How to talk to friends and family

It can be really tough when you are faced with an unknown condition such as an adverse effect from surgery or a medical device – It can feel very lonely and difficult to talk to friends and family about. Especially if it includes the word Vagina, as it does for some of us. However, it is important that our loved ones are informed so that we can reduce our isolation by sharing our experience.

We hope this leaflet you can print off may help.

you for trying to support me.

am suffering from Mesh injury as a result of a faulty medical device.

I had surgery for health reasons and have recently discovered that it has been adversely affecting my health for a while now in a number of ways.....

Please describe your pain or illness on the left, write your range of feelings accompanying the shock, or tick the boxes below that best explain your complications:

I have ongoing pain Hurts to go to the toilet I'm embarrassed I feel isolated and alone I feel like I am a burden It has been a huge shock to me to discover this and I am struggling I am depressed to come to terms with it. I am one of many men and women I am worried about the future Internationally who have been adversely affected by surgical mesh. I can't work because I am in too It has been used in some hernia repairs and for women suffering much pain I can't walk far or sit for long perifrom pelvic organ prolapses (POP) and stress urinary incontinence ods because of the pain (SUI), both are common after childbirth. П It hurts to have sex I am anxious about having further Sadly it is not an illness like Cancer where I have an official diagnosis surgery - where it is a condition that is well understood and acknowledged I am anxious about getting worse publicly, or where treatments options are readily available and I am scared accessible. It is an adverse reaction from a medical device made I have to get up to go to the toilet all from polypropylene and it is coming to light that this material ought the time not to have been used permanently in human bodies. It would seem I suffer from chronic urinary tract that the devices were not appropriately regulated and some were often misused and/or misplaced. I am feeling isolated and alone I am scared I will lose you as a since I found this out and find it tough to talk about. friend / partner As this is a sensitive subject for me I ask that if you would like further I can't control my bladder information about mesh injury, that you can visit the website I am relieved that I am not the only one this has happened to. meshinjured.info they have a friends & family support service. Thank