The Risk Reference Panel: a thematic analysis of a multidisciplinary forum for complex cases

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BACKGROUND

The assessment and management of the risk that a patient poses to themselves and others is a core duty for psychiatric services. There are several ways in which this is achieved, depending on the complexity of the case, the configuration of local services and resources available. In many cases, issues around risk are managed effectively by individual practitioners or teams on a day-to-day basis, using evidence-based guidance from national bodies such as the Royal College of Psychiatrists or the Department of Health, as well as locally agreed policies and procedures and clinical supervision. However, in some circumstances the risks posed may be exceptional or unclear, and individuals can engage in high levels of self-harm or antisocial behaviour. This can be analysed retrospectively by complex case reviews to reduce risk of recurrence, but prospective prevention or minimisation of harm is clearly preferable.

THE RISK REFERENCE PANEL

Multidisciplinary working is a key strategy for particularly risky cases, which can take various forms. From 2008 the Aneurin Bevan University Health Board Risk Reference Panel was developed by Dr Alan Wright, Forensic Psychologist, and author I.E., Forensic Psychiatric Nurse, as a local response to the public and professional anxiety surrounding exceptionally risky cases. The panel is intended to provide a pool of multidisciplinary expertise with representation from social work, nursing, forensics, intellectual disability and general adult psychiatry. To date, the panel has not included patient representation, but this could be considered in future. This paper offers an analysis of a way of addressing risk and management of complex cases by describing the types of cases that have been presented and the outcomes of panel meetings. A retrospective thematic analysis of the meetings was conducted.

AIMS AND METHOD

To describe the functions of the Aneurin Bevan University Health Board Risk Reference Panel and characterise the typical referrals presented and outcomes from the panel. A structured thematic analysis was performed on verbatim transcripts of 48 panel sessions.

RESULTS

The 79 codes identified were grouped into 16 subthemes. Four principal themes were identified: two characterising cases brought to the panel (childhood risk factors and current presenting difficulties) and two describing advice given (risk management and wider organisational issues). Quotations are given to illustrate cases presented and advice given by the panel.

CLINICAL IMPLICATIONS

The panel provides a valuable source of special expertise in the management of complex and exceptional cases where risk of harm is significant and clinical teams have ongoing concerns. This paper describes a model of peer-working to support staff and may further reduce the risk of harm associated with mental illness.

DECLARATION OF INTEREST

None.

KEYWORDS

Clinical governance; community mental health teams; forensic mental health services; qualitative research.
constitutes thematic analysis and its methodology. For our purposes, we have followed the model proposed by Braun and Clarke, which describes six discrete stages of thematic analysis.

The stages of the above method as applied to our analysis are described in the Appendix.

Discrete ideas (referred to in the process as ‘codes’) and broader groupings of these ideas into key patterns (‘subthemes’) were identified. The subthemes can then be grouped together to identify the main themes within the data.

**Data collection**

The Risk Reference Panel meetings are attended by a secretary who records what is said by typing verbatim as far as is possible; there is no audio recording of the meetings. These transcripts are then used to produce letters to the referring team, outlining what was discussed and the recommendations of the panel.

A pilot analysis was conducted whereby authors A.T. and G.J. independently applied the method described below to a sample of original transcripts for 8 of the 48 cases that had been presented to the Risk Reference Panel between May 2010 and April 2016. Ethical approval was not required as our study is a retrospective service evaluation where no change to the service was implemented and no service users were interviewed. All reported data are anonymised. Anonymised samples of the transcribed data, as well as examples of the codes generated from them and how they were recorded, are given in Boxes 1 and 2.

The pilot analysis generated an initial list of codes that were used as the basis for coding the rest of the transcripts, as well as demonstrating an acceptable consensus on what constituted individual codes. The remaining 40 transcripts were then coded, 20 each by authors A.T. and G.J., and the frequency of codes were recorded. The resultant database allowed the rest of the thematic analysis to be completed. A thematic map provided a diagrammatic representation of the various themes, subthemes and relationships between them (Fig. 1).

A feedback form allowed participants to indicate, via a Likert scale, the degree to which they agreed with a number of statements (outlined under ‘Feedback’ within Results) and to provide free-text feedback.

**Results**

Using this method, 16 subthemes were identified to group similar codes. The subthemes were further grouped into four principal themes: childhood risk factors, current presenting difficulties, assessment and management of risk to self or others and team or wider organisational issues. The first two themes encapsulate information presented to the panel during presentations by referrers; the latter describe recommendations made or issues identified by the panel.

The subthemes are organised by frequency of codes, referred to hereafter as occurrences, with absolute frequencies (i.e. the number of times a code occurred in the whole of the transcribed data, given as n) and the proportion of occurrences within that theme (given as a percentage). Percentages are rounded to the nearest 0.5%. There were a total of 522 occurrences of codes throughout the 48 transcripts analysed.
**Childhood risk factors**

The first theme contains codes pertaining to the history of the case, outlining risk factors pertinent to the current problem. There were four individual codes that accounted for 31 of the 522 occurrences (i.e., 6% of all occurrences). These were primarily adverse experiences that the person had endured in childhood, but also included childhood traits that are associated with subsequent antisocial behaviour.

The most common childhood adverse experience was being a victim of physical abuse (n = 14, 45% within theme) or sexual abuse (n = 11, 35%), followed by having been in care (n = 3, 10%). A history of aggressive behaviour as a child was also identified in a number of individuals (n = 3, 10%).

**Current presenting difficulties**

This theme accounted for the greatest proportion of individual codes, with 245 occurrences (47%). Five subthemes were identified within current presenting difficulties, namely nature of offences, problematic behaviour, factors affecting current presentation, police/criminal justice system involvement and established diagnoses.

The nature of offences subtheme explored the various types of offences committed, if applicable (only some of the cases presented had criminal convictions, hence their low frequencies). The offences captured in the transcripts included arson (n = 3, 1%), non-assaultive sexual offences (n = 3, 1% within theme), threats to kill (n = 1, 0.5%) and sexual assault (1, 0.5%).

The most common problematic behaviour was suicide attempts/self-harm (n = 32, 13%), which was often seen to be chaotic and instrumental. However, there were several cases where severe self-harm without suicidal behaviour was the primary problem. The next most frequent was a lack of engagement with services (n = 19, 8%). The remaining codes within the problematic behaviour subtheme were sexually inappropriate behaviour (n = 7, 3%), antisocial behaviour (n = 7, 3%), disruptive behaviour (n = 6, 2.5%), harassment of care-giving staff (n = 6, 2.5%), multiple contacts by phone (n = 6, 2.5%), absconding risk (n = 6, 2.5%), complaints to health board/Welsh Assembly Members/counsellors (n = 5, 2%), false allegations (n = 4, 1.5%) and litigation (n = 1, 0.5%). Although the most prevalent individual code was risk of self-harm or suicide, the combined frequency of codes relating to risk to others was around 1.5-times as high.

Another subtheme was factors affecting current presentation, which captured issues in the person’s life at present that were affecting their level of risk. Substance misuse was the most prominent among these (n = 27, 11%). Other codes within this subtheme were chronic pain/illness/physical issues (n = 7, 3%), social isolation (n = 7, 3%), multiple admissions to psychiatric hospital (n = 7, 3%), homelessness/moving area frequently (n = 6, 2.5%), family history of mental health issues (n = 5, 2%), admissions under the Mental Health Act (n = 5, 2%) and lack of insight (n = 5, 2%).

Less commonly, people had problems with anger (n = 3, 1%), bereavement (n = 3, 1%), sexuality (n = 3, 1%) and personality traits without a diagnosis of personality disorder (n = 2, 1%). There was police involvement in nine cases (3.5%) and the person was subject to the criminal justice system in 15 cases (6%).

Most of the cases presented to the panel had an established diagnosis, although within the 60 code occurrences in this theme, ‘conflicting/unclear diagnosis’ occurred 13 times (21.5%). These subthemes grouped the diagnoses by psychopathology.
'Developmental disorders' were the most common presentation and included autism spectrum disorder (n = 8, 13%), intellectual disability (n = 5, 8%), attachment disorder (n = 5, 8%), 'special needs' (n = 3, 5%) and attention-deficit hyperactivity disorder (n = 3, 5%). Emotionally unstable personality disorder (EUPD) was the only specified personality disorder diagnosis (n = 9, 15%) and the most common individual pathology. This was followed by psychotic illnesses, with 'treatment-resistant psychosis' accounting for seven cases (11.5%). Schizophrenia was only given as a named diagnosis in one case (1.5%). Lastly, affective/anxiety disorders included depression (n = 3, 5%), eating disorders (n = 2, 3%) and post-traumatic stress disorder (n = 1, 1.5%).

**Risk assessment and management**

There were 31 occurrences of codes (6%) related to an assessment of the level of risk posed to others by the individual. Three of these (9.5% within theme) were an observation that there was a poor understanding of the risks posed and that specific further information was required.

Much more of the output from panel meetings was regarding the management of specific risks to the person or others, with 95 code occurrences. Within this theme, four subthemes were identified: need for further assessments, specialist care plans, capacity/placement factors and responsibility of other services.

Most commonly, the panel advised that specific assessments would be beneficial to manage risk. In disorders that modulated risk, such as autism spectrum disorder, the panel would often suggest 'expert/specialist opinion required' (n = 17, 18%). If the risk was more general, then suggestions of which avenues to pursue included a need for HCR-20 (Historical Clinical Risk Management-20; an established actuarial tool for assessment of violence risk) (n = 3, 3%); legal advice (n = 4, 4%) and a need for more background information (n = 11, 12%) or updated formal risk assessment (n = 8, 8%).

Issues regarding the assessment of capacity, or the established lack of capacity, were also commonly cited. There were 14 occurrences (15%) of recommendations that a Mental Health Act assessment had not been considered and could be of use in containing the risks posed. Further to this, there were 13 recommendations (13.5%) that the acute problems should be managed as an in-patient.

Several of the cases indicated longer-term in-patient or other compulsory interventions were necessary, such as the need for low secure placement (n = 5, 5%) and management under guardianship (n = 2, 2%) or Protection Of Vulnerable Adults scheme (n = 2, 2%).

Team issues within the context of risk management were needed to improve engagement (n = 1, 1%), need to take positive risks (n = 1, 1%) and need for child protection training for team members (n = 1, 1%).

In eight cases (8%) the panel identified that responsibility for the patient's actions did not lie with health services as they were independent of mental health issues. These were divided into a recommendation that their behaviour be dealt with under the criminal justice system (n = 4, 4%), or an acknowledgement that although there is no criminal element to pursue in some cases, all appropriate actions to reduce risk have been considered and there is no more that the team can realistically do (n = 4, 4%).

Occasionally, the panel recommended specific management plans to manage particular disorders. Most frequently this was stated as a need for a specialist care plan (n = 24, 77%), incorporating advice from specialist assessments. The need for structure in the patient's life was identified (n = 3, 10%), as was the need for family work/family meeting (n = 1, 3%). More specifically, clozapine for treatment-resistant psychosis (n = 2, 6%) and dialectic behaviour therapy for EUPD (n = 3, 10%) were suggested.

**Team/wider organisational issues**

The final theme generated by the analysis contained 29 occurrences (0.5%) between five codes, and was subdivided into team working issues (n = 8, 27.5%), funding issues (n = 7, 24%), need for special staff/staff rotation (n = 6, 20.5%), need extra support for the staff involved (n = 5, 17%) and trust management issues/needs escalation (n = 3, 10%).

Team working issues captured differences within the team that may have hindered progress in the management of particular behaviours. For example, the panel acknowledged that because of splitting in the context of EUPD, it was necessary to establish a clear and consistent boundary narrative. However, outright disagreement within teams, over significant issues such as primary diagnosis and level of risk posed, was identified as an issue.

Funding issues identified that responsibility for funding was not always clear and had delayed the provision of appropriate placement. This was particularly relevant where the person had moved frequently between areas or was of no fixed abode.

The panel stressed that the staff who were dealing with particularly demanding individuals needed extra support themselves, which can often be neglected, and suggested rotation of staff to prevent burnout. In a small number of cases it was identified that the particular service or treatment indicated for a person's condition, such as dialectic behaviour therapy, was not available, and it was suggested that this was escalated on a trust level on the basis that this would reduce risk of harm.

**Feedback**

Feedback was overwhelmingly positive, with 20 out of 22 respondents (91%) indicating agreement or strong agreement with the following statements: ‘New ideas were generated in the session’, ‘The case feels safer as a result’, ‘I learned a great deal during the session’, ‘The questions I brought were answered’ and ‘I would recommend the process to a colleague.’

**Discussion**

The Royal College of Psychiatrists has highlighted a need for improved, tiered risk assessment and interventions on both a local and national level. A qualitative analysis of the content of Risk Reference Panel meetings has illustrated how it provides a potential model for services to address this need.
The panel is an innovative service model that was set up to provide a further level of support for teams who had been looking for support in managing very complex individuals with high-risk behaviour and has provided new direction and specialist advice in the management of these cases.

Most cases described physical or sexual abuse, with many experiencing multiple adverse childhood events (ACEs). These findings support the existing body of work on the relationship between multiple ACEs and subsequent morbidity and risk behaviours, such as that by Bellis et al., which showed that individuals with four or more ACEs were several times more likely to smoke, drink heavily, have low mental well-being and chronic health conditions. This also correlates with well-documented risk factors for self-harm, suicide and interpersonal violence. These data have more general implications on history-taking in risk assessment, with the inclusion of this background information crucial to the working of the panel. The potential for referrers to exaggerate the risk, both consciously and unconsciously, to encourage acceptance by the panel should be acknowledged.

Although a person’s risk of harm cannot be entirely removed, particularly in such complex cases, a thorough understanding of risks posed and their severity is crucial. The panel often identified areas where risk was not fully understood and suggested specific assessments or specialist opinion that would be beneficial. Having said this, the responsibility for risky behaviour, particularly in a capacitated patient, does not lie entirely with mental health teams and it is prudent to identify when all reasonable steps have been taken, or if involvement of the criminal justice system is more appropriate. Taking the case to such a panel allows an independent debate and makes this decision more defensible.

For patients who remain in the community, it is particularly difficult for teams to provide care and ameliorate risk if engagement is poor. Surprisingly, the meetings often highlighted that capacity assessments and use of the Mental Health Act beyond admission for brief assessment highlighted that capacity assessments and use of the Mental Health Act beyond admission for brief assessment had not been considered. This may reflect a lack of appropriate placements available to community teams, particularly long-term in-patient services when patients cannot be managed in the community.

This model provides a valuable multidisciplinary source of special expertise in the management of complex and exceptional cases, where risk of harm is significant and clinical teams have ongoing concerns. The format may not be appropriate for every service and there is also a place for improved training in formulation and management of patients with complex personality difficulties, but model could certainly be reproduced in other areas where clinical teams feel it would have a beneficial role, particularly to inform Care Programme Approach unmet needs, training needs within the staff workforce and service development needs.

Lastly, it is pertinent to consider alternatives to such a panel. As the heaviest burden appears to be personality pathology, these would primarily consist of improving training in managing patients with personality disorders. In some cases, it may be that this is the better and more economical option, although it has been acknowledged that the panel approach uses relatively few resources in terms of clinician sessions while providing an additional tier of support for community teams to reduce the risk of harm associated with mental illness.

Limitations
This study is only able to describe the function and output of the panel in question, rather than give a rigorous analysis of this model in comparison with other models, and there were no available data on matched people who were not referred. We were also unable to comment on the make-up and training of the teams referring into the panel; it is possible that the referrals represent professional anxiety that stems from sources other than the patient in question, such as gaps in senior management or reluctance to engage with complex individuals with personality pathology, although the panel meetings analysed cover a period of 6 years where it is likely that there would be some fluctuation in team make-up.

All referrals to the panel are made via a written referral letter, which details the history of the case, the assessments that have previously been conducted, a risk assessment and concerns that the referrers wish to address. As the panel will already be familiar with this information, some of it may not have been verbalised at the meetings and thus not captured by the transcription. However, it was noted that there was a verbal presentation of each case and that the relevant history was given.

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Appendix: Thematic analysis (Braun and Clarke) as applied to our study
1. Familiarisation with the data-set: All transcripts were read and re-read by authors A.T. and G.J., considering the application of the coding process to the data and noting initial observations.
2. Generating initial codes: Each transcript was read line by line and discrete ideas within the data were identified and systematically coded. Eight cases were coded by both authors. A.T. and G.J. and the remainder were divided in half between these two authors. Each occurrence of a code was recorded as a simple tally. For example, the history of the patient in
question would be given and each feature would be assigned a different code, such as ‘history of being in care’, ‘history of sexual abuse’, ‘diagnosis of schizophrenia’ or ‘poor engagement with the CMHT.’ Individual risks identified and management suggestions offered by the panel would all produce individual codes.

3. **Searching for themes:** Codes were reviewed and collated into potential themes. It was decided whether each code represented, for example, a demographic feature, a specific risk to his or her self or others, or a suggestion for managing a particular risk. All data relevant to each theme were identified, as some ideas could be given two distinct codes or fit into two or more themes.

4. **Reviewing themes:** Consideration was given to whether the identified themes adequately captured and collated the data in relation to both the individual coded transcripts (i.e. the narration of each Risk Reference Panel meeting) and the entire data-set (i.e. the overarching patterns of input and output for the Risk Reference Panel meetings as a whole). A ‘thematic map’ was generated.

5. **Defining and naming themes:** After satisfactory completion of stage four, each theme was analysed to refine its specific categorisation and the overall ‘story’ that the analysis has produced. Clear names and definitions were produced for each theme.

6. **Producing the report:** Writing the present report provided the final opportunity for analysis of the insights gained from the process. The analysis was used to address the initial research questions.

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