



One of our objectives is to campaign for improved services and standards of care for people with lymphoedema, including diagnosis and treatment. To strengthen our case it would be most helpful if we can show the prevalence and severity of lymphoedema among our members as an indication of the ongoing effect on them. We have the information sought below for a minority of our members and ask you to PLEASE reply to the questions below and return it to us a week before the open day if possible. Please be assured that we fully respect the requirements of the Data Protection Act and that any use we make of the data will be in the form of totals and percentages and will not be associated with any particular member.

Form with fields for personal details (Mr/Mrs/Ms, Address, First name, Surname, Sex, Date of birth, Home phone, Mobile phone, Membership number, Email), family history, diagnosis method, affected areas, severity, level of disability, hosiery, and therapist/treatment information.