

This picture doesn't tell the whole story...

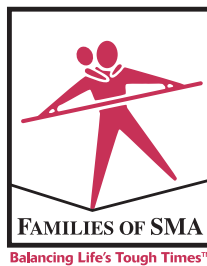
Spinal Muscular Atrophy

- 1 in 40 people are carriers.
- SMA robs individuals of their ability to control voluntary muscle movement.
- SMA has no age, gender, racial or religious boundaries.

There is no cure, but research provides hope.

This holiday season - give the gift of hope.

Although medical research is making incredible headway in the quest for a treatment and cure for spinal muscular atrophy, we're still losing children to SMA every day. Boys and girls throughout Canada and the world are losing their ability to crawl, walk, hold their head up, swallow and even breathe.



For more information visit www.curesma.ca or call **1-800-866-0016**

P.O. Box 22024, Brandon, Manitoba R7A 6Y9

To make a secure credit card donation please visit <http://www.canadahelps.org/search.aspx> and type "Atrophy" in the "Search terms:" field.

Give the gift of HOPE - make a donation to Families of SMA Canada.

Name: _____

Address: _____

City: _____ Province: _____ Postal: _____

email: _____

phone: _____



Thank you for your generosity!

I would like to make a donation of: \$25 \$50 \$100 \$250 \$other _____

Please make cheques payable to Families of SMA Canada.

Tax receipts will be issued for all donations over \$10.00