

My Child has Atopic Dermatitis

I am depressed and without hope, because it is so sad to see my baby suffer without knowing how to solve this problem.

My child doesn't understand why she has it.

When my son's flares are bad, we can't fully live our lives worrying about what he's touching, eating etc.

I avoid taking my son to play dates or out in public. We get stares and I feel like a neglectful mother as people pass judgement on my son.

I'm frustrated that our allergist, dermatologist and GP all have different theories as to why my son has it.

I can't always afford the medications for my daughter because my insurance does not cover them.



It's so very stressful. Every flare comes with anxiety over the immunosuppressant medication I'm using on my child at such a young age.

Always seems like we are fighting a losing battle. Prescription medications work only for the days we use them. If we go a few days with none, flare-ups come back quickly.