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## ***An Ode to Duchenne Muscular Dystrophy***

There are two threads X and Y  
And on the X-thread, an error arrives  
Mutating the 'DMD' gene  
Causing a condition progressive and mean  
07 September is the day to spread  
awareness about the same  
A condition we know as Duchenne  
Muscular Dystrophy, by name

A young boy hasn't walked up to  
18 months  
Your radar needs to beep!  
Super-raised blood CPK may be the  
first clue  
We need to delve deep

EMG and gene test may follow the line  
To cross the first hurdle  
And detect this condition in time

Screen their mums early, not late!  
Be wary,  
A single mutant copy can dilate their hearts,  
And cause a second child with DMD  
A matter of unease!

Monitor the kids up-and- close  
Allow good physiotherapy,  
Delay the muscles getting contracted and sore  
Timely steroids and breathing exercises  
Flu shots and more...

And how can we not mention  
Exon skipping, ataluren, CRISPR-Cas9  
and gene therapies ...  
Oh! The magical world of genomic  
possibilities!  
Wingardium Leviosa, Abra-ca-dabra  
'HOPE', we clinch on to you!  
Why let their bodies be wheelchaired?  
Let their dreams fly far!

Spread the word,  
The condition isn't all-that-rare  
We all need to stand up for these kids  
And show that we indeed,  
Dare to Care!