Population Name	Leicester, Leicestershire and Rutland + more distant CCGs
Population Size	One million
Report for 12 months ending	March 2014
Number of practices in the population	Unknown
Participating practices - number (%)	Unknown
Total number of patients from participating practices	68
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?	Yes- developed internally
% of PP patients identified in primary care	100%
% 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)	0%
% of routine or regular patients referred with 8 weeks of presentation by their GP (routine=persistent long term pain without significant progression)	0%
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	Variable
% of PP patients with a named co-ordinator of care	All patients will have a GP
% 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)	Unknown
% of PP patients who feel they have adequate methods to manage their pain	Unknown
Is there direct access to psychological services? If so, please describe.	? From where GP: access to IAPT Gynae: access to Clinical Psychology in hospital Pain Management Service: Direct access to Clinical Psychology for individual work and Pelvic Pain Group. Pelvic Pain Group advertised in Gynaecology and Genitourinary Clinics- but uptake not good
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-some of above
Are experiences of PP patients monitored? If so, please describe.	Medical care will be monitored if local patient

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?	No
Estimated cost of your service	approx £10,000/year for clinic and Pelvic Pain Group alone
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