

A small charity based in West Wales, established by a few patients, of this rare disease, coming together and sharing, the patients experience, of living with this "Life limiting" health condition.

Focusing on Ataxia and associated rare diseases and movement disorders, Allowing us to cater for a wider patient community, who might feel isolated, with a health condition that is generally un-heard of.

**Ataxia – "the most serious condition you`ve never heard of".**

**Ataxia is a Greek word meaning "lack of Order"**

**We aim to bring some order, to the lack of order.**

Although, most physical activity was very small, due to the (legacy of the Covid19) pandemic. Many events have been provided via on-line platforms.

Although, as we are more of an online charity, we were able to utilise, the advantage of a virtual presence, allowing us to keep interacting with the community.

During this year the theme for most events, was VIRTUAL (either via Zoom or Microsoft teams).

Events (on-line) 128

Meetings (F2F) 18 (Virtual) 4

Contribute to speaking, at virtual events 2

Supply text for media articles 2

Display/stand at 2 events

We hosted 2 volunteers during the year (adding to our social media presence), 1 volunteer relocated to Spain, but continued to contribute to our social media, by translating posts into Spanish and Catalan languages.

Interactions with local merchandise suppliers (for team Ataxia) 6

Our trustee and his wife, continue to make amazing links with other patient advocates, and other organisations, contributing to, and, conveying the patients perspective, including many creative writing and videos, poems, podcasts and speaking as patient advocates to many organisations.

Indeed, Louise, is quite an Ambassador, having had a book published, " Ruffling Feathers", from which she reads excerpts, from, at many Spoken Word, events. This continues to raise awareness of Ataxia, Neurology, and a positive outlook.

A highlight of the year was to be invited to the opening of the Genomic Centre in Cardiff

Which was officiated by the Health minister, and made the TV news, as a HUGE advancement in the Genetic/Rare disease arena.

Public liability Insurance

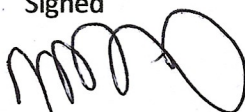
Markel London

Bank

Barclays 20-25-19

## Financials

Income		Expenditure	
Description	Amount	Description	Amount
In		Out	
Fundraising		Media materials	
Local Giving page	48.64	Just Print	150
Local Giving page	17.92	Get Stitched	200
Local Giving page	33.93	Get Stitched	165.4
Local Giving page	43.74	Get Stitched	168.99
Local Giving page	9.61	RareRevolution (charity membership)	65
Local Giving page	50.00		
Local Giving page	9.61	Local giving (use of,fundraising platform)	180
Local Giving page	19.42		
		Website (123)	117.54
		TSO Host (web. Hosting)	129.46
		Cheque to AT 23 balance	301.81
GAYL	15.03	postage to Volunteer	12.15
GAYL	15.39		
		Insurance	100.8
amazon	5.00		
amazon	15.96		
Total	284.25	>>>> MINUS	1591.15
End	1306.9		

Signed	full name	Position	date
	ALAD THOMAS	CHAIR.	16.1.25

Signed	full name	Position	date
D. Morgan	DAWN MORGAN	SECRETARY	16.1.25