

Faster Diagnosis Standards (FDS)

- What are FDS and what can we do to help targets be achieved?

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Clinical Specialist Sonographer

Presentation Outline

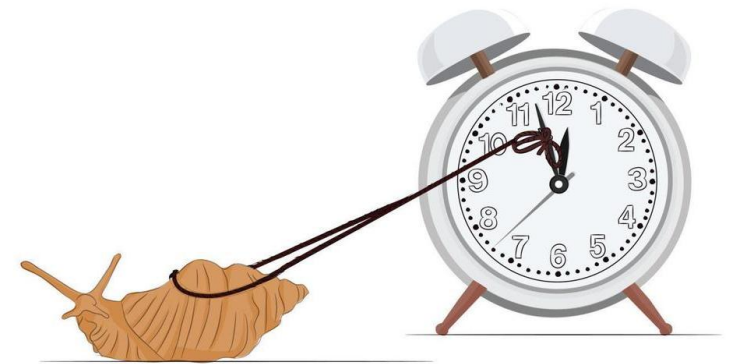


Humber Health
Partnership

- Background to FDS
- Best timed pathways
- Where we are with meeting targets
- Sonographers led services in FDS
- Some Hull examples
 - Urology – haematuria, prostate, testicular Ca
 - Ovarian Ca

Two week wait

- Consultation with specialist within 14 days of referral
 - Could be in person or telephone
- 93% target
 - 83% in Yorkshire
 - 75% in the rest of England



Problems

1. Onward referrals - Had to wait significant amounts of time for onward investigations, therefore had to wait longer to have cancer diagnosed or excluded
2. Treatment delays – wait time for results or having different tests on different days
3. No room for innovation – forgoing sophisticated tests to meet the targets



**“I want you to find a bold and innovative way
to do everything exactly the same way
it’s been done for 25 years!”**

Idea of FDS first floated in
2015

Too interested in the cinema?

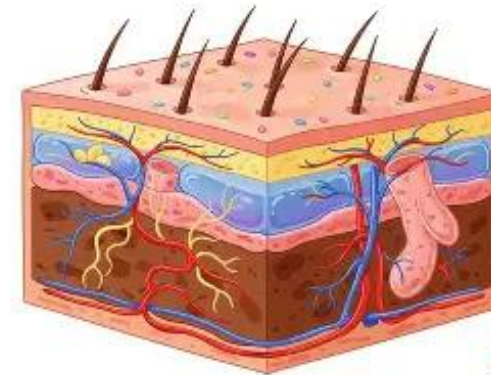
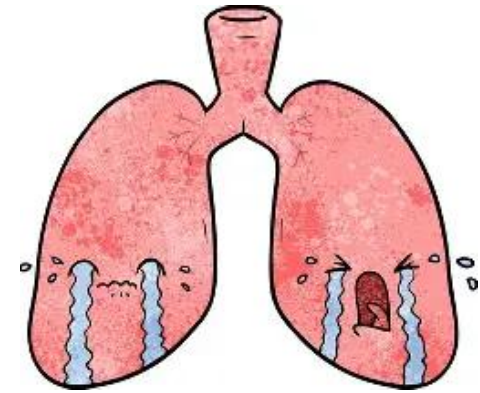


Personally I think apple are to blame...



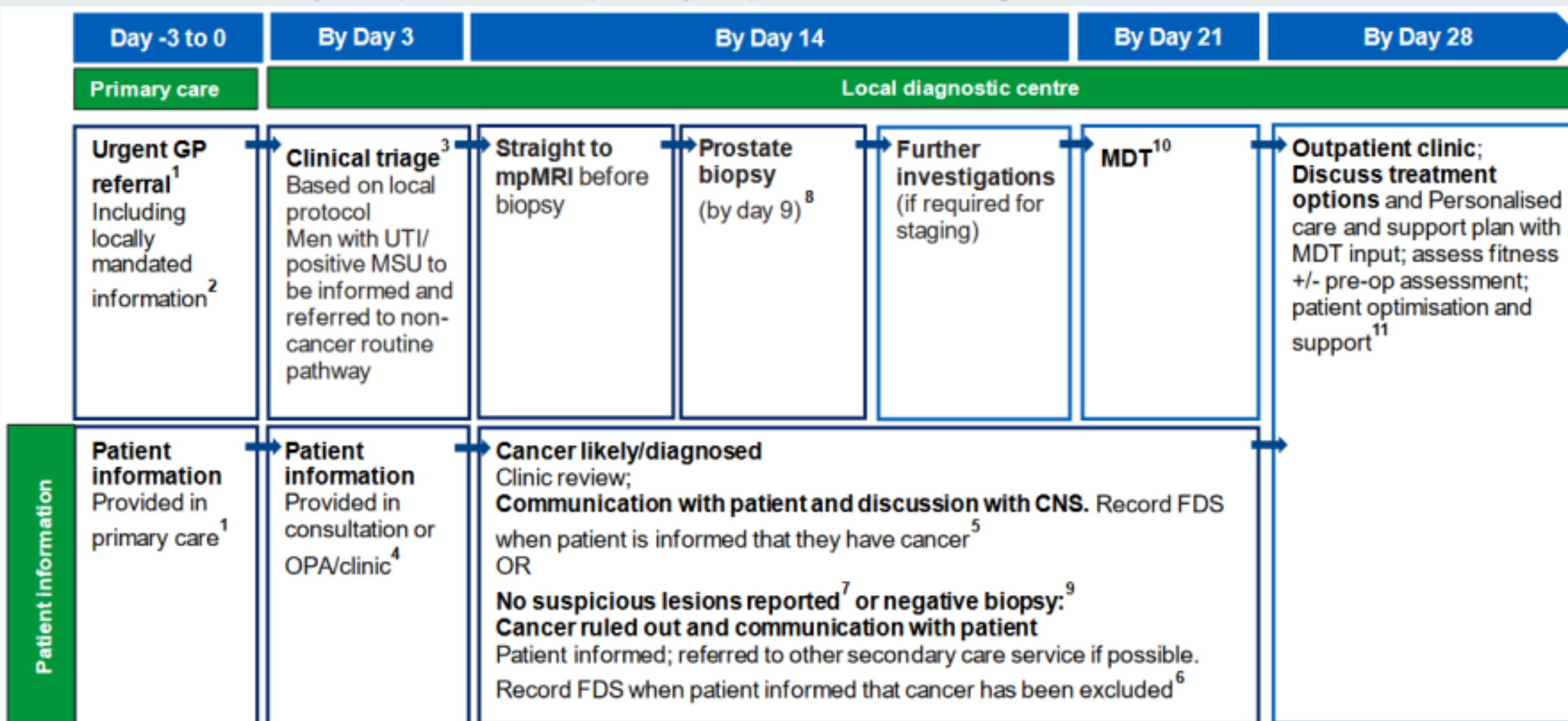
2018 – Best Timed Pathways

Outline best evidence structure and timing of various tumour specific pathways to meet FDS



Example..

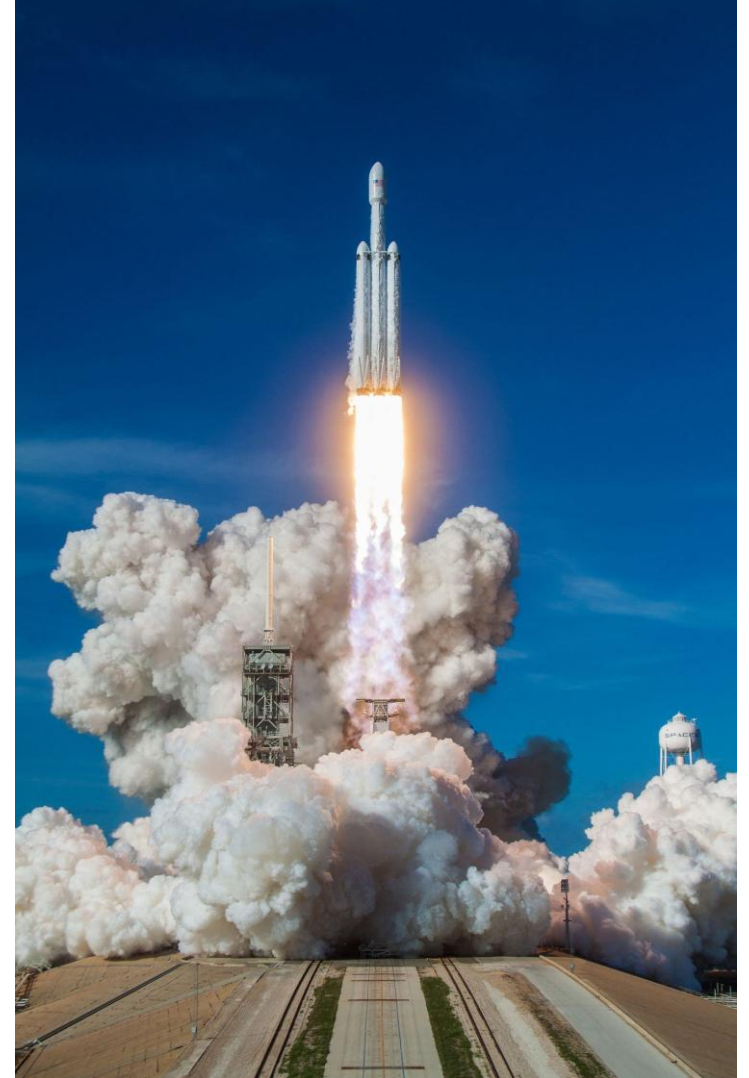
Prostate cancer - 28-day best practice timed pathway for prostate cancer diagnosis



This is a straight to test pathway using MpMRI. The 21-day pathway should be used when an immediate MRI is not required or is contra-indicated.

FDS Hard Launch 2021

- 2-week-wait for first appointment removed as NHS initiatives with Faster Diagnostic Standards Pathways.
- Nine previous performance targets streamlined into three key performance indicators
- Standard ensures patients will be diagnosed with, or have cancer ruled out, within 28 days of being referred urgently by their GP for suspected cancer.



3 Key Performance Indicators

28 Days Faster Diagnosis Standard

Diagnosis/Exclusion
of cancer by 28 days
of referral (80%)
(Prev 75%)

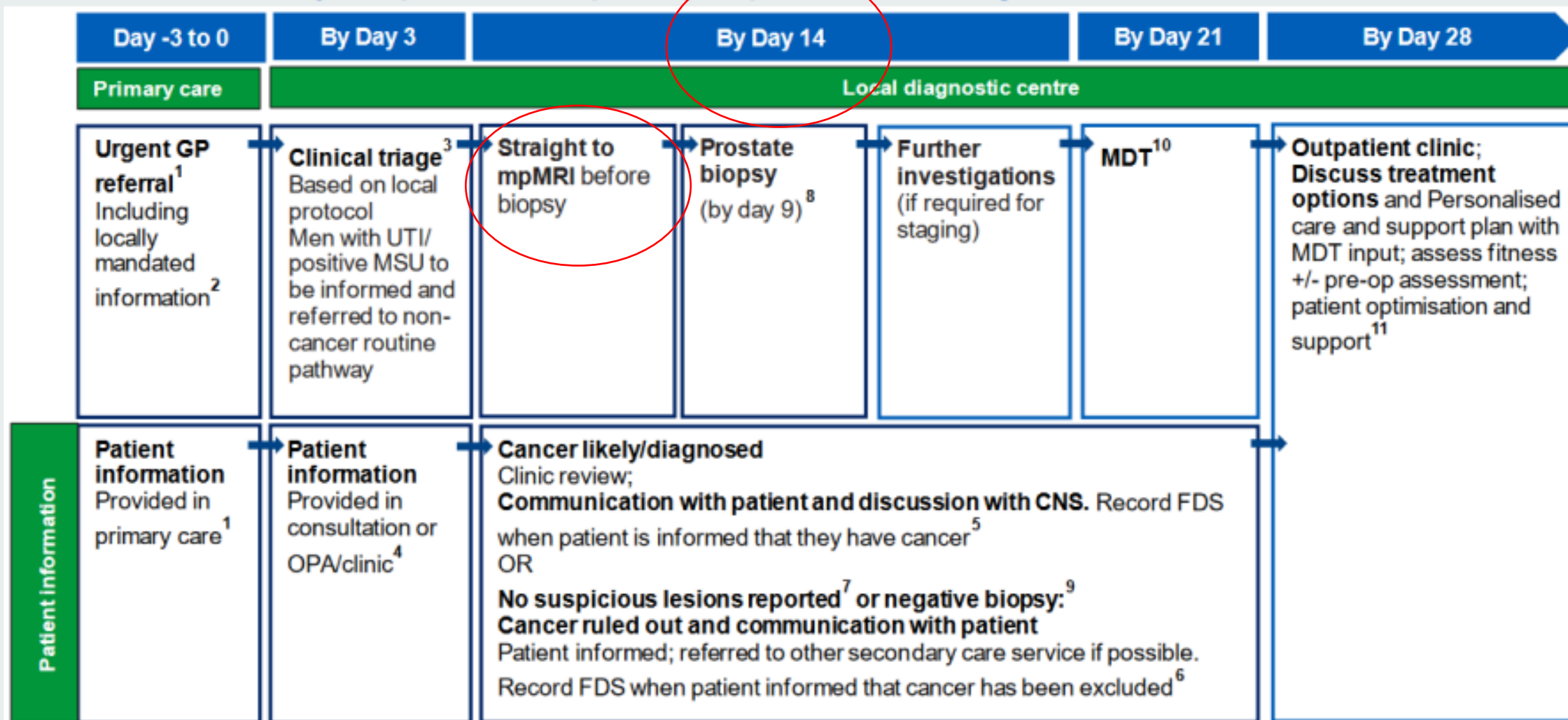
31 Day Treatment Standard

Start treatment within
31 days of a decision
to treat (96%)

62 Day Treatment Standard

Commence
treatment within
62 days of being
referred (85%)

Prostate cancer - 28-day best practice timed pathway for prostate cancer diagnosis

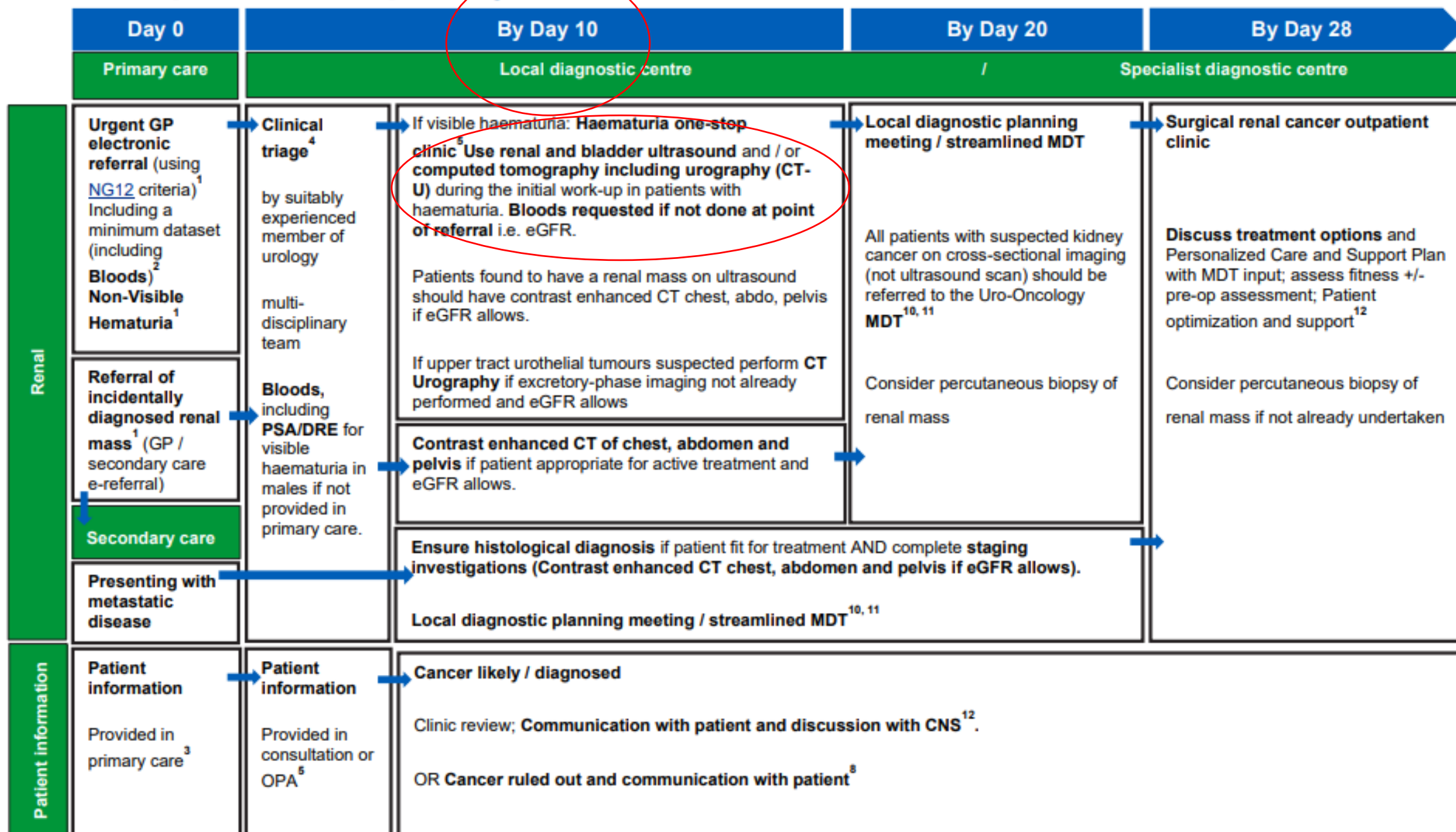


This is a straight to test pathway using MpMRI. The 21-day pathway should be used when an immediate MRI is not required or is contra-indicated.

Bladder best practice timed pathway

	Day 0	By Day 10	By Day 14	By Day 20	By Day 28
	Primary care	Local diagnostic centre		Specialist diagnostic centre	
Bladder and other Urothelial	Urgent GP electronic referral (using NG12 criteria) ¹ Including a minimum dataset (including Bloods) ² Non-Visible Hematuria ¹	Clinical triage ⁴ by suitably experienced member of urology multi-disciplinary team. Bloods , including DRE for visible haematuria in males if not provided in primary care. Offer PSA if abnormal DRE	If visible haematuria: Haematuria one-stop clinic ⁵ Use renal and bladder ultrasound and / or computed tomography including urography (CT-U) during the initial work-up in patients with haematuria. Once a bladder tumour has been detected, perform CT Urography in selected cases (e.g. – tumours located in the trigone, multiple- or high-risk tumours). AND Flexible Cystoscopy (can be omitted if straight to TURBT). ⁶	TURBT / Bladder Biopsy ⁷ (reported within 7 calendar days). Followed by CT of chest (if muscle invasive / advanced / metastatic bladder cancer and not yet done). In centres with capacity to follow an imaging guided pathway and sMDT agreement, if muscle invasive disease suspected at diagnostic flexible cystoscopy, arrange urgent MRI of bladder ⁹ If suspected upper urinary tract urothelial carcinoma electronically refer directly to sMDT ¹⁰ If suspicion of upper tract urothelial tumour: Consider Ureterorenoscopy +/- Biopsy if diagnostic uncertainty on imaging ⁷	If low risk non-muscle invasive bladder cancer, may remain in local MDT , for all others electronically refer to Specialist MDT ^{10,11} and specialist clinic appointment Bladder cancer clinic with histology results AND Discuss treatment options and Personalized Care and Support Plan with MDT input; assess fitness +/- pre-op assessment; Patient optimization and support ¹²
	Secondary care				
	Presenting with metastatic disease ¹		Ensure histological diagnosis if patient fit for treatment (TUR biopsies or biopsies from metastasis) AND complete staging investigations (CT of chest, abdomen and pelvis post contrast). Local diagnostic planning meeting / streamlined MDT . Refer appropriate cases directly to sMDT . ¹⁰	sMDT	
Patient information	Patient information Provided in primary care ³	Patient information / signposting Provided in consultation or OPA ⁵	Cancer likely / diagnosed Clinic review; Communication with patient and discussion with CNS ¹² . Record FDS when patient is informed that they have cancer OR Cancer ruled out and communication with patient ⁸		

Renal best practice timed pathway



Testicular best practice timed pathway

	Day 0	By Day 7	By Day 14	By Day 21
	Primary Care	Local diagnostic centre / Specialist diagnostic centre		
Testicular	Urgent Primary Care electronic referral (following NG12 criteria) ¹ Including a minimum dataset ² (including Bloods) ²	Clinical triage ⁴ by suitably experienced member of urology multi-disciplinary team.	Testicular one-stop clinic Clinical assessment, US ⁶ (if not done already) and Bloods (FBC, U&E's, LFT and AFP, bHCG and LDH tumour markers if not done already), followed by CT of CAP ⁶ if required (with hot reporting if available), if elevated markers or high-level suspicion of metastatic disease preservation where available. For small testicular masses which are indeterminate, where lesions are noted with infertility, bilateral tumours, or tumours in a single testis should be referred to supra regional MDT. 16–24-year-olds should be referred to Teenage and Young Adults Service. Offer sperm storage, request blood tests as required by fertility clinic. Provide sperm.	Patient informed if for inguinal orchidectomy and surgery completed. Discuss treatment options and Personalized Care and Support Plan with MDT input; assess fitness +/- pre-op assessment; Patient optimization and support ¹¹ Fertility Clinic
				Patients presenting with metastatic disease consider Supra-regional MDT review ¹⁰ AND Regional testis cancer clinic appointment (including outreach if provided)
Patient information	Patient information Provided in primary care ³	Patient information Provided in consultation or OPA ⁵	Cancer likely / diagnosed Clinic review; Communication with patient and discussion with CNS . Record FDS when patient is informed that they have cancer. ⁸ Discuss treatment options and Personalized Care and Support Plan with MDT input; assess fitness +/- pre-op assessment; Patient optimization and support. ¹² OR Cancer ruled out and communication with patient ⁸	

Gynaecology

Day -3 to 0	By Day 3	By Day 9	By Day 18	By Day 28
Primary care	Local Diagnostic Centre / Specialist Diagnostic Centre			
Pelvic examination / vaginal speculum and Urgent pelvic ultrasound / CT ¹ if available and appropriate based on NG12 and Ovarian NICE guidelines. Urgent referral ¹ Including minimum dataset ²	→ Clinical triage ³ by suitably experienced clinician Outpatient appointment by day 5 if not suitable for straight to one-stop clinic or CT ⁴	→ Straight to one-stop clinic (Ultrasound led) ⁶ Transvaginal Ultrasound if not carried out in Primary Care with same day hot reporting followed by Clinical Examination and/or LA Hysteroscopy or Pipelle Biopsy ⁷ (by day 12 if not carried out in same day clinic) followed by booking of MRI, if indeterminate US, significant clinical suspicion of lower genital tract cancer or biopsy indicating endometrial cancer or uterine sarcoma (may require 7 days between biopsy and MRI) ⁸ → Straight to one-stop clinic (Biopsy led) Clinical Examination and Biopsy, if visually suspicious lesions of lower genital tract ⁷ followed by booking of MRI, if indeterminate US, significant clinical suspicion of lower genital tract cancer or biopsy indicating endometrial cancer or uterine sarcoma ⁸ → Straight to CT ⁹ and Tumour markers CEA and CA19-9, and clotting	→ GA Hysteroscopy / Biopsy if LA not possible in one-stop clinic Triage of cases to local or specialist MDTs using agreed standard of care pathways to minimise MDT discussions ¹² required Pre-booked MRI carried out ⁸ → For significant risk of localised ovarian cancer: Formulation of management plan ¹⁰ and discuss at MDT ¹² in line with agreed protocols → For advanced ovarian cancer: MDT ¹² to discuss approach to biopsy including image guided or laparoscopic, followed by Biopsy ⁷	→ For endometrial cancer / uterine sarcoma: Thorax / Abdominopelvic CT ¹¹ for higher risk (high grade histology results) if not carried out in straight to CT clinic and/or Specialist MDT, if required ¹² and Formulation of management plan ¹⁰ → For cervical / vaginal / vulval cancer: Chest Abdominopelvic CT for assessing lymphadenopathy or PET-CT for higher risk (1b1 and above staging results) and/or Specialist MDT, if required ¹² and Formulation of management plan ¹⁰
Patient information Provided in primary care ¹	→ Patient information Provided in consultation or OPA / one-stop clinic ⁵	→ Cancer ruled out and communication to patient; Record FDS if patient informed that cancer has been excluded. Referred to other secondary care service, RDC or discharge ¹⁰ OR Cancer diagnosed; Outpatient Clinic and communication to patient Record FDS if patient is informed they have cancer. ¹⁰ Discussion with CNS and specialist MDT input, discuss treatment options, subject to staging testing being reported; Personalised Care and Support Plan discussion ¹⁰		→ Confirmation of treatment plan and Ongoing Personalised Care and Support Plan discussion with patient ¹⁰

Need for speed?



Figure 3: deaths from cancer (per 100,000 inhabitants), OECD countries, 2020

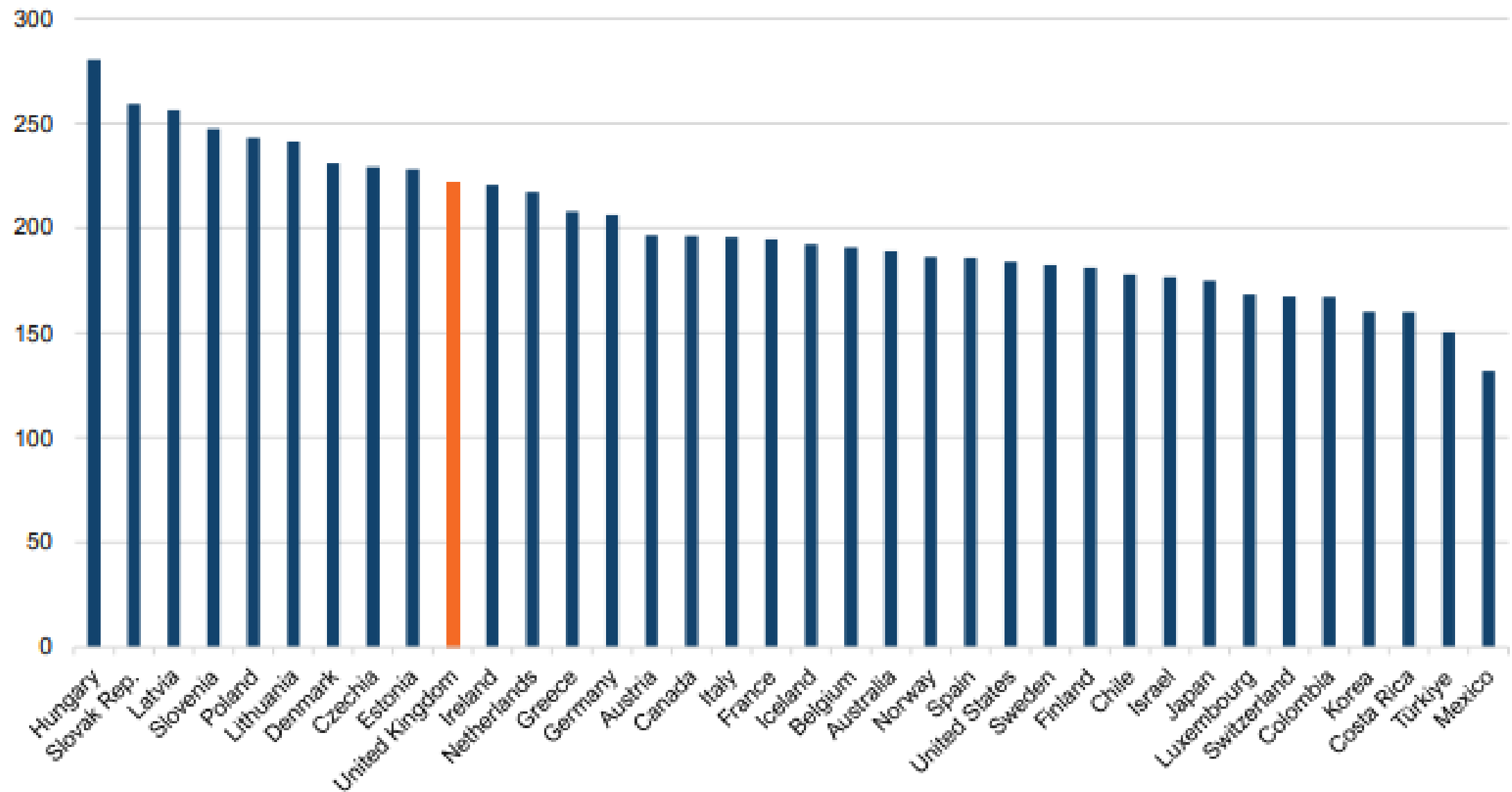
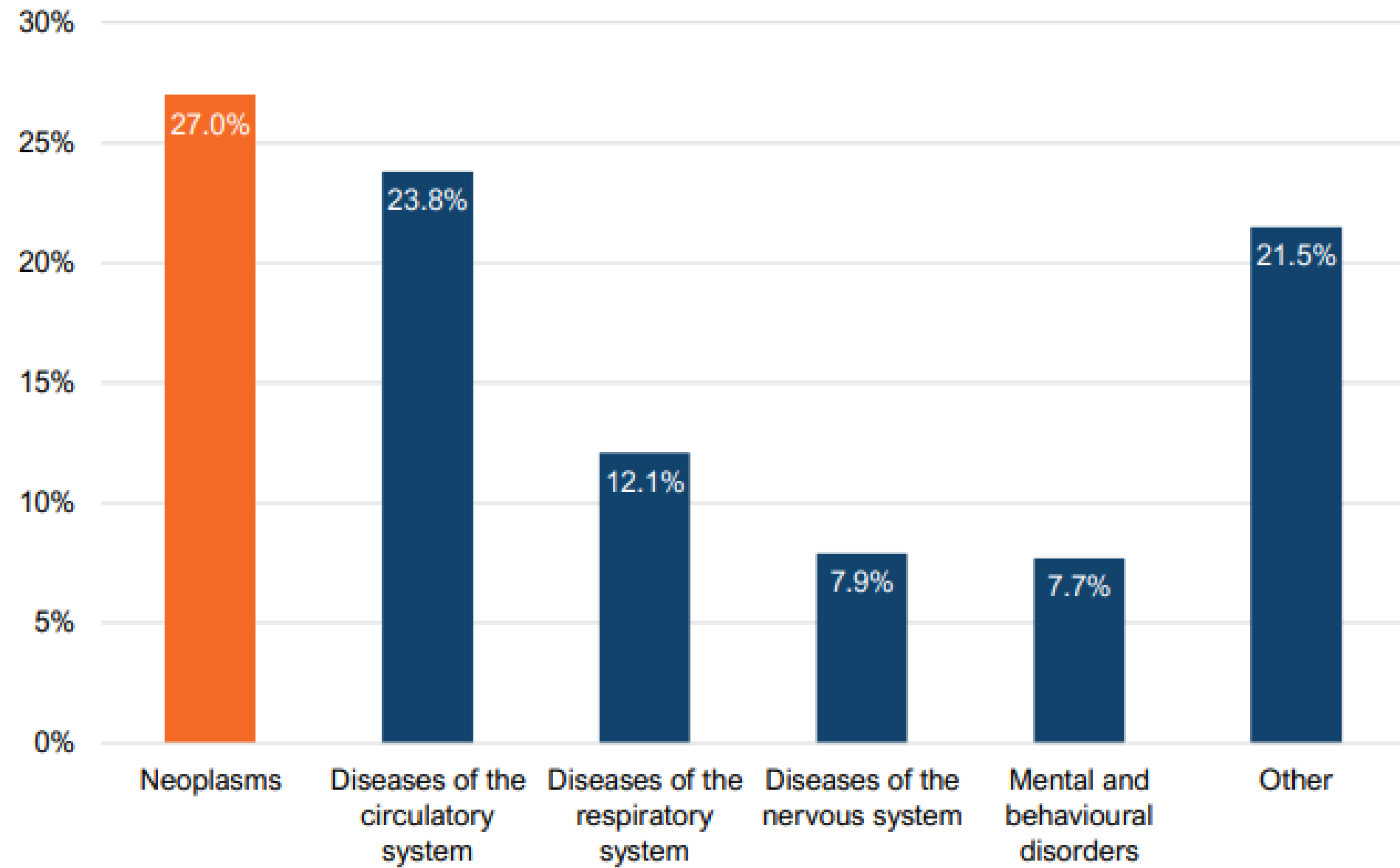


Figure 7: most common causes of death as a proportion of all deaths, 2024, England

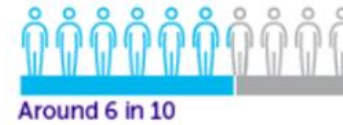


5 year survival
significantly
increased at
early stage

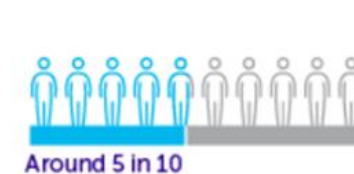
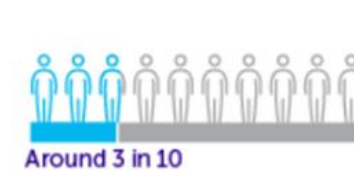
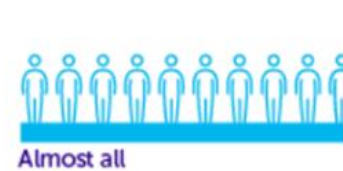
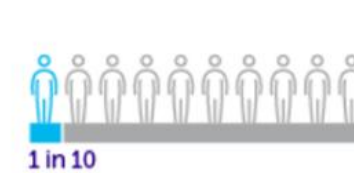
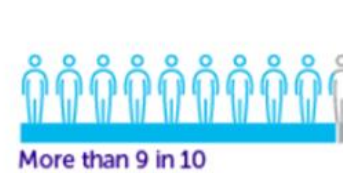
Cancer survival by stage at diagnosis

Proportion of people surviving their cancer for five years or more

Diagnosed at earliest stage



Diagnosed at latest stage

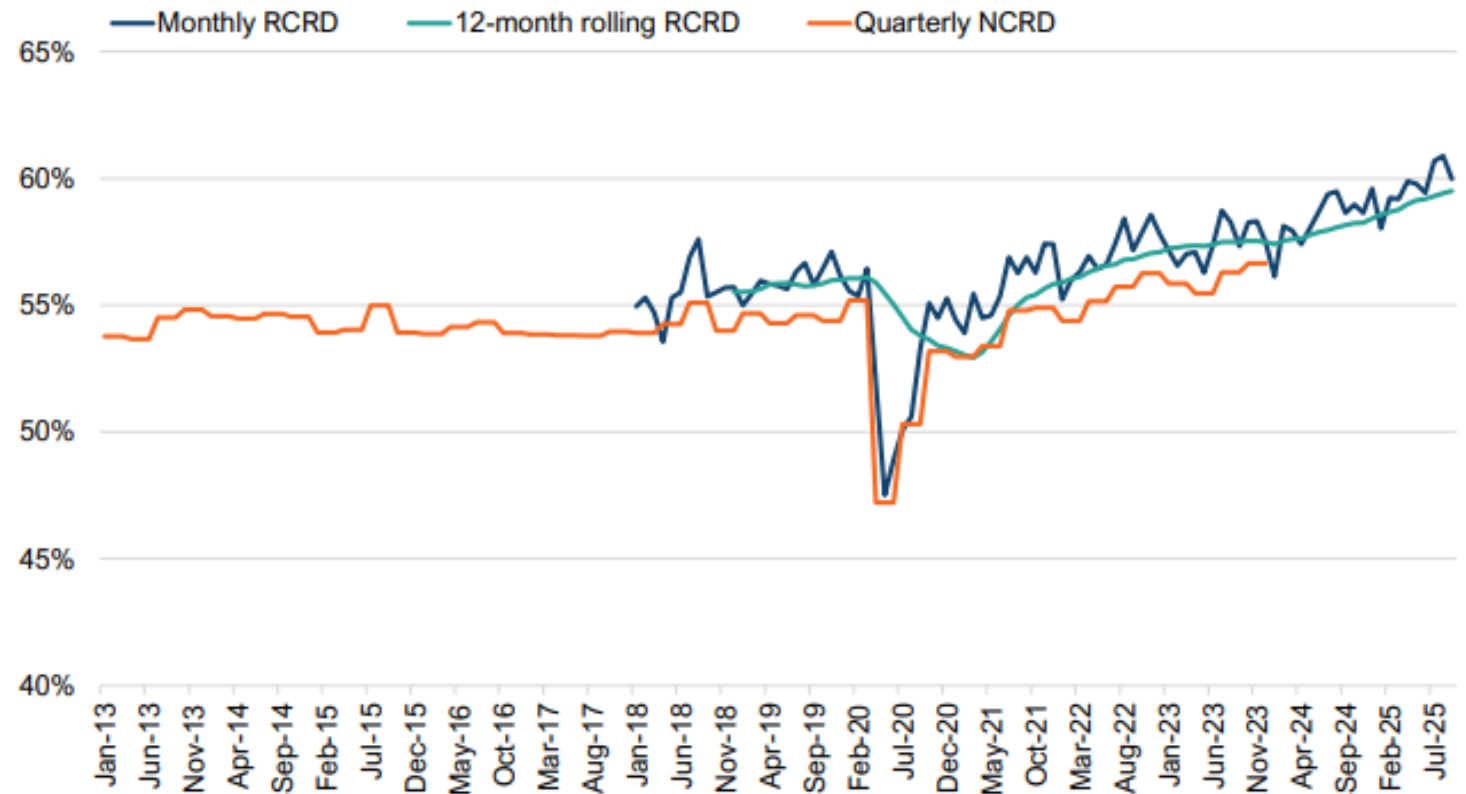


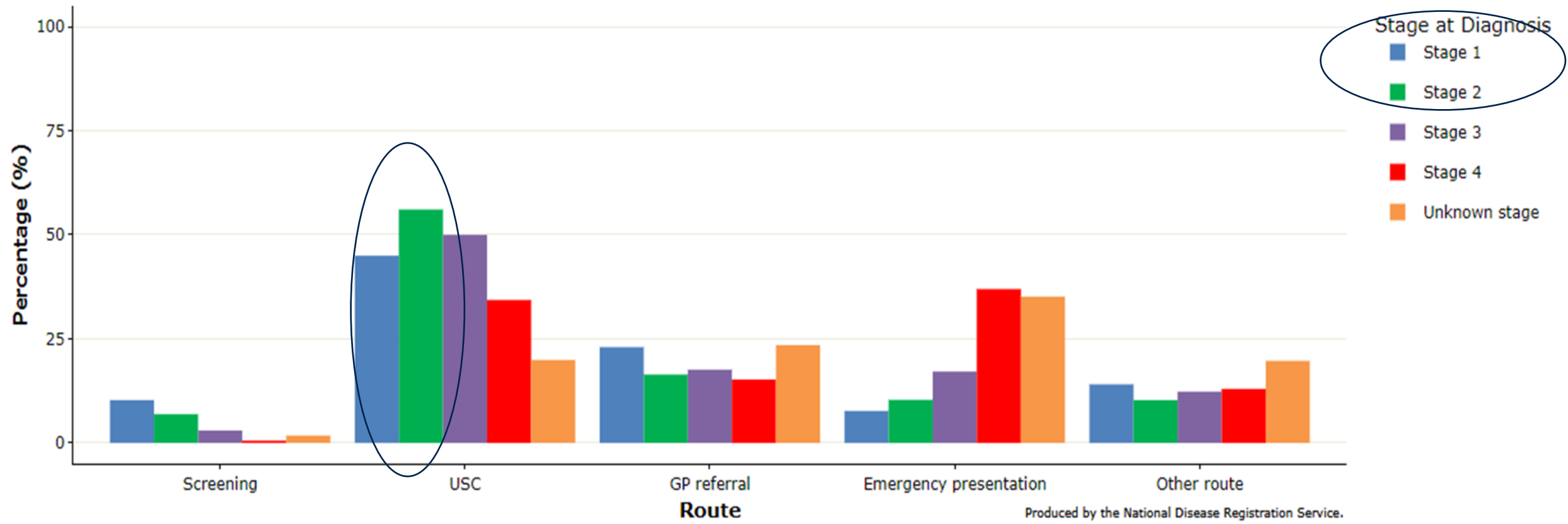
Earliest stage = stage 1; latest stage = stage 4.
Data for lung, bowel and breast cancer is age-standardised net survival for adults (aged 15 to 99 years) in England in 2015-2019 followed up to 2020. Data for prostate cancer is age-standardised net survival for adults (aged 15 to 99 years) in England in 2013-2017 followed up to 2018. Breast cancer data is for females only.
Source: Cancer survival in England, NHS Digital 2022.

Early diagnosis rates in 2024 and 2025 have been at their highest ever level.

Around 10,000 more people diagnosed at stages 1 and 2 since FDS became a performance target

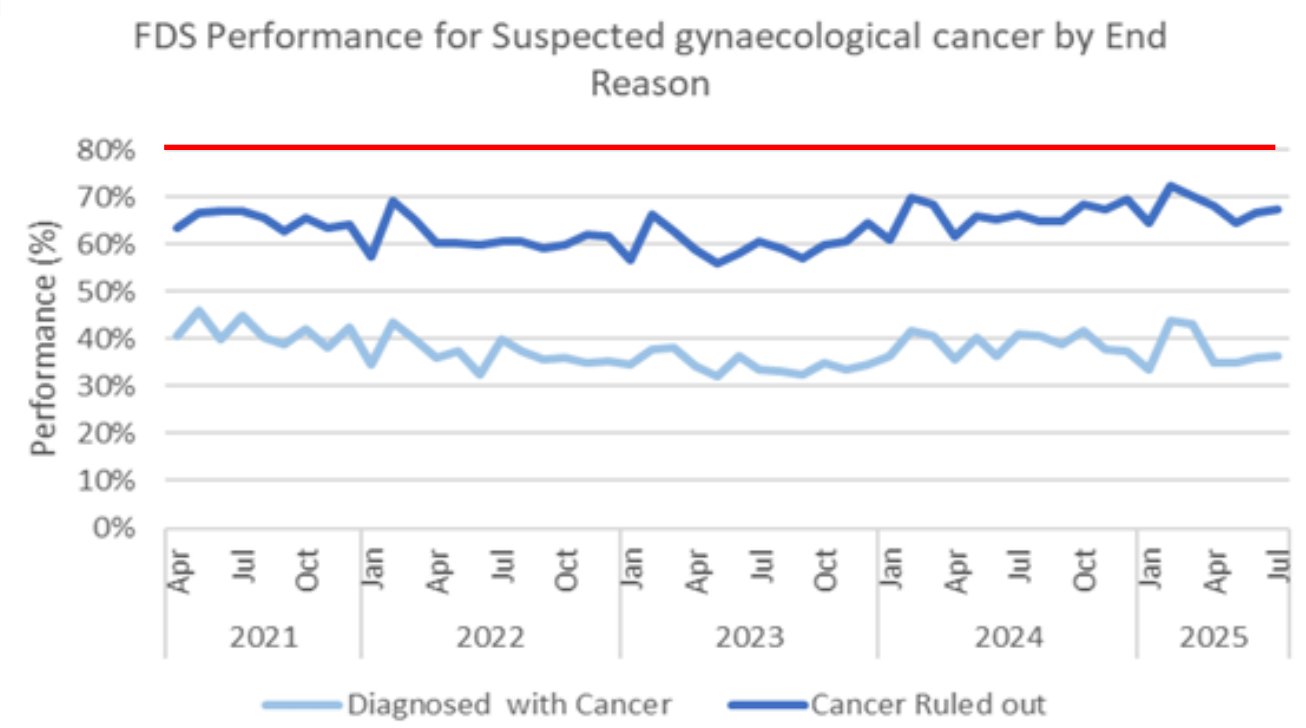
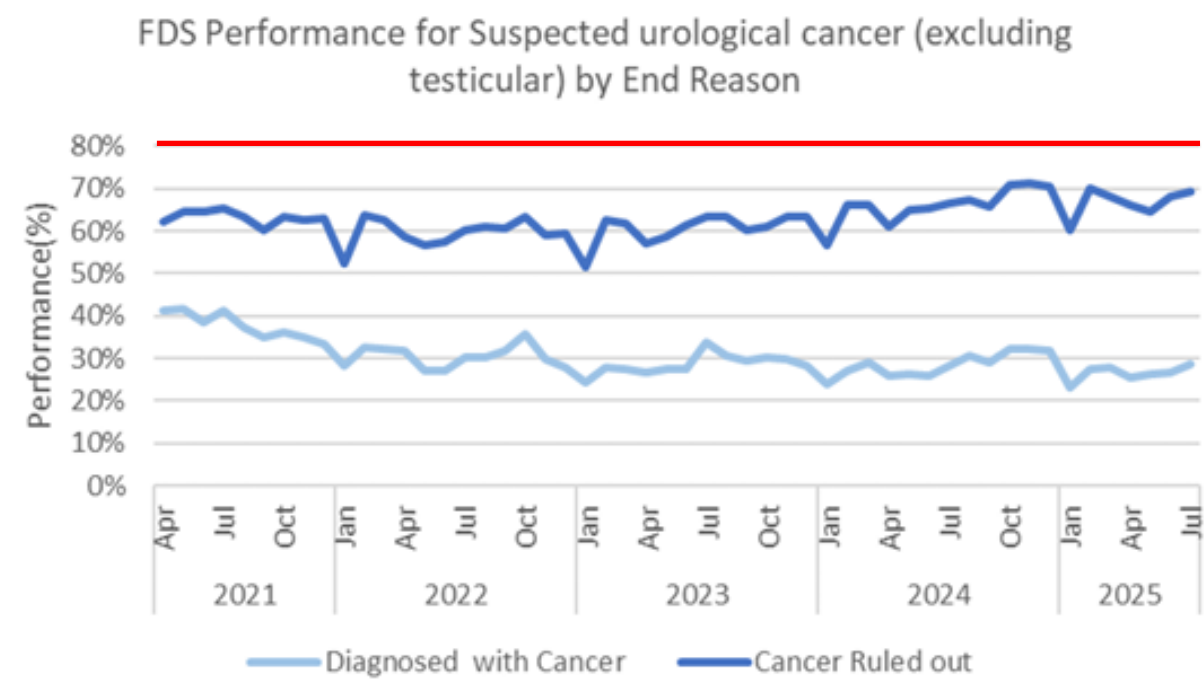
Figure 9: proportion of rapid cancer registrations (RCRD) and full cancer registrations (NCRD) diagnosed at stage 1 and 2, January 2013 to September 2025, England







[Cancer Waiting Time Statistics: Faster Diagnosis Standard: Statistics on End Reason](#)



Can we do more?



Sonographer led services

In our trust sonographers play a key role in achieving FDS

Sonographer role extension has been an underpinning part of the expansion of radiology services for over three decades.

Widely supported by radiologists and referring clinicians alike.

Evidence suggests they can provide holistic care, cut waiting times, ease staff pressures and increase job satisfaction.



Considerations

- Education and Training for your staff/team
- Protocols for training and competent sonographers
- Who will be the clinical lead and gold standard?
- Safety Considerations – written consent required etc.
- Are PGDs required?
- **Patients Journey through the system**
- Patient Information Leaflets
- Audit



Sonographer led services Hull

- Rapid Diagnostic Service
- Haematuria
- Prostate cancer
- Testicular cancer
- Ovarian cancer
- HCC (in development)



Hull urology cancer performance Sept 23 – Sept 24:

Diagnosis or cancer ruled out within 28 days of referral

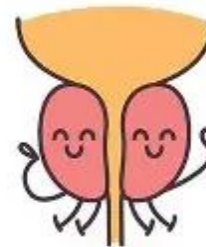
53.8% (75%) informed by day 28

One month (31-day) wait from diagnosis to first definitive treatment:

71.6% (96%) of people treated for urological cancers began first definitive treatment within 31 days of receiving their diagnosis

Two month (62-day) wait from urgent referral to first definitive treatment:

35.8% (85%) of people treated for urological cancers received first definitive treatment within 62 days of being referred



Haematuria



Haematuria – Traditional Pathway



Humber Health
Partnership

Various route referral into urology

*Urology GIRFT
recommended pathway**

One stop clinic – ultrasound &
cystoscopy*

60% were found to be normal

40% were having a CT examination,
regardless of the ultrasound findings

High risk patients have additional
CT Urogram

Long waits to enter clinics to find out
treatment options

Return for CT staging if bladder
cancer found

6% cancer detection rate on whole pathway

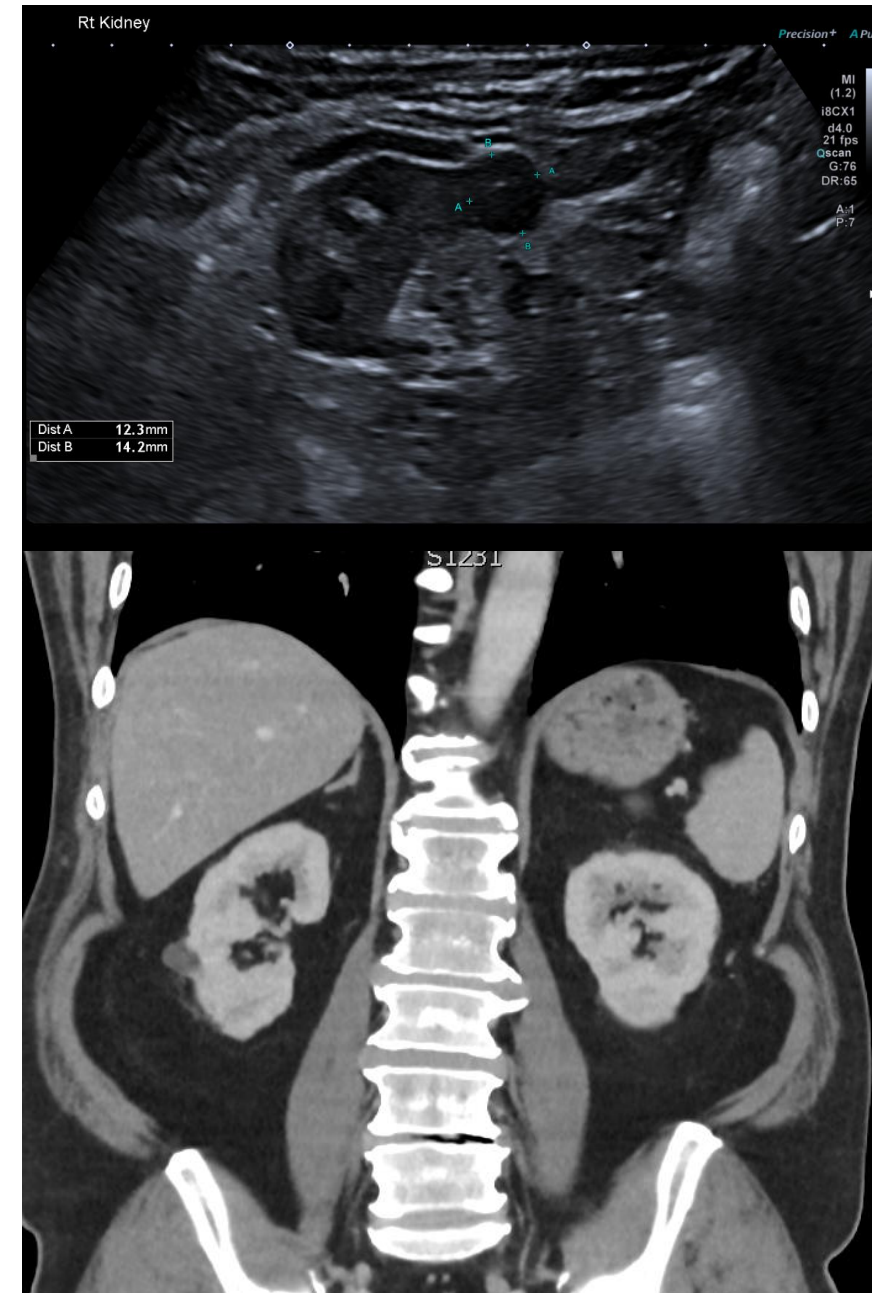
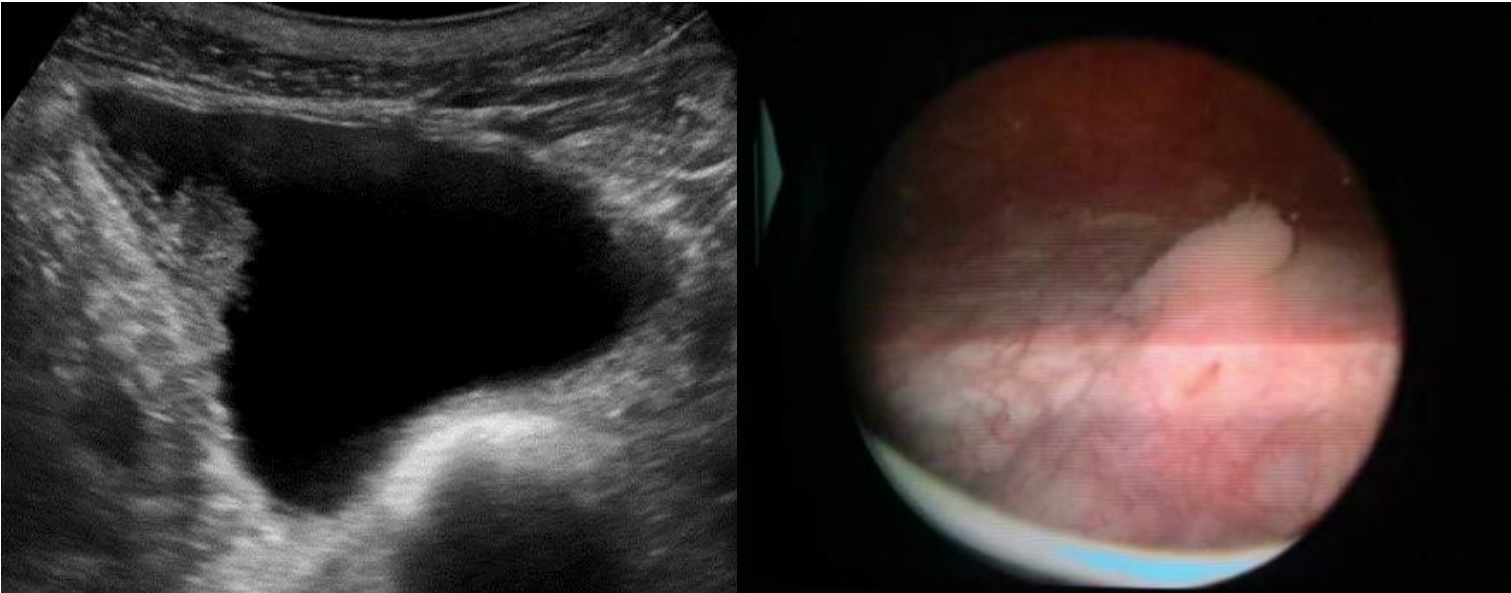
One-stop US imaging capacity

	Number of lists	Number of patients	
Potential capacity	26	312	
Lists cancelled by Urology including bank holidays	5	60	These spaces were back filled by general US patients
Lists not filled but ultrasound not informed	4	48	Some slots backfilled by ultrasound when became aware no clinic patients booked- This is dependent on clerical staff being available to contact patients by phone.
Actual slots used by Urology	17	204	
Patients booked		134	
Patients attended		101	
% potential capacity used		32.3%	

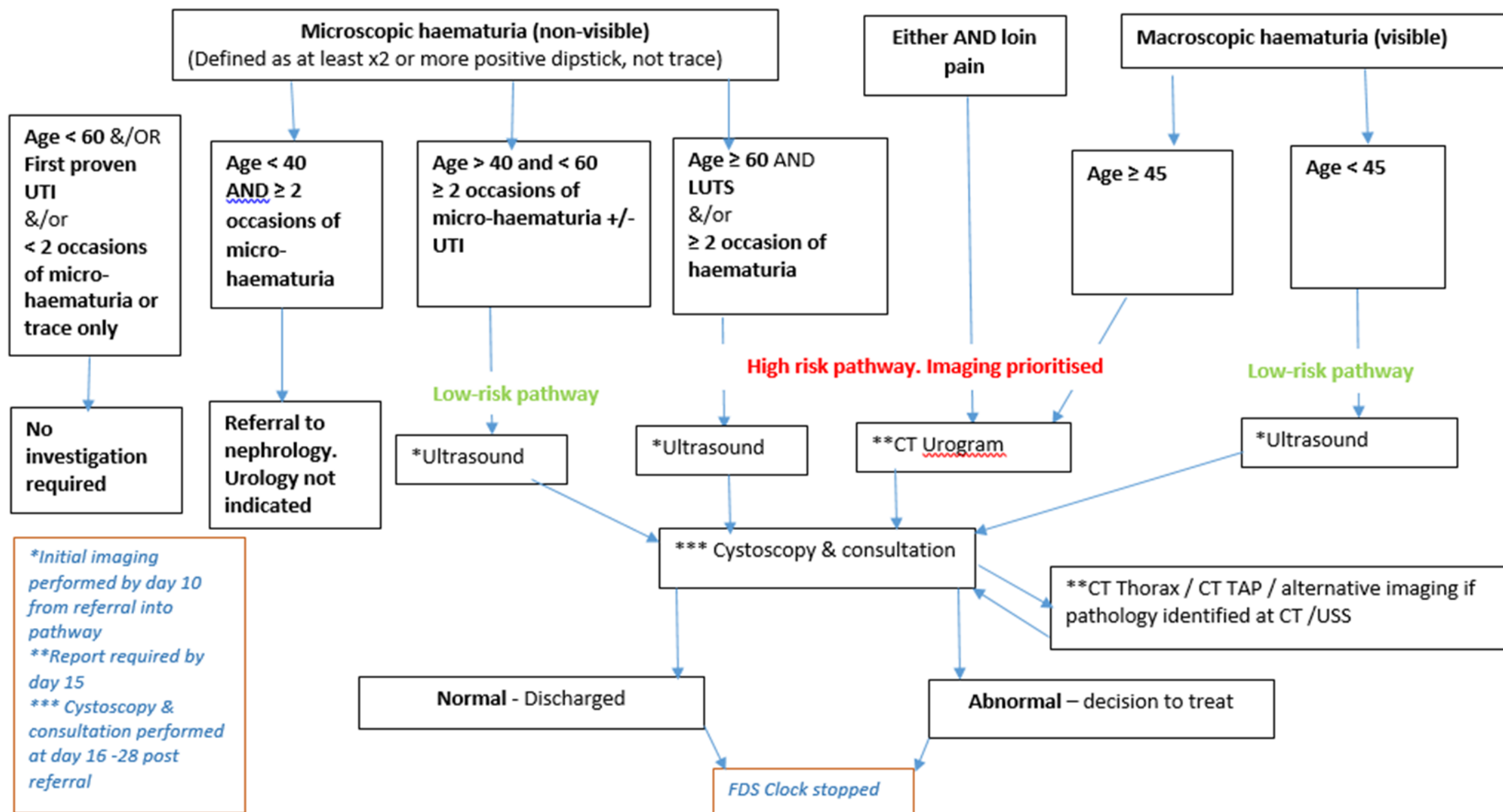
Traditional one stop

Actually, is the ultrasound needed?

And is a one stop the right way forward?



**Perform U&E in all patients at time of referral. Referrals with test not placed will be rejected.
Perform PSA if applicable, and with informed consent**



Haematuria – New Pathway

One route into
urology
Haematuria DOS

Cystoscopy arranged at
same time as first
consultation in urology

This date then placed onto
radiology request

Imaging
arranged prior to
cystoscopy
allowing time for
reporting

Decision to
treat/decision to
discharge given at
cystoscopy
appointment

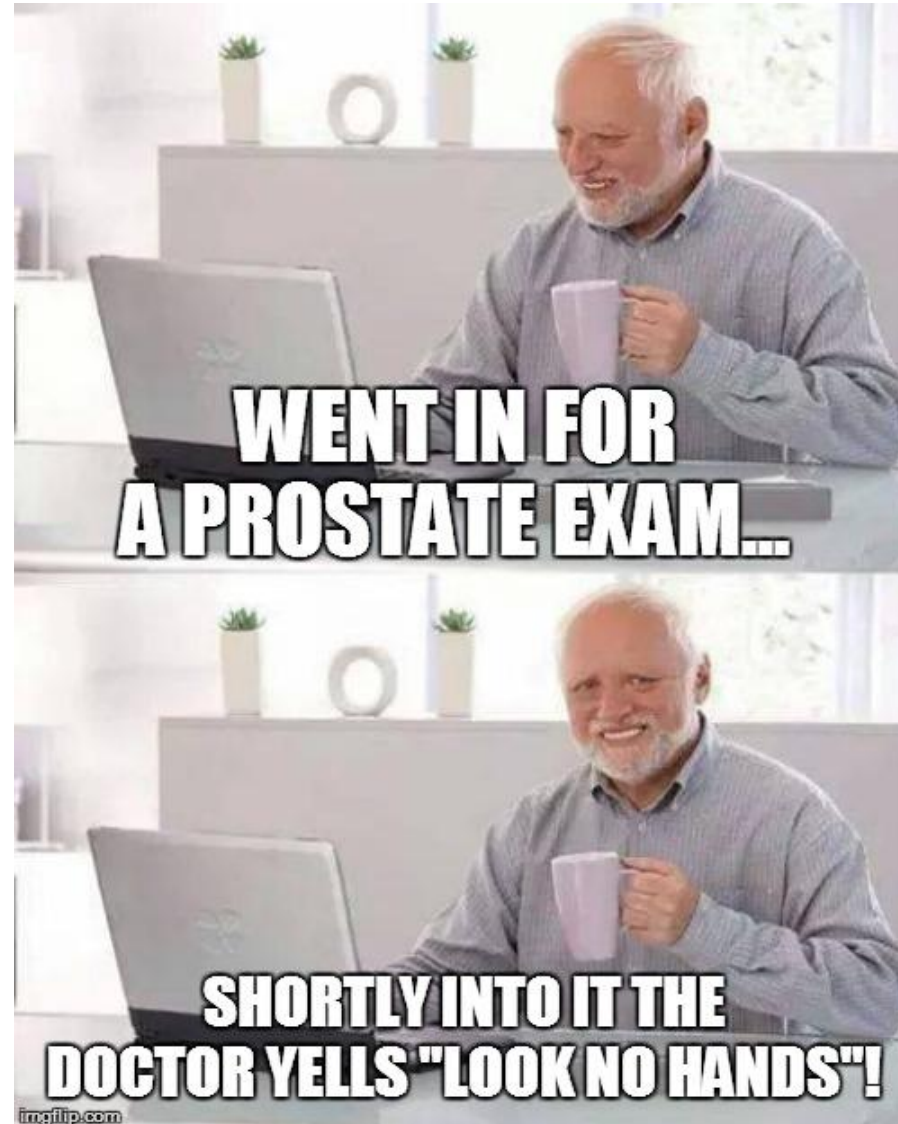
FDS pre pathway changes

Referral → patient informed of results = 65 day average

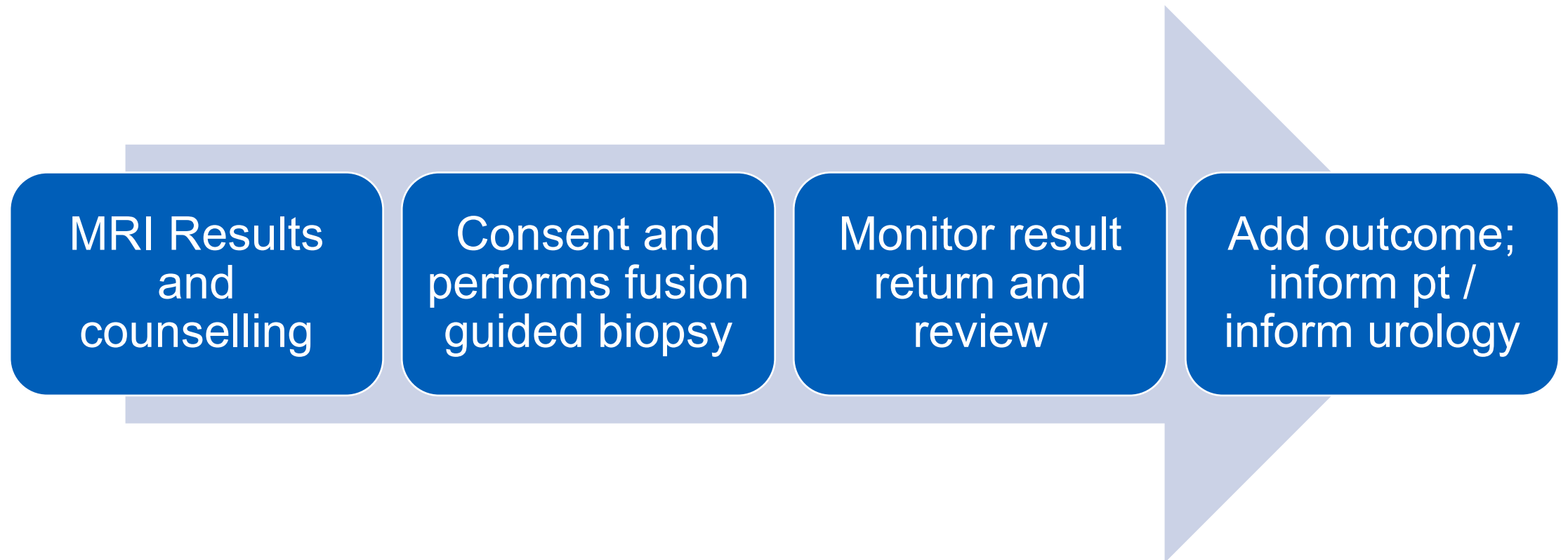
FDS post pathway changes

Referral → patient informed of results = 25 day average

Suspected Prostate Cancer



Sonographer role in PCa pathway



Prostate Outcomes

- Histology results emailed to prostate trackers
- Prostate trackers contact sonographer who performed biopsy
- Sonographer reviews histology, places outcome onto report & informs patient if benign
- Outcome emailed to urology & trackers
- Clock stopped if benign. MDT discussion if abnormal

Outcome Code	Description	Histology	Action
RPP1 Outcome1	Prostate Cancer	Gleason 3+3 and above	MDT discussion or clinical review
RPP 2 Outcome 2	Clinically significant Benign disease	High Grade PIN, ASAP	Not for MDT but follow up in urology. Sonographer contacts patient with results. Contact recorded and 2ww clock stopped
RPP 3 Outcome 3	Disparity between MRI and biopsy. ie PIRADS 4 and 5 on MRI	High Grade PIN, ASAP or benign disease	MDT Discussion required Sonographer contacts patient with results. Contact recorded and 2ww clock stopped
RPP 4 Outcome 4	Clinically insignificant Benign disease. (Includes low grade PIN, acute inflammation and prostatitis)		Not for MDT. Sonographer contacts patient with results. Contact recorded and 2ww clock stopped.

Prostate Summary

FDS pre pathway changes

Referral → patient informed of results

48 day average if positive

69 day average if benign

FDS post pathway changes

Referral → patient informed of results = 27 day average

Prostate cancer performance

Diagnosis or cancer ruled out within 28 days of referral

88.1% (80%) informed by day 28

One month (31-day) wait from diagnosis to first definitive treatment:

80.9% (96%) of people treated for urological cancers began first definitive treatment within 31 days of receiving their diagnosis

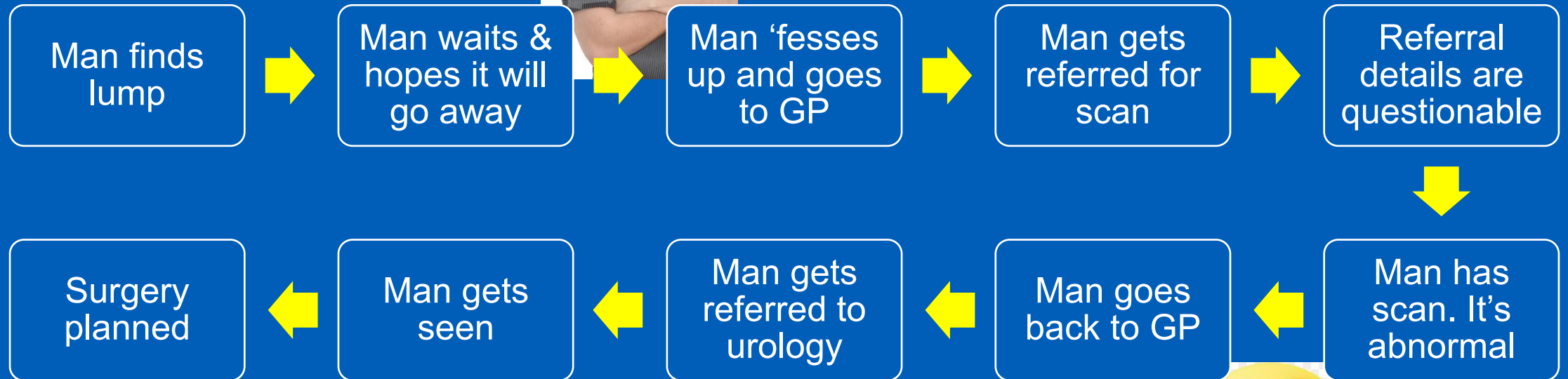
Two month (62-day) wait from urgent referral to first definitive treatment:

68.1% (85%) of people treated for urological cancers received first definitive treatment within 62 days of being referred

Suspected Testicular Cancer



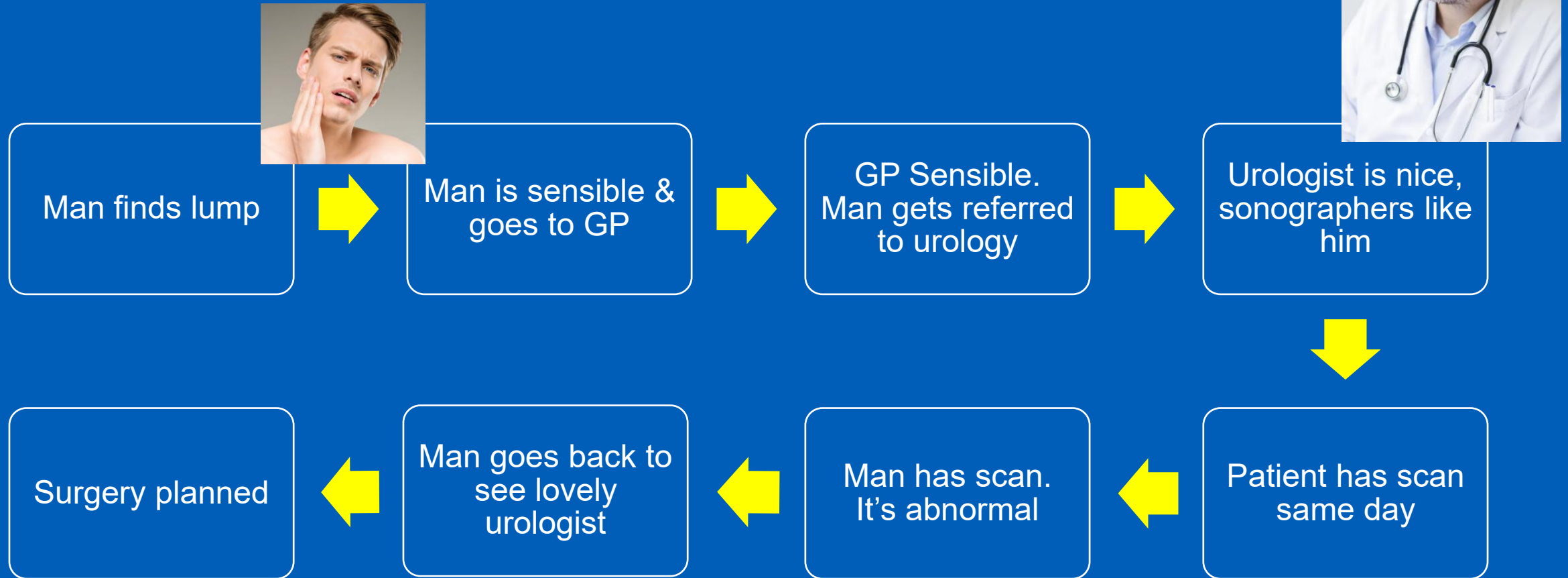
GP Pathway



Pathway ~ 66 days



Urology Pathway



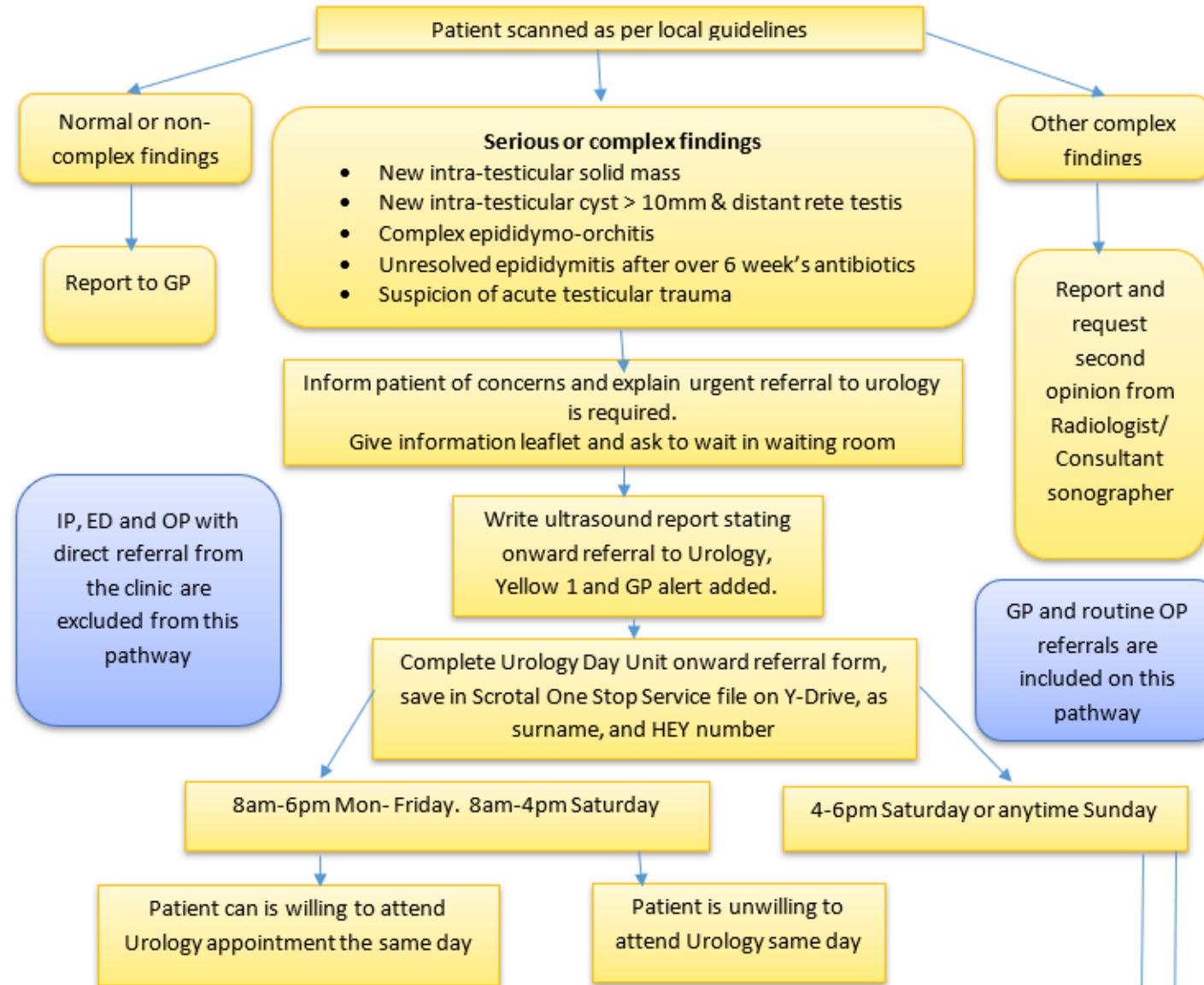
Common referral to treatment pathway **20 days**

Pathway considerations:

- Direct to urology is clinician and patient dependent
- Clinical information on US referral triage
 - Routine – 42 days (or more)
 - Urgent < 7 days
- Alert of US report; does it get read?
- Sonographers commonly faced with giving bad news
- Can we fast track?

Referrals from Primary

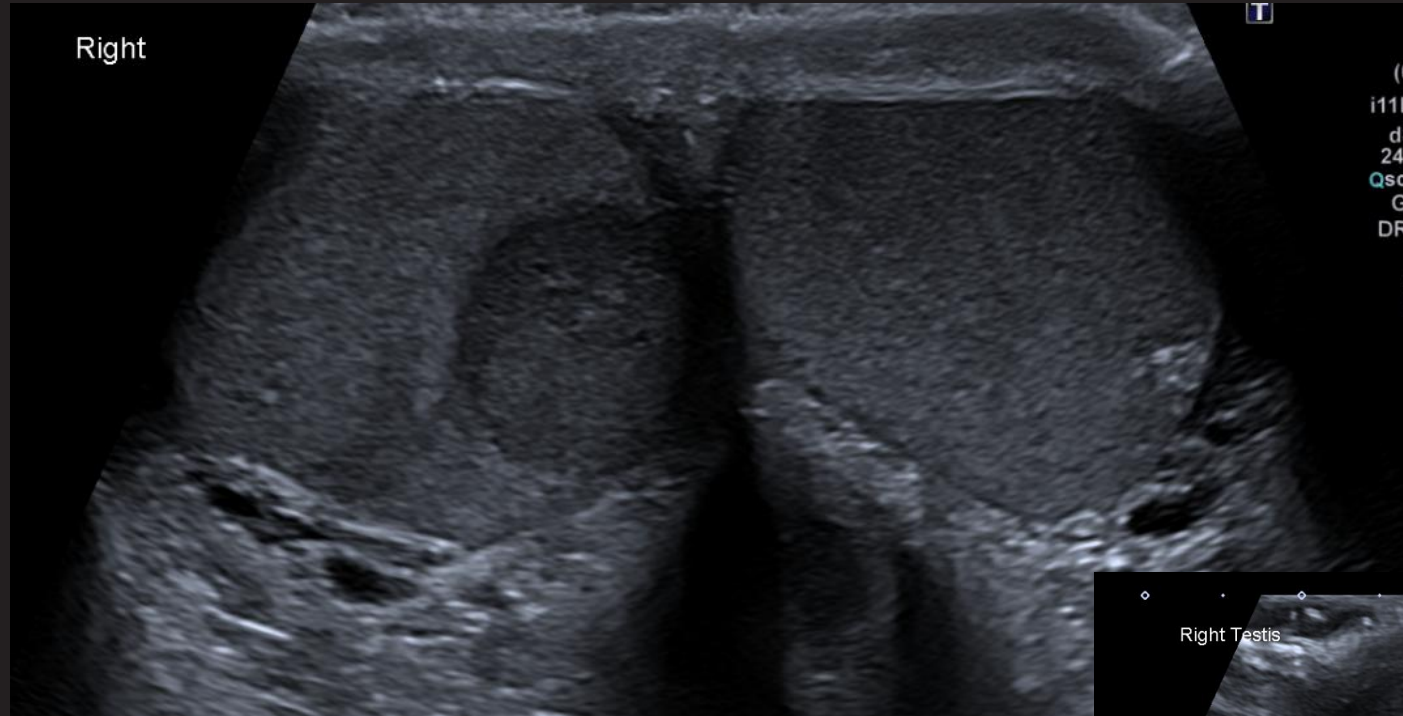
Flow Chart for Management of Incidental Significant and/or complex Scrotal Pathology



Testicular cancer pathway summary

- **Motivator:** Unnecessary steps & delays in GP pathway
- **Concern:** patients having verbal results but delays in accessing GP appointments for discussion & onward referrals
- **Solution:** sonographer led onward referral process
- **Result:** Quicker access to management and support for patients

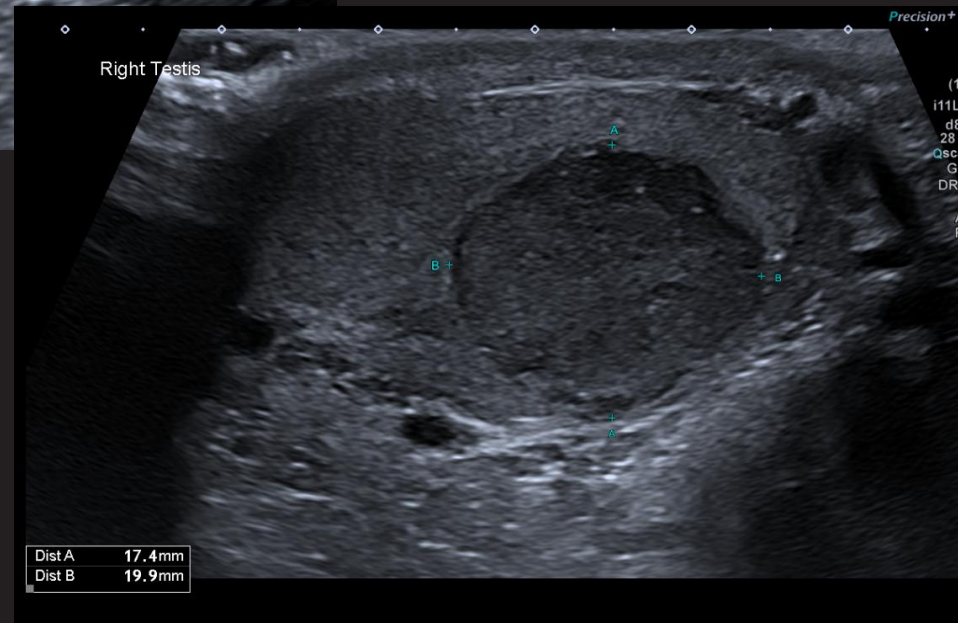
Case 2: 37 y/o. GP referral. Hard lump. Urgent scan at triage



Mixed germ cell tumour

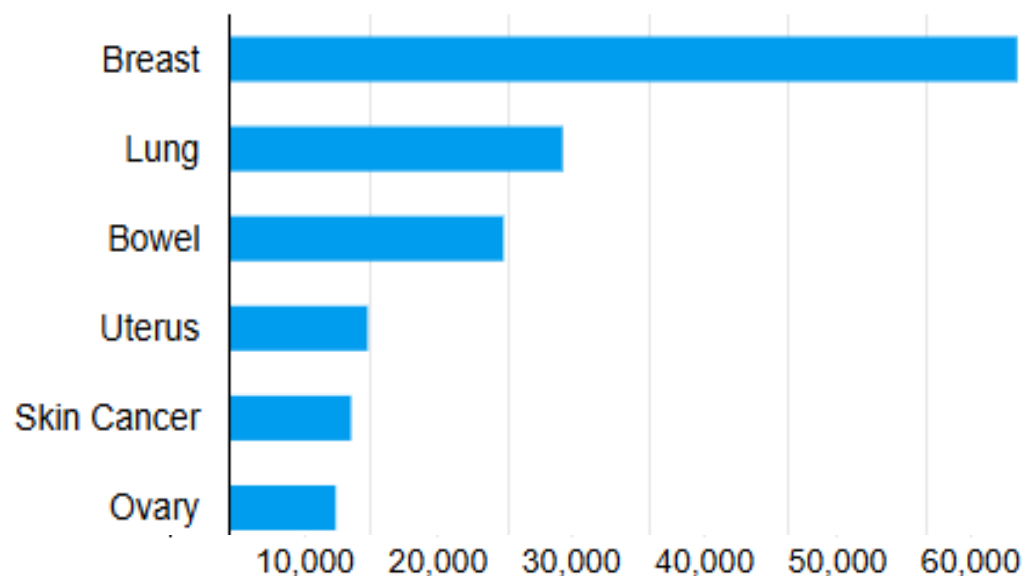
Referred from GP 13/10/2025.
Scan, sonographer direct
referred to urology day unit &
patient seen 14/10/25,
orchidectomy 25/10/2025

Total scan to treatment – 12 days



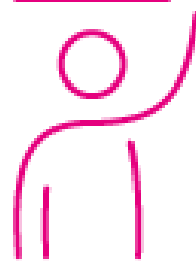
Suspected Ovarian Cancer





Survival

35%



Survive ovarian cancer for 10 or more years, 2013-2017, England

Survival

76%



Survive breast cancer for 10 or more years, 2013-2017, England

Survival

78%



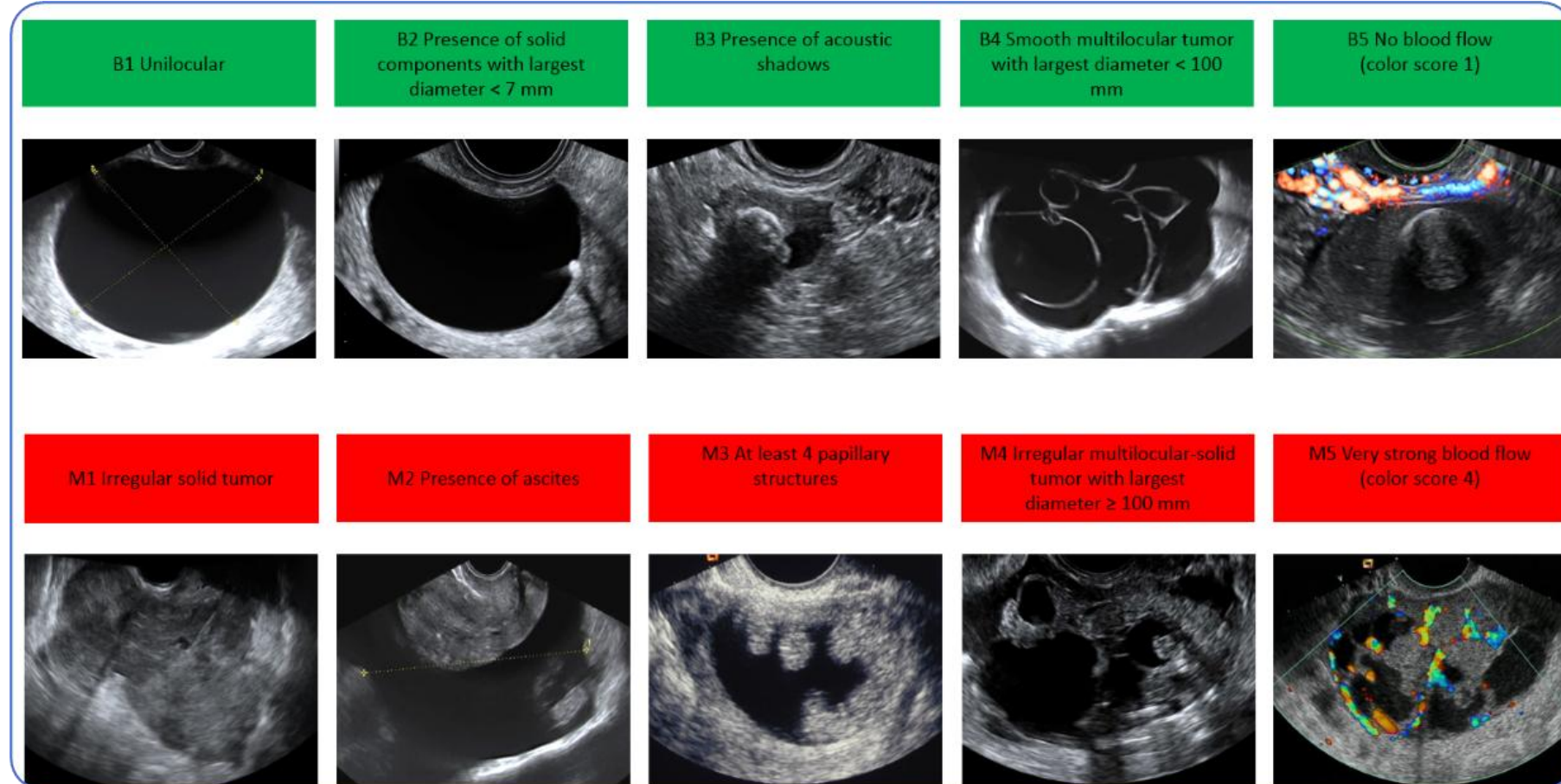
Survive prostate cancer for 10 or more years, 2013-2017, England

Suspected Ovarian Cancer – IOTA Simple Rules

When the simple rules applied conclusive in approximately 75% of adnexal masses (for malignant/benign)

25% indeterminate

Unable to determine tumour subtype



ADNEX

Diagnostic risk prediction model

3 clinical predictors

6 ultrasound predictors

Estimates the risk of five tumour types

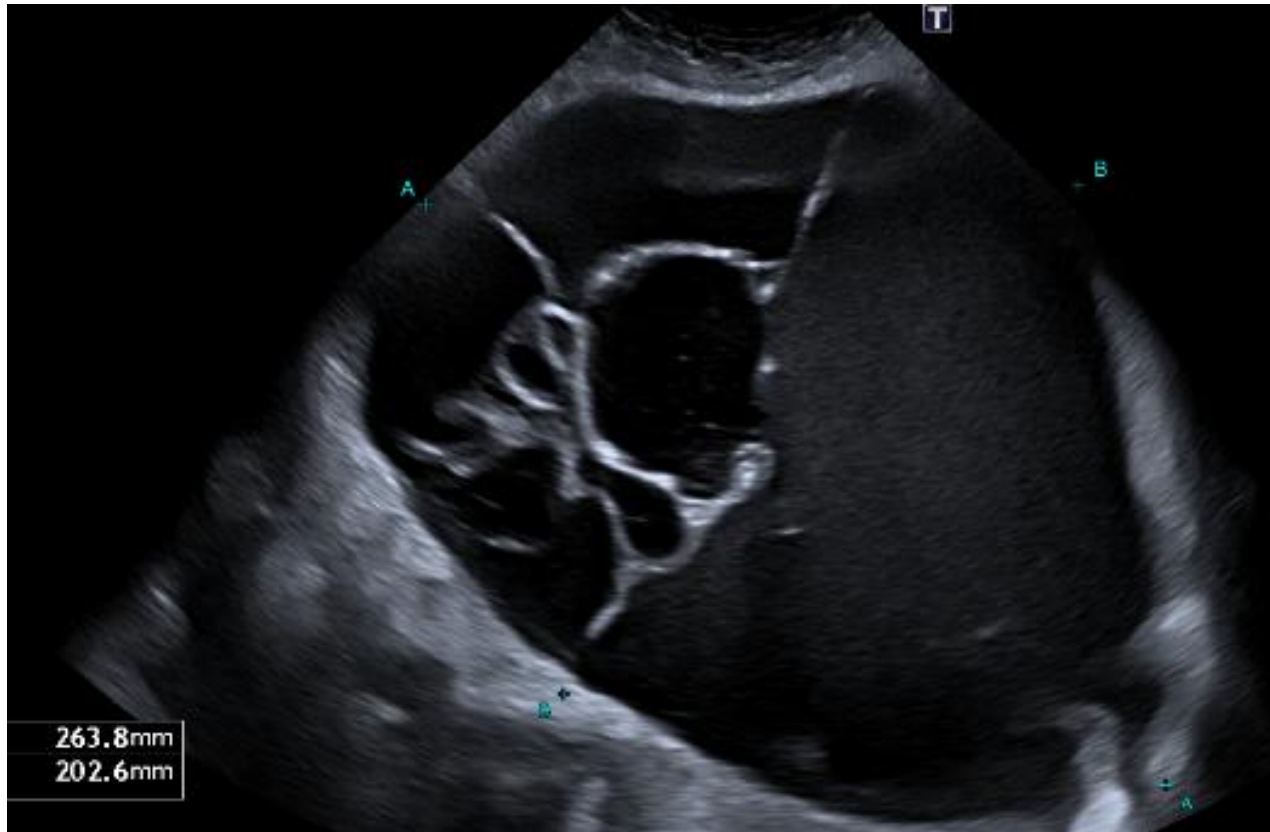
- Benign
- Borderline
- Stage I primary invasive
- Stage II-IV
- Metastatic

Clinical Predictors:

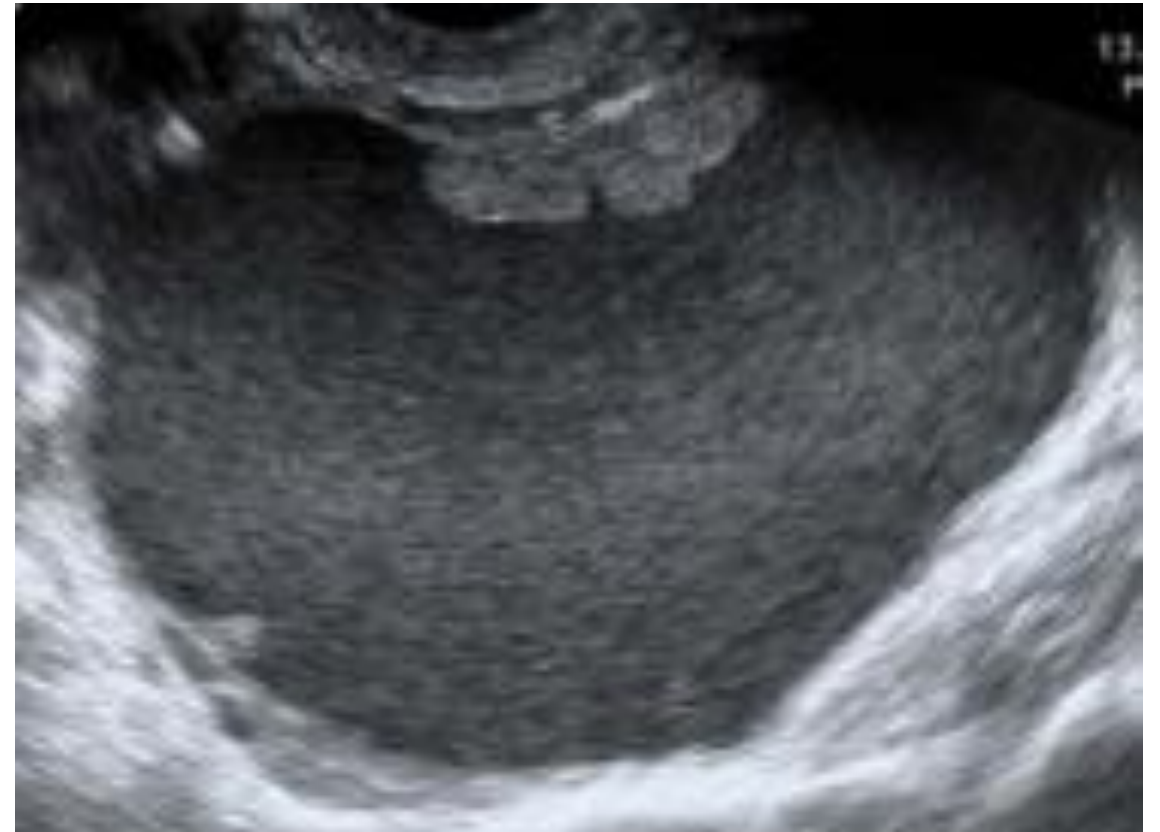
- Age (years)
- Serum CA-125 (U/mL)
- Type of centre to which the patient has been referred for ultrasound examination (i.e. oncology centre).

Ultrasound Predictors

Maximal diameter of the lesion (mm)



Number of papillary projections



Maximum diameter of solid tissue



Ascites



Presence of more than 10 cyst
locules



Acoustic shadows



How it looks:

Age



Oncology center

Is your center a tertiary referral center with a specific gynecological unit?

No Yes

Maximal diameter of the lesion

Largest of the diameters measured in three perpendicular planes



Maximal diameter of the largest solid part

Note: The maximal diameter of the largest solid part can never be larger than the maximal diameter of the lesion.



Cyst locules

Is the lesion multilocular or multilocular-solid with more than 10 cystic locules that are separated by septa?

0-10 >10

Number of papillations (papillary projections)

Papillations must be at least 3 mm. Note: If papillations are present the maximum diameter of the largest solid part cannot be 0!

None 1 2 3 >3

Acoustic shadow present

Presence of acoustic shadows is defined as loss of an acoustic echo behind a sound-absorbing structure

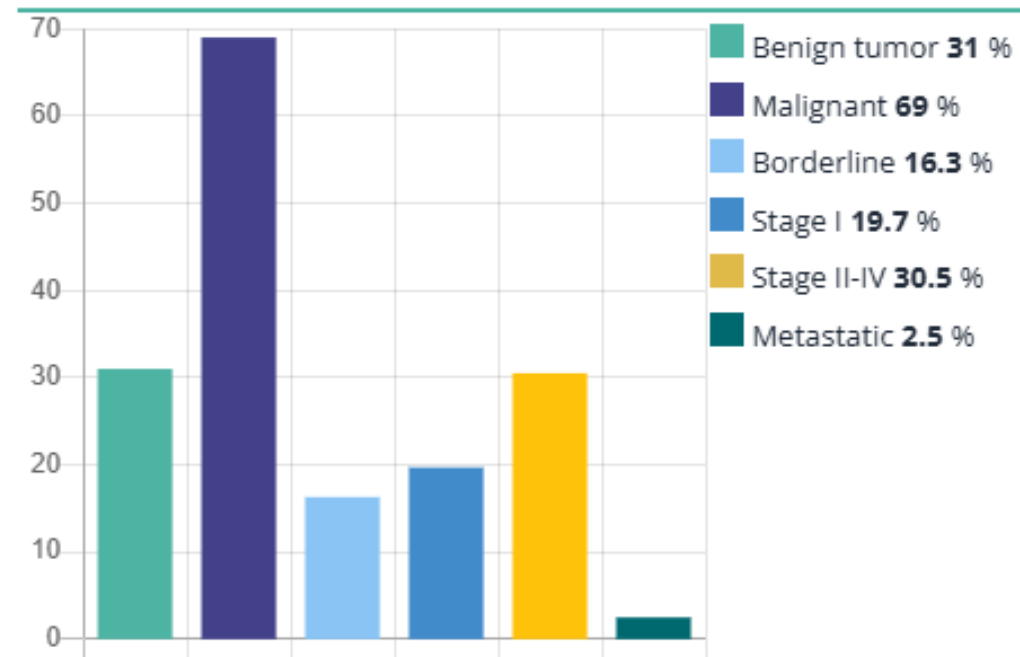
No Yes

Ascites present

Is there ascites outside the pelvis present?

No Yes

Serum CA-125



Pilot Aims:

- Performance compared to histology?
- Performance compared to MRI?

Could MRI capacity could be relieved with its implementation?

Could it help support FDS?

Step down indeterminate lesions that were benign

Escalate sinister masses

Results

ADNEX stage	Histology findings
2-4	FIGO 2B
Benign	Benign Cystadenoma
1	FIGO 1A
2-4	FIGO 3B
Benign	Cystadenoma
Stage 1	FIGO 1C
Benign	Cystadenoma
2-4	FIGO 2A
Benign	Cystadenoma
Borderline	Mucinous borderline tumour
2-4	Omental, peritoneal and ovarian involvement.
2-4	High grade tumour involving tubes, ovary and peritoneum
Borderline	Mucinous borderline tumour
Benign	Cystadenoma
Stage 1	FIGO 2B
Benign	Cystadenoma
Benign	Cystadenoma
2-4	Metastatic colon cancer
1	Primary uterine sarcoma

MRI result	Histology findings
FIGO 3A1	FIGO 2B
Cystadenoma – size of concern	Benign
Borderline	Borderline
Malignant/borderline	FIGO 1C
Benign	FIGO 1C
Cystadenoma	Cystadenoma
Unsure for MDT discussion	FIGO 2A
Cystadenoma	Cystadenoma
Cystadenoma	Cystadenoma
FIGO 3C2	Peritoneal, ovarian and omental involvement
? borderline ? benign	FIGO 2B
Cystadenoma	Cystadenoma
Malignant	Borderline
? borderline	Cystadenoma

ADNEX = 100% agreement
with histology

(n=21)

Results

Average 20 days from US to MRI

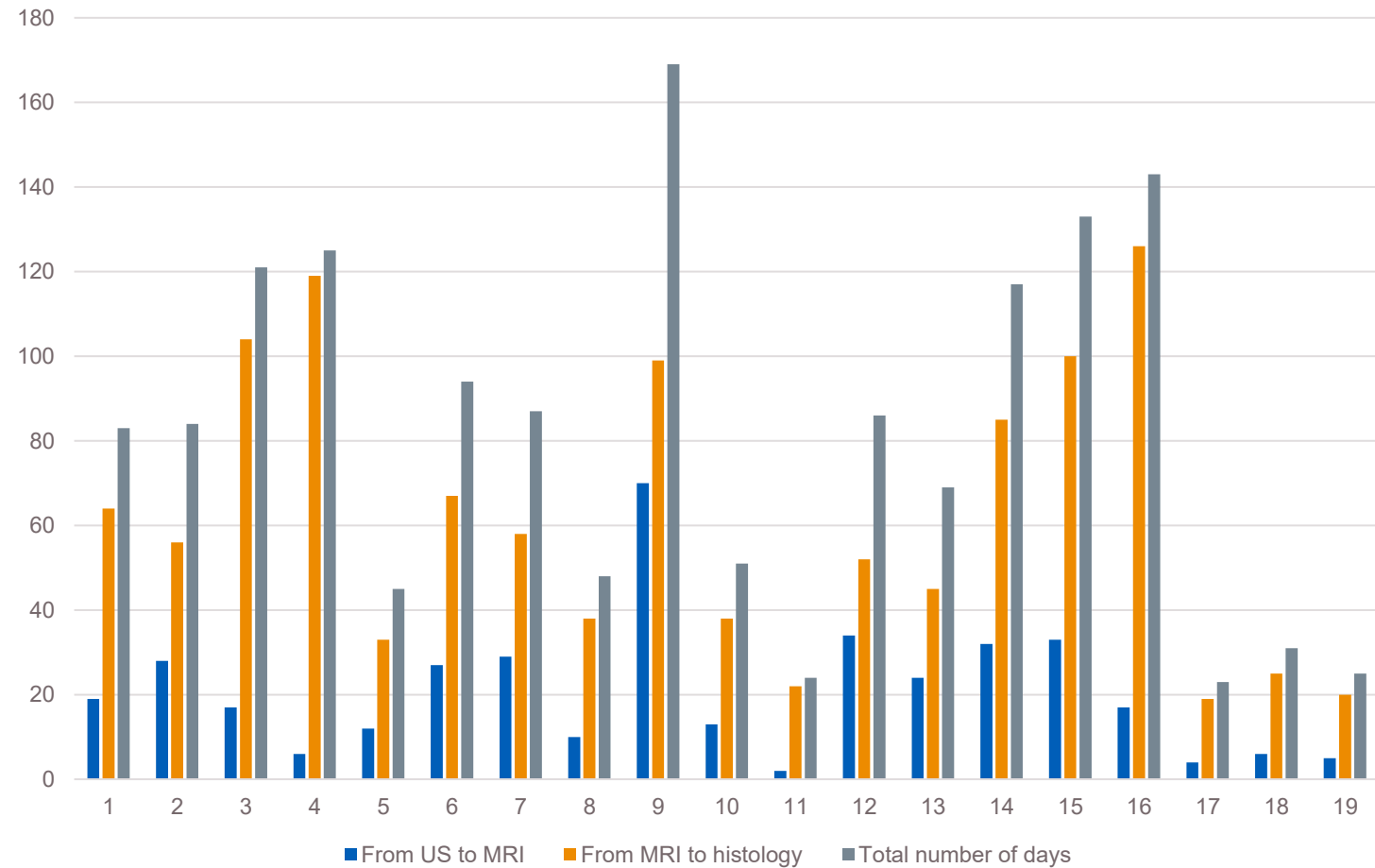
Average 61 days from US suspicion to removal

wait	Shortest wait	Longest
MRI/CT	2 days	70 days
Surgery	20 days	126 days

x2 of these radiology escalation - the two shortest waits

	2 days	5 days
MRI/CT	2 days	5 days
Surgery	20 days	25 days

Ovarian mass found at Ultrasound to Diagnosis in days

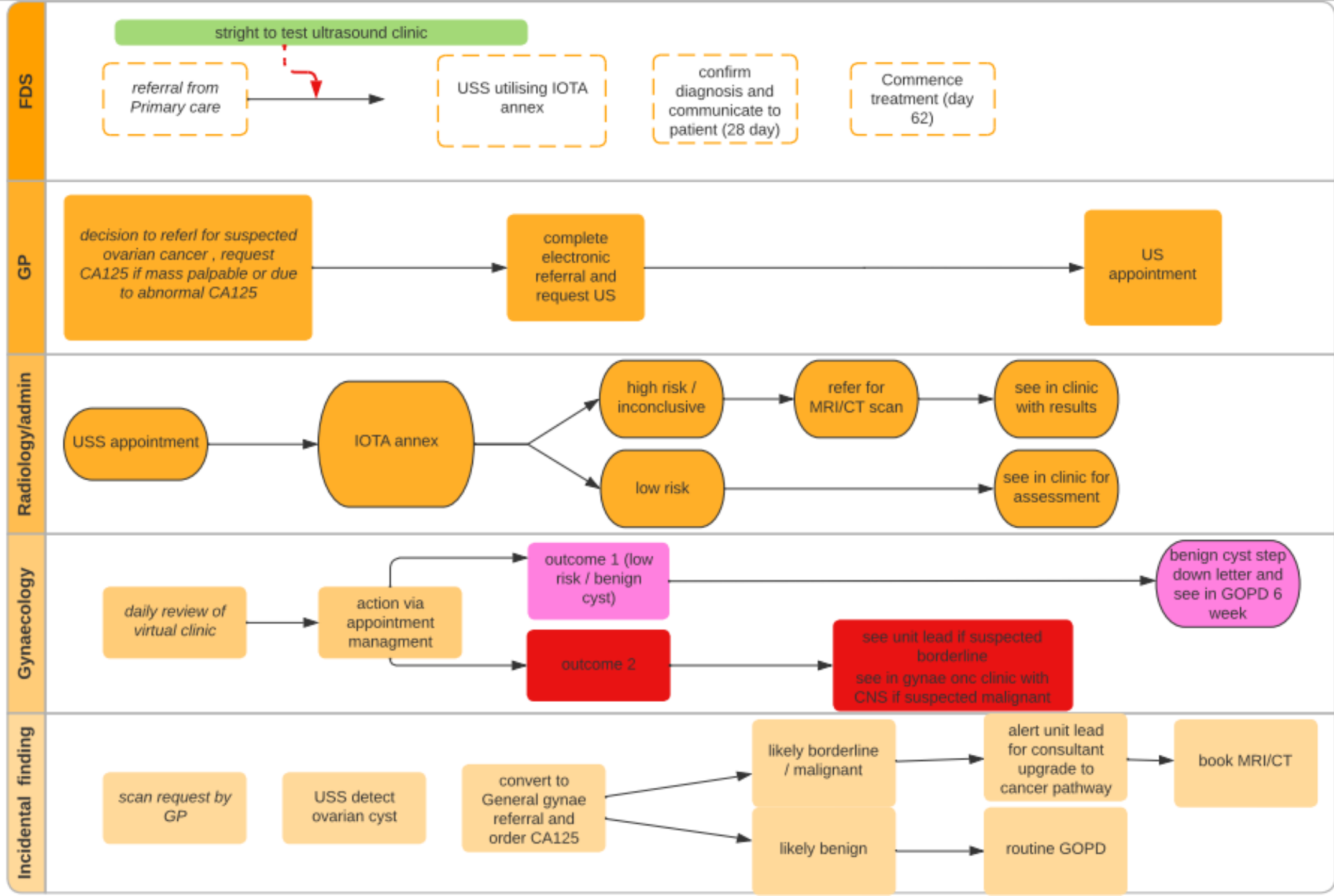


Results

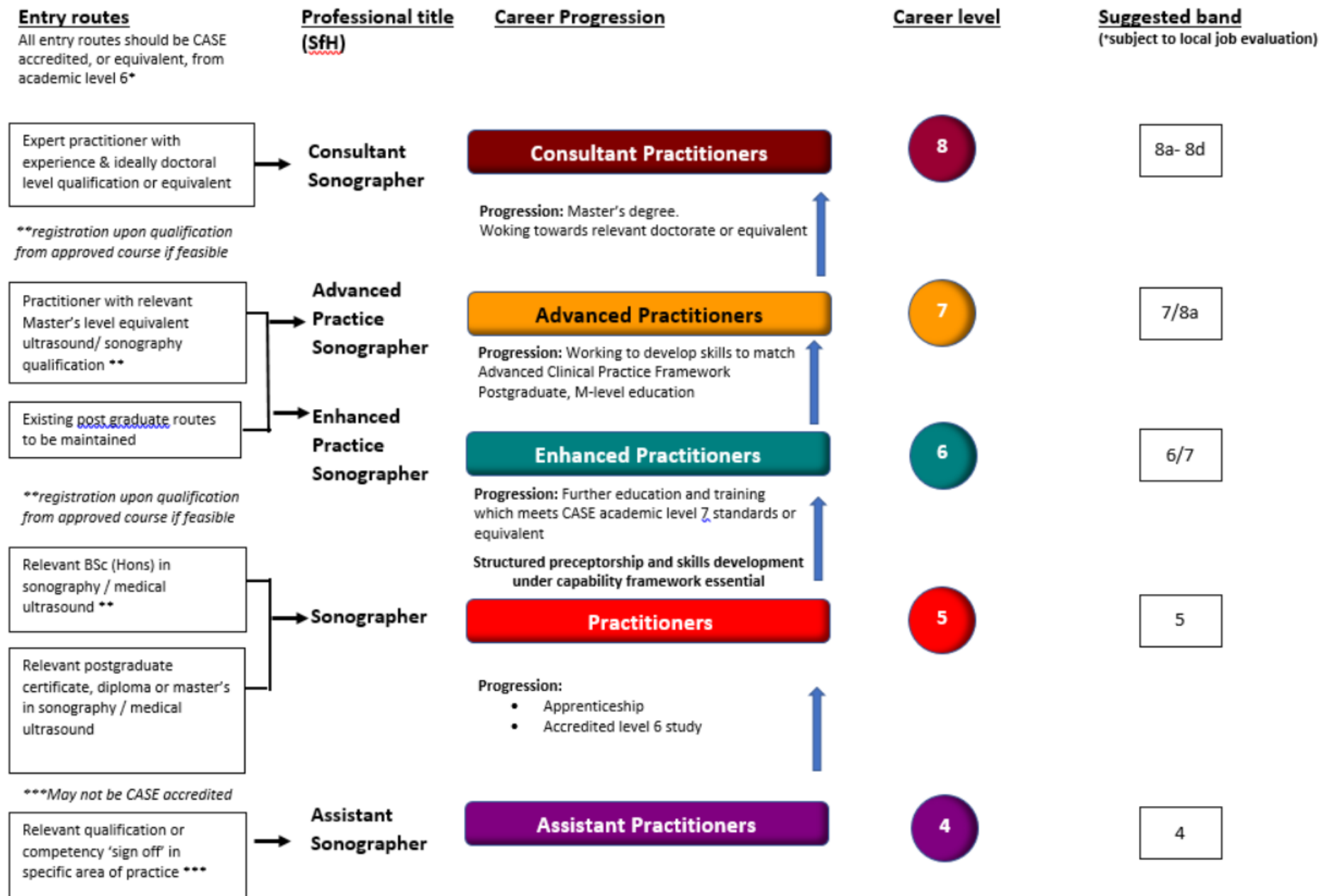
Escalation within radiology drastically reduced waiting

	Fast tracked case
MRI/CT	6 days
Surgery	19 days

x2 of these radiology escalation - the two shortest waits		
MRI/CT	2 days	5 days
Surgery	20 days	25 days



Outline Career and Progression Framework – Final V4. Updated March 2023



* Entry routes into sonography should be CASE equivalent i.e. the education and learning outcomes should meet CASE learning outcomes for the relevant academic level of study. See explanatory narrative for further details for non-CASE accredited awards.

Summary



Sonographers can have a positive impact on faster diagnostic pathways



Improves outcomes for patients and helps meet targets



Increase job satisfaction



Everyone is a winner!

Summary Continued

- Significant pressures are being faced with all FDS pathways and solutions are not always working harder, but we can work smarter.
- Don't underestimate the skill set of sonographers. Role extension and advanced practice goes beyond purely clinical acumen.
- Dynamic multi-disciplinary team approach, driven by sonographers, has led to tangible improvements in both patient care, and performance.

The background consists of several overlapping, curved shapes in two shades of blue. A lighter blue shape curves from the top left towards the bottom right, while a darker blue shape fills the rest of the frame, creating a dynamic, layered effect.

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