	
	Daiichi Sankyo	
	Solid Biosciences	
	Summit Therapeutics	
	FibroGen	
	NS Pharma	
	Pfizer	





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