

Population Name	Tees Valley including Hambeldon and Richmondshire
Population Size	approximately 1.5 million
Report for 12 months ending	April 2015
Number of practices in the population	Unknown
Participating practices - number (%)	Unknown
Total number of patients from participating practices	Unknown
Total number of patients identified with PP	Unknown
% of total patients identified with PP (prevalence):	Unknown
Was a Pelvic Pain questionnaire (e.g. IPPS) use for patients?	No
% of PP patients identified in primary care	Unknown
% 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)	Unknown
% of routine or regular patients referred with 8 weeks of presentation by their GP (routine=persistent long term pain without significant progression)	Unknown
Number of follow up PP patients in gynaecology, urology, psychology/counseling (IAPT, CBT), gastro-enterology, colo-rectal, pain/physiotherapy	Unknown
% of PP patients for whom a multidisciplinary meetings was held	Unknown
Average time from presentation to multidisciplinary meeting being held	Unknown
% of PP patients with a named co-ordinator of care	0
% 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)	0
% of PP patients who feel they have adequate methods to manage their pain	
Is there direct access to psychological services? If so, please describe.	No direct access but psychology is available via pain service
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.	Yes ad hoc in clinic. No formal arraignment
Are experiences of PP patients monitored? If so, please describe.	No formal monitoring for CPP. Patient experiences within the clinic assessed intermittently
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?	Non available
Estimated cost of your service	Unknown
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