

Population Name	Oxfordshire
Population Size	800000 plus tertiary referrals
Report for 12 months ending	2015
Number of practices in the population	82
Participating practices - number (%)	
Total number of patients from participating practices	
Total number of patients identified with PP	
% of total patients identified with PP (prevalence):	16%
Was a Pelvic Pain questionnaire (e.g. IPPS) use for patients?	yes from jan 2015. Short proforma developed for audit
% of PP patients identified in primary care	
% of urgent or semi-urgent patients referred within one month of presentation by their GP (urgent/semi urgent patients=severe undiagnosed or progressive pain with the risk of increasing functional impairment, generally of 6 months duration or more)	
% of routine or regular patients referred with 8 weeks of presentation by their GP (routine=persistent long term pain without significant progression)	
Number of follow up PP patients in gynaecology, urology, psychology/counseling (IAPT, CBT), gastro-enterology, colo-rectal, pain/physiotherapy	
% of PP patients for whom a multidisciplinary meetings was held	this is a multidisciplinary service. Not all patients seen by all elements of MDT but available to all
Average time from presentation to multidisciplinary meeting being held	average time from referral to clinic appt - 10-12 weeks.
% of PP patients with a named co-ordinator of care	100
% of PP patients for whom patient outcomes are reviewed every 6 months (being mindful of whether the patient should be monitored or discharged as self-managing)	100
% of PP patients who feel they have adequate methods to manage their pain	
Is there direct access to psychological services? If so, please describe.	yes - referral through pelvic pain clinic.Two counsellors. Also advice available through Dept of psychological medicine linking into community services.
Are patients given access to decision support tools (e.g. map of medicine, releavent websites, leaflets, local buddying and peer support groups)? If so, please describe.	range of leaflets, and book suggestions available. Websites including self help organisations frequently recommended
Are experiences of PP patients monitored? If so, please describe.	all follow up patients asked for feedback
Are local support groups or materials (e.g. shared decision aids, youtube videos, leaflets, etc.) developed with PP patient/care involvement? If so, please describe?	
Estimated cost of your service	
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