

Hello!



Piel de Mariposa
Butterfly Children

Martina (Huelva)

Photograph Campaign: Osos Preciosos de Fundación Carrefour



**Their skin is
as fragile as a butterfly's wing**



Butterfly Skin is a **rare disease** causing **extreme fragility of the skin**.
The fault lies in a **genetic mutation** that fails to “glue” the layers of the skin together meaning that **the slightest touch can cause a blister**. Even a simple embrace can cause harm.

Sadly, to date there is **no cure**.



**Take a moment to experience
the life of a Butterfly Child**

From birth **blisters and wounds continuously appear** all over their bodies.

As part of the treatment and as a form of protection the children have to undergo

long and painful bandaging routines every second day. In some cases management can be more complicated because the mucous membranes of the mouth, pharynx, oesophagus and eyelids can also be affected.

Everyday activities such as dressing, walking and eating can be challenging causing a high level of

disability and dependency.

The psychological effects of Butterfly Skin impact on **the whole family.**

Feelings of guilt, helplessness, anger and sadness can lead to depression, anxiety, fear and low self esteem.



**Supporting families
since 1993**

The **DEBTRA The Butterfly Children Charity** is an official **non profit and Public Interest Organisation**.

It was founded in Marbella by the parents of an affected baby. Their main aim was to **help and support other families** in the same situation. Twenty three years, our mission remains the same:

To improve the quality of lives for Butterfly Children and their families

Today we represent **more than 260 families** and consist of a team of **29 professionals**, 10 charity shops and more than 300 volunteers all inspired by the same mission.



Who is who?

Head office team



Manager



Nurses



Social Workers



Psychologists

Communications and Fundraising



Charity Shops



Admin Accounts





New-born Visits

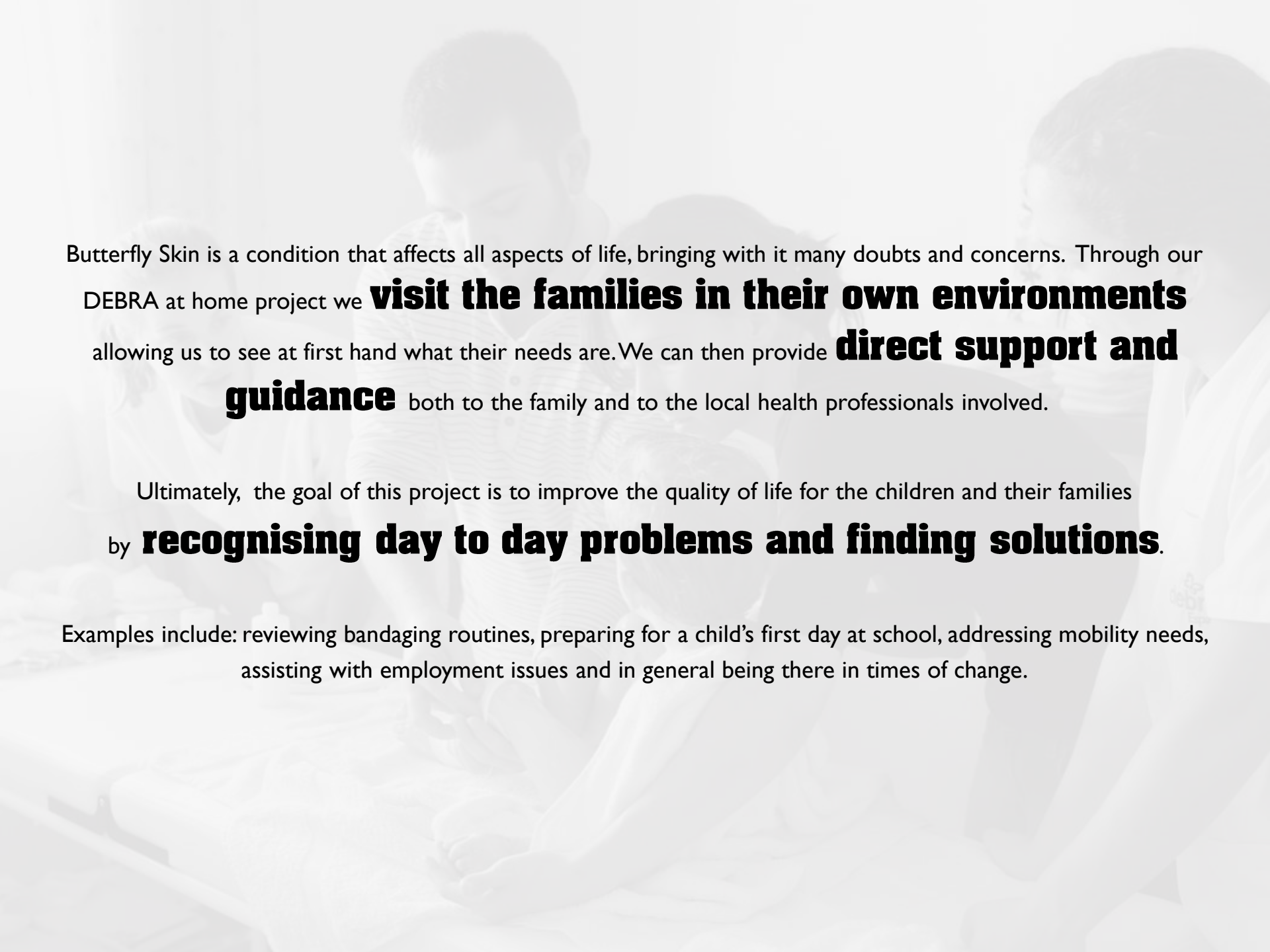
1 in every 227 people carry the faulty gene for Butterfly Skin unknowingly so the condition can arise in any family unexpectedly.

The birth of a child with Butterfly Skin is one of the **most traumatic situations** that a family can be faced with. During the first days and weeks the parents live with feelings of **disappointment, loneliness, uncertainty and fear.**

Our work at this time is highly influential. Through the expertise and experience of our Health Professional Team we aim to ease these feelings by **educating the family and their local health professionals** about the condition. This aids their understanding and assures them they are **not alone.**



DEBRA at home



Butterfly Skin is a condition that affects all aspects of life, bringing with it many doubts and concerns. Through our DEBRA at home project we **visit the families in their own environments** allowing us to see at first hand what their needs are. We can then provide **direct support and guidance** both to the family and to the local health professionals involved.

Ultimately, the goal of this project is to improve the quality of life for the children and their families by **recognising day to day problems and finding solutions.**

Examples include: reviewing bandaging routines, preparing for a child's first day at school, addressing mobility needs, assisting with employment issues and in general being there in times of change.

Dr. Raúl de Lucas
Head of Paediatric Dermatology



La Paz Hospital

Due to the rare nature of Butterfly Skin very few health and social professionals have the knowledge and experience of how to treat it.

For the last 2 years **our team has had a strong presence within the dermatology clinic** of La Paz Hospital in Madrid improving the level of care available and supporting its role as a specialist centre.

This year we are very pleased to announce that we have **extended this collaboration** by appointing one of our **nurses permanently to the clinic**. The other members of the team will continue to play a strategic role by visiting once a month to supplement the support, especially for the psychological requirements of the families.

We would like to replicate this experience in hospitals throughout Spain and have already started collaborations with hospitals in Seville and Barcelona and hope to see progress throughout this year.



National Meeting of Families



Every year we **unite families from all over Spain** in a “National Meeting”
an initiative that has been happening since the charity was created over 25 years ago.

“Mummy look, **I’m not the only alien!**” said one of Butterfly Children at his first National Meeting after seeing other children with the same condition .This comment alone reflects the importance of this project.

The meetings give the families the **opportunity to share experiences and learn** from each other as well as the specialist health professionals. It is a significant date in the diary for all those involved with the condition.



Estar Bien

Butterfly Skin is a **chronic condition** that affects the child for life. Our team of specialist health professionals are **always available** to the families and their local health professionals **to help them cope with the difficulties** that arise along the different stages of life.

Examples include:

HEALTH

- Advice on materials and **methods of bandaging.**
- Recommending local and regional professionals with experience.
- Supporting the genetic diagnostic process.

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SOCIAL

- Assessing and assisting with **disability and dependency.**
- Advice and assistance with the financial benefits process.
- Addressing schooling and employment issues.

...

PSYCHOLOGICAL

- Help with the initial **impact of the diagnosis.**
- Coping mechanisms for the individual and the family.
- Support for the main carers.

...



**Collaborations
and synergies**

We are **members of**

and we **collaborate with**



Internationally we are:

Members of the European Organisation of Rare Diseases **EURORDIS**.

Founders and members of the International network “**DEBRA International**”.

Members of the International network for the Health Professionals of Butterfly Skin “**EB Clinet**”.

In addition, we also work with other countries who do not have an association in place. We help them to create their own DEBRA especially the Latin American countries.





Our Claims

Our claims concentrate on **3 basic requirements** of life in the developed world:

1

Access to **specialised health care** through the designation of Reference Hospitals. Here, care from highly experienced professionals will be given regardless of their postcode.

2

For many years we have been campaigning for **free and easy access to bandaging materials**. Last year we achieved the first step towards this by receiving an agreement from the government and local health authorities to guarantee this right. However we know that not all the families are benefiting from it yet, so this claim remains open.

3

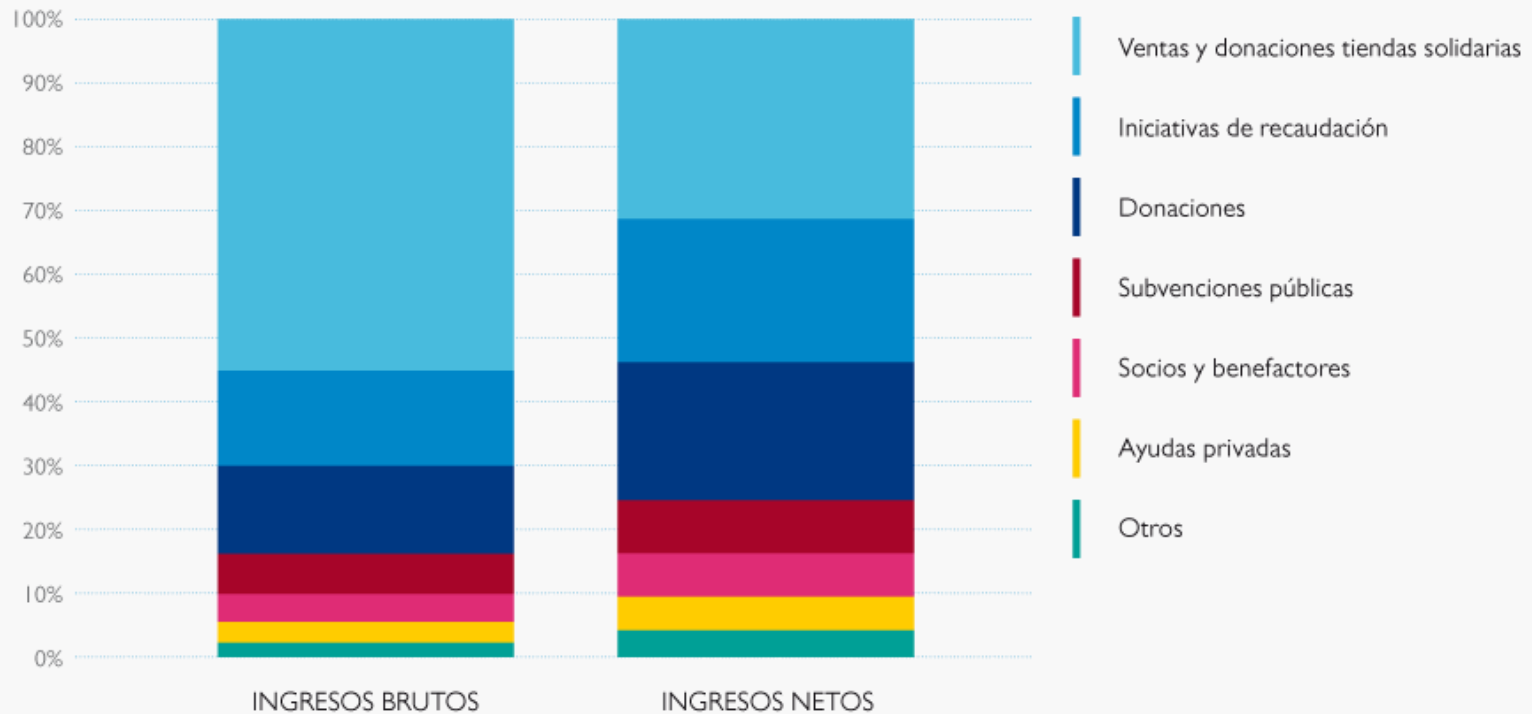
Families face problems such as disability, dependency, social exclusion and financial difficulties. These do not adhere to the social benefits criteria that exist today therefore the **social support and resources** available need to adapt accordingly.

Our projects are in place to compensate for the failings of the Public Health System. Once these have been resolved we can then dedicate more resources to areas such as research and awareness.

A large number of monarch butterflies are scattered across a clear, bright blue sky. The butterflies are in various stages of flight, with some showing their characteristic orange and black wings. The overall scene is one of natural beauty and movement.

Your help gives us wings

To carry out our work comprehensively we rely on the **generosity of individuals and companies** as well as our own fundraising methods such as our **charity shops**.



*Sources of Income 2016

Less than 10% of our funds come from the government.

Whether you are a **company or** an **individual** there are many ways that **you can collaborate** to help build better lives for the Butterfly Children and their families.

Here are some examples...and we always welcome news ideas!

Become a Benefactor

**Visit or donate to a
charity shop**

**Take part in one of
our events**

Organise an event

Finance a project

**Donate €1/month
Teaming.net**

**Advertise in our
publications**

La Risa es el Alimento del Alma



**Our shops
give us wings**



The **Butterfly Children Charity Shops** sell high quality items at incredible prices. All funds raised support the projects and services of the charity. You can **collaborate by donating, volunteering or shopping** in your local shop.



Supporting the Butterfly Children Charity Shops you are helping the families living with Butterfly Skin.



Who is who? Charity Shops



Jorge



Alberto

Madrid Lavapiés

C/ Embajadores 35, Local 5
28012 Lavapiés (Madrid)
911 960 539
lavapiésmadrid@pieldemariposa.es



Belén

Madrid CC Alcalá Norte

CC Alcalá Norte. C/ Alcalá 414, Local 24
28027 Madrid
914 071 320
madrid@pieldemariposa.es



Ana

Sevilla

C/ Feria, 145
41002 Sevilla
609 569 052
sevilla@pieldemariposa.es



Antonia

Marbella San Pedro de Alcántara

Avda. Pablo Ruiz Picasso 46
29670 San Pedro de Alcántara (Marbella)
637 744 217 / 686 962 422
sanpedro@pieldemariposa.es



Eileen

Marbella Benavente

C/ Jacinto Benavente 12
29601 Marbella (Málaga)
658 464 227
marbella@pieldemariposa.es

Piel de Mariposa

Tienda solidaria Charity shop



Barcelona

Carrer d'Olla 100
08012 Barcelona
626 56 21 44
barcelona@pieldemariposa.es



Núria

Valencia

C/ Dolores Marqués 30
46020 Benimaçlet (Valencia)
963 616 963
valencia@pieldemariposa.es



Ana Lucía



Fina

Valencia Patraix

C/ Escultor Vicente Rodilla, 5
Bajo 2 (.dcha) 46007 Valencia
627 803 078
patraixvalencia@pieldemariposa.es



Lucía

Urbanización La Marina

C/ Luis de Góngora, local 37, Urb. La Marina
03177 San Fulgencio (Alicante)
616 076 072
lamarina@debra.es



Mary

Rojales

Urb. Pueblo Bravo
03170 Rojales (Alicante)
966 718 134
rojales@debra.es



Anne

Málaga

C/ Héroe de Sostoa 80
29002 Málaga
655 667 669
malaga@pieldemariposa.es



Carmen

Your help gives us Wings
Thank you!





www.butterflychildrencharity.com

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