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Pelvic floor dysfunction: A scoping study exploring current service provision in the UK, interprofessional collaboration and future management priorities

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Disclosures

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Abstract

Objective

Pelvic floor dysfunction(PFD) has a significant socioeconomic and healthcare cost. This study aimed to investigate current service provision for PFD in the UK, highlighting any gaps and areas for improvement to inform future service improvement.

Methods

A three phase design comprised a scoping literature review, consultation survey with frontline practitioners from four key professional groups and an overarching synthesis. An interpretative analytical framework was informed by the concepts of interdisciplinary and interprofessional collaboration.

Results

Empirical evidence on PFD service provision is limited. No overarching strategic approach to PFD as a single clinical entity in the UK was identified. Two hundred and forty three medical, nursing and physiotherapy practitioners from different clinical subspecialties participated in the survey. Access and availability to services, models of delivery and individual practice vary widely within and across the disciplines. Time restrictions, mixed professional attitudes, lack of standardization and low investment priority were identified as major barriers to optimal service provision. Five overlapping areas for improvement are highlighted: access and availability, team working and collaboration, funding and investment, education, training and research, public and professional awareness.

Conclusions

Current services are characterized by a fragmented approach with asynchronous delivery, limited investment and poor interprofessional integration. An improved service delivery model has the potential to improve outcomes through better interdisciplinary collaboration and efficient use of resources.

Key words. Female, pelvic floor dysfunction, services, interdisciplinary care, collaboration

<u>What's known</u>

PFD comprises symptoms ranging from bothersome to debilitating that are challenging to patients and healthcare professionals in primary and secondary care and have substantial NHS healthcare cost implications. PFD is a multi-disciplinary issue that concerns a wide range of healthcare practitioners. Traditionally, its management has followed a clinically compartmentalised and fragmented approach that potentially is failing to address multicompartmental symptomatology and resulting in an incomplete resolution of the problem.

What's new

This study has highlighted gaps and inconsistencies in current service provision for women with PFD. There is under utilization of primary care resources and variable access and availability to specialist care. Management pathways are further compromised by circuitous routes of referral, treatment and diagnostic delays, duplication of effort and inappropriate referrals to specialist services. New ways of working with improved Interdisciplinary service provision have the potential to improve outcomes of care.

Text Revised and reduced in length

Background

Pelvic floor dysfunction (PFD) presents an increasing clinical challenge and financial burden to the National Health Service. Against a background of rising public expectations, technological advances, increasing prevalence of predisposing factors such as obesity, diabetes and, an ageing population it is predicted that future demand for PFD-related healthcare will rise by 50% over the next 30 years, potentially stretching current service capacity^{[1].} PFD comprises symptoms that are associated with anatomical and functional disturbances of the lower urinary tract and micturition, lower gastrointestinal tract and defaecation and/ or sexual function^[2,3]. These range from being bothersome to debilitating and are challenging to patients and healthcare professionals in primary and secondary care. Although PFD is common among women of all ages, the risk of developing symptoms rises steadily in middle and older age affecting an estimated 40% of women aged between 65-70 years increasing to 50% in those over 80^{[4].} It is estimated that than one in ten older women will undergo surgery for PFD with as many as 30% of these requiring further surgery for symptom recurrence^{[5].} Traditionally management of PFD has followed a clinically compartmentalized and often fragmented approach that may, due to variable access and availability to integrated services, be failing to address multi-compartmental symptomatology and resulting in an incomplete resolution of problems. This study presents the findings of preliminary study undertaken to investigate current service provision for women seeking treatment for PFD in the UK. Data gathered from a range of healthcare practitioners working in different specialty areas and mapped against relevant policy and documentary evidence is used to enhance a scoping literature review. The professional groups studied include nurses, doctors and physiotherapists with a specific interest in women's health.

Aims and Objectives

The primary aim of this scoping study is to investigate and synthesize evidence on current service provision in the UK. The secondary aim was to highlight priority areas for change to inform future service improvement and research. The following objectives were refined to aid the identification and interpretation of evidence;

- Identify the range, nature and extent of services
- Determine models of service provision and levels of interdisciplinary practice
- Ascertain the experiences, views and needs of frontline service practitioners in relation to current and future PFD service provision

Methods

A three phase structured design comprised:

(1) Concurrent scoping literature review and analysis of relevant research, documentary and policy evidence

(2) Consultation with frontline practitioners

(3) Overarching synthesis

This phased approach was deemed appropriate in the light of the complexity of the topic area and limited existing research evidence. The main strengths of a scoping study lie in its ability to extract the essence of a diverse body of evidence and give meaning and significance to a topic that is both developmental and intellectually creative. The concepts of interdisciplinary and inter-professional collaboration provided an interpretative analytical framework for this study^{[6].} The underlying principles of sharing, partnership, power, interdependency, and process were used to guide data extraction, appraisal and synthesis. The primary research questions posed were; *What type of service provision is available to women seeking treatment for pelvic floor disorders and what level of inter-professional collaboration exists*?

(1) Scoping literature review strategy

Our approach followed established methods for a scoping literature review^[7,8]. The review process comprised: establishment of the analytical framework, definition of key concepts and search terms, identification of key data sources, charting of the evidence, expert consultation and a summary analysis of the findings. Eight electronic bibliographic databases were searched including Cochrane Database of Systematic Reviews. These searches were supplemented by the pursuit of grey literature retrieved through secondary references, hand searching non-index linked publications and, advice from experts in the field. A systematic search strategy was used based on keywords comprising explicit single and combined terms relating to the topic (for example pelvic floor and pelvic floor dysfunction/disorders, incontinence and continence as free text and related indexed terms) and, the target population (women). A second layer of keywords such as multidisciplinary, interdisciplinary, integrated service and inter-professional collaboration were then used to refine the search. To ensure comparability and applicability of research findings, we focused our review on publications specific to services for women with pelvic floor disorders. Additional limitations were employed to include only material that was published in the English language and reported on service provision from 1990 to 2009 to ensure current relevance. Policy documents, reports and position statements relevant to the management of women with PFD in the UK were also sought using the following databases; - Department of Health National Institute for Health and Clinical Excellence(NICE) (government agencies) the Health Care Commission (healthcare agencies), Royal Colleges and professional bodies and, charitable associations such as The International Continence Society.

(2) Consultation

To elicit the expert views held by UK healthcare practitioners already engaged at the frontline of practice and enhance the findings of the scoping literature review, a questionnaire survey was conducted among a purposive sample of health care professionals. These were initially sought through contact with the UK membership of a wide range of groups and associations including; The International Continence Society, Association of Coloproctology of Great Britain and Ireland, UK Royal College of Nursing's Continence, Stoma and Gastroenterology and Nurse Practitioner Forums, Women's Health Chartered Physiotherapists(ACWHP), Royal College of General Practitioner Academic faculty groups in addition to local meetings and seminars. The questionnaire was piloted among a convenience sample of eight specialist practitioners. Following minor revision the final versions were made available for completion. The questionnaires comprised a series of open and closed questions that sought information about individual professional roles and practices, range of symptoms assessed, management and treatment strategies employed, pathways of referral within and across specialties, use of existing national guidelines/protocols and treatment outcome measures. The following final free text question, asked to each of the professional groups, canvassed opinions and recommendations on service improvements; "In order to provide optimal services for women with pelvic floor disorders, particularly to women with co-existing symptoms relating to pelvic organ prolapse, bowel, bladder and sexual function, what service developments/improvements do you think are required?"

It is not uncommon to use more than one method of purposive sampling techniques and in this phase three specific types were used;

- critical case sampling to elicit participants who exemplified key characteristics of PFD service provision, for example those who had published/lectured on PFD
- snowball sampling to determine other suitable participants; and
- maximum variation sampling that sought a range of participants from different disciplines to ensure that our data was reflective of a diversity of experiences.

Each respondent received a copy of the study information so that they could decide in advance whether they wished to participate. A prepaid-reply envelope was provided to those returning completed questionnaires by surface mail. Consent was accepted if the

 questionnaire was completed and returned. Two follow up reminders were made to non respondents at 4 and 8 week intervals.

(3) Data Synthesis

An interpretative analysis of the survey, documentary and literature data was undertaken by the PI and then discussed among the researchers. Summary descriptive statistics and manifest content analyses were the primary tools used for categorising and identifying emerging themes. The findings are synthesized and presented in the form of a narrative summary. The literature and documentary evidence was managed in a word program with quantitative survey data managed and analysed using SPSS[™] version 11 (SPSS inc, Chicago,II USA).

Ethical consideration

The UK National Research Ethics Service declared this study a service evaluation that did not require further ethical review by local Ethics Committees.

Results

Only 36 papers of over 2000 publications potentially focused on aspects of service provision and were selected for further review. Twenty one were considered eligible for inclusion ^[9,10,11,12,13,14,15,16,17,18,19,20,21,22,23,24,25,26,27,29,30]. Five (published across 7 publications) provided a holistic functional perspective of the evaluation and management of PFD and potential benefits of collaborative, multidisciplinary approaches in relation to both outpatient and surgical service delivery ^[10,11,14,17,19,22,24]. The remainder focused on service provision in terms of just one aspect of PFD (urinary incontinence ^[9,12,13,15,16,18,20,21,23,25,26,27,29,30] or pelvic organ prolapse^[15]. These papers tended to focus on the heterogeneity of access, appropriateness and treatment provision, service inconsistency and fragmentation, the importance of multidisciplinary team work and the need for greater service integration. Unlike other countries such as the United States of America ^[31] and Australia ^[32] who have developed National Women's Health policies to ensure that the planning and delivery of health services best meets the needs and improves health and well being of women in their respective populations, no national overarching strategy for women's health and PFD in particular, was identified for the UK. Although numerous sources providing a wide range of commissioning guidance's, treatment pathways and protocols for specific aspects of PFD, were identified (Table 1).

The frontline survey highlighted a number of issues associated with the provision and delivery of pelvic floor services. However, in the absence of a national association or

database of practitioners with a special interest in PFD and the decision to use a variety of sampling techniques, final response rates were unable to be calculated. The two most difficult professional groups to access, despite positive liaison with representatives of their respective professional forums, were the GP's and nurse specialists. In light of a very small national GP response(26) the steering committee agreed that the focus of further efforts to gather the views of GP's should be placed on establishing a representative cohort. This involved recruiting GP's from 56 practices in one large, urban Primary Care Trust. GP's from 19(32%) of these practices responded.

Following reflective discussion between the researchers analysis of the survey data focused on three key dimensions (i) demographics and professional characteristics of practitioners (ii) management of PFD and models of healthcare delivery and, (iii) priority areas for improvement.

(i) Demographics and professional characteristics

 We recruited 243 practitioners including 95(39%) specialist physiotherapists, 63(26%) medical consultants, 45(19%) general practitioners (with a cohort of 19 from one PCT) and, 40(16%) specialist nurses, from a variety of clinical subspecialties (Table 2). They were geographically spread across the ten Strategic Health Authorities in England with a small number responding from services based in Scotland(10), Wales(6), Northern(8) and Southern Ireland(6) and, the Channel islands(1) (Figure 1). Eighteen (45%) of the nurses worked exclusively within primary care with the exception of 2 general practice-based, nurse practitioners, the remainder were continence nurse advisors, 16(40%) within secondary care(5 colorectal, the remainder in urogynaecology) and 6(15%) worked across primary and secondary care(providing continence care). Of the 95 physiotherapists, 22 (23.1%) worked in primary care(6 worked in dedicated continence services with 1 in private practice). 48(50.5%) in secondary care, 25(26.3%) provided services across primary and secondary care. All declared a specialist interest in women's health physiotherapy with 13 having full membership of the clinical interest group the Association of Chartered Physiotherapists in Women's Health(ACPWH). Three provided extended-scope practitioner services. The medical consultants worked in District General, Secondary and Tertiary Hospitals. The proportion of time estimated to be devoted to PFD was 60-100% by urogynaecologists, \leq 20% by the urologists and colorectal surgeons and less than 1% of the GP's practice.

(ii) management of PFD and models of healthcare delivery

management

The management of PFD varied widely across the disciplines and between practitioners. Medical consultations were greatly restricted by time compared to specialist nurses and physiotherapists who were able to arrange longer appointments (45-60 minutes) and generally involved additional symptom severity and quality of life assessment measures and pelvic examination. Of the GP's only 2 recommended a second appointment for further assessment, examination and discussion. Four referred directly either to a hospital consultant, women's health physiotherapist or continence nurse advisor rather than manage symptoms themselves. The initial treatment strategies described by the group collectively, were broadly similar. First line management comprised lifestyle interventions, diet and fluid manipulation, bowel and bladder retraining, evacuation techniques, pelvic floor muscle exercises, vaginal pessaries and pharmaceutical therapies prescribed as required. Generally the guidelines for Urinary Incontinence(UI)^{[33],} Faecal Incontinence(FI)^[34] Pelvic Organ Prolapse^[35] were followed however, when asked about the use and usefulness of guidelines in clinical practice, the practitioners views were not universal. Only 3 GP's and 12 consultants perceived clinical guidelines as very useful while 23 GP's and 38 consultants found them moderately useful. The remainder either had not read any guidelines or did not find them useful. For nurses and physiotherapists, clinical guidelines and treatment protocols provided the framework for their practices.

models

Four distinct models of healthcare delivery were identified; single practitioner-led, multidisciplinary within-specialty, across-specialty and cross boundary (Figure 2). Nearly 50% (including the GP's) managed PFD as single practitioners. The majority of the remainder followed a within-specialty model and used a combination of different service approaches. Only 33 practitioners worked in across-specialty services and just 2 (working in continence) provided cross boundary services between primary and secondary care. Within these three models many practiced formalized triage systems developed to facilitate direct referral pathways to single specialist-led, combined and/or joint multidisciplinary outpatient clinics and diagnostic services. There was one example of a consultant-led outreach clinic in primary care and four secondary/tertiary centres provided dedicated PFD outpatient services. Attendance at regular multidisciplinary team(MDT)-style meetings tended to be confined to practitioners within individual clinical subspecialties in secondary care. The attendance by nurses was less than half 16(40%) and only just over half among physiotherapists 49(51.5%). The main reason given for non-attendance was 'no meetings', limited time to attend and logistical difficulties. Complex cases were discussed by 43(65%) consultants at infrequent, ad hoc meetings. With the exception of one consultant surgeon who managed all pelvic floor problems as a single specialist surgeon, the majority of consultants preferred a single specialty approach referring patients requiring further surgery for other pelvic compartmental symptoms on to surgical colleagues as appropriate. Only 23(36.5%) of the surgeons participated in combined operative procedures.

(iii) priority areas for improvement

One hundred and ninety nine (81.8%) practitioners provided views on how services could be improved. Of the 44 that declined to offer an opinion, over half were medical staff; comprising 15/45 (33.3%) GP's, 16/63 (25.3%) consultants (11 colorectal, 2 urology, 3 urogynaecology), 8/95 (8.4%) of the physiotherapists and 5/40 (11.1%) nurses. Using a qualitative methodological approach to the analysis of this data, five overlapping themes identified as priority areas for improvement, were identified;

- 1) Access and availability to specialist services
- 2) Team working and collaboration
- 3) Funding and investment
- 4) PFD education, training and research
- 5) Public and professional awareness

There was consensus that access and availability to specialist care was being potentially compromised due to the low priority and lack of a coherent strategy to PFD. While the perspectives of this theme differed slightly between professional groups, it was generally agreed that circuitous routes of referral, duplication of effort and treatment delays presented particular obstacles to optimal service provision. For example, when symptoms failed to respond to treatment, few specialist nurses and physiotherapists, particularly in primary care, had direct referral access to consultants or specialist diagnostic investigations. Local arrangements commonly required patients be referred back to their GP's for further assessment in order to secure an onward referral. In secondary care similar frustrations were also described in relation to direct referral access to continence advisors, WHP's and hospital services were variable. While for some, local arrangements were good for others access was reported as more difficult for several reasons including; lack of direct referral arrangements to specialist investigations or 'one/two stop' specialist PFD clinics and, uncertainty about the availability or location of local specialist nurse or physiotherapy services.

Improving inter-professional collaboration was a key issue and supported by numerous recommendations. These included making better use of existing resources to improve communication and coordination between different service fragments, development of a national database, patient-held documentation and, wider involvement of specialist nurse/physiotherapists working in advanced roles to facilitate services and improve earlier

 access to treatment. Practitioners generally concurred that services needed to shift with the development of more primary care-directed service models that would not only encourage a more synchronous and seamless service approach but, would also have the advantage of freeing up limited specialist secondary/tertiary care resources for patients with more complex problems.

There was universal agreement that greater funding and investment in staff, equipment and training was required if services were to improve and new ways of working were to be developed. Numerous recommendations to achieve standardized approaches to care were made in relation to improving opportunities for inter-professional education and training within and across primary and secondary care settings although specific educational needs tended to differ across the groups.

The need to raise professional and public awareness of PFD as a significant women's health problem was also highlighted and a number of proposals were made that would potentially enhance current services including the provision of more primary care-based and school based educational sessions to promote healthy pelvic floor practices, ante and post natal drop in clinics, self help support groups, better advertising of services and the development of local health service websites, information leaflets and posters.

Discussion

This study suggests that a combination of external (macro) and internal (meso) organizational and, personal (micro) factors are contributing to a fragmented, highly variable and poorly integrated approach to PFD service delivery in the UK.

At a 'macro' level, the strategic policy approach to women's health care in the UK differs significantly from that of other countries ^[31,32,36] where through synchronized approaches to cross referral and team working, models such as the Calgary Pelvic Floor Disorders Clinic (currently being evaluated in Canada) ^[37] and the Maimonides Centre for Pelvic Floor Dysfunction in the USA ^[24] have developed. It could be argued that although the majority of women's health services in the UK are well provided through local services, from the views expressed by the respondents in this study, there is a suggestion that the tendency to subsume or 'piggyback' women's health issues within other areas of health policy could result in the multiple causes and consequences of PFD being inadequately addressed. This may also explain some of the service inconsistencies we identified and support evidence that outcomes in relation to the management of multi-compartmental pelvic floor symptoms are suboptimal^{[2,38].} Another factor to consider at the macro level relates to targets. These have become an essential element in the UK Government's plans for modernizing the NHS (DH

 2000)^[39] and despite efforts to move away from a target-driven to a "high quality" service (DH 2008)^[40] initiatives like the 18 week commissioning pathways remain important service indicators^{[41].} We contend that although such initiatives have gone a long way to improving patient healthcare they are not without their critics particularly when managing the time consuming and often complex problems associated with PFD that require a plural rather than singular approach. Our findings indicate that in patients with mulitple symptoms the efficacy of treatment can be limited if a single specialty service position is consistently endorsed.

At the 'meso' level we found that the majority of practitioners describe broadly similar approaches to care with a high level of autonomy in the provision of services. However, as patients often enter and re-enter the system via many routes and consult with many different practitioners along the treatment pathway, there is potential for patients to be exposed to duplication and delay in treatment (Figure 3). Although there is evidence supporting the clinical and cost effectiveness of more specialist nurse and physiotherapist practitioner involvement ^[9,13,30] confirmation of inter-professional collaboration is limited. We suggest that this may provide an explanation for the predominance of the partitioned, 'within specialty' model approach identified in this study that, alongside the lack of a standardized approach to managing PFD as a single clinical entity, inhibits service innovation and the development based on locally available expertise of robust, effective integrated services. Another factor to consider at this level relates to the use of and adherence to clinical guidelines. Our findings concur with previous studies^[42] where adherence by practitioners to guidelines developed for the identification and management of bowel, bladder or sexual function, may be only partial. The major reasons highlighted in our study include; insufficient time, lack of available resources especially staffing numbers limited or no access to specialist 'one-stop MDT services'. The recommendations made by practitioners for new service configurations that include more specialist services being made available within primary care, would appear to offer a solution to this problem. In addition it would facilitate greater opportunities for interdisciplinary professional development and support recent recommendations made by The Department of Health^[43]. We propose that by creating a more synchronized, collaborative, inter-professional approach between GP's, primary care-located triage hubs and specialist centers, the potential benefits already apparent in other specialties^[44] will become evident.

Finally, at the practitioner 'micro' level, several factors emerged that emphasized links between the pressures on consultation/treatment time and poor resourcing, inadequate knowledge and lack of understanding of the expertise of colleagues and efficacy of treatments. Previous research has shown that patients presenting with PFD are likely to

suffer complex groups of problems which if not routinely enquired about may be underappreciated and lead to inadequate treatment^{[2,3,4].} In addition, the propensity for duplication of effort for assessment and conservative therapies as patients bounce around the system, are further indications that optimal care is being compromised. Our findings suggest that differences in professional attitudes, experiences and knowledge alongside a lack of government directive and financial investment in PFD in the UK has contributed to a degree of professional territorialism and may account for the 'therapeutic nihilism' reported in some studies ^[45]. Recommendations from practitioners for more 'protected consultation time', 'ring fenced' clinical arrangements, more inter-professional educational opportunities in addition to improved specialist staffing levels were identified as key areas for improvement.

Conclusion

The assessment and management of PFD is time consuming and involves the input of a wide variety of healthcare professionals. This study has highlighted areas which will be of interest to practitioners, providers and commissioners involved in the future planning and delivery of women's health services. Although our survey data cannot be interpreted as representative of the wider views of each professional group because they are based on a self-selected, purposive sample with unknown response rates, we contend that this study affords an opportunity to gain valuable insight into the current views and recommendations of key frontline practitioners working across the different subspecialties. Given the impact of PFD on the health and wellbeing of women and the eminently treatable nature of symptoms regardless of age, early access to specialist health care is important. Raising awareness of optimal service provision for the prevention and management of PFD should be a health policy priority. This study provides for the first time, baseline evidence that exposes a number of barriers hindering the development of specialist integrated services. Our analysis reveals that availability and access to formalized interdisciplinary PFD care in the UK is highly variable. We hypothesize that new service configurations will facilitate greater service collaboration and break down traditional professional boundaries across primary and secondary care and ensure more effective and efficient use of resources. Improving interdisciplinary education and training would also strengthen relationships and change practitioner's views on the benefits of combined management approaches. This will not only lead to improved understanding of the complexity of female PFD but also expose more patients to the range of therapeutic benefits available including pelvic floor health promotion initiatives helping to prevent pelvic floor complications developing later in life, contributing to healthier ageing and well being and improving women's participation in health decision making and health management.

Authors Contribution

All authors have approved the final draft for submission and had access to the study data. Specific contributions were made in the following areas: Kathy Davis was responsible for the conception and design, data collection, analysis and interpretation of this scoping study. In addition, she was responsible for the preparation, drafting and critical revision of the manuscript. Devinder Kumar participated on the data interpretation, preparation and critical revision of the manuscript for important intellectual content and contributed to statistical analysis of the data. Martyn Wake assisted in the design of the study, facilitating the acquisition of data from general practitioners and critical review of the final drafts of the manuscript.

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Table 1

Summary of key documentation & evidence based practice guidelines identified to support best practice in the assessement and management of PFD

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Table 2

Summary of practitioners by principle specialty across the four disciplines

Specialty	TOTAL	Specialty	TOTAL
	n (%)		n (%)
1. Consultants		2. Specialist Nurses	
a) Urogynaecology & Gynaecology	17 (27)	a) Urogynaecology	5 (12.5)
b) Obstetric & Gynaecology	5 (7.9)	b) Obstetrics	1 (2.5)
c) Colorectal	26 (41.3)	c) Colorectal	6 (15.0)
d) Urology	14 (22.2)	d) Urology	1 (2.5)
e) unspecified	1 (1.6)	e) Continence	23 (57.5)
		f) Primary care Nurse Practitioners	4 (10.0)
	Total =63		Total=40
3. Physiotherapists		General Practitioners	
a) Women's health (O & G including Urogynaecology)	31 (32.6)	GP's from one PCT	19 (42.2)
b) Women's health and continence	29 (30.5)	Other GP's	26 (57.8)
c) Continence	14 (14.7)		
d) Women's health and general physio	12 (12.6)		Total = 45
e) Mixed- clinical provision across subspecialty eg.	6 (6.4)		
urology, general surgery and medicine			
f) unspecified	3(3.2%)		
	Total = 95		

Figure 1 Respondents by Geographical Location





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Figure 2

Models of healthcare delivery



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Figure 3

Current service pathways for investigation & management of PFD

