Substantially lower lung cancer incidence with TARGIT-IORT

(targeted intraoperative radiotherapy) compared with whole breast radiotherapy in the TARGIT-A randomised trial for breast cancer: 25-year update

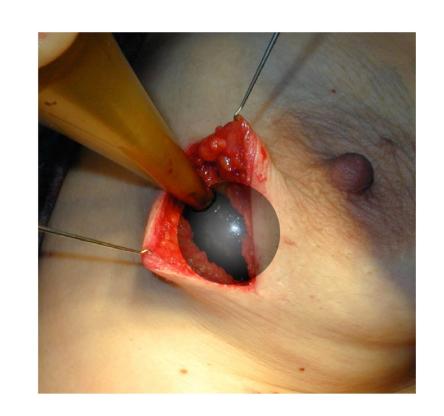
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and reduce travel and cost.

Background External beam whole breast radiotherapy (EBRT) given after breast conserving surgery inevitably leads to carcinogenic irradiation of nearby vital organs such as the lungs. TARGIT-A randomised trial (recruited from 2000-2012) found that TARGIT-IORT during the initial lumpectomy is as effective

as EBRT in controlling breast cancer.





Over 50,000 breast cancer patients in 260 centres from 38 countries have been treated with TARGIT-IORT



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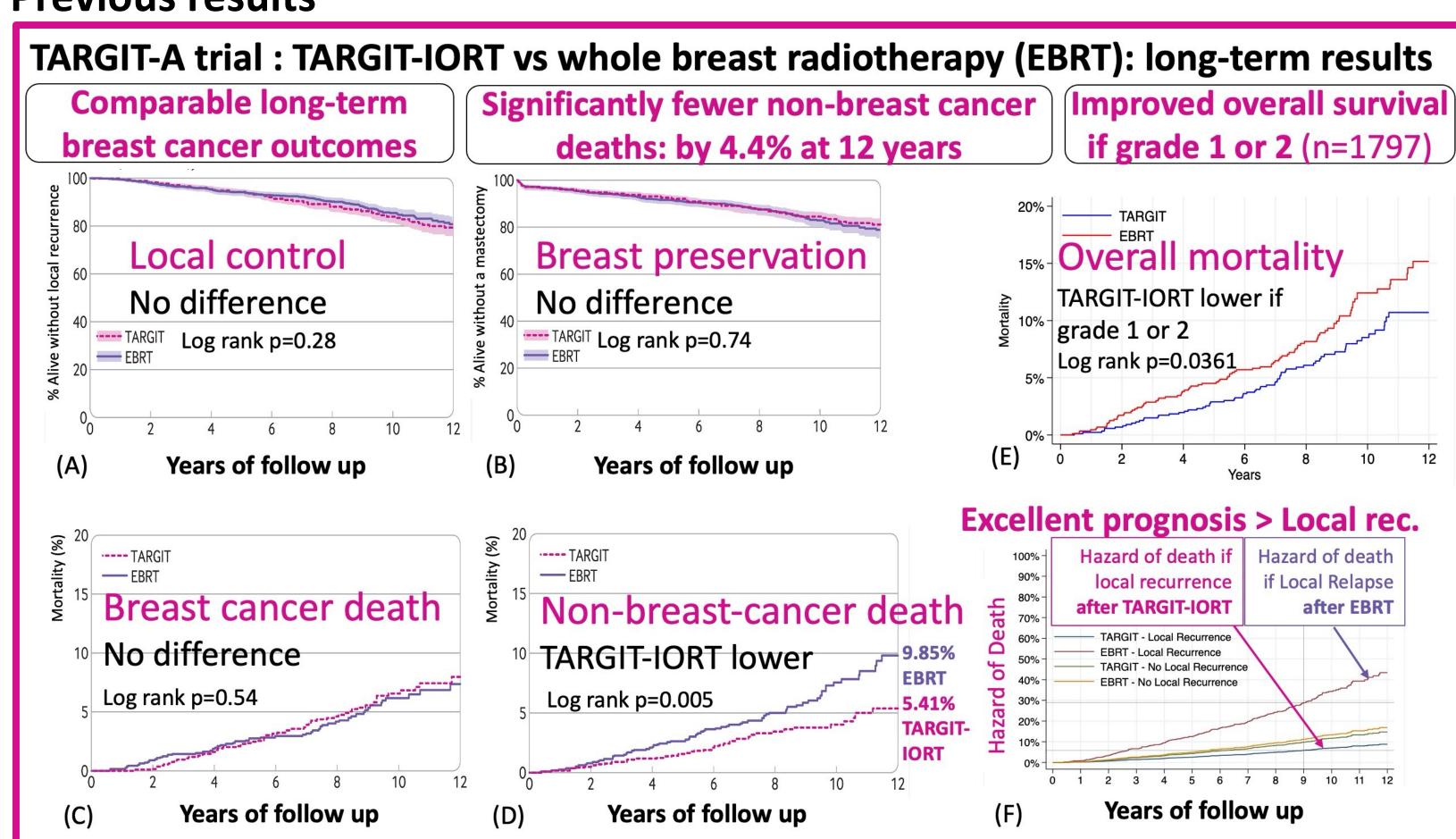
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- Significantly fewer deaths from non-breast-cancer causes with TARGIT-IORT vs. whole breast radiotherapy (EBRT).
- TARGIT-IORT conferred an overall survival benefit in patients with grade 1 or 2 cancers: 12-year mortality reduction from 15% to 10.5%

TARGIT-IORT reduced pain, improved cosmetic outcome, quality of life,

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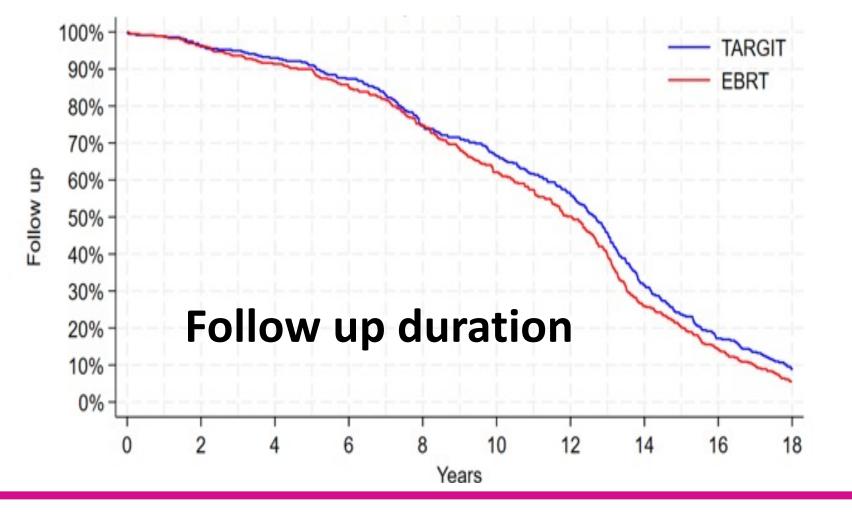
Previous results



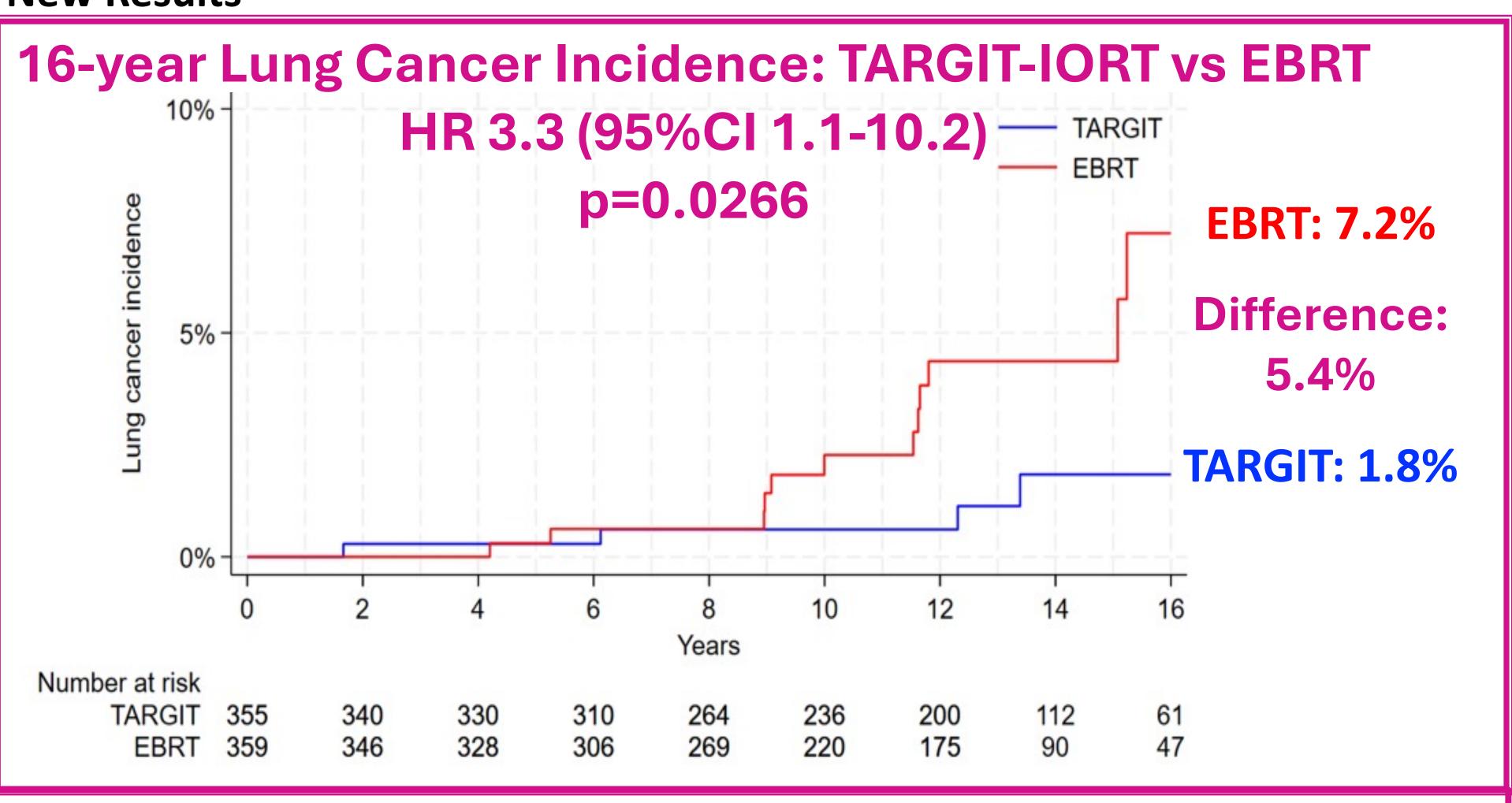
Material and methods

- We collected long term data about health status and new cancer diagnoses of UK patients from the TARGIT-A randomised trial, using direct patient contact, & NHS Digital data.
- We compared lung cancer incidence between patients randomised to TARGIT-IORT vs EBRT.

N.B. The duration of follow-up for patients receiving EBRT was slightly shorter vs TARGIT-IORT (p=0.0399). In other words, some cases of lung cancer in the EBRT group may have been missed, so the benefit of targeted intraoperative radiotherapy may have been underestimated.



New Results



Headline results

- Significantly more lung cancer with EBRT vs TARGIT-IORT HR 3.3 (95%CI 1.1-10.2)., p=0.0266
- 16-year incidence of lung cancer: EBRT: 7.2% vs. TARGIT: 1.8% Reduction with TARGIT-IORT = 5.4% (95%CI 0.3 -10.5)

50,000 breast cancer patients could avoid getting lung cancer by taking TARGIT-IORT

*An estimated 920,000 breast cancer patients worldwide are suitable for TARGIT-IORT during lumpectomy, annually.

Using the 5.38% reduction in lung cancer risk that we have observed, if TARGIT-IORT were to be made accessible to these patients, then 49,496 (95%CI 5500-134320) of these patients would be spared the diagnosis of a lung cancer during their follow up.

Conclusions

- With very long-term follow data from of a large TARGIT-A randomised trial, we found a substantial increase in lung cancer incidence with EBRT vs TARGIT-IORT.
- It is a tragedy when women who outlive breast cancer then succumb to this frequently lethal radiation-induced lung cancer, which is avoidable by using TARGIT-IORT during lumpectomy instead of post-operative EBRT.
- These new data further mandate* full discussion about benefits of TARGIT-IORT vs. EBRT with patients, including reduction in lung cancer incidence, before their surgery, so they have a choice to take it during their initial surgical excision.

*Discussion about TARGIT-IORT as an option to EBRT is mandatory as per GMC guidance and the UK law (UK Supreme Court. Montgomery v Lanarkshire Health Board. 11 Mar 2015.

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