

Hi there,

We're organizing a living with EPP webinar on Sunday, August 11th at 20.00 CET/14.00 EST.

Please share the information or this link with your members and contacts:

<https://www.rareconnect.org/en/community/porphyria/forum/topic/living-with-epp-webinar>

To access the webinar room on August 11th:

[http://eurordis.emea.acrobat.com/rare\\_disease\\_communities/](http://eurordis.emea.acrobat.com/rare_disease_communities/)

Enter as a Guest, no password required. Connect a headset, or turn up your speakers to hear people living with EPP and their families speak about their experiences.

If you have any questions, please let me know. If you are interested in telling your story on the webinar, we still have a space available.

Best,  
Rob