



1



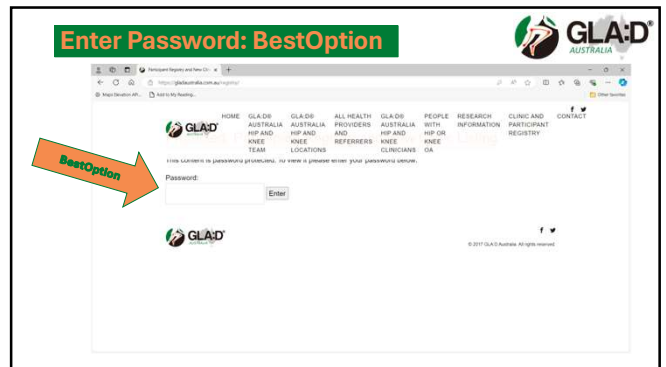
2

Go to the GLA:D Australia website, www.gladaustralia.com.au



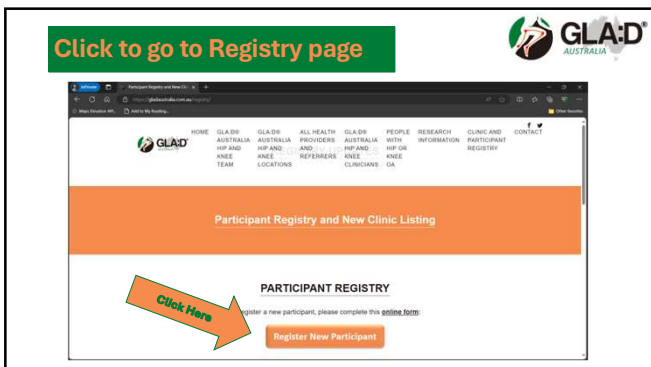
3

Click the tab labelled 'Clinic and Participant Registry'



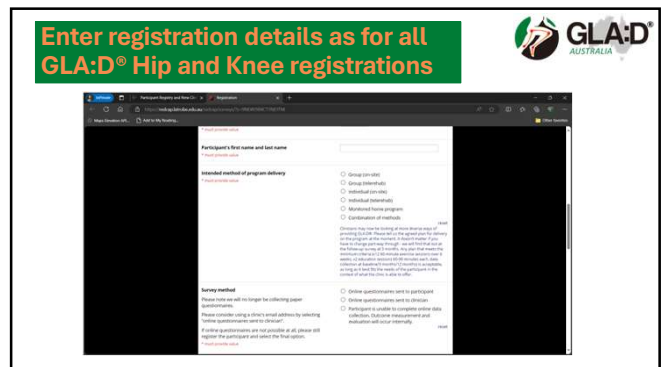
4

Use the password 'BestOption' to access the registry page



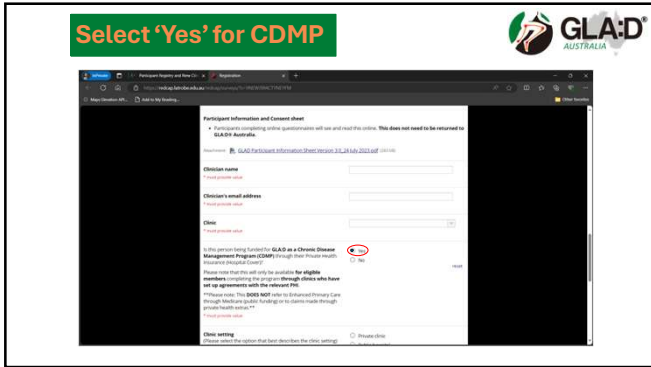
5

Click the button to 'Register New Participant'

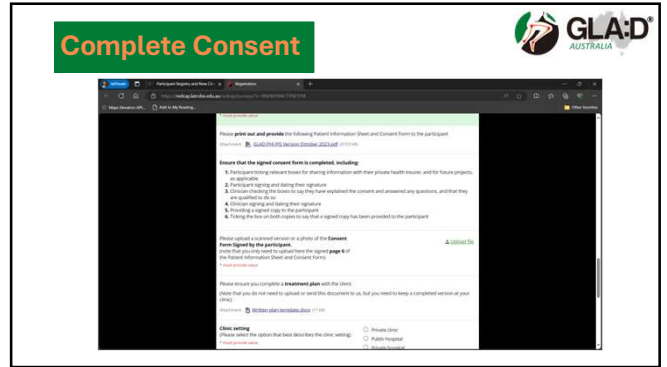


6

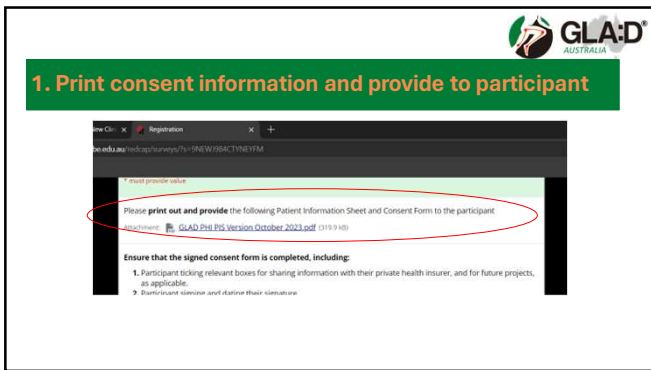
Complete the registration form, just as you would for any GLA:D Hip and Knee participant registration



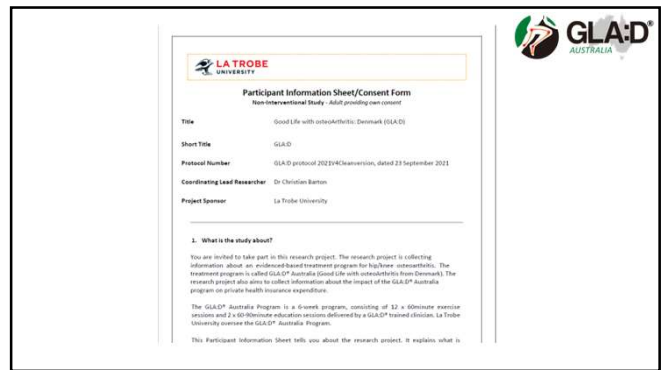
7 When you select 'yes' to the CDMF question, some further fields will open up below which will help complete the consent



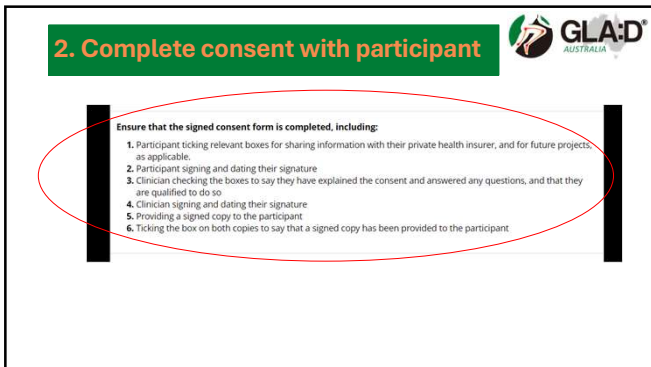
8 First, download and print the form. If you have copies already available, make sure it is the current version of the information sheet and consent form.



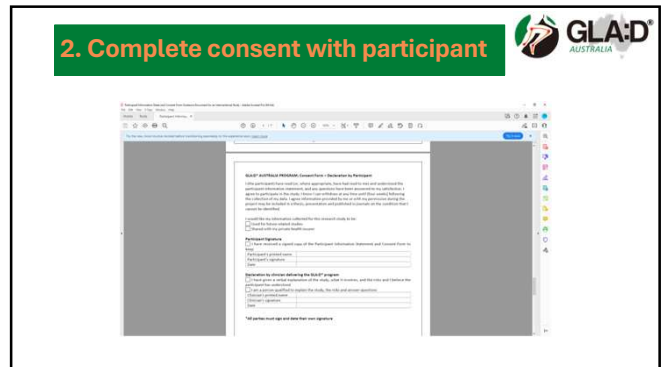
9 Give the form to the participant to read; the clinician is responsible for making sure the participant has read it or had it read to them, and that they understand the information.



10 This is what the front of the Participant Information Sheet and Consent Form looks like. See the end for further information.



11 The next step is to make sure that all the relevant fields in the form are completed:



12 The signature page is page 6.

13

First, the participant needs to select whether they consent to sharing their data with their insurer, and/or being used for future studies. They need to write their name, sign and date. A copy must be provided to the participant, and the relevant box for that ticked to confirm this has happened.

14

Then the clinician completes their declaration - that they have explained the information to the participant, and that they are the right person to do so. They also need to write their name, sign and date.

15

Only page 6, with the signatures, needs to be uploaded. Don't forget to give a copy to the participant. Do not upload withdrawal of consent forms here. If the participant doesn't consent to having their name confirmed with the insurer, they should indicate that on the form. If they do not consent to sharing their name with GLA:D Australia, do not register them.

16

There is a prompt in the registration form to complete a written treatment plan as part of meeting your provider obligations, and the template is there for your to download if needed. Keep that for your own records - don't provide it to us.

The Participant Information Sheet and Consent Form outlines the GLA:D Program and all that's involved with it, just like the Participant Information Sheet for the main GLA:D Program. It has some extra content that covers what else will happen to CDM participants' data; that is, that GLA:D Australia will confirm their name with their private health insurer, and that we will provide their de-identified aggregated data (data that can't identify them in any way) as part of reports to the private health insurer. This private health insurer will use that information to facilitate payment of clinicians, and to help inform decisions about continuing this type of funding. Providing consent by signing this form will give the participant's name to GLA:D Australia, and then to their health insurer. Once they receive their baseline questionnaire, they can still decide whether or not to provide GLA:D Australia with their data through the questionnaires.