What is a ‘serious’ genetic condition? The perceptions of people living with genetic conditions

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INTRODUCTION

As an era of targeted genetic testing gives way to global mainstreaming of high output genomic sequencing, important decisions need to be made regarding which genetic variants, associated with which conditions, are considered ‘worth knowing’ [1]. In addition to the strength of genotype/phenotype correlations, the availability of effective treatment/prevention and significance of the genetic findings to wider family members [2], the ‘seriousness’ of conditions has been pivotal to policy decisions governing the uses of genetic screening [3]. The notion of ‘seriousness’ has been employed to guide professionals [4, 5] and patients [6] in determining which incidental findings should be returned from whole genome sequencing and which variants should be included on population level genetic screening panels [3, 7]. It is also employed to govern access to selective pregnancy termination and technologies such as pre-implantation genetic diagnosis (PGD). However, what ‘counts’ as a serious condition remains contested, with no clear legal or social definition [8, 9].

Building on guidelines produced by the American College of Medical Genetics [10], various attempts have been made to operationalise the concept of ‘seriousness’ by researchers [6, 11–13], authoritative bodies [14], and working groups [15] in order to support clinical and policy decision-making. These attempts often draw on the views of professionals (e.g. obstetricians, genetics professionals, paediatricians), to delineate core ‘dimensions’ of seriousness (e.g. age of onset, lifespan, variability of symptoms, effective treatment). Dimension thresholds are then assigned to higher order categories (e.g. ‘mild’, ‘moderate’, ‘severe’, or ‘profound’) [6], onto which genetic conditions are then mapped. The resulting taxonomies [5, 6] are attractive in their ability to synphon complex conditions into discrete manageable categories and transform unfamiliar rare disease names into accessible lay language. Designed to streamline reproductive genomic decision-making, they enable patients to decide whether to receive genetic results for groups of similarly serious conditions, with the ultimate goal of facilitating informed decision-making [6], and to guide decisions regarding the inclusion of conditions into expanded carrier screening programmes. However, concerns remain regarding their applicability to unpredictable, fluctuating, and spectrum conditions [5], as well as their neglect of non-clinical factors (e.g. social/environmental contexts) [9, 16]. Moreover, genetics professionals indicate a preference for genetics patients to co-produce definitions of seriousness [8].

The importance of including the perspectives of people living with genetic conditions in understanding the subjective nature of ‘seriousness’ is increasingly acknowledged [7, 17–21]. Indeed, there are examples in the literature of people with disabilities evaluating their condition and quality of life (QoL) differently—and often more positively—than others around them do, suggesting differences in condition appraisal between those with and without lived experience [2, 22–24]. Disability focused research, however, has been slow to infiltrate the literature on reproductive genetics, despite the fact that people living with the...
conditions that screening programmes could potentially identify, arguably have the most accurate and intimate understanding of life with a genetic condition [20, 25].

This paper examines the perspectives and experiences of adults living with highly contrasting genetic conditions—including those that are treatable (haemophilia, thalassaemia), those which are life-limiting (cystic fibrosis, spinal muscular atrophy), those involving cognitive (fragile X syndrome) and/or physical impairment (spinal muscular atrophy, thalassaemia) (Table 1). Of these, Thalassaemia is the only condition routinely screened for prenatally in the UK. Perceptions of the ‘seriousness’ are established through an analysis of 45 in-depth interviews and 469 surveys with adults diagnosed with these conditions, that explore their perceptions of suffering, QoL, health, and attitudes towards prenatal screening (PNS) (and selective termination) for their condition. Mixed methods techniques are employed to combine the two datasets. The use of mixed methods allows a detailed exploration of participant attitudes, whilst also including a large number of participants, which is particularly beneficial when analysing such complex and nuanced topics. Finally, participant reflections on condition seriousness are positioned in the context of the broader literature and existing taxonomies [26], in order to highlight their relevance and necessary contribution to future applications of reproductive genomic medicine.

**METHODS**

The data for this study were collected as part of a larger sequential mixed methods study exploring the attitudes of people living with genetic conditions towards different types of population screening programmes (pre-conception, PNS, newborn). The analysis presented here focusses on the concept of seriousness among people with SMA, thalassaemia, CF, haemophilia (men/women analysed separately), and FXS, with the addition of people with VWD and FXTAS/FXPOI. Detailed methods can be found in [16, 27, 28]. Ethical approval was granted by the Biomedical and Scientific Research Ethics Committee and the Health Research Authority (17/WM/0231 01/08/17). All participants were informed of the study aims and potential data uses. Interview participants signed a consent form, completion of surveys (anonymously) was deemed a proxy for consent.

**Qualitative data collection and analysis**

Forty-five interviews were conducted between March 2017 and January 2019. Thirty-five participants were recruited through condition support groups and 10 through an NHS CF clinic. Interviews were face-to-face (n = 14), by telephone (n = 29) or email (n = 2), as determined by participants’ preferences. Participants were asked about their daily lives with their condition and their attitudes towards screening. Names/identifiers were removed during (verbatim) transcription and replaced with pseudonyms. Open coding (Nvivo 11) identified broad descriptive themes from which subthemes were developed. A conceptual cross-cutting framework was created and organised into a hierarchical framework, utilising an iterative analytic process driven by the data (constant comparison method) with reference to the literature. This was conducted until data saturation was achieved.

**Quantitative data collection using surveys**

A core set of questions was developed from the qualitative data (with assistance from condition support groups) with amendments to ensure applicability for each condition. As the survey was originally designed to explore attitudes to hypothetical future screening programmes, questions relating to PNS were not asked of thalassaemia participants due to the pre-existence of a thalassaemia PNS programme in the UK. Surveys (paper/online) were completed between December 2017 and November 2019. Questions were designed to explore attitudes towards screening, but also current health/wellbeing, QoL, suffering, social support, PNS, termination for own condition and the consequences of this (Table 2). Participants responded using five-point likert scales, which were condensed for analysis into positively (agree/strongly agree) or negatively (disagree/strongly disagree) balanced answers and those that were neither (neither/don’t know). Current health and wellbeing responses were condensed into poor/very poor/fair versus good/very good. Statistical differences were determined using Chi squared or Fisher’s Exact (where assumptions of Chi violated) (IBM SPSS Statistics 27). Comparisons across all participants

| Condition Name | Description |
|----------------|-------------|
| Spinal Muscular Atrophy (SMA) | Neuromuscular disorder. SMA1 is generally fatal in infancy. SMA2 (onset 6–18 months) typified by ability to sit but not walk, many attain a near-normal lifespan. SMA3 and SMA4 (childhood and adult onset, respectively), typified by progressive muscle weakness, lifespan unaffected. |
| Fragile X Syndrome (FXS) | Most common cause of inherited intellectual disability. Severity and combination of symptoms varies widely (e.g. can include ADHD and epilepsy). Males are more severely affected (X-linked) |
| Fragile X-associated Primary Ovarian Insufficiency (FXPOI) and Fragile X-associated Tremor/Ataxia (FXTAS) | Affects some FXS ‘premutation’ carriers: FXPOI (women), causing reduced fertility and early menopause; FXTAS (men and women), a late-onset (>50 years) progressive neurodegenerative disorder. |
| Thalassaemia | Affects haemoglobin production. Only condition for which universal PNS is available in the UK. Individuals with lifetime access to blood transfusions/and chelation therapy can achieve a near-normal lifespan. Comorbidities include heart, bone, and endocrine problems, diabetes, cirrhosis, gallstones and reduced fertility. |
| Cystic Fibrosis (CF) | Most common inherited life-limiting disease in European populations (UK life expectancy 47 years). Newborn screening offered (UK) since 2007, with cascade carrier screening and PNS for affected families. Thickened epithelial secretions cause progressive damage to the lungs, intestine, liver, kidneys, pancreas, and reproductive organs. Respiratory failure causes over 80–90% of deaths, but comorbidities include malabsorption, digestive tract cancers, diabetes, renal disease, osteoporosis, cirrhosis, and fertility problems. |
| Haemophilia | Haemophilia A is the most common and severe form, affecting mainly men (X-linked). Current life expectancy on average 15 years less than the general population. Associated with spontaneous bleeding (in joints/soft tissues/intestine/brain), leading to acute pain, long-term disability, or death. |
| Von Willebrand disease (VWD) | Most common inherited clotting disorder. Affects men and women. Types 1 & 2 generally less severe than haemophilia (lifespan unaffected), type 3 is equivalent to severe haemophilia A. |
Table 2. Summary of attitudes of 469 survey respondents.

|                     | SMA2 | SMA3 | SMA4 | Haemophilia Women | Haemophilia Men | VWD | Cystic Fibrosis | Thalassaemia | FXS | FX/TAS/POI |
|---------------------|------|------|------|-------------------|----------------|-----|----------------|--------------|-----|-----------|
|                     | (N = 28) | (N = 31) | (N = 23) | (N = 29) | (N = 151) | (N = 77) | (N = 21) | (N = 61) | (N = 18) | (N = 30) |
| **N**               | %    | %    | %    | %    | %    | %    | %    | %    | %    | %    |
| **Current health and wellbeing** |       |       |       |       |       |       |       |       |       |       |
| Bad/Vb              | 0.0% | 2.0% | 10.5% | 3.0% | 16.7% | 6.9% | 24.6% | 16.1% | 13.7% | 7.0% | 4.8% |
| Fair                | 23.1%| 7.6% | 36.8% | 11.0%| 61.1% | 17.2%| 53.5% | 35.6% | 20.2%| 26.3% | 7.3% |
| Good/Vg             | 76.9%| 8.5% | 65.2% | 22.0%| 22.2% | 75.9%| 22.2% | 48.3% | 43.2%| 35.6% | 16.1%|
| **It is possible to have a good quality of life** |       |       |       |       |       |       |       |       |       |       |
| Agree (N = 28)      |       |       |       |       |       |       |       |       |       |       |
| Disagree            | 0.0% | 2.0% | 6.0%  | 0.0% | 0.0%  | 4.7% | 3.9%  | 1.4%  | 0.0% | 0.0% | 0.0% |
| DK/Neither          | 7.1% | 5.0% | 16.0% | 17.4%| 10.3% | 19.0%| 9.2%  | 19.0% | 9.2% | 19.0%| 5.9% |
| **Condition causes people to suffer** |       |       |       |       |       |       |       |       |       |       |
| Agree               | 92.9%| 23.1%| 76.7% | 15.0%| 65.2% | 26.9%| 89.7% | 87.3% | 86.8%| 81.0%| 80.0%|
| Disagree            | 0.0% | 23.1%| 23.1% | 23.1%| 23.1% | 23.1%| 23.1% | 23.1% | 23.1%| 23.1%| 23.1%|
| DK/Neither          | 7.1% | 5.0% | 16.0% | 17.4%| 10.3% | 19.0%| 9.2%  | 19.0% | 9.2% | 19.0%| 5.9% |
| **People/families well supported by society** |       |       |       |       |       |       |       |       |       |       |
| Agree               | 76.9%| 21.4%| 13.3% | 1.4% | 52.2% | 31.0%| 27.3% | 35.1% | 31.0%| 27.5%| 16.1%|
| Disagree            | 17.6%| 17.6%| 17.6% | 17.6%| 17.6% | 17.6%| 17.6% | 17.6% | 17.6%| 17.6%| 17.6%|
| DK/Neither          | 6.5% | 1.0% | 1.0%  | 1.0% | 1.0%  | 1.0% | 1.0%  | 1.0%  | 1.0% | 1.0% | 1.0% |
| **I support pre-natal screening** |       |       |       |       |       |       |       |       |       |       |
| Agree               | 55.6%| 52.6%| 65.2% | 31.0%| 27.3% | 35.1%| 17.2% | 31.0% | 50.0%| 59.0%| 58.8%|
| Disagree            | 44.4%| 47.4%| 34.8% | 69.0%| 72.7% | 64.9%| 82.8% | 49.9% | 40.0%| 41.2%| 41.2%|
| DK/Neither          | 9.9% | 9.9% | 9.9%  | 9.9% | 9.9%  | 9.9% | 9.9%  | 9.9%  | 9.9% | 9.9% | 9.9% |
| **PNS leads to fewer people existing who could have fulfilling lives** |       |       |       |       |       |       |       |       |       |       |
| Agree               | 67.9%| 21.4%| 13.3% | 4.3% | 63.6% | 44.8%| 42.7% | 52.4% | 52.4%| 43.8%| 55.2%|
| Disagree            | 32.1%| 78.6%| 86.7% | 95.7%| 56.4% | 57.3%| 47.3% | 47.6% | 47.6%| 44.8%| 44.8%|
| DK/Neither          | 6.0% | 6.0% | 6.0%  | 6.0% | 6.0%  | 6.0% | 6.0%  | 6.0%  | 6.0% | 6.0% | 6.0% |
| **Would be loss to have fewer people with condition** |       |       |       |       |       |       |       |       |       |       |
| Agree               | 64.3%| 26.7%| 26.7% | 35.4%| 37.1% | 30.7%| 38.1% | 38.1% | 38.1%| 38.1%| 38.1%|
| Disagree            | 35.7%| 73.3%| 73.3% | 64.6%| 62.9% | 61.9%| 61.9% | 61.9% | 61.9%| 61.9%| 61.9%|
| DK/Neither          | 1.0% | 1.0% | 1.0%  | 1.0% | 1.0%  | 1.0% | 1.0%  | 1.0%  | 1.0% | 1.0% | 1.0% |
| **PNS enables informed decision making whether to have child** |       |       |       |       |       |       |       |       |       |       |
| Agree               | 71.4%| 83.3%| 86.2% | 71.5%| 62.2% | 81.0%| 62.5% | 80.0% | 80.0%| 80.0%| 80.0%|
| Disagree            | 28.6%| 16.7%| 13.8% | 28.5%| 37.8% | 19.0%| 19.0% | 20.0% | 20.0%| 20.0%| 20.0%|
| DK/Neither          | 7.1% | 5.0% | 16.0% | 17.4%| 10.3% | 19.0%| 9.2%  | 19.0% | 9.2% | 19.0%| 5.9% |

Grey shading signifies question not asked of group.

PNS prenatal screening. Vb/Vg very bad/very good. Dk don't know.
Mixed-methods data analysis of the concept of seriousness
Quantitative findings from the surveys were cross-referenced with related conceptual themes from the qualitative data during mixed methods analysis to offer context and greater depth of understanding of statistical findings. Representative qualitative excerpts of key findings presented in this paper were selected during this mixed methods analysis.

RESULTS
There were 469 survey returns and 45 interview participants. Women were over-represented (Table 3), except in SMA4 (survey) and haemophilia (survey and interview) groups. Participants were predominantly of white British ethnicity, except for the thalassaemia group (Asian/Arab, or white European). Survey participants in CF, thalassaemia, SMA2 and FXS groups tended to be younger than the other groups.

Current health and wellbeing, QoL, and suffering
Participants generally reported good current health, with very few stating they had bad or very bad health/wellbeing (Table 2). Interview participants highlighted how their positive health status could come as a surprise to other people. Samantha (32, CF), who was on the waiting list for a lung transplant, commented, "I think people are surprised when I say I'm doing fine and feel ok most days. Yes, I will have been doing nebulisers all morning and night but that's my normal. Perhaps my normal is different to other people's, but it's my normal, you know?" This sense of being in good health, despite the perceptions of others, was mirrored across conditions. Participants with SMA2 reported the highest levels of good/very good current/health welfare (76.9%). Mark (25, SMA2), was living in University halls supported by care workers at the time of interview. He was a full-time wheelchair user and used a C-PAP machine to support his breathing. He reflected, "You see, I see myself as really healthy, actually. I don't often get colds, or have any serious breathing issues or chest infections...I've only been hospitalised twice my whole life...[...]. The SMA part is a disability, but it doesn't impact much on my health actually. It doesn't make me ill as such, or in much pain...it's just the way my body is. I don't think it's a contradiction to say you're both healthy and disabled."

Although not commonly reported, bad or very bad current health/wellbeing was most frequent in participants with VWD (17.1%), SMA4 (16.7%), men with haemophilia (16.1%), and people with FXSTAS/POI (15.4%). Due to their infrequency, these responses were combined with 'fair' for pairwise analyses. Participants with SMA4 had the highest levels of bad/fair (77.8%) and lowest levels of good health/wellbeing (22.2%) (Tables 2 & 4), which is striking as SMA4 is clinically the mildest form of SMA with late onset (usually after 35 years). Helen (45) described the dramatic changes in her life and health brought about by SMA4; "I'll never forget the day I was diagnosed...[...]. I was 37, just got all my kids in school and had just started in a new job...[...]. It's been downhill from there really, I'm 45 now and I still walk, but...with a frame now. I couldn't keep up with the job and had to chuck it in which was devastating and that hit us...financially. I've just...lost my independence really. Most days I'm so weak I struggle with everything now, it's a terrible illness...So I would say my health is pretty bad now." Helen incorporated her SMA into her evaluation of her own health, viewing it as an 'illness'. In contrast, Mark drew a conceptual distinction between his disability—which had been a comparatively stable presence since infancy—and his health, which he interpreted as being the absence of illnesses, pain and suffering.

Nearly all participants agreed that it is possible to have a good QoL with the condition they lived with; however, in a similar pattern to that observed with health, participants with SMA2 were most likely to agree (92.9%) and participants with adult onset conditions least likely to agree (SMA4 65.2%, FXSTAS/POI 67.9%). This difference between early and later onset conditions was also evident within the qualitative dataset. James (46, haemophilia) had received regular treatment with blood products from childhood and through this, contracted Hepatitis C at age 10. The hepatitis was successfully treated, but James's haemophilia led to arthritis, prompting him to resign from his active job. He reflected, "Quality of life...is what you make it really. Yes, I've got my issues to deal with, but so does everyone, just in different ways...although some people will tell you you need your health to have a quality of life, I think I'm living proof that you can have it anyway. As long as you have positive relationships, you know, you have things to look forward to, you have things to do, that's all you really need." The qualitative data revealed that participants, such as James, who had lived with their condition since birth or childhood—even though his condition had changed over time and had implications for health as well as disability—were more likely to define QoL as a combination of factors external to themselves, such as social relationships, interests and activities, which haemophilia did not preclude. This was in stark contrast to participants with later onset conditions, such as Mary (74), diagnosed with FXTAS in her early 60s who commented on the onset of her condition, "My quality of life deteriorated considerably. Everything you take for granted in life gets pulled out from under you, like driving, shopping, cooking a meal, even basic things like drinking a hot drink without spilling it all over yourself...you shake constantly and feel tired constantly...[...]. and when you can't do those things no more—there isn't much quality left is there?". For Mary, health, QoL and FXTAS were inextricably bound together. Having once enjoyed life, Mary's FXTAS was a restrictive, unwelcome and unexpected burden, prompting a re-evaluation of her lifestyle and identity, akin to what Bury has referred to 'biographical disruption' and re-integration [29].

Differences between early and later onset conditions were less pronounced, however, within perceptions of suffering. Almost all of those with CF (90.5%) agreed that CF causes suffering (Table 2), which was significant compared with SMA2 (25.0%), thalassaemia (59.0%), FXS (58.8%) and FXSTAS/POI (53.6%) (Table 4). After CF, the greatest proportion of participants reporting suffering were those with haemophilia (men 78.5%, women 75.9%), SMA3 (70.0%) and SMA4 (72.7%). This seeming contradiction, that genetic conditions cause suffering, but that a good QoL can be achieved, was expressed by the majority of participants within the qualitative data. Johara (36, thalassaemia) commented, "Thalassaemia is hard...I am ill a lot, I get tired, and I have restrictions in my life because of it...but that doesn't mean that you can't have a good life...[...]. I'd always choose to be here, even with thalassaemia."

Societal influences
Across all conditions, fewer than 50% of respondents felt well supported by society. Men with haemophilia most frequently (although still only 46%) felt well supported, and respondents with SMA3 (13.3%) and SMA4 (4.3%) least frequently felt supported. Within the qualitative data, links between societal support and lived experience were made explicit, with many participants emphasising that the social consequences of their condition—and its interface with their environmental context—could be as disabling as the physiological effects of a genetic condition, as highlighted by Olivia (37, SMA2), "I don't much think about having SMA, what I think more about is the access issues. As long as society is accessible, I'm not restricted...But if I can't get into a building, if I can't get around...yeah, that's when I'm disabled." Indeed, within the survey, of the 27 participants who felt that good QoL was not possible with their condition, 77.8% felt unsupported by society (compared with 35.8% of the 385 people
Table 3. Demographics of 469 survey and 45 interview respondents.

| Variable                        | SMA2 Survey | SMA3 Survey | SMA4 Survey | Haemophilia Women Survey | Haemophilia Men Survey | VWD Survey | Cystic Fibrosis Survey | Thalassaemia Survey | FXS Survey | FXTAS/POI Survey |
|---------------------------------|-------------|-------------|-------------|--------------------------|------------------------|------------|------------------------|---------------------|------------|---------------------|
| Gender                          |             |             |             |                          |                        |            |                        |                     |            |                     |
| Female                          | 20 (71.4%)  | 17 (68.8%)  | 14 (66.7%)  | 31 (68.8%)               | 17 (68.8%)             | 2 (8.3%)   | 17 (68.8%)             | 14 (66.7%)          | 31 (68.8%) | 17 (68.8%)          |
| Male                            | 8 (28.6%)   | 8 (31.5%)   | 6 (27.3%)   | 14 (31.5%)               | 10 (41.7%)             | 4 (16.7%)  | 10 (41.7%)             | 9 (29.0%)           | 14 (31.5%) | 15 (33.3%)          |
| NA                              | 0           | 0           | 0           | 0                        | 0                      | 0          | 0                      | 0                   | 0          | 0                   |
| Age (yrs)                       |             |             |             |                          |                        |            |                        |                     |            |                     |
| Under 35                        | 11 (39.3%)  | 3 (12.0%)   | 1 (4.6%)    | 14 (31.5%)               | 14 (56.9%)             | 4 (16.7%)  | 14 (56.9%)             | 12 (46.4%)          | 14 (31.5%) | 10 (21.3%)          |
| 35–55                           | 14 (50.0%)  | 8 (32.0%)   | 16 (72.7%)  | 21 (47.8%)               | 9 (35.4%)              | 5 (20.8%)  | 12 (47.8%)             | 17 (62.1%)          | 28 (61.5%) | 13 (27.1%)          |
| Over 55                         | 1 (3.6%)    | 1 (4.0%)    | 2 (9.1%)    | 2 (4.4%)                 | 1 (3.9%)               | 0          | 0                      | 0                   | 0          | 0                   |
| Ethnicity                       |             |             |             |                          |                        |            |                        |                     |            |                     |
| Asian/Arab                      | 1 (3.6%)    | 0           | 1 (4.6%)    | 1 (2.3%)                 | 0                      | 0          | 0                      | 0                   | 0          | 0                   |
| Black                           | 0 (0.0%)    | 1 (4.0%)    | 0           | 0                        | 0                      | 0          | 0                      | 0                   | 0          | 0                   |
| White British                   | 26 (92.9%)  | 7 (28.0%)   | 16 (71.4%)  | 36 (77.7%)               | 31 (121.4%)            | 4 (16.7%)  | 31 (121.4%)            | 25 (91.4%)          | 30 (68.8%) | 23 (48.9%)          |
| White Other                     | 1 (3.6%)    | 2 (8.0%)    | 1 (4.6%)    | 1 (2.3%)                 | 0                      | 0          | 0                      | 0                   | 0          | 0                   |
| Religious faith                 |             |             |             |                          |                        |            |                        |                     |            |                     |
| No                              | 10 (35.7%)  | 8 (32.0%)   | 8 (36.4%)   | 18 (40.0%)               | 15 (58.6%)             | 4 (16.7%)  | 14 (58.6%)             | 12 (43.8%)          | 17 (37.2%) | 14 (29.1%)          |
| Yes                             | 17 (60.7%)  | 10 (40.0%)  | 12 (53.6%)  | 27 (58.0%)               | 9 (34.0%)              | 5 (20.8%)  | 19 (72.0%)             | 21 (76.1%)          | 23 (50.0%) | 18 (37.2%)          |
| Educational level               |             |             |             |                          |                        |            |                        |                     |            |                     |
| No                               | 11 (38.3%)  | 6 (24.0%)   | 6 (27.3%)   | 17 (37.2%)               | 15 (58.6%)             | 4 (16.7%)  | 14 (58.6%)             | 12 (43.8%)          | 17 (37.2%) | 14 (29.1%)          |
| Yes                             | 26 (88.9%)  | 19 (76.0%)  | 16 (72.7%)  | 47 (98.0%)               | 52 (200.0%)            | 5 (20.8%)  | 48 (192.0%)            | 35 (125.0%)         | 52 (110.0%)| 42 (87.2%)          |
| NA                              | 0 (0.0%)    | 0           | 0           | 0                        | 0                      | 0          | 0                      | 0                   | 0          | 0                   |

NA = did not answer or 'prefer not to say.'
Table 4. Significant differences in attitudes between survey respondents of different condition groups (Pairwise comparison, significant differences $p < 0.05$ in bold, trends of $p \leq 0.10$ also included).

### Current health and wellbeing

| Condition          | SMA2 | SMA3 | SMA4 | Haem Women | Haem Men | VWD | Cystic Fibrosis | Thalassaemia | FXS | FXTAS/POI |
|--------------------|------|------|------|------------|----------|-----|----------------|--------------|-----|-----------|
| SMA2               |      |      |      |            |          |     |                |              |     |           |
| SMA3               |      |      |      |            |          |     |                |              |     |           |
| SMA4               |      |      |      |            |          |     |                |              |     |           |
| Haem Women         |      |      |      |            |          |     |                |              |     |           |
| Haem Men           |      |      |      |            |          |     |                |              |     |           |
| VWD                |      |      |      |            |          |     |                |              |     |           |
| Cystic Fibrosis    |      |      |      |            |          |     |                |              |     |           |
| Thalassaemia       |      |      |      |            |          |     |                |              |     |           |
| FXS                |      |      |      |            |          |     |                |              |     |           |

- It is possible to have a good quality of life
  - SMA2
  - SMA3
  - SMA4
  - Haem Women
  - Haem Men
  - VWD
  - Cystic Fibrosis
  - Thalassaemia
  - FXS

- The condition causes people to suffer
  - SMA2
  - SMA3
  - SMA4
  - Haem Women
  - Haem Men
  - VWD
  - Cystic Fibrosis
  - Thalassaemia
  - FXS

- People with condition and families well supported by society
  - SMA2
  - SMA3
  - SMA4
  - Haem Women
  - Haem Men
  - VWD
  - Cystic Fibrosis
  - Thalassaemia
  - FXS

- I support prenatal screening
  - SMA2
  - SMA3
  - SMA4
  - Haem Women
  - Haem Men
  - VWD
  - Cystic Fibrosis
  - Thalassaemia
  - FXS

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Table 4 continued

| Current health and wellbeing | SMA2 | SMA3 | SMA4 | Haem Women | Haem Men | VWD | Cystic Fibrosis | Thalassaemia | FXS | FXTAS/POI |
|-----------------------------|------|------|------|------------|----------|-----|----------------|--------------|-----|----------|
| Haem Men                    |      |      |      |            |          |     |                |              |     |          |
| VWD                         |      |      |      |            |          | 0.052|                | 0.094        |     |          |
| Cystic Fibrosis             |      |      |      |            |          |     |                |              |     |          |
| Thalassaemia                |      |      |      |            |          |     |                |              |     |          |
| FXS                         |      |      |      |            |          |     |                |              |     |          |

Pregnancy screen will lead to fewer people existing who could have lived fulfilling lives

| Current health and wellbeing | SMA2 | SMA3 | SMA4 | Haem Women | Haem Men | VWD | Cystic Fibrosis | Thalassaemia | FXS | FXTAS/POI |
|-----------------------------|------|------|------|------------|----------|-----|----------------|--------------|-----|----------|
| SMA2                        | 0.016*|      |      |            |          | 0.075|                |              |     |          |
| SMA3                        |      |      |      |            |          |     |                |              |     |          |
| SMA4                        |      |      |      |            |          |     |                |              |     |          |
| Haem Women                  |      |      |      |            |          |     |                |              |     |          |
| Haem Men                    |      |      |      |            |          |     |                |              |     |          |
| VWD                         |      |      |      |            |          |     |                |              |     |          |
| Cystic Fibrosis             |      |      |      |            |          |     |                |              |     |          |
| Thalassaemia                |      |      |      |            |          |     |                |              |     |          |
| FXS                         |      |      |      |            |          |     |                |              |     |          |

Loss to society to have fewer people with condition

| Current health and wellbeing | SMA2 | SMA3 | SMA4 | Haem Women | Haem Men | VWD | Cystic Fibrosis | Thalassaemia | FXS | FXTAS/POI |
|-----------------------------|------|------|------|------------|----------|-----|----------------|--------------|-----|----------|
| SMA2                        | 0.012*|      |      |            |          | 0.042*|                |              |     |          |
| SMA3                        |      |      |      |            |          |     |                |              |     |          |
| SMA4                        |      |      |      |            |          |     |                |              |     |          |
| Haem Women                  |      |      |      |            |          |     |                |              |     |          |
| Haem Men                    |      |      |      |            |          |     |                |              |     |          |
| VWD                         |      |      |      |            |          |     |                |              |     |          |
| Cystic Fibrosis             |      |      |      |            |          |     |                |              |     |          |
| Thalassaemia                |      |      |      |            |          |     |                |              |     |          |
| FXS                         |      |      |      |            |          |     |                |              |     |          |

Pregnancy screen enables informed decision about having child with condition

| Current health and wellbeing | SMA2 | SMA3 | SMA4 | Haem Women | Haem Men | VWD | Cystic Fibrosis | Thalassaemia | FXS | FXTAS/POI |
|-----------------------------|------|------|------|------------|----------|-----|----------------|--------------|-----|----------|
| SMA2                        | 0.012*|      |      |            |          | 0.042*|                |              |     |          |
| SMA3                        |      |      |      |            |          |     |                |              |     |          |
| SMA4                        |      |      |      |            |          |     |                |              |     |          |
| Haem Women                  |      |      |      |            |          |     |                |              |     |          |
| Haem Men                    |      |      |      |            |          |     |                |              |     |          |
| VWD                         |      |      |      |            |          |     |                |              |     |          |
| Cystic Fibrosis             |      |      |      |            |          |     |                |              |     |          |
| Thalassaemia                |      |      |      |            |          |     |                |              |     |          |
| FXS                         |      |      |      |            |          |     |                |              |     |          |

Pregnancy screen will prevent unnecessary suffering

| Current health and wellbeing | SMA2 | SMA3 | SMA4 | Haem Women | Haem Men | VWD | Cystic Fibrosis | Thalassaemia | FXS | FXTAS/POI |
|-----------------------------|------|------|------|------------|----------|-----|----------------|--------------|-----|----------|
| SMA2                        | 0.012*|      |      |            |          | 0.042*|                |              |     |          |
| SMA3                        |      |      |      |            |          |     |                |              |     |          |
| SMA4                        |      |      |      |            |          |     |                |              |     |          |
| Haem Women                  |      |      |      |            |          |     |                |              |     |          |
| Haem Men                    |      |      |      |            |          |     |                |              |     |          |
| VWD                         |      |      |      |            |          |     |                |              |     |          |
| Cystic Fibrosis             |      |      |      |            |          |     |                |              |     |          |
| Thalassaemia                |      |      |      |            |          |     |                |              |     |          |
| FXS                         |      |      |      |            |          |     |                |              |     |          |
who thought a good QoL was possible, $\chi^2 = 23.63$ $p < 0.001$), suggesting that social factors play an important role in negative lived experience.

**Attitudes towards PNS and pregnancy termination**

There were high levels of support for PNS from people with FXS (88.2%), CF (81.0%), SMA3 (80.6%), SMA4 (77.3%), and FXTAS/POI (76.7%) (Tables 2 & 4). Men with haemophilia and people with VWD showed lower levels of support (57.9% & 60.0%) but were more likely to select don’t know/neither (28.3% & 32.0%) than disagree (13.8% & 8.0%). People with SMA2 were least in favour of PNS (55.6%), and most likely to select disagree (40.7%). Enabling informed-decision making about whether to have an affected child (62–86% across groups) appeared to be a key driver of support for PNS. Isabelle (26, FXS), commented, “I don’t think it’s [PNS and pregnancy termination] a problem as it’s the parents’ choice. I think they have a right to choose not to have a child with fragile X if they don’t want to. It’s important to have full choice and...be fully aware.” Other interview participants highlighted the importance of fully committed parents to successfully raise a child with additional support needs, seeing PNS as a way to offer a 'way out' (David, 47, CF) to parents who feel they could not cope. Despite this, there was little support for selective pregnancy termination for their condition (<15% agreement for most conditions). The exceptions to this were FXTAS/POI and SMA3 groups (46.4% and 33.3%, respectively), and also SMA4 (27.3%, not significant). The interview data highlighted the complexity of this issue for people living with genetic conditions. Ilona (23, VWD) reflected, “I support it [PNS], but I don’t think it’s necessary suffering.” Claire (30 years, SMA2), commented, “I don’t think SMA causes suffering... Perhaps the parents suffer because they were thinking their child would be fine....Certainly I don’t know anyone with type II who would say they’re suffering... I think that’s perhaps just an idea about disability that people put on us ... unnecessarily, you know [laughs] because we’re fine!”

**DISCUSSION**

This paper is, to the best of our knowledge, the largest to explore attitudes and experiences relevant to the concept of condition ‘seriousness’ and PNS/selective termination exclusively from the perspectives of 514 adults living with clinically ‘serious’, and widely contrasting, genetic conditions. When set in the broader literature, all of the included conditions (with the exception of haemophilia which wasn’t subject to the algorithm) were classified as ‘severe’ using Lazarin et al’s taxonomy [5, 13], a system designed to classify conditions to simplify reproductive decision-making for prospective parents. In addition, all five are included on the pre-conception population screening panel as part of the McKenzie’s Mission pilot (Australia), based on their seriousness, treatment availability and whether an ‘average’ couple could reasonably be expected to want to avoid having a child with that condition [3]. One key finding of this study is, however, that whilst experiencing a clinically serious condition, the vast majority of participants reported both good health and the possibility of having a good QoL. Assuming that health/wellbeing/QoL are core dimensions of a condition’s seriousness, these findings indicate notable differences between the lived experience of affected adults and the way genetic conditions are defined and understood in the context of reproductive screening programmes. This
divergence illustrates the importance of including the views and lived experiences of those directly affected by genetic conditions in ethical debates surrounding the scope, and practical implementation of screening programmes. More specifically, debates around which conditions are considered ‘serious’ enough to qualify them as candidates for screening, as well as those around the adequacy and quality of informed consent for screening and any subsequent (prenatal) diagnosis. This is particularly important in relation to rare genetic conditions where public awareness, and knowledge of their lived realities, is typically low.

It has been suggested that people with genetic conditions might use different frames of reference for assessing their own health/wellbeing vis-à-vis those without genetic conditions [23, 24], and there was evidence within the dataset of this e.g. Samantha, CF. Indeed, physical and psychological adaptation to the condition, including its incorporation within personal identity (e.g. Mark, SMA2) was found to play a significant role in the way that health and QoL were interpreted. Participants who did not see their condition as part of themselves were more likely to view it as an illness, separate to their sense of self, that impacted their health and QoL (Mary, FXTAS).

Those with later onset conditions were more likely than those with early onset to report poorer health and consider a good QoL impossible. Previous research suggests that age of onset has a critical role to play in positive adaptation to a condition, and the likelihood of its incorporation into personal identity [30]. Indeed, those with later onset conditions were more likely to experience ‘biographical disruption’ [29] (Helen, SMA4), and associated forms of identity re-formulation around a new health state. This contrast was most clearly demonstrated in relation to SMA, where participants with milder, later onset types (SMA3/4), consistently reported poorer health/wellbeing than those with SMA2, suggesting a lack of correlation between clinical definitions of condition seriousness and lived realities for these adults. Late onset conditions have traditionally been excluded from reproductive screening programmes [26], often being interpreted as less serious (by virtue of the number of condition-free years) [31], yet, our data suggest that those with later onset conditions experience unique challenges in adjusting to their condition, irrespective of clinical seriousness.

The majority of participants perceived poor societal support, which was particularly evident in the minority who did not consider a good QoL to be possible. The influence of social and environmental factors on disease seriousness has been vastly under-explored in the context of reproductive genetics. As Kleiderman et al. [9] outline, the concept of ‘serious disease’ cannot be understood outside of social/environmental contexts—including societal/cultural influences on the interpretation of serious disease, health, and disability, which vary by time and place. However, the social and cultural context in which the condition will be experienced is not accounted for within taxonomies of condition seriousness.

The complexity of the views of people with genetic conditions, their largely positive lived experiences and supportive attitudes towards PNS, but also their resistance to the associated practice of selective termination, all emerged from this study. Participants highlighted that suffering and high quality of life can co-exist without contradiction. Their widespread support for PNS, when interpreted on its own, masks the deeply ambivalent views they held towards reproductive genetic technologies, and the value they still assign to future lives with their condition [18]. When designing taxonomies capable of classifying condition seriousness, future work may usefully take into account this complexity of lived experience, particularly in the context of the emergence of new therapies (e.g. nusinersen for spinal muscular atrophy) that may modify the nature of those experiences for future generations. Mixed methodologies, capable of capturing not only lived experience, but also the way that participants value different aspects of that experience, and balance the relative harms and benefits of screening (e.g. discrete choice experiments, Q methodologies) may prove to be particularly useful in this regard given the multifaceted nature of this topic. Indeed, the age of onset of the condition, its intersection with perceptions of health, disability and identity, and the social and environmental context in which it is experienced, were all found by this study to be fundamentally important to the way that genetic conditions were lived, yet have all been under-explored in debates around condition seriousness. These factors were found to override clinical markers of disease seriousness in their influence on lived experience, suggesting a need for a much wider range of factors and perspectives to be taken into account when determining what constitutes a serious condition. This is particularly important in an age of expanded genomic screening, wherein the conceptual apparatus used to define condition seriousness determines not only access to genomic screening programmes, but also the range of reproductive options available once a condition is found; ultimately influencing who does, and who does not, come into the world.

LIMITATIONS
The study questions included in this analysis were designed to assess attitudes towards hypothetical screening scenarios rather than perceptions of seriousness per se. For this reason, not all of the survey questions were asked of every group (e.g. participants with thalassaemia were not asked about prenatal screening). These data were therefore missing. The self-selecting nature of the sample introduces limitations, in that those who participated may have been in better health, and have more positive perspectives than those who opted not to. Despite this, participants represented a broad range of experiences/clinical severities within/across conditions and the included questions were all highly relevant to an analysis of seriousness. Adults with FXS were, however, underrepresented in the final sample. This was in spite of efforts to remove barriers to interview participation to allow for learning differences (e.g. adapted interview schedule/alternative interview methods, supportive adult present). As some of the reported data date back to 2017, this could also be considered a limitation of the study, given the pace of change, particularly in the field of expanded carrier screening in the international arena. However, as this study offers insights into the lived experiences of adults with a broad range of genetic conditions, their perceptions of their condition’s ‘seriousness’, and ultimately its prevention, the findings retain a high degree of relevance to debates underpinning the inclusion of conditions on such carrier screening panels (e.g. [7]), which are set to continue, even as such programmes begin to be implemented and/or expanded.

DATA AVAILABILITY STATEMENT
The datasets generated and analysed during the current study are not yet publicly available but will be deposited at the end of the study. However, they are available from the corresponding author on reasonable request.

REFERENCES
1. Schneider JL, Goddard KAB, Davis J, Wilford B, Kauffman TL, Reiss JA, et al. “Is It Worth Knowing?” Focus Group participants’ perceived utility of genomic pre-conception carrier screening. J Genet Couns. 2016;25:135–45.
2. Boardman FK, Young PJ, Warren O, Griffiths FE. The role of experiential knowl-edge within attitudes towards genetic carrier screening: a comparison of people with and without experience of spinal muscular atrophy. Health Expect. 2018;21:201–11.
3. Kirk EP, Ong R, Boggs K, Hardy T, Righetti S, Kamien B, et al. Gene selection for the Australian Reproductive Genetic Carrier Screening Project ("Mackenzie’s Mission"). Eur J Hum Genet. 2021;29:79–87.
4. Grody WW, Thompson BH, Gregg AR, Bean L, Monaghan K, Schneider A, et al. ACMG position statement on prenatal/preconceptional expanded carrier screening. Genet Med. 2013;15:482–483.
5. Arjunan A, Bellerose H, Torres R, Ben-Shachar R, Hoffman JD, Angle B, et al. Evaluation and classification of severity for 176 genes on an expanded carrier screening panel. Prenat Diagn. 2020;40:1246–57.
6. Leo MC, McMullen C, Wilford BS, Lynch FL, Reiss JA, Gilmore MJ, et al. Patients’ ratings of genetic conditions validate a taxonomy to simplify decisions about preconception carrier screening via genome sequencing. Am J Med Genet A. 2016;170:574–82.
7. Thomas LA, Lewis S, Massie J, Kirk EP, Archibald AD, Barlow-Stewart K, et al. Which types of conditions should be included in reproductive genetic carrier screening? Views of parents of children with a genetic condition. Eur J Med Genet. 2020;63:10075.
8. Wertz DC, Knoppers BM. Serious genetic disorders: can or should they be defined? Am J Med Genet. 2002;108:29–35.
9. Kleiderman E, Ravitsky V, Knoppers BM. The ‘serious’ factor in germline modification. J Med Ethics. 2019;45:508–13.
10. ACMG Newborn Screening Expert Group. Newborn screening: toward a uniform system-executive summary. Pediatrics. 2006;117:5296–307.
11. Chokoshvili D, Janssens S, Vears D, Borry P. Designing expanded carrier screening panels: results of a qualitative study with European geneticists. Pers Med. 2016;13:553–62.
12. Wilford BS, Goddard KA. It’s complicated: criteria for policy decisions for the clinical integration of genome-scale sequencing for reproductive decision making. Mol Genet Genom Med. 2015;3:239–42.
13. Lazarin GA, Hawthorne F, Collins NS, Platt EA, Evans EA, Haque IS. Systematic classification of disease severity for evaluation of expanded carrier screening panels. PLoS ONE. 2014;9:e114391.
14. Edwards JG, Feldman G, Goldberg J, Gregg AR, Norton ME, Rose NC, et al. Expanded Carrier screening in reproductive medicine-points to consider; a joint statement of the ACMG, ACOG, NGGC, PQF and SMFM. Obstet Gynecol. 2015;125:653–62.
15. Molster CM, Lister K, Metternick-Jones S, Baynham G, Clarke AJ, Straub V, et al. Outcomes of an international workshop on preconception expanded carrier screening: some considerations for governments. Front Public Health. 2017;5:25. https://doi.org/10.3389/fpubh.2017.00025.
16. Boardman FK, Clark C, Jungkurt H, Young PJ. Social and cultural influences on genetic screening programme acceptability: a mixed-methods study of the views of adults, carriers, and family members living with thalassemia in the UK. J Genet Couns. 2020;29:1026–40.
17. Roadhouse C, Shuman C, Anstey K, Sappleton K, Chitayat D, Igignani E. Disability experiences and perspectives regarding reproductive decisions, parenting, and the utility of genetic services: a qualitative study. J Genet Couns. 2018;27:1360–73.
18. Boardman FK, Hale R. How do genetically disabled adults view reproductive decision? Impairment, identity, and genetic screening. Mol Genet Genom Med. 2018;6:941–56.
19. McNeill A, Lewis R, Freeth M. Views of adults with 22q11 deletion syndrome on reproductive choices. Am J Med Genet A. 2020;182:1284–7.
20. Paul DR. Imagining life with a genetic disorder: the challenge of evaluating health states that exist from birth. OBM Genet. 2021;5:11.
21. Nijmeger SCM, Conijn T, Lakeman P, Henneman L, Wijburg FA, Haverman L. Attitudes of relatives of mucopolysaccharidosis type III patients toward preconception expanded carrier screening. Eur J Hum Genet. 2020;28:1331–40.
22. Albrecht GL, Devlieger PJ. The disability paradox: high quality of life against all odds. Soc Sci Med. 1999;48:977–88.
23. Ubel PA, Loewenstein G, Schwarz N, Smith D. Mismatching the unimaginable: the disability paradox and health care decision making. Health Psychol. 2005;24:557–62.
24. Lacey HF, Fagerlin A, Loewenstein G, Smith DM, Riis J, Ubel PA. Are they really that happy? Exploring scale recalibration in estimates of well-being. Health Psychol. 2008;27:669–75.
25. Petersen A. The best experts: the narratives of those who have a genetic condition. Soc Sci Med. 2006;63:32–42.
26. Kornjgiebel DM, McMullen CK, Amendola LM, Berg JS, Davis JV, Gilmore MJ, et al. Generating a taxonomy for genetic conditions relevant to reproductive planning. Am J Med Genet A. 2016;170:565–73.

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AUTHOR CONTRIBUTIONS
FB designed the study, conducted data collection and undertook the qualitative analysis. CC analysed the statistical data. The authors collaboratively undertook the mixed methods analysis, data integration and drafted the manuscript.

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COMPETING INTERESTS
The authors declare no competing interests.

ETHICS APPROVAL
Ethical approval was granted by the Biomedical and Scientific Research Ethics Committee and the Health Research Authority (17/WM/0221; 01/08/17). All participants were informed of the study aims and potential data uses. Interview participants signed a consent form, completion of surveys (anonymously) was deemed a proxy for consent.

ADDITIONAL INFORMATION
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