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Systematic Review: Psychological/psychosocial interventions for the families of gender diverse youth under 18 years old

Background:

The aim of this paper is the systematic review of psychological/psychosocial interventions for gender diverse youth under eighteen years of age and their families, based on the published protocol: PROSPERO 2020 CRD42020163995.

Methods:

A search strategy was developed using key terms. An electronic literature search was completed using the following data bases (OVID MEDLINE; EBSCO CINAHL; ProQuest MEDLINE; OVID PsycINFO). Only studies published in English between 2001-2021 were included. This review is based on PRISMA guidance. Studies meeting inclusion criteria were quality appraised using the Mixed Methods Assessment Tool (MMAT).

Results:

8,405 studies were independently screened. Four studies met the inclusion criteria for the study. Parents of transgender youth attended between one and eleven psychological/psychosocial group interventions. Parents reported reduced isolation and increased knowledge, which enabled them to advocate for their young person's needs. Psychological/psychosocial group interventions were creating challenges in terms of group processes, with some parents dominating interactions. Psychological/psychosocial group interventions were positive for parents, but no outcomes were collected for transgender young people.

Conclusion:

More research is required to understand the role of group facilitators, the optimal group size and the number of psychological/psychosocial intervention sessions required.

Keywords: words: systematic review, young people, children, families, transgender, gender dysphoria, gender diversity, psychological, psychosocial, therapy, interventions

Background

Transgender or gender diverse young people have a higher risk of poor mental health outcomes compared to their cis gender (non-transgender) age peers (Connolly *et al.* 2016; Reisner *et al.* 2015). This is not necessarily surprising given the multitude of challenges gender diverse adolescents' face, in trying to cope with the mismatch between their growing physical bodies and their internalised sense of self. Transgender adolescents are four times more likely to have depression compared to their cis gender peers (Peterson *et al.* 2017). Clinically, young people with depression can present as sad, hopeless, irritable and tired with loss of interest in previously enjoyable activities (Thapar *et al.* 2012). Depression can impact negatively in terms of school, home and leisure activities. In 2007, Grossman and D'Augelli

suggested that 58% of transgender adolescents receive a diagnosis of depression, of these, 51% experience suicidal ideation and 32% attempt suicide. These figures are alarming and highlight the need for a greater understanding of the relationship between adolescent transgender identity and mental health, in order to find ways to support young people and to reduce their risk of suicidality.

Increased risk of poor mental health outcomes has been associated in the literature with minority stress (Hendricks and Testa, 2012). In 2003, Meyer described the minority stress model, in which stigma, prejudice and discrimination impact negatively on mental health. While this concept was originally developed to understand the impact of a sexual minority status, it can be applied to a range of minority populations. It is worth noting, that many young gender diverse people, belong to more than one minority group. Aside from their transgender identity, they could be part of a sexual, ethnic or disability minority group, with overlapping areas of stigma and discrimination. Minority stressors can be described as distal (environmental), interactive (expectation of external threat) and internalised (internalised negative attitudes) (Meyer, 2003). While minority stressors can cause poor mental health, this is not a given. In a recent systematic review in 2021, Tankersley et al. identified risk and resilience factors for mental health in transgender young people. Risk factors were reported as: (1) physical and verbal abuse, (2) exposure to discrimination, (2) social isolation, (3) poor peer relations, (4) low selfesteem, (5) weight dissatisfaction and (6) age, with older adolescents identified at increased risk (Tankersley et al. 2021). Resilience promoting factors in the systematic review included: (1) parent connectedness, (2) social support, (3) school safety and (4) the ability to use the preferred name (Tankersley et al. 2021). Parent connectedness in this context refers to the closeness between the transgender young person and their parent or caregiver.

Looking in more detail at parent connectedness and family relationships could identity resilience promoting behaviours, which support gender diverse young people. Positive relationships between young people and their parents in some of the reviewed studies, promoted resilience in young people who were seeking gender affirming treatments and those who were not. Gower *et al.* (2018) reported lower odds of depression, suicidality and suicide attempts in young people with greater parental connection. In 2013, Simons *et al.* emphasised the positive association of parental support with a higher quality of life for transgender adolescents, highlighting the value of interventions, which promote parental support. Psychosocial or psychological interventions involve therapies or actions used to help gender diverse young people assimilate healthily with their families into society. While there is research evidence to highlight the positive impact of family acceptance on young people's mental health (Olson *et al.*2016), little is known about how to best provide psychological and psychosocial support to families of gender diverse young people. This systematic review aims to address this current gap in knowledge.

The Review

Aims

- To review the evidence on psychological and psychosocial support for families of gender diverse youth.
- To review established interventions/therapies for families of children and young people exploring gender diversity
- To determine the effectiveness of the above interventions/therapies in improving family relationships and mental wellbeing for transgender young people

Design

Systematic reviews have been described as the gold standard among reviews, collecting research in a systematic, transparent and reproducible way (Davis *et al.* 2014). High quality systematic reviews can avoid unnecessary primary research (Gough *et al.*2015). In transgender healthcare, research studies are frequently conducted with small sample sizes in dispersed specialist centres. Gender diverse individuals have been the subject of a range of research studies, with some transgender individuals reporting research participation fatigue (Ashley, 2021).

Protocol and registration

This systematic review is based on the published protocol: Psychological/psychosocial interventions for gender diverse youth under eighteen years of age and their families: a systematic review: PROSPERO 2020 CRD42020163995. The protocol states that risk of bias assessments will be completed using the ROBIS tool (Whiting *et al.*2016). On further evaluation, the ROBIS tool was found to be unsuitable for this type of systematic review as it focuses on assessing risk of bias in systematic reviews rather than primary research studies. This is a deviation from the original protocol, with the Mixed Methods Appraisal Tool (MMAT) developed in 2018 by Hong *et al.* chosen to assess risk of bias in its place. As the protocol had been published in 2020, the inclusion period for studies has been adjusted from (2000-2020) to 2001 to 2021 to capture the last 20 years of research.

The eligibility criteria are clearly stated in the systematic review protocol in terms of the population, interventions, comparators and outcomes. The population are families of gender diverse youth, defined as children or adolescents under the age of eighteen years of age with a parent or caregiver. Interventions are defined as any psychological or psychosocial intervention, offered to gender diverse youth and their families. If studies include comparisons between groups or within groups,

comparators are defined as other types of psychological, psychosocial interventions or no treatment. Outcomes in this review include the identification of relevant interventions and effectiveness of these interventions.

Search methods

An electronic literature search was conducted using OVID MEDLINE; EBSCO CINAHL; ProQuest MEDLINE; OVID PsycINFO. Only studies published in English from 2001-2021 were included in the search. The last twenty years were used as the period for the assessment of the most recent subject material, which is likely to include or relate to the specified population. The search strategy was developed based on key terms. Search terms were reviewed with guidance from the university subject librarian. The search strategy terminology was adjusted slightly in keeping with search terminology of each data base.

Example of search strategy

- (1) exp Family/
- (2) exp Parents/
- (3) siblings mp.
- (4) (famil* or parent* or sibling* or brother* or sister* or father* or mother* or grandparent* or grandmother* or grandfather* or guardian* or spouse* or adopt*) mp.
- (5) 1 or 2 or 3 or 4
- (6) transgender persons mp.
- (7) Exp transsexualism/
- (8) (transgender* or transsexual* or gendervariant* or gender divers* or gender incongruent* or non-binary or gender dysphor* or genderqueer or queer or genderfluid) mp.
- (9) 6 or 7 or 8
- (10) 5 and 9
- (11) Limit 10 to yr="2001-2021"
- (12) (intervent* or treatment* or therap* or education* or support*) mp.
- (13) 11 and 12

Search outcome

During the data collection process, the protocol authors independently screened titles and abstracts identified through the electronic searches and compared them against the inclusion criteria. The authors accessed full text studies if the titles and abstract met the inclusion criteria or if there was uncertainty.

Both reviewers screened full text articles and made decisions about the inclusion of the study. Any disagreements were resolved through further discussion.

Quality appraisal

Assessment of the methodological quality is the key to ensuring that systematic reviews are trustworthy, valid and reliable (Harden and Gough, 2012). The MMAT version 2018 was developed to focus on reporting, conceptual and methodological aspects of mixed methodological studies (Hong *et al.*2019), highlighting potential bias in each area. The MMAT tool 2018 was piloted prior to use. Both authors independently followed the algorithm for study categories and completed the appropriate sections of the MMAT (Hong *et al.*2019) for each included study. Disagreements were resolved through further discussion.

Data abstraction

Abstracting the quantitative element of a study is relatively easy as findings are based on means, odds ratios or other numerical concepts. In qualitative studies, direct quotes from participants constitute the raw data of a study and it can be difficult to identify key concepts or summaries in some studies, in particular if the authors have only described and summarised what participants have reported (Thomas and Harden, 2008). In this review, information contained in the results section of each paper was extracted. This included qualitative and quantitative elements. Data from each study was independently extracted and the main findings of each study were summarised under the headings: (1) author/date, (2) inclusion/exclusion, (3) sample size, (4) participant demographics, (5) study methodology, (6) intervention, (7) method and (8) key findings. Tables 1-4 provide a summary of each included study.

Synthesis

Synthesising qualitative research is a complex area, with ongoing debates about appropriate methods and whether qualitative data is too specific in terms of context, time and participants to be generalisable (Thomas and Harden, 2008). This has led to suggestions, that qualitative synthesis de-contextualises findings (Sandelowski and Barosso, 2006). Thematic analysis of primary qualitative data follows a structured process: (1) familiarisation with the data, (2) initial line by line coding, (3) sorting of initial codes into broader themes, (4) review of emerging themes and (5) renaming and refining codes (Braun and Clarke, 2006). Thematic synthesis has been developed to analyse secondary data, following a number of steps: (1) line by line coding, (2) the development of descriptive themes close to the primary studies and finally (3) generation of new hypothesis (Thomas and Harden, 2008).

Due to the small number of included studies, there was limited data to analyse. Qualitative data analysis was thematic and were possible thematic synthesis steps were followed. Data was initially described on the basis of the PICO criteria- families of gender diverse youth (population), psychological/psychosocial support group (intervention) and outcomes. As none of the studies included a comparison group, the comparator aspect was removed. The limited quantitative data is reported separately.

Results

Study selection

Overall 8,405 studies were independently screened. By the screening of titles and abstracts 8,292 studies were excluded, while 113 studies were screened for further detailed analysis, from which 109 studies did not meet the eligibility criteria. Only four studies met inclusion criteria for the review. Figure 1 outlines the study selection process based on PRISMA guidance. The following results are organised and expanded on the basis of the PICO criteria. Quality appraisal of each study was conducted using the MMAT tool (Hong *et al.*, 2019). This highlighted potential risks of bias in each study, in terms of for example missing data (Caldarera *et al.*, 2021) and missing standard deviations (Di Ceglie and Thümmel, 2006). In other studies it was impossible to know whether there was coherence between data sources and analysis (Menvielle and Darryl, 2010) or whether the risk of non-response bias was low (Hillier and Torg, 2019). As these studies represent the limited available research available on family interventions for transgender youth, the quality of included study was accepted as basis for this review.

Population

All included studies were focused on treatment seeking transgender children and young people who were involved with specialist gender services, or waiting for input from specialist gender services. Services were based in the United States (US) (Hillier and Torg, 2019; Menvielle and Hill, 2010), Italy (Caldarera *et al.*2021) and Great Britain (Di Ceglie and Thümmel, 2006). Services catered for large catchment areas, ranging from citywide (Hillier and Torg, 2019), regional (Caldarera *et al.*2021), national (Di Ceglie and Thümmel, 2006) and international covering 34 states in the US and six countries (Menvielle and Hill, 2011). Participating parents had children and young people between the ages of four and seventeen years six months old, with only one study (Hillier and Torg, 2019) including parents of young people up to the age of twenty two years of age. This study included nine parents of eighteen to twenty two year olds. As it was impossible to separate the experiences of this small subsection, the study was included in the review. Parents (including step, foster, adoptive and legal guardians) of children and adolescents were offered access to the psychological/ psychosocial support group interventions regardless of the age of their young person, which meant grouping parents of children and adolescents together. For one study (Menvielle and Hill, 2011), young people had to meet clinician

assessed DSM-IV-TR (American Psychiatric Association, 2000) criteria to be included in the study. This is not to suggest that young people in other studies did not meet the respective DSM criteria, but acceptance to the service was seen as sufficient for participation. Participation in the studies was voluntary, but in one study (Menvielle and Hill, 2011), parents were paid \$37.50 for taking part in the study. Di Ceglie and Thümmel (2006) offered reimbursement of travel costs through a charitable donation to participants, who were not eligible to have travel costs re-reimbursed through other means.

Intervention

Most of the included studies provide details about the group intervention offered to parents, apart from Hillier and Torg (2019), who only make reference to the support group, without providing specific details. Groups were led by clinical staff, some of whom had other types of contact with the families in the studies. Hillier and Torg's (2019) study was conducted by two parents who were attending the monthly support groups, while the other studies were conducted by clinical staff who were leading the support groups. The definition of intervention differed across studies. Participants in the studies by Di Ceglie and Thümmel (2006) and Caldarera et al. (2021), attended at six monthly parent groups (Di Ceglie and Thümmel, 2006) and up to eleven monthly parent groups (Caldarera et al. 2021). The other studies specified one or more attendance at the monthly support group (Hillier and Torg, 2019) and a range of different interventions on their own, or in combination, such as monthly parent groups, listery (internet forum) contact and website access (Menvielle and Hill, 2010). Parent groups lasted between sixty to ninety minutes (Caldarera et al. 2021; Hillier and Torg, 2019; (Di Ceglie and Thümmel, 2006). No details about the group length were available for Menvielle and Hill (2010). The group formats were therapeutic, with some educational elements (Di Ceglie and Thümmel, 2006). Parent groups were closed in two studies (Di Ceglie and Thümmel, 2006; Caldarera et al. 2021) with consistent group membership and open in the other two studies.

Comparators

None of the studies included comparison of the intervention with treatment as usual or other interventions.

Outcomes

Studies used a range of quantitative and qualitative outcome measures for parents. None of the measures used were standardised tools. In terms of feedback surveys, studies included quantitative elements, based on closed questions and open ended questions, which invited parents to include free text, which was thematically analysed. Menvielle and Hill (2011) was the only study which was based on interview data.

Quantitative outcomes

Two studies (Di Ceglie and Thümmel, 2006; Caldarera *et al*.2021) contained the same questions in their survey, in terms of helpfulness of the group, meeting other parents, reduced isolation and parents having a changed approach as a result of the group. This was deliberate as the second study was based on previous research. Both studies used very similar interventions, offering a closed therapeutic group with similar sample sizes in their studies. A comparison between studies, highlights a mean difference of 0.25 (helpfulness of group, 0.4 (meeting other parents), 0.38 (reduced isolation) and 1.35 (changed approach as a result of the group). Mean scores for Di Ceglie and Thümmel (2006) were marginally higher for all items, but no standard deviations were reported. Data was collected after 6 months (Di Ceglie and Thümmel, 2006) compared to after 12 months (Calderera *et al*.2021).

Hillier and Torg (2010) collected information about the importance of the support group for family members on a Likert scale containing four categories (ranked from not particularly helpful to single most important). In this study authors reported proportions in each category, with a majority of 35/48 (73%) indicating importance or critical importance in terms of the support group.

Theme 1- Reduced shame and isolation: Participants reported reduced shame about their child or young person's gender identity (Caldarera et al.2021) and felt less isolated (Caldarera et al.2021; Di Ceglie and Thümmel 2006; Hillier and Torg, 2019). Listening to others also reduced anxiety (Hillier and Torg, 2019). Participants reported the value of emotional support, in particular the opportunity to build friendships with other families (Menvielle and Hill, 2011). Participants in Hillier and Torg's (2019) study expanded on the concept of friendship, describing other participants as friends or chosen family. Parents took the role of supporters to other parents who had less experience with their transgender child (Hillier and Torg, 2019).

Theme 2- group processes: For some participants, the group format itself brought challenges, which differed in terms of the group set up. In the open group set-up by Hillier and Torg (2019), some participants felt the group had become less personal and overcrowded. It is unclear what group sizes were offered by the service. In the closed group set-up by Di Ceglie and Thümmel (2006), some participants reported that other group members dominated interactions. Some participants were critical of the lack of racial diversity in the group (Hillier and Torg, 2019), while group formats in other services were set up specifically for interracial families (Menvielle and Hill, 2010). The logistics of attending the group were reported as difficult by some participants due to physical distances, requiring overnight stays for some (Hillier and Torg, 2019).

Theme 3- Looking into the future: Participants in Di Ceglie and Thümmel's (2006) study, valued talking to others about different approaches in managing their children and different outlooks into the future. Participants in Menvielle and Hill's (2010) study reported that taking part in the group allowed them a window into the future, preparing them for what might be ahead of them. Combining parents of children and young people at different developmental and transition stages in a group also brought challenges, with some parents presenting a terrifying and gloomy picture of the future (Di Ceglie and Thümmel, 2006) to others. This experience was shared by participants in (Hillier and Torg, 2019) who requested a splitting of groups by age, feeling overwhelmed after having to listen to other parents talk about surgical procedures when they were only beginning to get used to different pronouns.

Theme 4- increased understanding of transgender young person's experience: Participants reported an increased understanding about their child's experience in terms of navigating their development in the context of gender incongruence (Di Ceglie and Thümmel), while others reported gaining knowledge in terms of medical, legal and political aspects of parenting a transgender child (Hillier and Torg, 2019). Parents in the study by Caldera *et al.* (2021) reported the importance of increased awareness of difficulties faced by gender diverse young people and the legal issues they may face.

Theme 5- parents as advocates for transgender youth: Participants in Menvielle and Hill (2010) reported feeling more open to discuss difficult topics with their child, with the group intervention serving as a reference point. In the study by Caldarera et al. (2021), participants reported that mutual learning created better insights into different ways of coping with the same situation, while also decreasing parental sense of guilt. In some of the studies, parents made specific reference to the benefit of support from clinical staff in the group, which some described as sympathetic and professional (Di Ceglie and Thümmel, 2006; Caldarera et al.2021). For participants, having access to a network of knowledge from other parents and clinicians increased their knowledge about specific doctors and health insurance coverage, enabling them to effectively advocate for their children (Hillier and Torg, 2016).

Discussion

Family support is a known protective factor against health risks for transgender youth (Simons *et al.*2013; Olson *et al.*2016). This review reported on group interventions for parents of transgender youth. The total sample of participants in the review was 114, highlighting the limited available amount of research studies in this area. None of the included studies measured outcomes in terms of transgender youth and it is unclear if any transgender young people benefited from parents attending the group interventions. Two studies (Di Ceglie and Thümmel, 2006; Caldarera *et al.*2021) completed an

evaluation of a structured closed group intervention, indicating benefits for parents in terms of increasing their understanding, meeting other parents, feeling less isolated and changing their approach to managing their child's gender diversity. As parents and their young people received other supports from services, it is unknown whether the group intervention created the reported change. It is unclear whether the context of the group interventions based in specialist gender services impacted on the reported outcomes by parents. Clinicians in specialist gender services are put in the position of gate-keepers to physical interventions. Lev (2009) argues that the gate-keeping role impacts on the building of healthy therapeutic relationships. It unknown whether facilitation of the groups by clinical staff was viewed by parents as part of a wider assessment process for their children's eligibility to access gender affirming interventions and if this influenced their feedback about the groups.

Studies highlighted the need to address potential negative group dynamics (Di Ceglie and Thümmel, 2006; Hillier and Torg, 2019), in terms of parents having different needs, based on the developmental stages of their children and the risk of some parents dominating discussions. This highlights the need to explore the role of group facilitators and whether groups should be managed by clinical staff, parents or both. Group size and group membership were also crucial aspects for some parents. While participants attended between one and eleven group sessions in the studies, it is not clear whether the number of group attendances is important for parents. Lawlis *et al.* (2020) completed a survey of parental preferences for group interventions, which highlighted that the majority of parents liked monthly meetings, with a group size of between six and ten people, with the majority of parents happy to attend the clinical service base for the group as long as travel time to the group was between eleven and twenty minutes. This highlights many common barriers relevant to all types of group attendances in terms of time and convenience. Some parents reported reduced need for the group intervention over time (Hillier and Torg, 2019), which could be directly related to the positive impact of the intervention or due to natural development processes in the child which reassured the parent.

Measuring the impact of group interventions is complex and none of the studies in the review used before and after measures. It is therefore unknown whether group interventions impacted on parenting practices or family dynamics. Brown *et al.* (2020) in their review of family relationships of transgender youth, differentiate between general family support and gender-identity specific support. All included studies offered a gender-identity specific group intervention to parents. Brown *et al.* (2020) argue that while both types of family support can be protective, it is unclear if the need for general family and gender identity specific support varies within family members.

The studies by Menvielle and Hill (2010) and Hillier and Torg (2019) focused a group intervention as part of a range of activities of an affirmative service programme. The use of incentives by Menvielle and Hill (2010) and the participation of the researchers as parents in the group (Hillier and Torg, 2019) may have introduced recruitment bias. As all the studies were based in specialist gender

services, group interventions may have been offered to parents who were at least somewhat supportive of the gender diversity of their children. This could have excluded parents who did not support their children's gender diversity or who were at the very early stages of their child, questioning their gender identity. As there were no comparison groups, this cannot be established. It is also not clear whether clinical staff input or parental input into the groups made a difference to the experience, as groups tended to be facilitated by clinical staff, but parents reported benefits in meeting other parents. Studies highlighted risks and benefits of incorporating parents of children and adolescents but it is unclear whether parents of young children would have received greater benefit without hearing the potential frightening aspects of adolescence and whether parents of adolescents gained no benefit themselves hearing about childhood experiences, they had already managed.

Conclusion

This review highlights the limited available research in terms of family interventions which could increase resilience in transgender youth and protect them from potential adverse mental health outcomes. This review compares and summarises available evidence of group family interventions, which show a positive impact on parents in terms of reducing their isolation and meeting their needs for knowledge in managing their transgender young person.

Looking at the needs of families of gender diverse youth highlights that they experience marginalisation alongside their young person. From a minority stress theory point of view, parents are struggling to meet the needs of their transgender young person in a family and community context which could be unwelcome or even hostile. The group interventions for many parents reduced their isolation and created a new network of supportive adults. Whether this in turn has enabled parents to create more psychologically healthy and supportive home environments for their children is unclear. There is a need for further research to investigate whether families require general family interventions or specific gender identity focused family interventions.

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Table 1: Study 1: Caldarera et al. (2021)

Author/ date	Caldarera, A.M., Davidson, S., Vitiello, B. and Baietto, C., 2021. A psychological support group for parents in the care of families with gender diverse children and adolescents. <i>Clinical child psychology and psychiatry</i> , 26(1), pp.64-78.					
Inclusion/ Exclusion	Psychological support group for parents of gender diverse children and adolescent who were attending the service.					
Sample size	14 parents of children between 8 and 17 years old. Full data available for 11 parents only.					
Participant demographic s	Parents of gender diverse children between the ages of 8 to 17 years who spoke Italian fluently. Parental education levels ranged from high school to university masters level.					
Study methodology	Data collection: Stage 1 (after 6 months- T1) – parents completed short form about relevant topics Stage 2 (after 12 months/ end of study-T2)- parents completed feedback					
Into many ortions	questionnaire					
Intervention	11 monthly closed psychological support group (group therapy and psycho educational techniques), lasting 90 minutes, over 12 months, for parents whose children were attending the gender identity service. The group was led by clinical staff at the gender service.					
Method	Data collection at: 6 months – short form indicating main relevant topics 12 months- repeat of short form & semi-structured feedback questionnaire Thematic analysis					
Key findings	-Importance of sharing experiences with other parents -gender identity as a process, which evolves over childhood and adolescence -child/ young person's relationships within and outside the family -Evolution of themes across time- parents taking a more complex perspective on gender diversity and needs of their young person; parents more able to deal with uncertainty related to process of child's gender development.					

Table 2 Study 2: Di Ceglie and Thümmel (2006)

Author/ date	Di Ceglie, D. and Thümmel, E.C., 2006. An experience of group work with					
	parents of children and adolescents with gender identity disorder. Clinical					
	Child Psychology and Psychiatry, 11(3), pp.387-396.					
Inclusion/	Parents of children or young people who were assessed and were involved					
Exclusion	in treatment at the service.					
Sample size	10 parents of seven children between 7 and 17 years old.					
	Data available for 7 participants (couples returned one joint questionnaire)					
Participant	Two couples, five mothers and one aunt of gender diverse youth. No					
demographic	information about parental education levels.					
S						
Study	Evaluation questionnaire after 6 months- end of the study.					
methodology						
Intervention	6 monthly closed psycho dynamically informed support group, lasting 90					
	minutes, over 6 months, for parents whose children were attending the					
	gender identity service. The group was led by clinical staff at the gender					
	service.					
Method	Data collection at: 6 months – short form indicating main relevant topics					
	Thematic analysis of qualitative responses.					
Key findings	-Reduced sense of isolation through meeting other parents with children					
	similar to their own					
	-Helpful to learn about different outlooks and approaches to dealing with					
	their children.					
	-Least helpful- tendency for some parents to dominate; difficulty					
	understanding the way staff explained things; lack of parents whose child is					
	of similar age; focus of group on changing parental perception rather than					
	treating gender dysphoria					

Table 3- Study 3: Hillier and Torg (2019)

Author/ date	Hillier, A. and Torg, E., 2019. Parent participation in a support group for families with transgender and gender-nonconforming children: "Being in the company of others who do not question the reality of our experience".					
	Transgender Health, 4(1), pp.168-175.					
Inclusion/	Parents of gender diverse youth who are receiving care at one of the					
Exclusion	participating services and who are referred to the support group by a					
	member of staff. Participation in at least one monthly psychological					
	support group.					
Sample size	48 parents/ guardians of 48 children between 5 and 22 years old.					
Participant	Participants were at least 18 years old and were biological, step, foster,					
demographics	adoptive or legal guardians of gender diverse youth attending either of the					
	two services.					
Study	Online cross-sectional survey through REDCap of parents who participated					
methodology	in at least one monthly support group identified through email listsery or					
	secret Facebook group.					
Intervention	Thematic analysis of qualitative content.					
	Monthly psychological support group, lasting 90 minutes, for parents					
	whose children were attending the gender identity service. The group was					
	led by two parents of transgender children. While parents attended their					
	support group, their children and siblings of their children attended separate					
	support groups facilitated by clinical staff.					
Method	Quantitative analysis of closed ended questions using SPSS v. 25.					
	Qualitative themative analysis to identity patterns.					
Key findings	-Emotional support for the whole family					
	- children were able to develop friendships with other transgender children					
	through their support group					
	-parents gained knowledge about medical, legal, political aspects of					
	parenting trans child from other parents					
	-limitations- parent group perceived by some as too large, overcrowded,					
	less personal, not racially diverse, no other children similar in own to own					
	child.					

Table 4- Study 4: Menvielle and Hill (2010)

M 11 E 1III DD 2010 A CC 21 1 C C 21					
Menvielle, E. and Hill, D.B., 2010. An affirmative intervention for fan					
with gender-variant children: A process evaluation. Journal of Gay &					
Lesbian Mental Health, 15(1), pp.94-123.					
Parents affiliated with the service whose child met GID criteria (assessed					
through clinician telephone or live interview).					
42 parents (26 couples, 16 parents) of 31 children between the ages of 4 to					
17.5 years.					
Parental involvement with the service ranged between one month to 7					
years. Involvement ranged from freceiving listserv emails to attending					
monthly parent group over 2 years). Parents were 80% White, 10%					
Hispanic, others identified as Black or multiracial. Participants lived					
throughout the US.					
One off telephone interviews. Participants were paid \$37.50 for their					
participation.					
Participants had between 1 month to 7 years involvement with the service,					
ranging from monthly in-person parent group attendance to accessing					
listserv resources online (which included the parent guide). Children of					
parets who attended the in-person group had access to a child group.					
Semi-structured open-ended telephone interviews lasting between 20 to 80					
minutes. Parents were recruited from the in-person parent group at the					
centre or from the email listsery. Data was analysis of verbatim interviews					
using NVIVO 7.					
-increased understanding of gender identity and the ability to educate other					
people					
-accessing information about future challenges					
-reluctance to participate for fear of stigmatization					
-negative emotions (sadness and loss at confronting reality					
-affirmation					
-for children accessing the group- finding like-minded peers					
-coping with social exclusion					

Table 5 - Risk of Bias based MMAT version 2018 (Hong et al. 2019)

Study	Study design	Methodological quality criteria	Responses		
	8		Yes	No	Can` t tell
Study 1 Caldarer a et al. (2021)	Quantitativ e non- randomise d	Participants representative of target population	V		
		Measurements appropriate regarding outcome and intervention	1		
		Complete outcome data		V	
		Confounders accounted for in design and analysis			1
		Intervention administered as intended	1		
Study 2 Di	Quantitativ e	Sampling strategy relevant to research question	1		
Ceglie and	descriptive	Sample representative of target population	V		
Thümme		Measurements appropriate			$\sqrt{}$
1 (2006)		Risk of non-response bias low	$\sqrt{}$		
		Statistical analysis appropriate			$\sqrt{}$
Study 3 Hillier and Torg (2019)	Quantitativ e	Sampling strategy relevant to research question	1		
	descriptive	Sample representative of target population	1		
		Measurements appropriate	V		
		Risk of non-response bias low			
		Statistical analysis appropriate			
Study 4 Menviell	Qualitative	Approach appropriate to research question	V		
e and Darryl		Methods adequate to address research question	V		
(2010)		Findings derived from data	1		
(2010)		Results substantiated by data	Ì		
		Coherence between data sources,	Ì		
		collection, analysis and interpretation			



Figure 1: PRISMA Flow Diagram (Moher et al. 2009)

