

HOW CAN YOU CONTRIBUTE?

Carers/family/support people - Share your story

We are always looking to share stories of positivity and growth.

Provided below is a guideline for you to write about your experience of supporting or 'caring' for someone who lives with borderline personality disorder (BPD).

Your story can be incredibly impactful and supportive for other people who may be experiencing challenges while supporting someone living with BPD. Share the 'gems of wisdom' you have learnt in your own journey. As your story and the story of your family member are interwoven it is absolutely essential that your 'someone' is aware that you are participating and has approved what you have written.

BEFORE YOU CONTRIBUTE:

Sharing your story safely is essential!

- You should have a good support structure available before you share your experiences – it can be confronting and attract attention you might not expect or be able to control. We highly suggest having a mental health professional you can work with available if needed.
- Many people are negatively affected by key words and themes. These might include experiences of child abuse, sexual assault, self-harm and suicide. Whilst it is your right to acknowledge them, it is not appropriate in our spaces to detail or recount them as stories. This is unhelpful to your healing and the healing of others. These specific stories are best expressed in safe and secure environments with trained mental health professions who know how to respond and validate them. Hearing about detailed events of abuse and pain of others can cause distress to others and yourself. Re-experiencing trauma is common for people sharing difficult events in their lives – we need to keep everyone safe, especially you.
- Your contribution when told safely is important, valid, and impactful. Making sure that your contribution is fully heard is key, and this is why we suggest describing the feelings and actions you took to overcome your challenges as opposed to what held you back or hurt you. This is the information that others are lacking and need to hear to heal.
- Our audience is largely people with lived experience, their carers and supporters and those who work with people living with BPD – It is important that your contribution allows everyone to find solutions to challenges and provide deeper insight into what helps. Many people with lived

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experience know what 'bad treatment' looks and feels like – what they might not know is what 'good treatment' is like – share what worked!

- By sharing your valuable experience, you will be supporting others and they may want to contact you about this. Make sure you have strong safe boundaries, professional support and let us know if this becomes an issue.
- We may ask you to re-structure or focus your content to make sure your story is effective and strong. Your contribution feedback will come from people with extensive experience in lived experience advocacy, so please keep in mind that they have a strong knowledge of the complexities of sharing and are trying to support you.
- If you are feeling unsure, you can contact our team and speak to one of our experienced consumer or carer advocates, we are here to support you to have a voice.

Format for sharing your story:

- Word document or PDF
- Up to 500 words
- Please refrain from using profane language, naming family members, friends, clinicians or places of treatment/services/hospitals.
- Please remember to not include graphic details or traumatic or distressing experiences.

Things you need to know before sharing

- We will only share your first name, and how you identify. For example: Mahlie (Lived Experience Advocate). Alternatively, we can use a pseudonym.
- Nothing ever goes away on the internet – even if we try to make it go away, we have little control. We also cannot force other agencies or websites to remove or unshare your information once it is placed in the public domain.
- Your story may be shared and re-shared – and we cannot monitor this!
- Your story is about you and should not mention other people in your life – if you are a carer this means you must not identify the person you support *without their expressed permission*. As a carer/support person please consider how you identify yourself to not breach the privacy of the person you are speaking about.
- You will be asked to sign a disclaimer form before your content is shared and you have the final say on the content's editing.
- You have rights and should always use them when needed! (Contact the Foundation for more information.)
- To read an additional great resource about telling your story – visit <https://wayahead.org.au/telling-your-story/>

These questions are suggestions to guide your writing:

1. Was receiving a diagnosis positive for the person you care for?
Was it positive for you/other family?
2. What could have made the experience of receiving a diagnosis better?
3. What would you like carers who are struggling to accept the diagnosis of BPD to know?
4. What would you like carers who are facing challenges to support someone with BPD and feeling overwhelmed to know?
5. What good things did clinicians and mental health professionals do to support you?
6. Are you receiving support in your caring role?
7. What support did you receive/ access?
8. What can clinicians and mental health professionals offer to better support you?
9. What self-care strategies work for you?
10. With the wisdom of hindsight what 'words of wisdom' would you offer carers struggling to support someone with BPD?

The person I support has given me their permission to share,

Disclaimer:

I, [full name], confirm that all information provided in this submission is true and accurate to the best of my knowledge. I have obtained the consent of any third parties discussed therein. I indemnify the Australian BPD Foundation ('the Foundation' or 'us') for any loss, damage or claims arising in relation to this article, whether published by us or any other party, including third parties. I acknowledge and accept that the Foundation is under no obligation to publish my submission or any part of it.

SignatureDate...../...../.....

Email:.....

Phone:.....

