

WHAT IS OVERDIAGNOSIS AND HOW CAN REAL-WORLD DATA HELP US UNDERSTAND THE HARMS OF THE PROSTATE CANCER DIAGNOSTIC PATHWAY?

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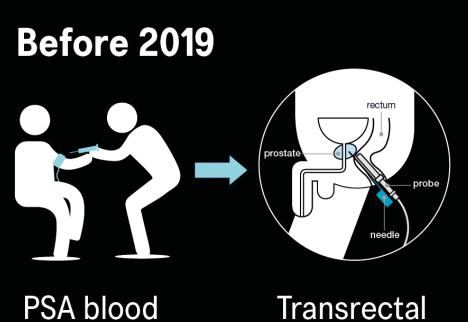
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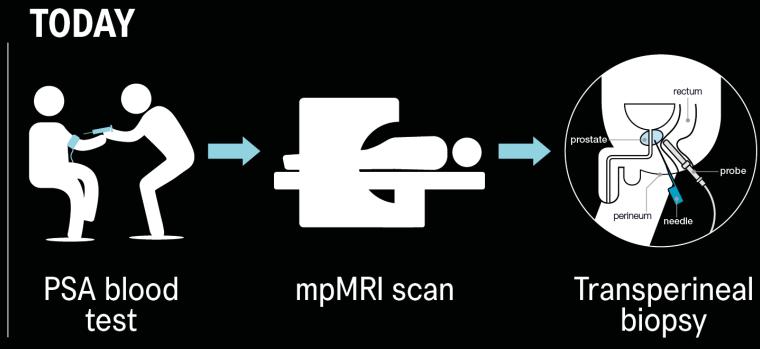


biopsy





test







- Used in the 3 largest prostate cancer screening trials: CAP, ERSPC,
 PLCO.
- Following this pathway, ERSPC demonstrated a 20% reduction in prostate cancer specific mortality for men who were screened vs men who were not screened at 16 years follow up.
- ERSPC and CAP proved that prostate cancer screening saves lives.
- ERSPC and CAP also showed that screening is associated with harms: unnecessary biopsy, sepsis, overdiagnosis, and overtreatment.





- Some cancers grow so slowly (or not at all), that they don't cause any symptoms or harm if left untreated.
- These are known as clinically insignificant or indolent cancers.
- Overdiagnosis happens when these harmless cancers are found through screening or tests.
- It can lead to psychological and physical harms, including unnecessary worry and overtreatment.





- Overdiagnosis ≠ overtreatment.
- Overtreatment is a consequence of overdiagnosis.
- It happens when a man receives unnecessary treatment for a prostate cancer that would not have caused any harm if left untreated.
- Some of the physical harms associated with overtreatment including urinary incontinence, erectile dysfunction, fatigue, bowel issues, and higher risk of infection.





- It is difficult to measure overdiagnosis and overtreatment we can only estimate which cancers are more or less likely to progress or cause harm.
- In the UK, men diagnosed with low-risk prostate cancer are offered active surveillance.
- Active surveillance involves regularly monitoring prostate cancer to check for any signs of growth or progression.
- It helps avoid or delay treatment until it's really needed.
- It helps reduce overtreatment.





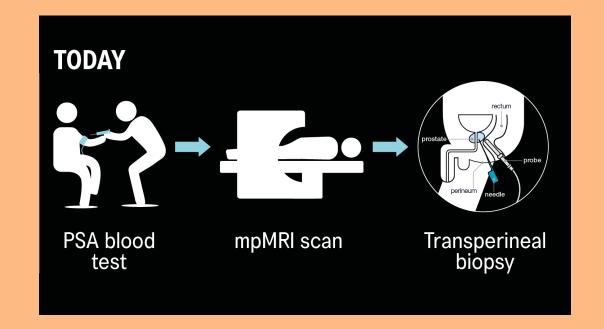
- The UK is a world leader in active surveillance.
- 8% of men diagnosed with low-risk localised cancer in England underwent radical treatment 12 months after their diagnosis.
- We assume the remaining 92% choose active surveillance as their first treatment choice.

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- Pre-biopsy mpMRI reduces the number of men who have an unnecessary biopsy, and clinically insignificant prostate cancer diagnoses.
- 40% (and rising) of prostate biopsies are now transperineal – this has significantly reduced the risk of sepsis.





REAL-WORLD DATA RESEARCH

- We analysed data from 16 hospitals in London and the South West of England to measure how many men experienced harm after a PSA test under the current prostate cancer pathway.
- We compared this to older data from the CaP and ProtecT trials, to see whether the harms from PSA testing have reduced over time.



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Possible harm outcome after a PSA test	pre-MRI (CaP/ ProtecT)	Pre-MRI (UK clinical practice)	Current UK pathway model	Current pathway RAPID (London) RWD	Current pathway SW England RWD
% of men having a PSA test who have a biopsy showing no cancer	9.46%	9.34%	6.51%	2.33%	1.52%
% of men having a PSA test who have a biopsy showing insignificant cancer	3.83%	3.30%	1.11%	0.73%	0.83%
% of men having a PSA test who suffer sepsis	0.10%	0.21%	0.07%	0.02%	No data

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WHAT DOES THIS MEAN?



Advances in technology have reduced the risk of harm when being tested for prostate cancer by a massive...

79%

*compared to the diagnostic pathway before MRI.

Today,

90%

fewer men develop sepsis after a PSA blood test

*Reduced from 0.21% in the diagnostic pathway before MRI to 0.02% in current UK clinical practice.

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KEY MESSAGES



- Our study provides the first quantitative estimates of reduction in harm after a PSA test.
- We have confirmed that the current UK prostate cancer diagnostic pathway is safer and more accurate.
- Our research makes it clear that NHS guidelines are out of date.
- With this new evidence, we believe men at highest risk must be proactively informed of their risk and given the choice of a PSA test.





THANK YOU!

