

Spina Bifida and Hydrocephalus Association of Southern Alberta

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We are delighted to extend our warmest welcome to you as a new member of our Spina Bifida and Hydrocephalus Association. As we strive to support individuals and families affected by Spina Bifida (and related hydrocephalus), it is important for us to maintain accurate records and ensure that our resources are effectively allocated to meet the needs of our members. To this end, we kindly request your cooperation in completing the following documentation.

Pls mail or scan a copy of this completed form to the address at the top of this form.

First & Last name of person with Spina Bifida / Hydrocephalus:

_____ DOB (dd/mm/yyyy): _____

I confirm that the above-named person has been a permanent resident residing in Southern Alberta since (dd/mm/yyyy) _____, residing at the following address (street address, city, postal code):

_____ Date of Spina Bifida & related Hydrocephalus Diagnosis (dd/mm/yyyy): _____

Parent / Legal Guardian / Adult with SB/H applying for membership (print full name):

Signature: _____ Date (dd/mm/yyyy): _____

Myelo Clinic Representative (print full name): _____

Signature: _____ Date: (dd/mm/yyyy): _____

Neurosurgeon or Physician confirming Spina Bifida a/o related Hydrocephalus diagnosis:

(print full name): _____

Office Contact #: _____

Signature: _____ Date: (dd/mm/yyyy): _____

* I acknowledge that any false information provided may impact the services and support that the association can offer. This form allows us to better tailor our programs and services to meet the specific needs of individuals living with Spina Bifida in Alberta a/o related Hydrocephalus in Southern Alberta.

Sincerely, Program Coordinator, SBHASA

Disclaimer *We can only support individuals with diagnosed Spina Bifida a/o related Hydrocephalus. People who have conditions like SB/H are welcome to apply for a Friends & Others Membership located on our website.

Our Mission: To improve the quality of life of all individuals with Spina Bifida a/o related Hydrocephalus and their families through awareness, advocacy, education, and support.