CANCER SCREENING







INTRODUCTION What is screening? Wilson's screening criteria Testing apparently healthy people for signs of a Condition is an important health problem with a natural history that is understood and a recognisable latent or disease. UK National Screening committee (NSC) decides early symptomatic stage. Continuous case finding and accepted treatment which is what to screen for. more effective if started early. UK population screening - cervix, bowel, breast. UK targeted screening - lung (55-74, ever Clear policy on who to treat. Diagnosis and treatment cost-effective. smoked).

| CERVICAL SCREENING | | |
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| Who is screened and how often? | How does it work | Special situations |
| All ♀ (registered ♀ at birth regardless of gender identity) who still have their cervix. England/NI 3-yearly 25-49 then 5-yearly 50-64. Scotland/Wales 5-yearly 25-64. HIV +ve - annually 25-64. Stop at 65 unless recent test +ve | Primary HPV screen - if -ve, back to normal recall. +ve HPV → cytology → +ve cytology → colposcopy. -ve cytology → test 1y later → if this happens 3 times then colposcopy. Some evidence for self-take HPV | HPV can transmit via non-penetrative sex; screen if any sexual contact. Learning disabilities – individual decision based on risks/benefits. Trans man – screen if has cervix, will need manual recall if gender marker male on notes. |
| or haven't had test since age 50, in which case can offer one >65. | swab as first step, rolling out first among smear non-responders. | Perimenopause – 6/52 vaginal oestrogen first if test is painful. |

| BOWEL AND BREAST SCREENING | | | |
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| Bowel | Breast | | |
| • Age 50 – 74 (England, Scotland), 51-74 (Wales), | • All ♀ (registered ♀ at birth regardless of gender identity), | | |
| 60-74 (NI), ≥75 can request if wanted in | 3-yearly mammogram 50 – 70. | | |
| England and Scotland only. | First invite sometime between 50-53. | | |
| Faecal immunochemical test (FIT) 2-yearly. | Trial ongoing re effectiveness from 47 and to 73. | | |
| +ve FIT → colonoscopy. | Some concerns about overdiagnosis – <u>information for</u> | | |
| +ve threshold ~120μg/g (screening), 10 μg/g | <u>public</u> on gov.uk site. | | |
| (symptomatic) therefore still send FIT with | High-risk ♀ more often and/or with MRI and | | |
| symptoms even if recent negative screen. | frequency/modality adjusted for breast density. | | |

| symptoms even if recent negative | S | frequency/modality adjusted for breast density. | |
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| OTHER CANCERS Prostate No national programme but guidance for GPs to do PSA if requested and aged >50 or +ve FH. This is unresourced. PSA specificity & sensitivity poor, doesn't differentiate between slow growing and aggressive cancers. Some evidence that MRI screening is sensitive and picks up aggressive cancers even in those with normal PSA. Ongoing research which NSC will review in a few years, including about genetic test done on saliva. | High-risk bowel Refer to genetics if one find degree relative diagnosed or two people who are findegree relatives to each cand at least one is a first-relative to your patient, diagnosed at any age. Screening may involve on regular colonoscopy. Prophylactic surgery e.g. swith UC or certain mutati No guidance for pancreas patients can enrol in EUR study if 2 first-degree relative to general surgery e.g. swith pancreatic cancer. | High risk breast rst- I < 50, st- other degree degree o 1 first-degree relative ♀< 40, bilateral < 50, ♂ any age. o 2 first or 1 first and 1 second- degree any age. o 1 first or second degree breast and 1 first or second degree ovarian at any age (at least one first degree). o ↓ referral threshold: FH bilateral / ♂ breast cancer, ovarian cancer, Jewish ancestry (≥1 Jewish grandparent → NHS BRCA test in England), sarcoma | |
| | | FH breast cancer. | |