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- Leaving a Legacy to The Scleroderma Society
- Localised Scleroderma
- Physiotherapy & Scleroderma
- Pulmonary Hypertension in Scleroderma
- Scleroderma & the Fingers
- Scleroderma & the Foot
- Scleroderma of the Heart
- Scleroderma in Kidneys
- Scleroderma of the Lung
- Scleroderma, Not Just Hard Skin
- Scleroderma Oral Problems
- Scleroderma & Pregnancy
- Scleroderma in Young People
- Sexual Problems in Women with Scleroderma
- The Gastro-Intestinal Tract
- The Skin in Scleroderma

Order our free booklet

(Please tick or enter number of copies required)

- Understanding & Managing Scleroderma

Federation of European Scleroderma Associations (FESCA)

The Scleroderma Society is a founder member of FESCA and this enables us to establish and maintain contact and co-operate with organisations like ours from all over Europe. We participate in European Scleroderma Awareness Day, which takes place every year on 29th June.

Visit www.fesca-scleroderma.eu for further information.

Data Protection

The information you have provided will be held securely on a computer system and will be used by the society to keep in contact with you. The Scleroderma Society is registered with the Data Commissioner for the purposes of Data Protection. The trustees will not pass any information you have provided on this form to any third party, or to any other member without your express agreement. If you wish to join our scheme for putting members in contact, we will send you the necessary agreement which you will need to sign.

The Scleroderma Society is a company limited by guarantee and not having a share capital. Registered in England, company no. 01704872. Trustees K Fligelstone, S Saunderson, S Hoare, A Thorpe, R Dodds, S Holloway (secretary). Registered address 37 Warren Street, London W1T 6AD



Membership Application Form

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The Scleroderma Society

Founded in 1982 to help people with scleroderma and their families, the society came about through the initiative of doctors who specialise in this field and because of their concern that, since the condition is comparatively rare, there are only a small number of these people in any one area.

The Society is based in Chichester, with local groups in areas where there is sufficient interest and membership to support one.

What we do

- Support people with scleroderma
- Increase awareness of scleroderma among the medical profession & general public to improve early diagnosis & prognosis
- Fund medical & scientific research

Why become a member?

- Contact other people with scleroderma and share experiences
- Learn more about scleroderma from our leaflets, booklet and newsletter
- Use our 0800 telephone helpline
- Take part in our web forum and discuss scleroderma with people from all over the world
- Get involved with fundraising to help us fund research
- Become a volunteer helper
- Join a local group
- Attend our annual conference

How much does it cost?

UK Membership costs £5.00 per annum or £20.00 for 5 years. Membership renewal is due on 1st April and we will send a reminder.

Life membership is available for a one-off payment of £100.00.

How can we contact you?

Title: *(Mrs. Mr, Miss, Ms, Dr, Other)* _____

Forename: _____

Surname: _____

Address: _____

Postcode: _____

Telephone: _____

Email: _____

These statements are optional

(Please tick)

- I have scleroderma
- I would like to be in contact with other members
- I would like to receive the newsletter by email

You must sign this guarantee

I guarantee to pay a sum of not more than £1 in the event of the Society winding up with debts while I am still a member, or within one year after I cease to be a member, for payment of the debts and liabilities of the Society contracted before I ceased to be a member.

Signature: _____ Date: _____

Please select your payment

(Please tick)

- £5.00 for 1 year (*£10.00 overseas*)
- £20.00 for 5 years (*£40.00 overseas*)
- £100.00 Life membership (*£200.00 overseas*)
- I include a donation of £_____ (*optional*)

Please Gift Aid your payment

If you are a UK tax payer and you would like us to claim Gift Aid on your payment, please ensure you have completed your name, address and postcode above and tick this declaration.

- I am a UK tax payer. Please claim Gift Aid on my payment.

How to make your payment

(Please tick one)

- Pay by cheque made out to The Scleroderma Society
- Pay online using a credit or debit card
(See the link on our website, www.sclerodermasociety.co.uk)
- Pay by BACS direct to our account from your online bank account
- Pay by Annual Standing Order

Send your completed form & cheque to

The Secretary
The Scleroderma Society
PO Box 581
Chichester
PO19 9EW

Our bank details for Standing Order & BACS payments

CAF Bank Ltd
Account No: 00016089
Sort Code: 40-52-40

Please use your name as the reference so we can identify the payment.

Information you may wish to receive

We do not pass any of your contact information to other organisations, but occasionally we receive information from other organisations, for example patient support groups in other countries, which we think may be of interest to you. If you would like to receive this carefully selected information from us, please tick here