

BREAST CANCER SUPPORTIVE CARE

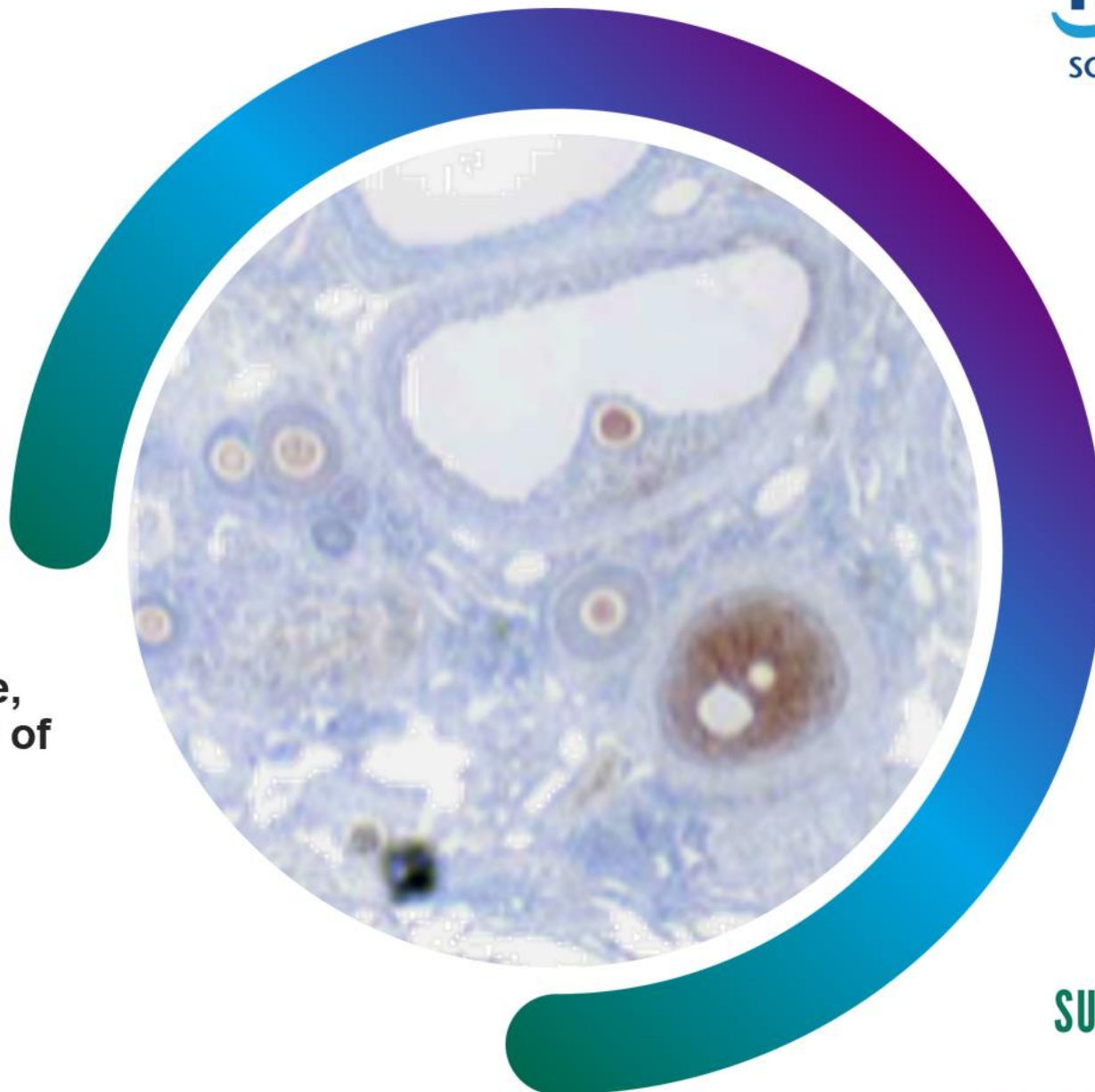
EDUCATIONAL SYMPOSIUM



FERTILITY & ITS PRESERVATION IN BREAST CANCER

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Edinburgh.



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**BREAST
CANCER
SUPPORTIVE
CARE**

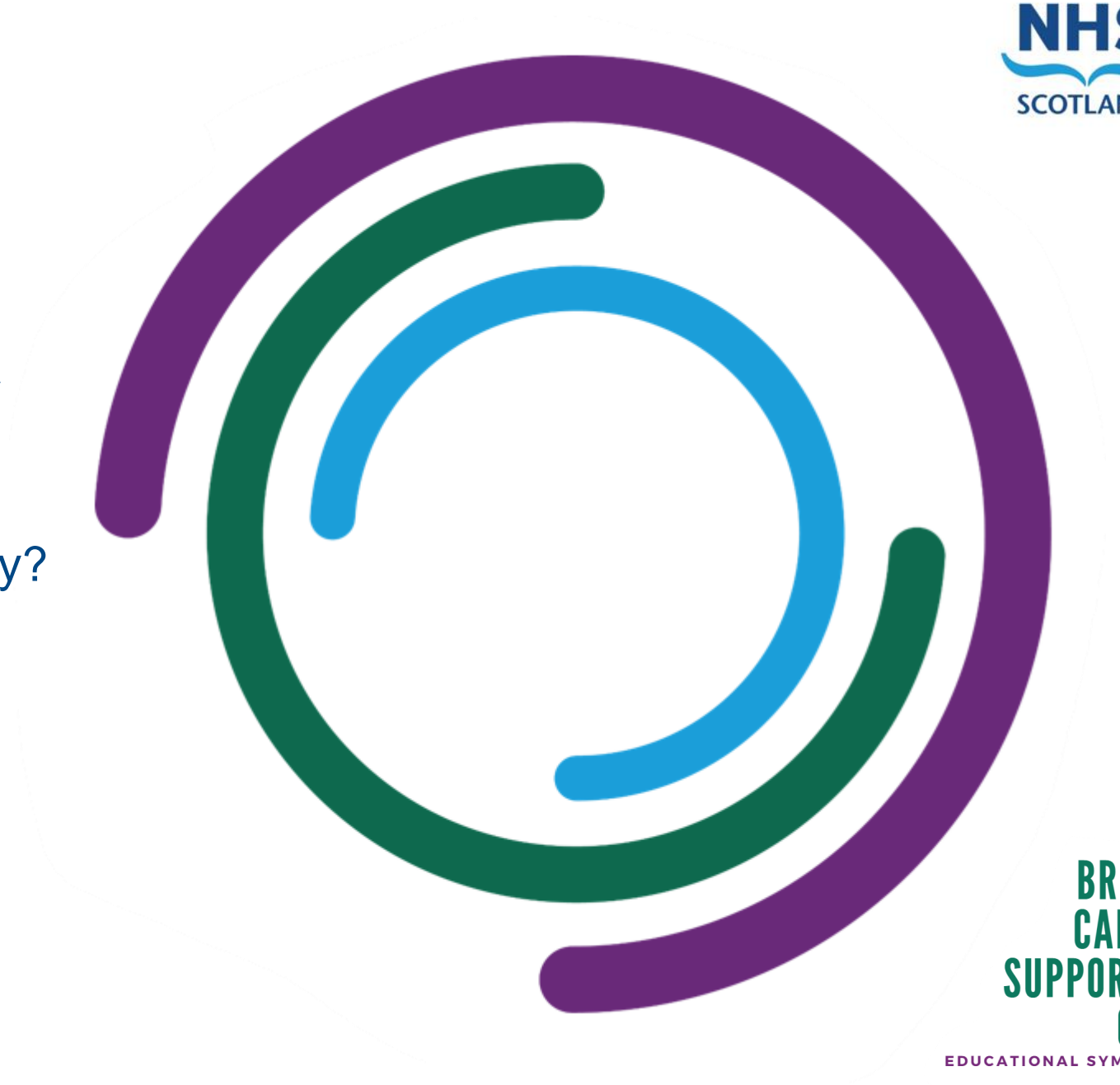
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Questions

Will I be able to have a baby after treatment?

Will it be safe to have a pregnancy?

What can I do to increase my chances of being able to have a baby afterwards?



Chemotherapy: immediate and late effects on the ovary

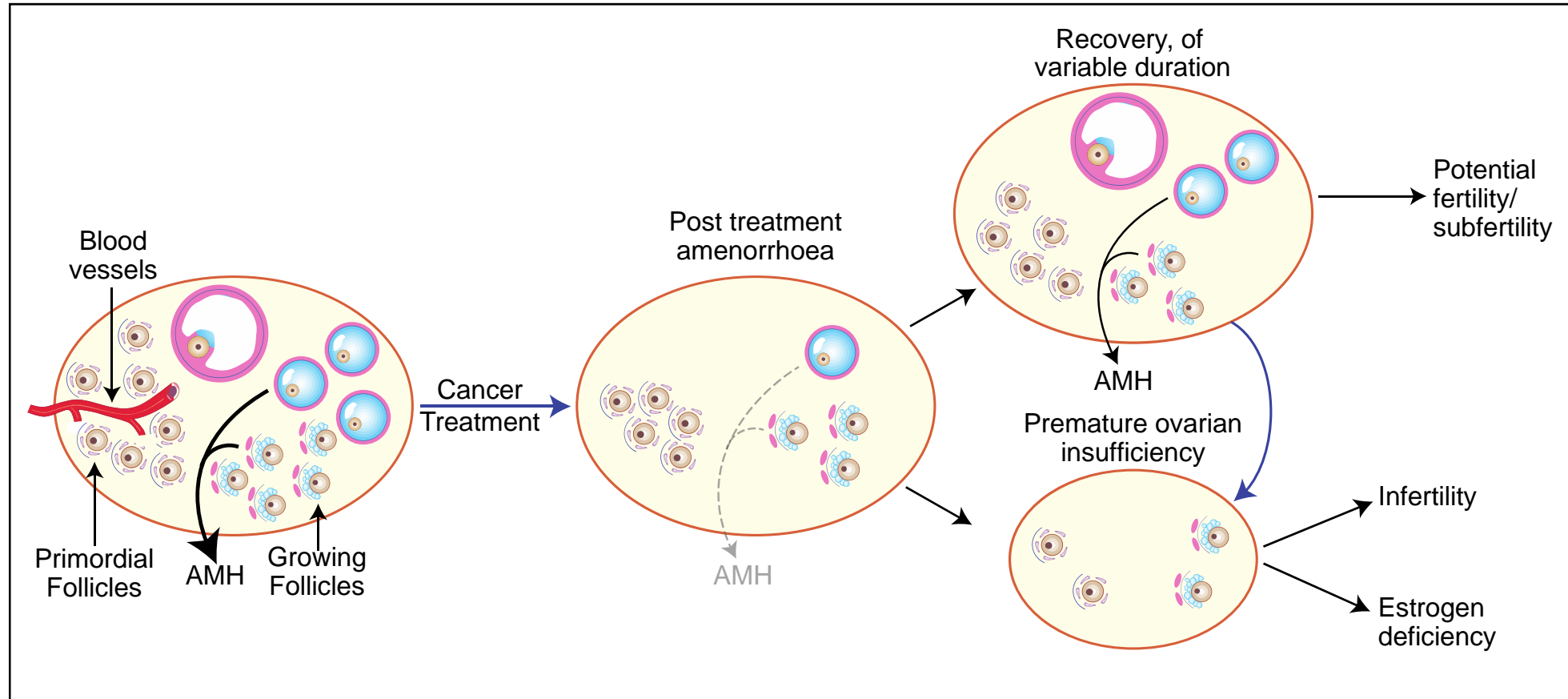
Depletion of growing follicles

Himelstein-Braw R, Peters H and Faber M (1978)
Morphological study of the ovaries of leukaemic children.
Br J Cancer 38, 82-87

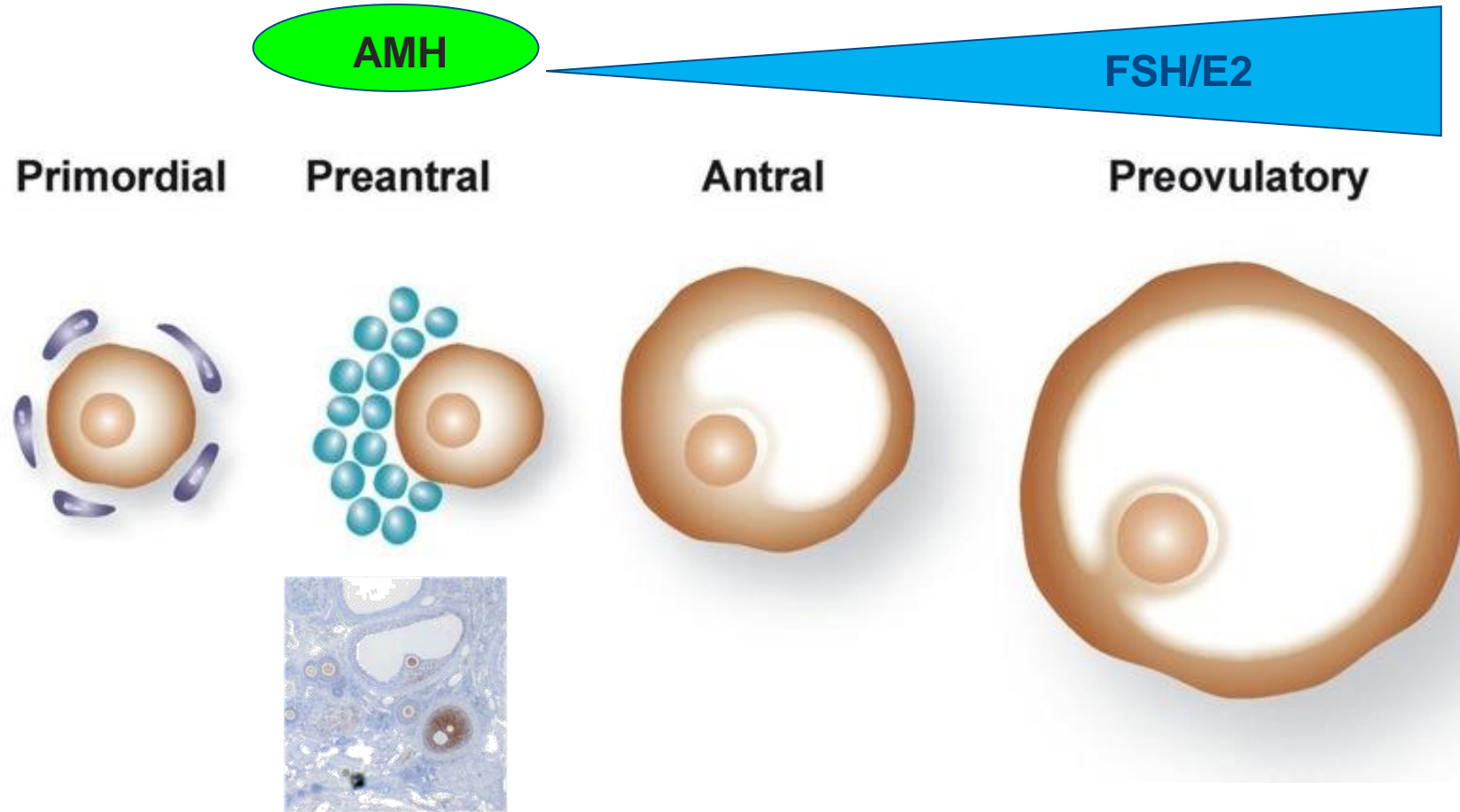
Premature ovarian failure

Chapman RM, Sutcliffe SB and Malpas JS (1979)
Cytotoxic-induced ovarian failure in women with Hodgkin's disease. I. Hormone function.
JAMA 242, 1877-1881

Effects of cancer therapy on the ovary



AMH as key biomarker of the ovarian reserve



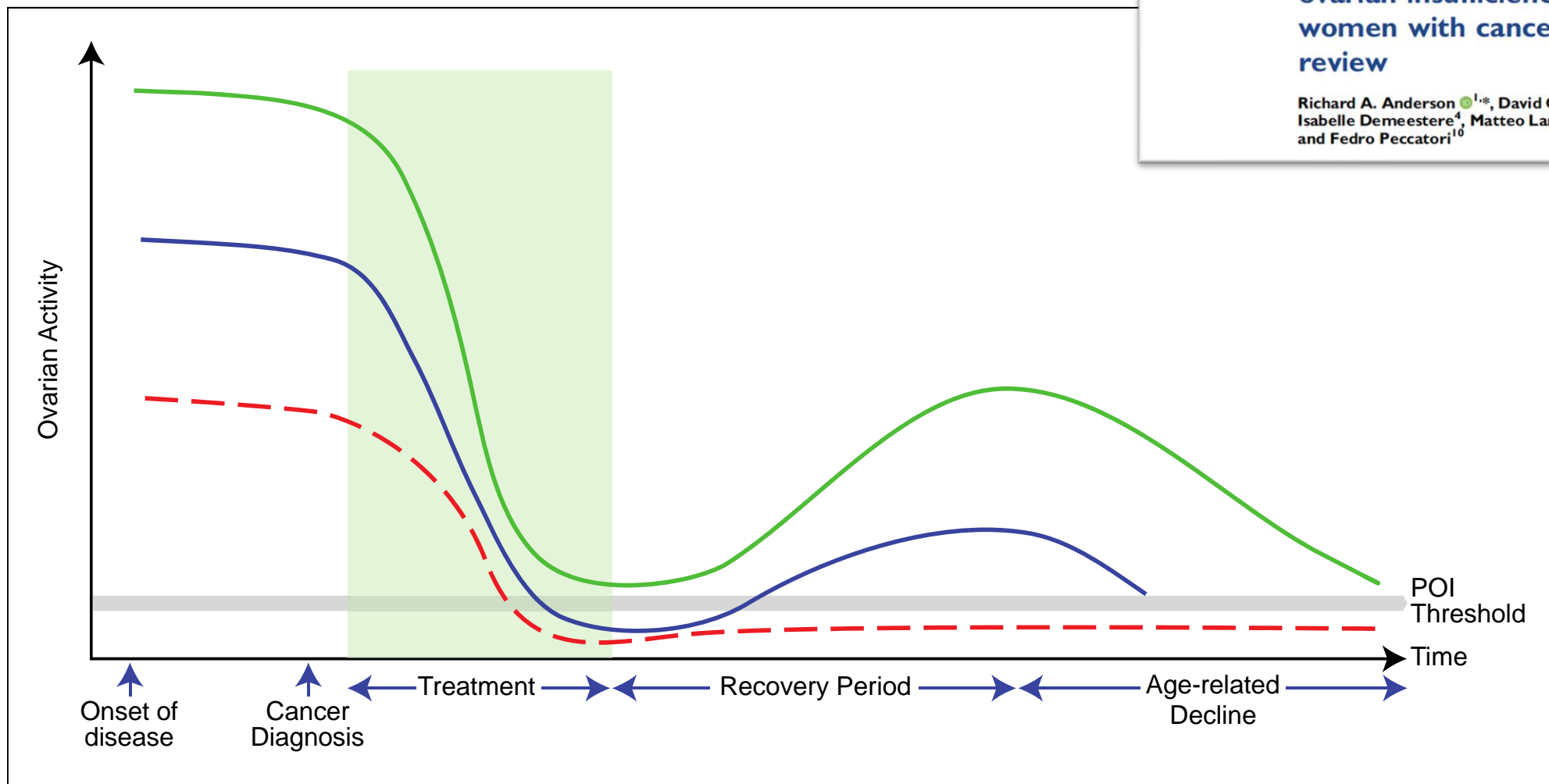
Identify and predict ovarian damage

Human Reproduction Update, pp. 1–18, 2022
<https://doi.org/10.1093/humupd/dmac004>

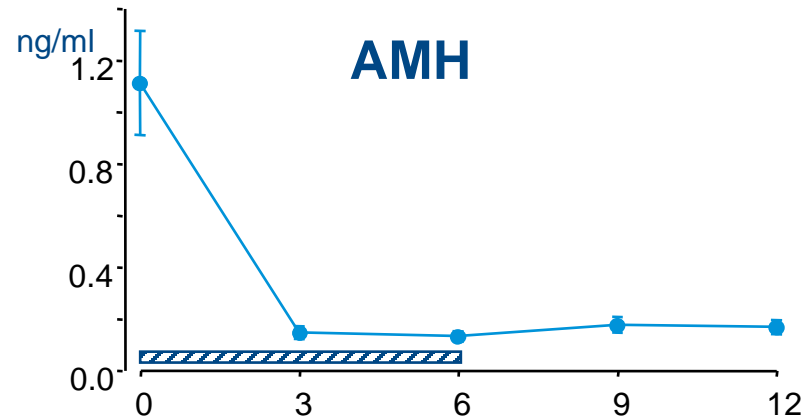
human
reproduction
update

Anti-Müllerian hormone as a marker of ovarian reserve and premature ovarian insufficiency in children and women with cancer: a systematic review

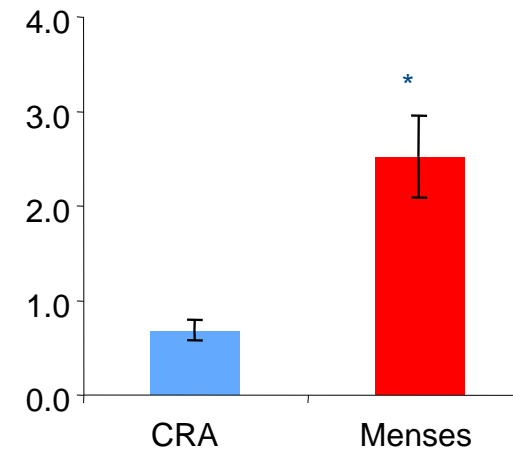
Richard A. Anderson^{1,*}, David Cameron², Florian Clatot³,
Isabelle Demeestere⁴, Matteo Lambertini^{5,6}, Scott M. Nelson^{7,8,9},
and Fedro Peccatori¹⁰



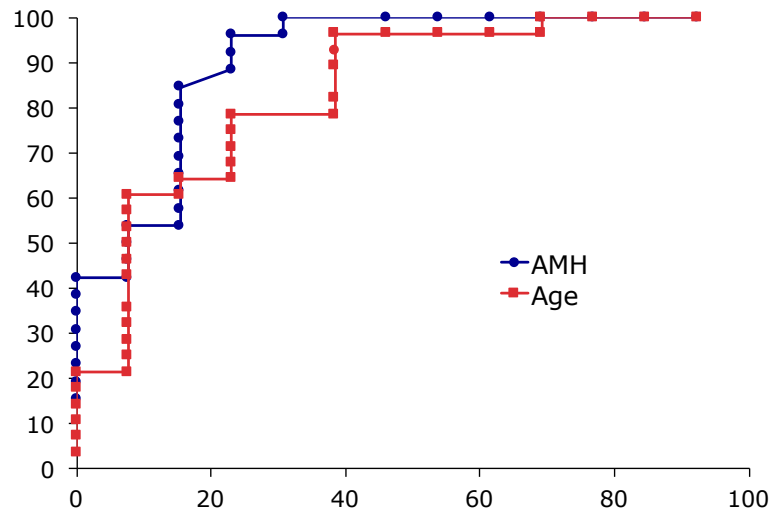
Pretreatment AMH predicts long-term ovarian function in eBC



Pretreatment AMH



Prediction
of amen



AMH at diagnosis of early breast cancer is higher in women still having menses 5 years later

The impact of cancer on subsequent chance of pregnancy: a population-based analysis

**Richard A. Anderson^{1,*}, David H. Brewster², Rachael Wood³,
Sian Nowell^{4,5}, Colin Fischbacher³, Tom W. Kelsey⁶,
and W. Hamish B. Wallace⁷**

¹MRC Centre for Reproductive Health, Queen's Medical Research Institute, University of Edinburgh, 47 Little France Crescent, Edinburgh EH16 4 TJ, UK ²Scottish Cancer Registry, Information Services Division, NHS National Services Scotland, 1 South Gyle Crescent, Edinburgh EH12 9EB, UK ³Information Services Division, NHS National Services Scotland, 1 South Gyle Crescent, Edinburgh EH12 9EB, UK ⁴eData Research & Innovation Service (eDRIS), Information Services Division, NHS National Services Scotland, 1 South Gyle Crescent, Edinburgh EH12 9EB, UK ⁵Farr Institute Scotland, School of Computer Science, University of St. Andrews, St. Andrews, Fife KY16 9SS, UK ⁶Department of Paediatrics, Royal Children's Hospital, Park Road, Edinburgh EH10 6DT, UK ⁷Department of Paediatrics, Royal Children's Hospital, Park Road, Edinburgh EH10 6DT, UK

ORIGINAL ARTICLES: EPIDEMIOLOGY

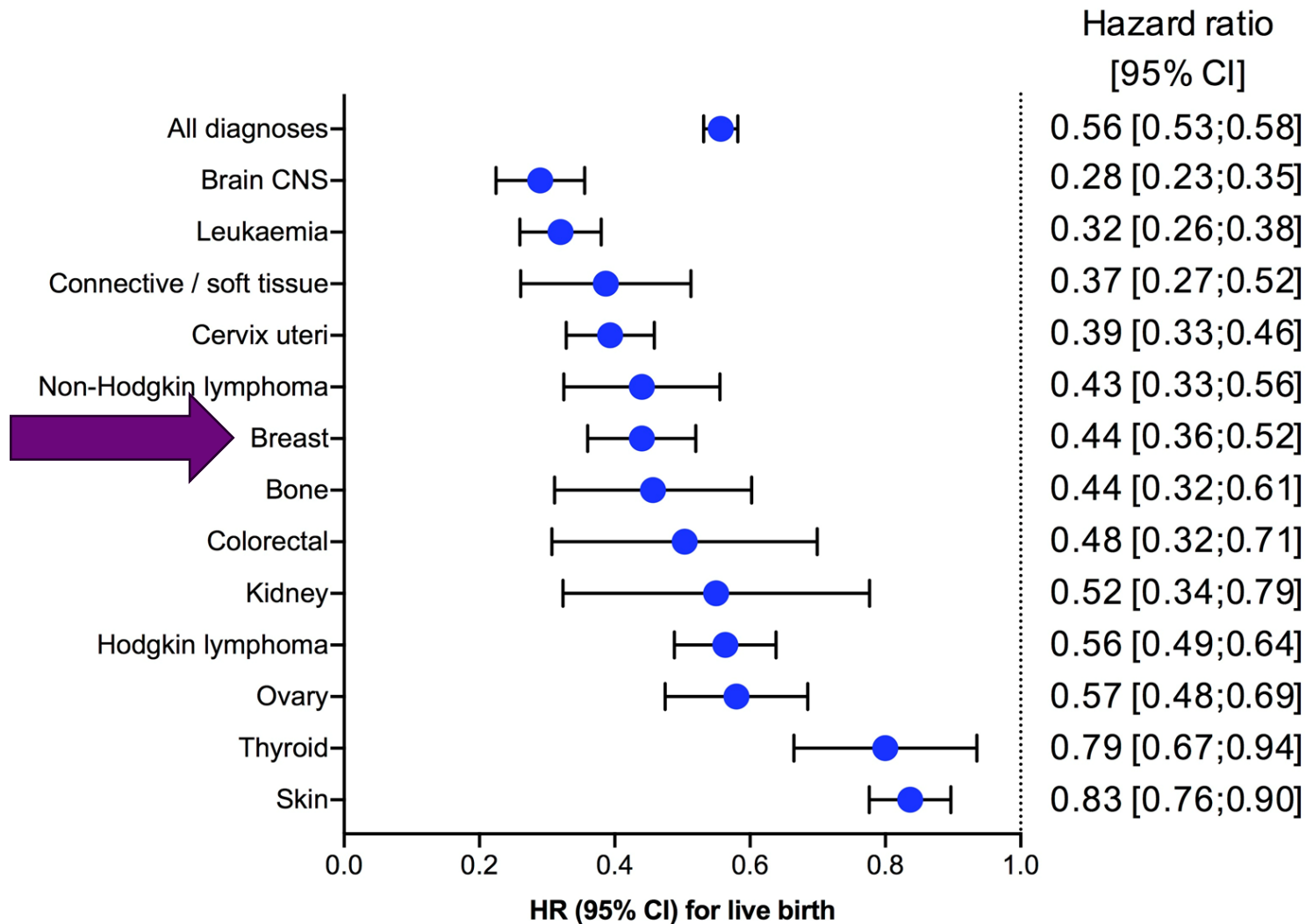
 Check for updates

Family size and duration of fertility in female cancer survivors: a population-based analysis

Richard A. Anderson, M.D., Ph.D.,^a Tom W. Kelsey, Ph.D.,^b David S. Morrison, M.D.,^c
and W. Hamish B. Wallace, M.D.^d

Human Reprod, 2018

Cancer (treatment) reduces the chance of live birth



Scottish population data:

New cancer diagnosis

1981-2012

N=10,267 with no prev pregnancy

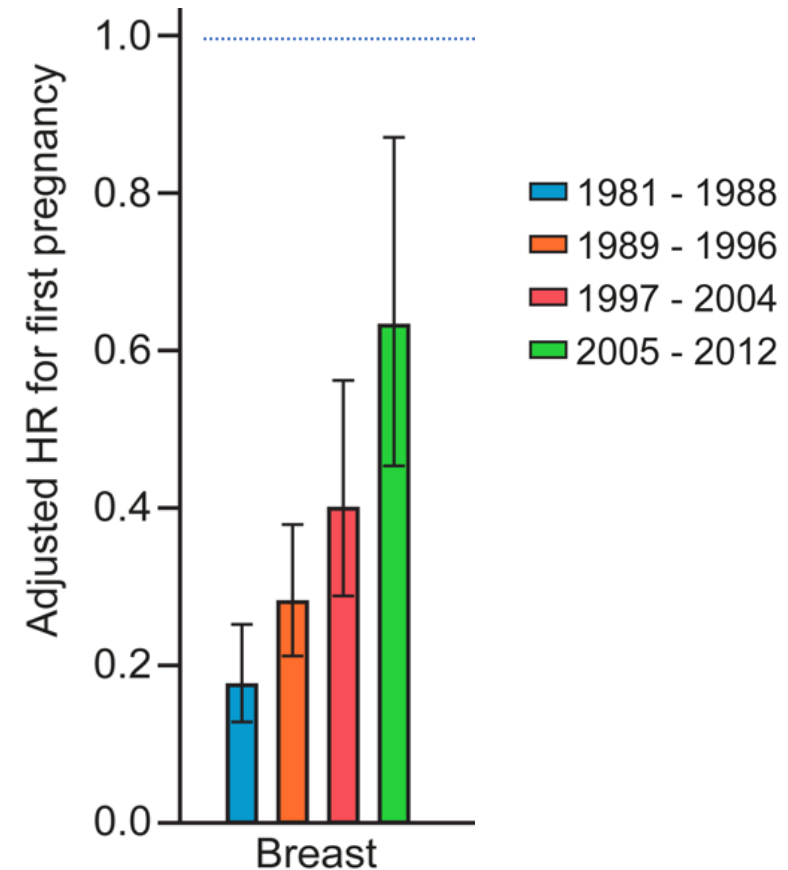
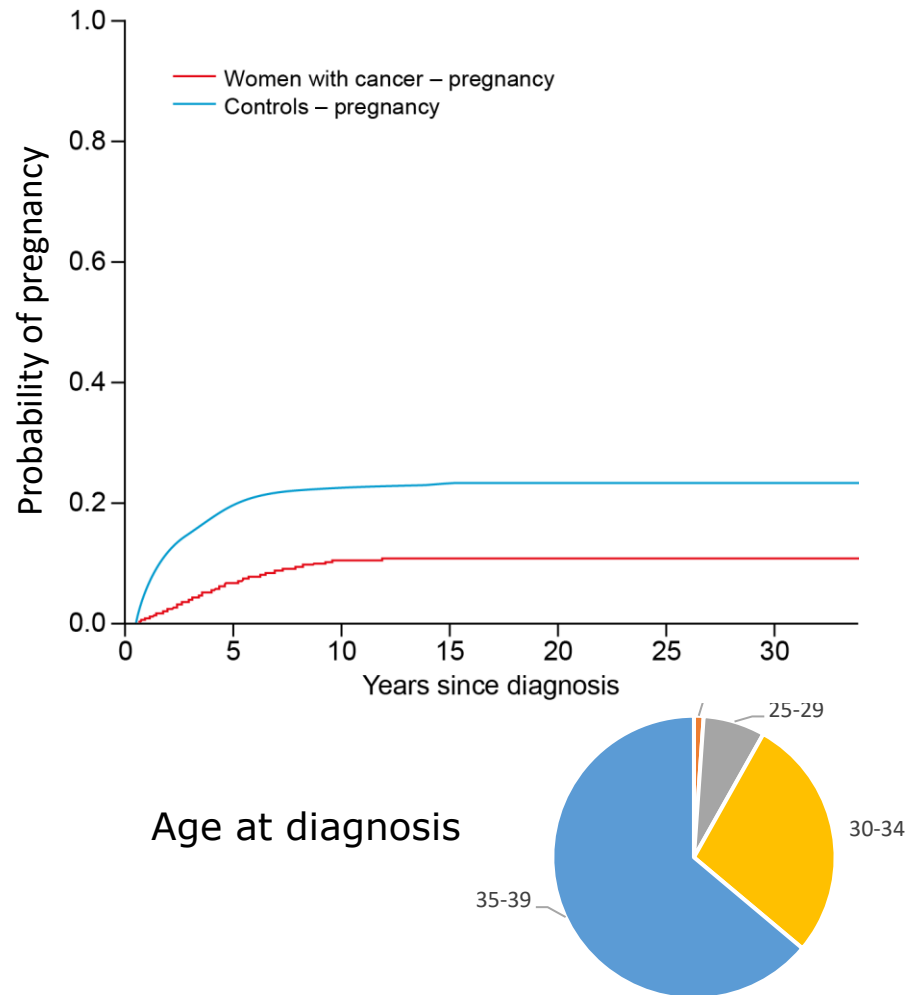
With 3x controls

Matching:

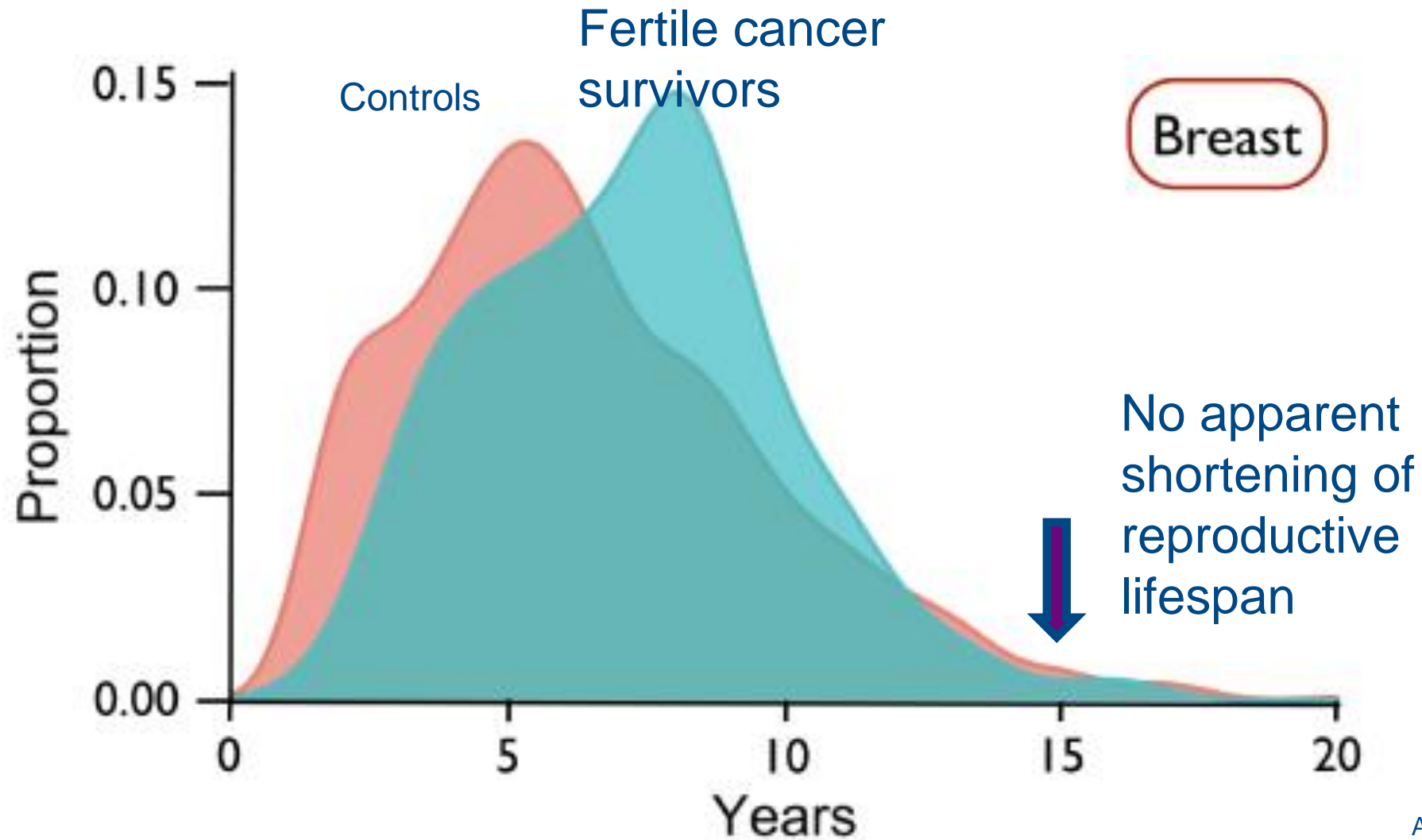
Age

Period of diagnosis

Impact of breast cancer on fertility: Scottish national data



‘Fertile lifespan’ after breast cancer



Original Research

Survival after breast cancer in women with a subsequent live birth: Influence of age at diagnosis and interval to subsequent pregnancy

Richard A. Anderson ^{a,*}, Matteo Lambertini ^{b,c}, Peter S. Hall ^d,
W. Hamish Wallace ^e, David S. Morrison ^f, Tom W. Kelsey ^g

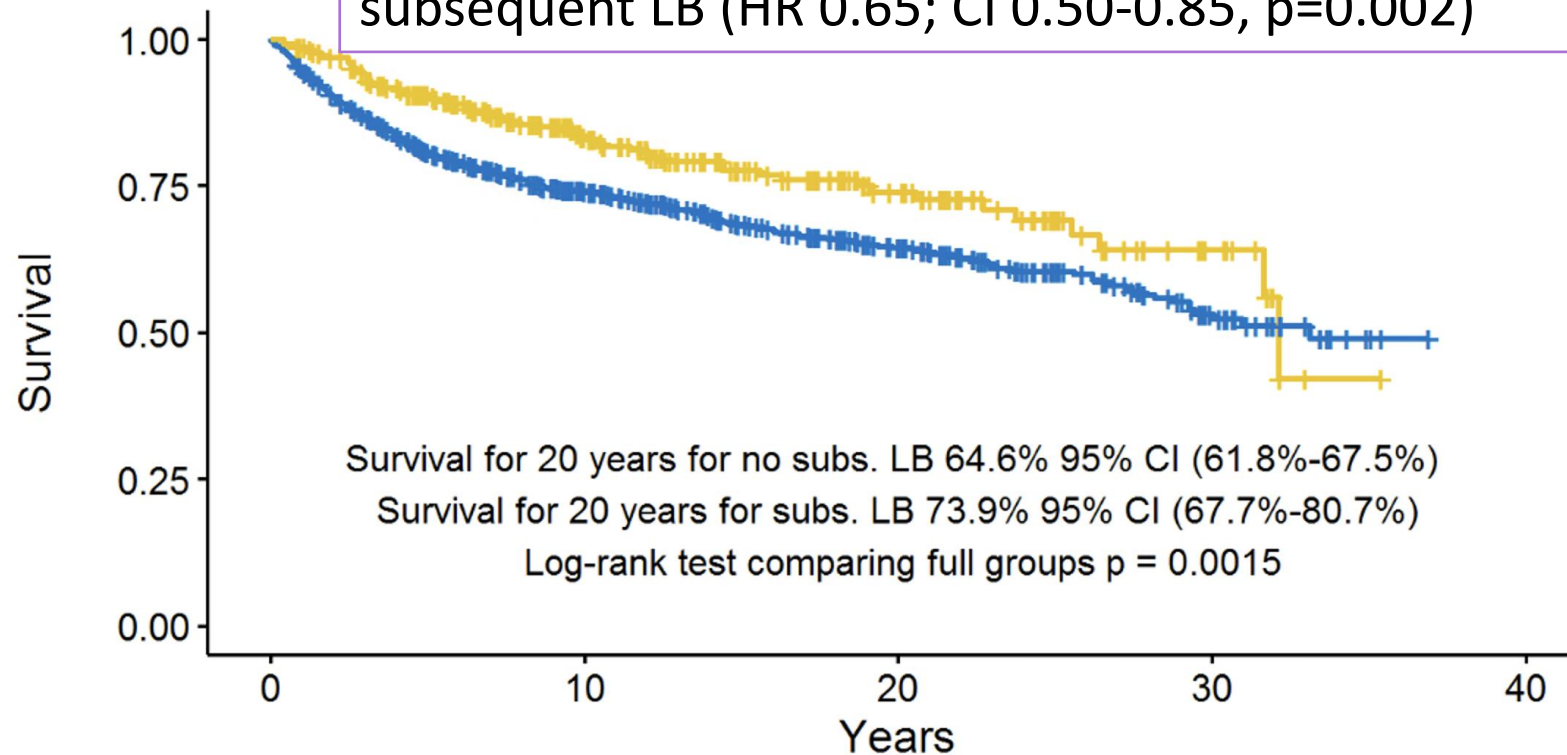


5181 with breast cancer,
290 women with SLB

No differences in stage
distribution, EP status,
age

After breast cancer, does pregnancy impact survival?

Overall survival was increased in women who had a subsequent LB (HR 0.65; CI 0.50-0.85, $p=0.002$)



Number at risk

no SLB	1682	789	320	59	0
SLB	290	151	65	13	0

Subgroup analyses

Pregnancy prior to diagnosis

Age at diagnosis

<> 5 years interval between diagnosis and LB

ER status (positive or negative)

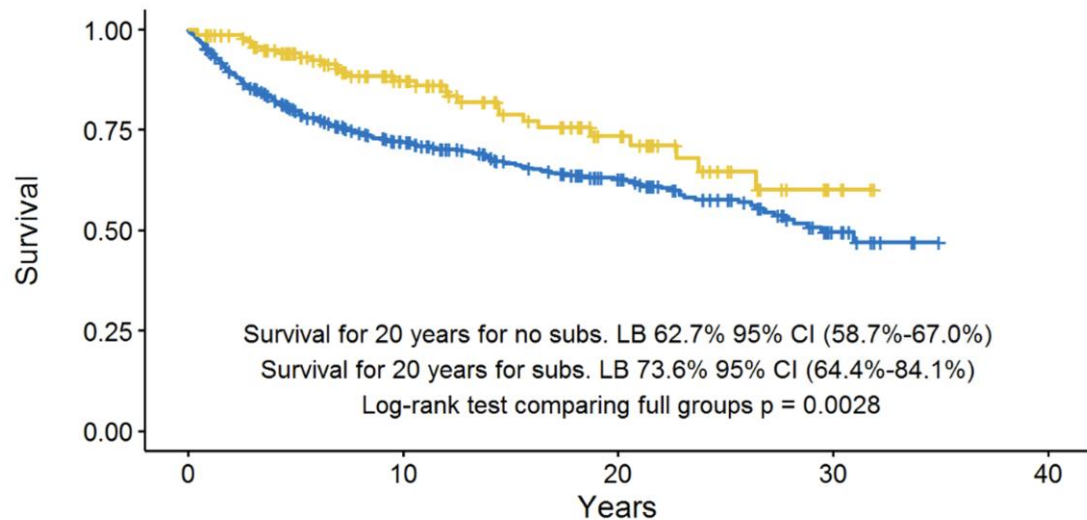
Tumour stage

Known treatment with chemotherapy

Period of diagnosis

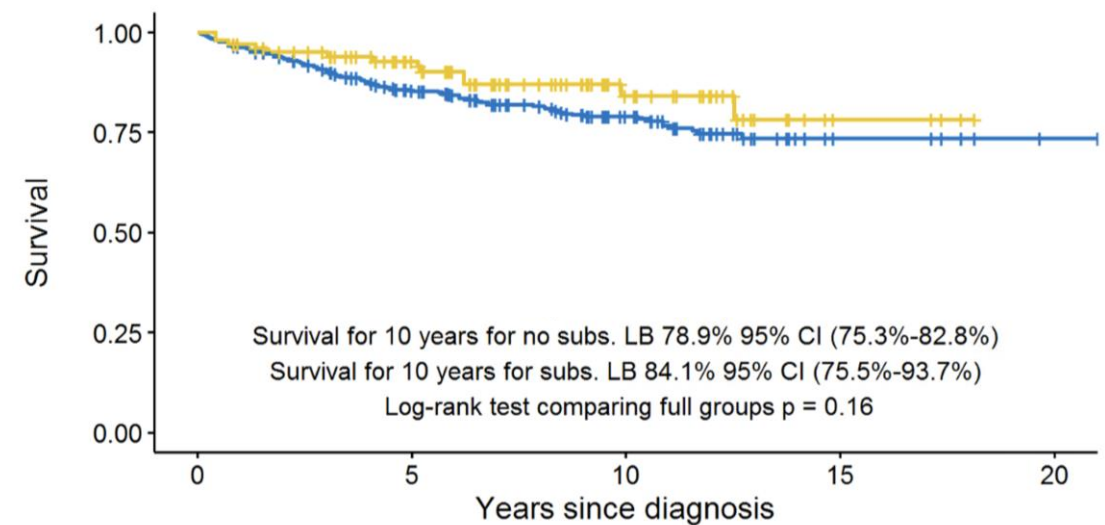
Women with no previous pregnancy, and ER+ve

No prev preg: HR 0.56 (0.38-0.82, p=0.003)



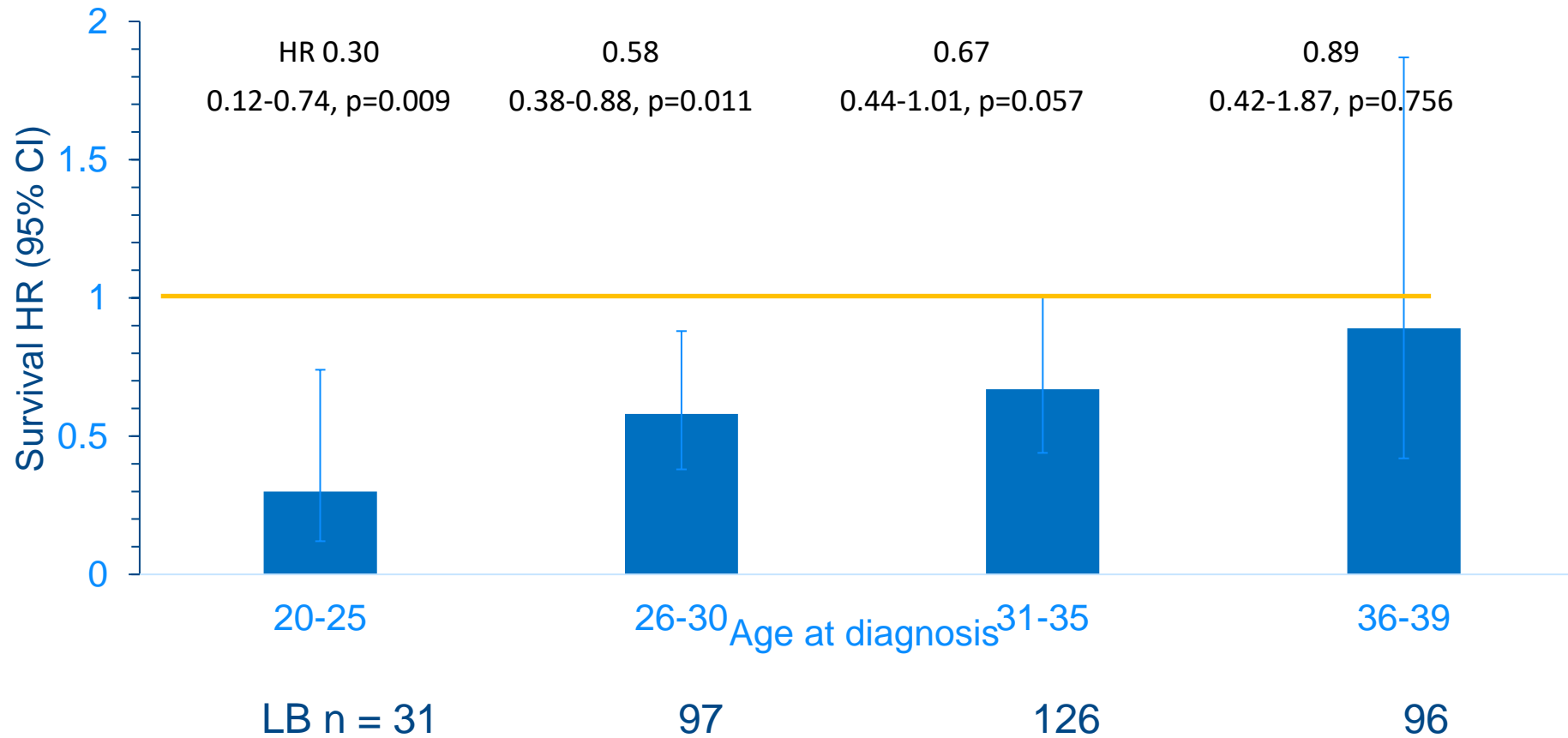
Number at risk					
no SLB	794	366	159	27	0
SLB	142	75	33	5	0

ER Positive HR 0.66 (0.37-1.18, p=0.160)



Number at risk					
no SLB	612	389	168	22	4
SLB	102	71	27	4	0

Age at diagnosis



POSITIVE trial: interruption of endocrine treatment for pregnancy

≤42 years of age

stage I, II, or III disease

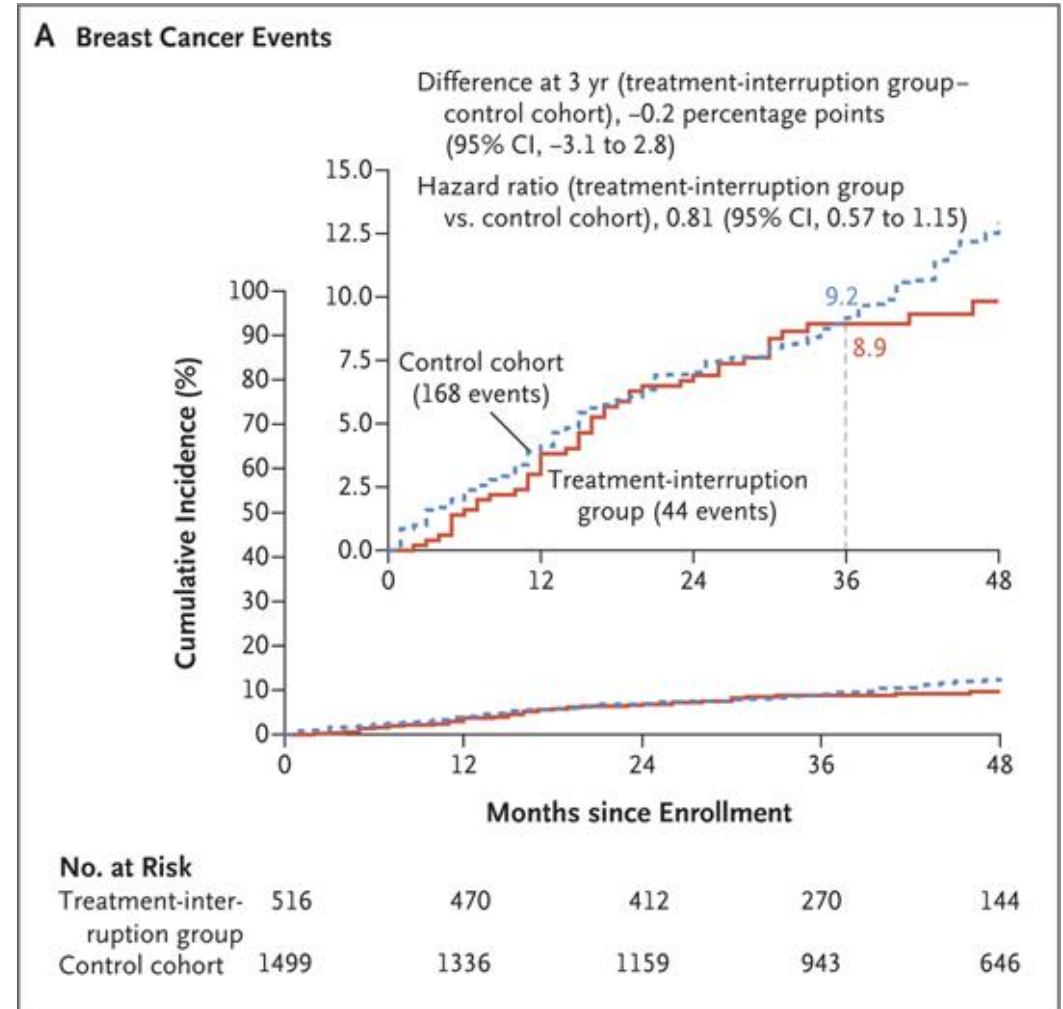
had received adjuvant endocrine therapy for 18 to 30 months

desired pregnancy

Controls: similar but no wish for pregnancy

N=497, 74% pregnant, 64% had LB

Analysis at 1638 patient-years



Acknowledgements

David Cameron and colleagues at Edinburgh Breast Centre
Peter Hall

Hamish Wallace (paed oncology)
Tom Kelsey (mathematician, St Andrews)



REGISTER HERE

UK Fertility Preservation Conference

May 22-23 2024 - Edinburgh, UK

NOW ACCEPTING

UKFP2024 abstract submissions

Submit by April 11th 2024

Enquiries to UKFP-2024@ed.ac.uk



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FEMALE FERTILITY PRESERVATION IN BREAST CANCER

Maya Chetty

Consultant in Reproductive Medicine,
Edinburgh Fertility Centre, NHS Lothian.



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Fertility Preservation Guidelines



Breast Cancer Care Survey 2018

254 responses

Women < 45 years old

21% did not have fertility discussion with their cancer team



“When I asked about having children I was made to feel like I was wasting their time and basically ignored as they felt not important.”

“I’m in a same sex relationship and it was unfortunately overlooked by my team. I had to ask about fertility treatment as when initially diagnosed I was advised I would be having chemo. This was later reversed due to the Oncotype test but they just assumed fertility was not an issue for me being in a same sex relationship which I found very upsetting.”

“I was offered fertility treatment after my lumpectomy and before I started chemo. I was provided a lot of information by [my hospital], where I received my fertility treatment. [My hospital] gave me brilliant service and care, they were amazing. It was quick and painless for me. I had 20 eggs taken and 18 successfully stored”

“I had a fantastic experience - my surgeon discussed fertility treatment and referred me without me even mentioning that it was an area of concern. It's appalling that this isn't the default level of treatment for all women!”

Breast Cancer Care wants to see every breast clinic have a process for referring women promptly to a fertility specialist. This referral shouldn't depend on local in vitro fertilisation (IVF) funding arrangements.

Fertility Preservation Provision in Scotland



- Generous and equitable NHS funding
- Service provided by 4 NHS IVF centres: Glasgow, Edinburgh, Dundee and Aberdeen
- Funding by Health Board
- Referral from secondary care
- Commitment to seeing patients +/- starting treatment ASAP and maintaining close communication with cancer team

Access Criteria

Diagnosis	Subfertility with an appropriate cause of any duration - all couples OR Unexplained subfertility of two years – heterosexual couples Unexplained subfertility following six cycles of donor insemination – same sex couples
Residency	Live within Lothian or the Borders.
Stable relationship	Couple have been co-habiting (at the same address) in a stable relationship for a minimum of two years.
Sterilisation	Neither partner has had voluntary sterilisation (even if reversed).
Biological child	At least one partner must have no living biological child. Same sex couples who have previously conceived through assisted conception and are both named on the resulting child's birth certificate are not eligible for future treatment.
Body Mass Index (BMI)	The prospective birth mother must have a BMI above 18.5 and below 30.
Smoking	Both partners must not smoke, vape or use nicotine for at least three months before referral for treatment and couples must continue to be non-smoking/vaping and nicotine free during treatment.
Alcohol and Drugs	Both partners must not use illegal and abusive substances. Both partners must be Methadone free for at least one year prior to referral for treatment and continue to be Methadone free for the duration of treatment. Neither partner should drink alcohol prior to or during the period of treatment.
Age	If a couple is referred for IVF/ICSI treatment where the intended birth mother is aged 39, the couple is unlikely to receive more than one NHS funded cycle. Couples where the intended birth mother is over the age of 42 years old will not be eligible for any NHS funded treatment.
Previous treatment	Any frozen embryos stored at another clinic in the UK must be transported to EFC before NHS treatment can commence. If the previous treatment was not NHS funded, the transport will be at the patient's cost.
Welfare of the Child	Satisfy the requirements Welfare of the Child Assessment.

Condition or treatment causing imminent threat to fertility

Not applicable at time of storage

Individual has no children

BMI <35

Not applicable at time of storage

Age < 40

Not applicable at time of storage

Egg or embryo freezing

One cycle of egg/ embryo storage (counts as one of max three NHS funded cycles)

Option of self-funding or private treatment (cost £4-6k and ongoing storage fee)

Fertility counselling always offered

Storage for up to maximum of 55 years

Can nominate named partner for posthumous use

Full access criteria apply at the point of use

Possibility of pre-implantation genetic testing (PGT)

Emergency IVF

Treatment can start ASAP

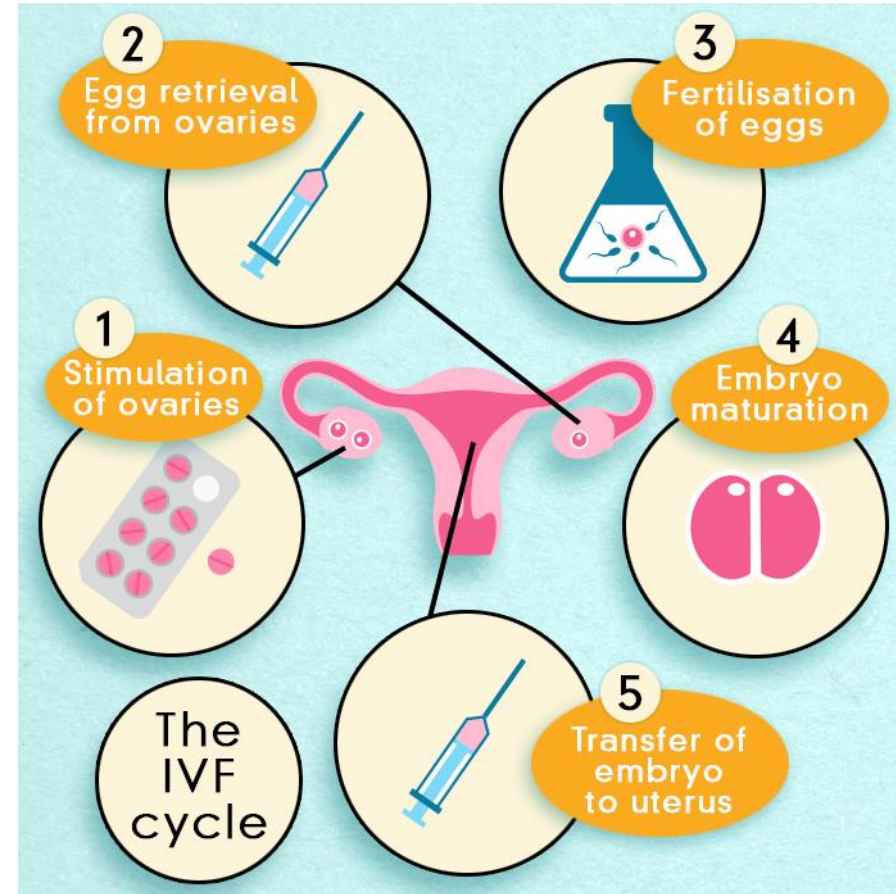
Takes 2-3 weeks start to finish

Risks: operative, anaesthetic, OHSS

Eggs v embryos

Thaw cycle

Chance of live birth approx 4-5% per egg frozen,
30-35% per embryo frozen



Controlled ovarian stimulation

1. Random start
2. Antagonist cycle
3. Higher dose FSH
4. Agonist trigger
5. Co-treatment with anti-oestrogen

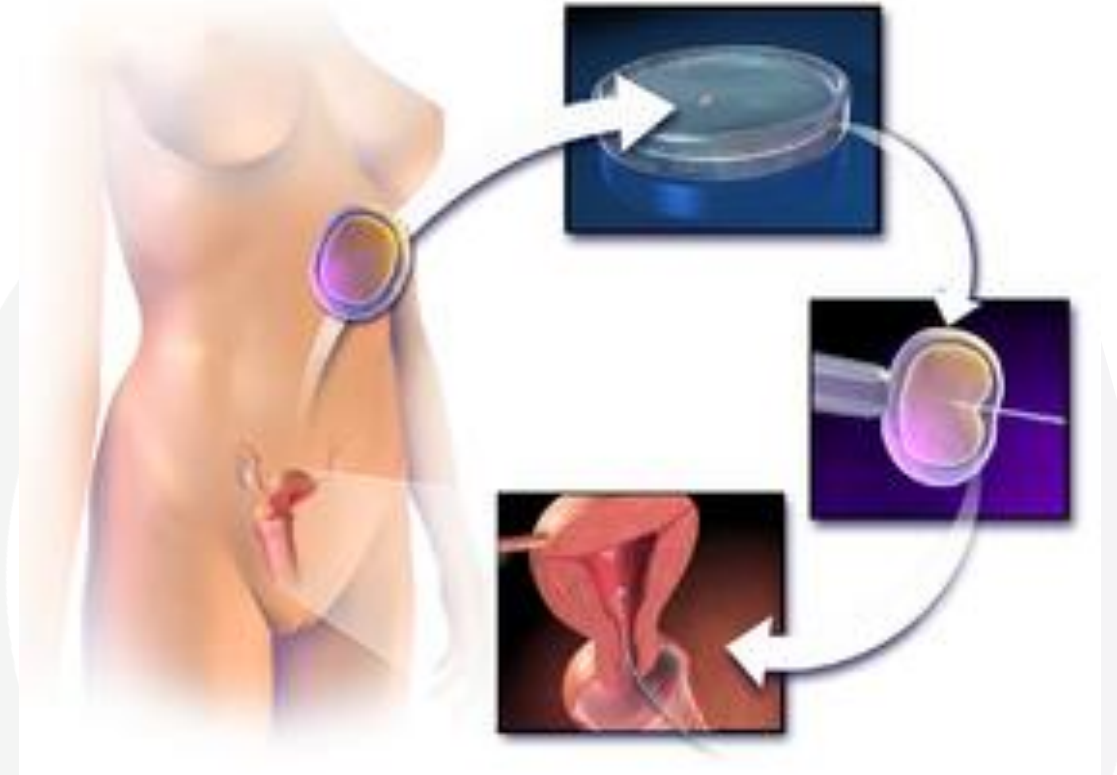


- Minimize duration
- Minimize duration and OHSS
- Optimise ovarian response
- Minimise OHSS
- Oestrogen sensitive tumours

Egg or embryo freezing



Image fertilitysolutions.com.au

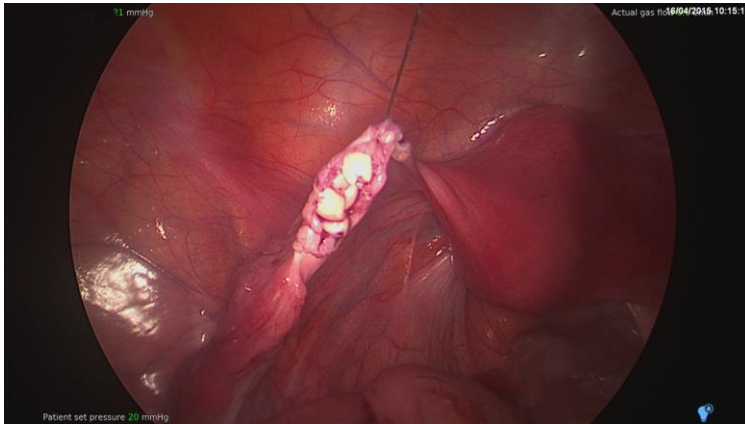


Embryo cryopreservation

established option
But....



Ovarian tissue cryopreservation



Alternative option when time not adequate for ovarian stimulation or other contraindications

Age < 35

Requires laparoscopy for tissue harvesting and re-implantation

First patient in Edinburgh in 1993

1mm thick, 1 ovary, strips v oophorectomy

Slow freezing (vitrification 2 babies in Japan)

Laparoscopic replacement

- Surgical placement and technique

- Risk of malignant contamination

- Chance of live birth ? approx 25-40%

More than 200 live births

Edinburgh data

- 76 referrals since 2015
- Increasing numbers of referrals and patients opting for fertility preservation
- Mainly white women
- Increasing numbers of random starts and agonist triggers
- Small numbers of OTC

Thanks to Cheryl Dunlop and Richard Anderson



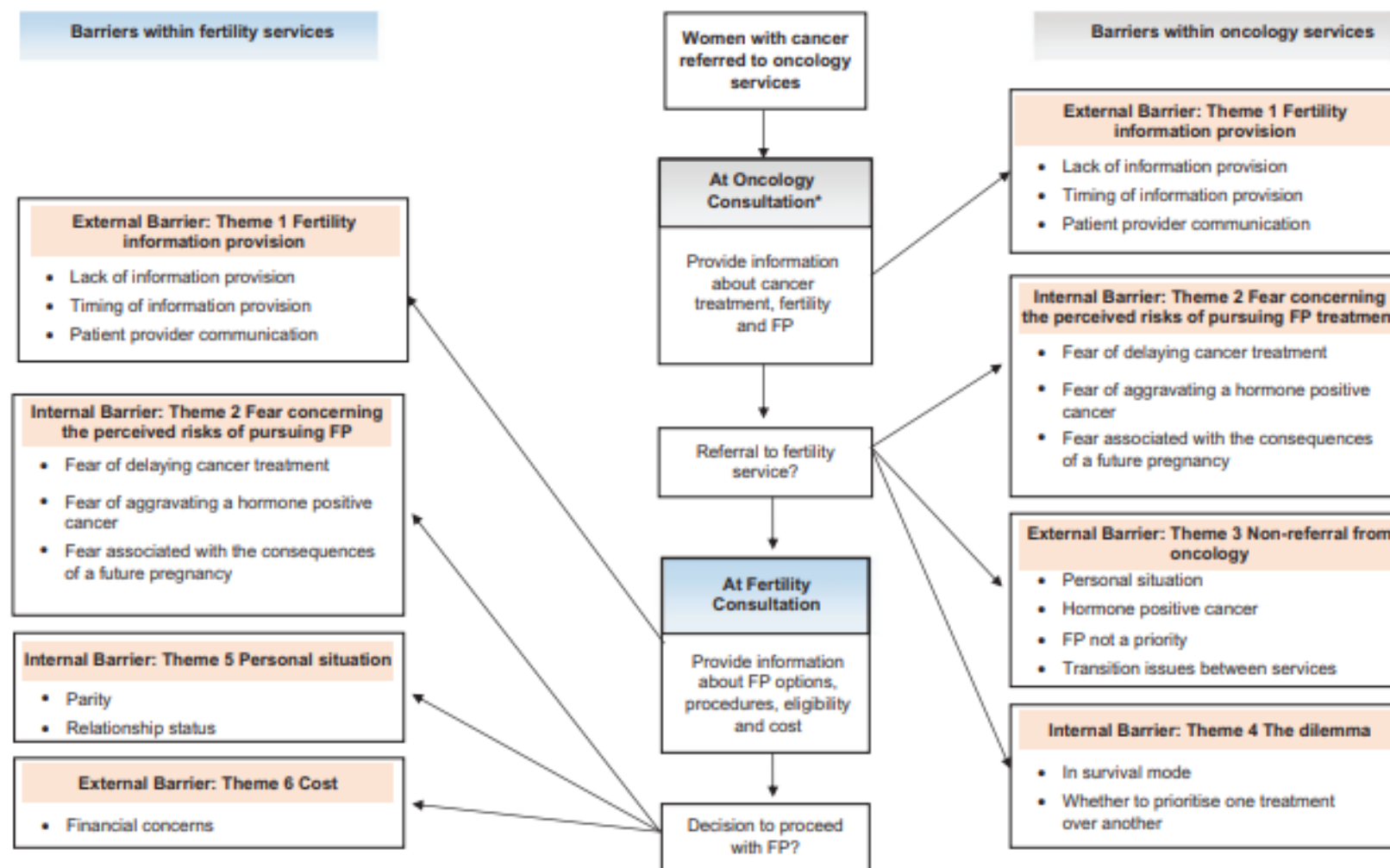


“I do remember thinking it was just totally unfair that guys could freeze their sperm and there really were no options for women.”

“If the situation were ever similar to that which it is for young males, which is to say... freeze your sperm... I think it would be outstanding. But secondarily, I think it also gives a large measure of hope and expectation to someone at the front-end of it.”

Nieman et al. Fertility Preservation and Adolescent Cancer Patients: Lessons from Adult Survivors of Childhood Cancer and Their Parents. Cancer Treat Res. 2007 ; 138: 201–217.

Overcoming barriers



* Consultation where fertility preservation is first discussed at either diagnosis or cancer treatment planning stage

Endorsed decision aid



[Cancer Fertility and Me | Helping you make the right decision for you https://cancerfertilityandme.org.uk/](https://cancerfertilityandme.org.uk/)

Questions

Scan or click the QR code to ask a question:



TOPIC:
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