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diagnosis and

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~~you probably DON'T  
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During the End of Life  
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Care of The Patient  
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the End-of-Life Period  
Respiratory failure  
associated with ALS is  
mainly due to  
respiratory muscle  
weakness. Respiratory  
symptoms usually  
develop late in the  
disease process and in

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conjunction with  
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direct effect on the lung

but the mechanical

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the patient with  
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sclerosis during the end  
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practitioners, neurology

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Loss of motor neurons  
in the cortex, brainstem  
and spinal cord is the  
hallmark of motor  
neuron  
disease / amyotrophic

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lateral sclerosis (MND / ALS), resulting in weakness of limbs, respiratory and bulbar muscles and eventually death from respiratory failure in the majority of patients. Around 20% – 30% have bulbar symptoms at onset—this is less common in younger patients, but affects more than 40 ...

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complications ...  
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Management Of ALS:  
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Amyotrophic lateral sclerosis (ALS) is a debilitating, progressive disease with degeneration of motor neurons in the brain and spinal cord causing weakness, muscle atrophy, fasciculations and spasticity.<sup>1</sup> Onset in

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the limbs, with  
extremity weakness and  
impairment in mobility,  
is the most common  
presentation, occurring  
in about 70% of  
patients.<sup>2</sup> Bulbar onset  
with oropharyngeal  
muscle ...

Canadian best practice  
recommendations for  
the management ...  
in amyotrophic lateral

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sclerosis also has no direct effect on the lung but the mechanical respiratory system is significantly involved also affects all the major muscle groups of the mechanical respiratory system

- 1 upper airway muscles abnormal swallowing and cough
- 2 expiratory muscles inadequate cough and
- 3 inspiratory muscles

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symptoms usually  
develop late in the  
disease process and in  
conjunction with  
extremity or bulbar  
muscle involvement  
other disease processes  
such as

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Hospice Nurses Journal  
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2008;10(4):214-215. Lee  
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Bartlett, 2009. Page 75

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