

Information sourced from Tourettes Action

SYMPTOMS/CHARACTERISTICS

Tics (involuntary sounds and movements) are the main characteristic. There are other tic disorders, but for tourette's, multiple motor tics and one or more vocal tics must be present for at least 12 months, though not always concurrently.

Tics can be simple (head jolting, grunting) or complex (touching objects/people, repeating phrases).

Most people associate TS with swearing (coprolalia), but only 15%-20% have this.

can

help

Many people with TS experience a physical premonitory feeling / urge feeling a tic. Recognising this suppress tics for a while, but often makes it worse. It's only reduced by performing the tic. For some, the tic has to be performed multiple times before it 'feels right'.

Many people with TS experience a physical premonitory feeling / urge preceding a tic. Recognising this feeling and ignoring it can help suppress tics for a while, but often makes it worse. It's only reduced by performing the tic. For some, the tic has to be performed multiple times before it 'feels right'.

Tic attacks are bouts of severe, continuous, non-suppressible and disabling tics that can last from minutes to hours.

Up to 85% with TS experience co-occurring symptoms:

ADHD OCD

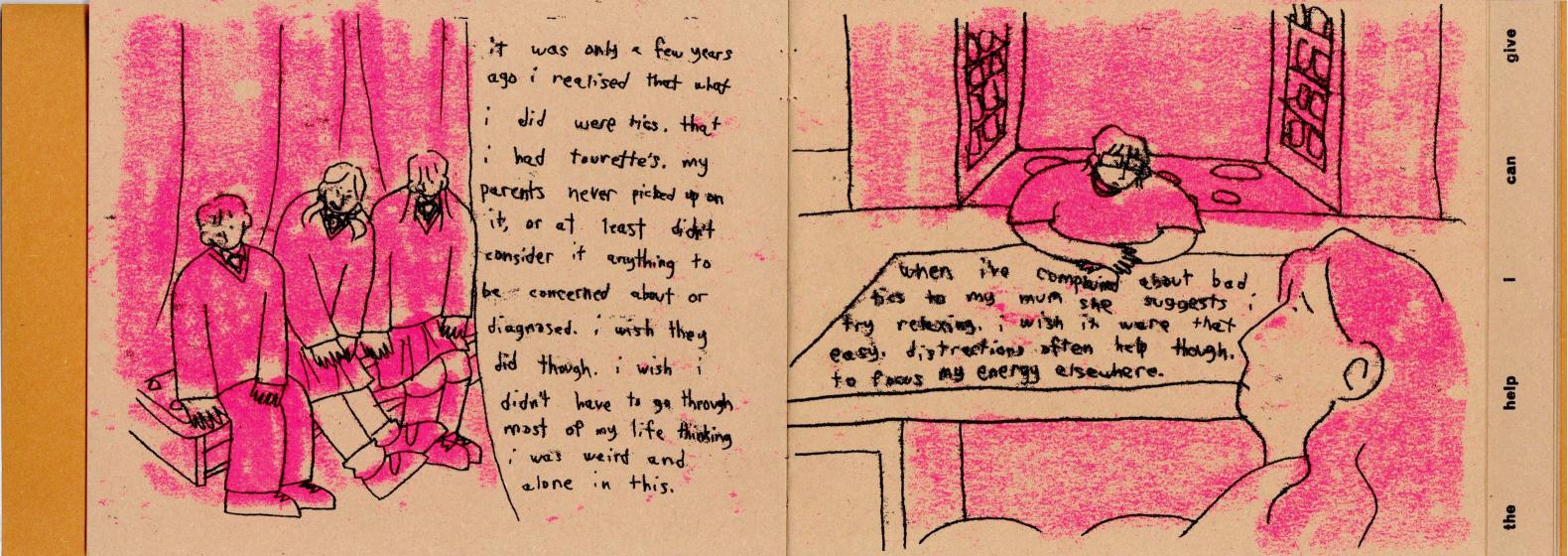
> **Anxiety** Anger

Executive functioning difficulties Sensory processing difficulties

Depression

Autism spectrum disorder Sleep problems







you have tourettes? ricouldn't even tell as much in front of People.
a masking thing i guess.

it always makes me feel weind when people say they can't even tell i have tourette's while i'm grateful to have it less severe than others and so can live my life relatively unchallenged, it just makes me doubt myself is my tourefte's even bad enough to telk about ? it affects me, but usually in private, i don't went People to think i'm making a big deal out of nothing or faking, being aware of how People may perceive me affects my tics. Some 'private' ties i never do in public, some Public ties i namely do in private.

give

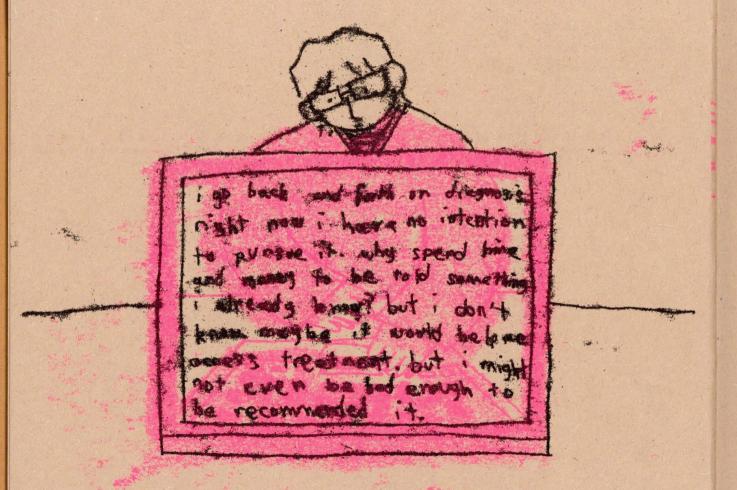
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THINK

YOU

HAVE

TOURETTE'S?

putting a name to something is a good first step, the validation i felt when my experiences lined up with the description of 'tis' on the NHS website was a great relief.

it's your choice if you want to be diagnosed many people with TS were diagnosed as a child, but if, like me, you weren't, consider what a diagnosis could do for you.

talk to your support system and if you decide on diagnosis, talk to your GP who can refer you to a specialist. Townstes Action also has a list of specialists throughout the UK.

Tourettes Action has really good resources and information, and is often what i refer to.

sometimes i look at the touretter teg on tumble to read posts of people seeking out other people with 15 ean be really comforting and make talking about their own experiences or feel less alone. there is raising awareness, sometimes, read thoso basically no mainstream representation same posts over and over there is much of TS, but there are some comfort in knowing you're not alone. that people documentas their experiences you're part of an unknowingly large online. a youthber it recommend group who understand. is Zara Beth. DELL 0000 (P)

THINGS THAT CAN HELP

seeking specialised advice from a professional is best if your ties are bad, but tips to manage them can be found online and through others with TS.

distrecting yourself by concentrating on something else can work well to reduce ties for a while. Hings like:

excercise & sport listening to/making music stim toys making art if a certain tic is bothering you, try redirecting it by doing something else. i can sometimes calm my abdominal tensing tic by humming.

ties are worsened by stress and tiredness, being aware of what worsens your ties means you can avoid or be prepared for environments that may exacerbate them.

for more painful ties like banging or julting, you can protect yourself with padding and help the pain with braces and/or a hot water bottle. seek out mobility aids and any other disability accommodations if you think they would help.

TS is a disability and there's no shame in asking for help.

this is my perspective and experience. everyone is different.



@batsanrats