



Dear new member,

Thank you for contacting **CORPAL**. We are a small charity made up of parents with children affected by Agenesis of the Corpus Callosum and/or Aicardi Syndrome.

We hold our annual general meeting and conference in April/ May time, generally in the London area. On these occasions we invite speakers to talk on relevant issues concerning ACC. This event is always a good time to meet up with other families and share experiences. We also hold other regional meetings.

We generally send out newsletters at least twice a year. We would love to hear from you if you would like to include any information that could be of interest to our members, e.g. your experiences with ACC, forthcoming events, or even just a photo.

If you are interested in receiving information about the annual event and our newsletter, **please fill in the attached form** and return to us, if this is not received we cannot guarantee you will be notified.

We must stress the committee members **are not medically qualified**, therefore are not able to give comment on any medical issues, although we can help you by guiding you to find the right professional person who should be able to help you further.

We are here to offer you support and send out further information as it becomes available, please be sure to inform us of your **email address** if you have not already done so. Do not hesitate to contact us if you need to discuss anything further.

I hope you find the information pack useful.

Yours sincerely

All at Corpal.