

## Modelling – "Bald is Beautiful"

Niamh Woods

So, I was asked to model for the company Ally\_alopecia, sponsored by BooHoo! It was amazing working with all these beautiful girls and hearing their stories. It was also a very nerve-racking experience, it was the first time I had ever stepped outside without my wig on.

I was born with a condition called ectodermal dysplasia which caused my alopecia. My skin disorder has been its own rocky journey but right now I am just going to talk about hair loss.

I always knew I was going to lose my hair eventually and I always avoided the topic and I used to tell myself that it wasn't even happening to the point I was in complete denial even though I had massive obvious bald patches. It had got to the point where I was scared of wearing wigs because it would be accepting the fact that it was actually happening, and it would be such a massive change that everyone would realise. However, in year 11 I started to think about wearing wigs because prom was coming up and I wanted to look nice.



The day I wore my first wig was a scary day, but it improved my confidence in so many ways! After that point I didn't hide or avoid the fact I have alopecia, I'd say I'm pretty upfront about it now because I genuinely couldn't care, I'm not mad, I'm not sad, I actually see it as some kind of blessing as it gives me the opportunity to be creative, I never get bored of my hair because I can switch it up whenever I want, more than once a day if I really want to! However, I had never really wanted to go outside without my wig on. This wasn't because I'm ashamed of how I look without it on, it's just because I'm comfortable wearing wigs and I enjoy wearing them! I am proud of who I am, and I am proud of the journey I have been on.





Please head over to www.niamhwoods.wordpress.com to read Niamh's blog - "Miss Wiggy".

## Supporting a normal lifestyle

Ectodermal Dysplasia Society (Registered Charity No. 1089135). Disclaimer: Any views or opinions are made by the author in good faith. No liability whatsoever is accepted by the author or the Ectodermal Dysplasia Society. Recipients should make their own additional enquiries of medical and other relevant authorities before acting on these views. The use of a product name does not constitute a recommendation or endorsement by the author or the Society.