

**ESTRO**
2026**15-19 May 2026****Stockholm, Sweden**

ESTRO 2026

AMEND

CONTINUE**Submission ID**

3766

Author approval (read-only)

I confirm that this abstract has been approved by all authors.

Accuracy of the abstract (read-only)

I accept responsibility for the accuracy of the abstract. I understand that no changes to the abstract text are possible after the submission deadline of 12 November 2025 (23h59 CET).

Author list (read-only)

I confirm that all authors contributing to the abstract are properly included in the author list. I understand that no changes to the author list are possible after the 1 February 2026.

Abstract presentation (read-only)

I certify that, if the abstract is selected, the information reported is for the exclusive presentation in the scientific programme and will not be presented at any commercially-sponsored satellite symposia during the congress.

register to attend and present at the congress.

Embargo of abstracts (read-only)

I understand that abstracts are embargoed until 1 month prior to the congress at which point abstracts are published in the abstract book and online searchable programme. Should you wish for an abstract to be embargoed until the congress, please email abstracts@estro.org by 28 February 2026.

Abstract verification (read-only)

I accept to check the abstract overview provided in the submission confirmation email and ensure all abstract details are correct.

Contact person (read-only)

I accept to be the contact person for all correspondence related to the abstract and to share relevant information with the co-authors.

Topic (read-only)

Physics: Inter-fraction motion management and daily adaptive radiotherapy

Keywords (read-only)

Synthetic-CT, Adaptive, Breast

Request for Digital Poster only

I want this abstract to ONLY be considered for a Digital Poster. N.B. I understand that this abstract will then NOT be considered for a live presentation (Proffered Paper, Mini-Oral, Poster Discussion).

Abstract title (read-only)

Diffusion-based synthetic-CT generation from breast CBCT

Clinical affairs, TheraPanacea, Paris, France

Audrey Duran a.duran@therapanacea.com

AI engineering, TheraPanacea, Paris, France

Quentin Spinat q.spinat@therapanacea.com

AI engineering, TheraPanacea, Paris, France

Pierre Olléon p.olleon@therapanacea.com

AI engineering, TheraPanacea, Paris, France

Sami Romdhani s.romdhani@therapanacea.com

AI engineering, TheraPanacea, Paris, France

Nikolaos Paragios (Presenting) n.paragios@therapanacea.com

CEO, TheraPanacea, Paris, France

Pauline Maury pauline.maury@gustaveroussy.fr

Department of radiation oncology, Institut Gustave Roussy, Villejuif, France

Pascal Fenoglietto pascal.fenoglietto@icm.unicancer.fr

Department of radiation oncology, Institut du Cancer de Montpellier, Montpellier, France

Purpose/Objective (read-only)

Cone-beam CT (CBCT) imaging is routinely employed in radiation therapy for patient positioning, but its limited image quality, spatial resolution, and field of view restrict its broader clinical use. Extending its role beyond positioning could accelerate the adoption of adaptive radiotherapy. To achieve this, improvements enabling organ-at-risk delineation, dose calculation, and treatment replanning are essential. This study clinically evaluates an AI-based solution that generates synthetic-CTs (sCTs) from CBCT images, addressing major technical constraints and demonstrating the potential to fully exploit CBCT for adaptive radiotherapy in breast cancer management.

Material/Methods (read-only)

.Training a 3D multimodal translation model for medical imaging presents several challenges, including the large computational burden of volumetric data, the scarcity of high-quality paired datasets, and the need to preserve quantitative fidelity for clinical use. To overcome these, a 3D latent diffusion model was developed. Its variational autoencoder (VAE) component achieves strong compression with high reconstruction quality and computational efficiency. The diffusion model was first trained in an unsupervised manner on unpaired data, then fine-tuned for cross-modality translation using limited paired samples. A ControlNet-based conditioning strategy was employed to achieve anatomically consistent and quantitatively reliable image synthesis. The unsupervised model was trained on approximately 34000 CT scans (50% pelvis, 37.5% head and neck, 12.5% thorax) to ensure anatomical diversity, followed by fine-tuning with 248 breast cancer patients (1126 paired datasets).

Results (read-only)

Median relative differences in DVH parameters for target volumes between sCTs and wCTs were minimal (0.09–0.19% - Table 1), with the largest difference observed for D98% (0.19%). Median gamma pass rates reached 99.74% (2%/2mm) and 99.96% (3%/3mm) with a 1% low-dose threshold (Table 2), confirming excellent dosimetric agreement.

Structure	DVH parameter	Median relative difference (%)
Target volumes	Dmax [Gy]	0.16
	Dmean [Gy]	0.09
	D2% [Gy]	0.10
	D5% [Gy]	0.10
	D50% [Gy]	0.10
	D95% [Gy]	0.17
	D98% [Gy]	0.19

Gamma index	2%/2mm			3%/3mm		
Low dose threshold	1%	10%	20%	1%	10%	20%
Median	99.74	99.40	99.12	99.96	99.89	99.86

Conclusion (read-only)

This study demonstrates the feasibility and accuracy of AI-based CBCT-to-CT conversion for adaptive breast radiotherapy. The resulting sCTs provide dosimetric precision comparable to conventional CTs, supporting their integration into a comprehensive adaptive workflow including organ-at-risk delineation, dose simulation, and replanning to enhance personalised and efficient radiotherapy.

Conflict of Interest Disclosure (read-only)

- ☒ I do not have a Conflict of Interest to disclose.
- ☐ I do have a Conflict of Interest to disclose.

Permission to publish (required)

In submitting this abstract, you give ESTRO permission to publish it in the congress abstract book, a supplement to the Green Journal and online in various forms (searchable programme, media library, congress platform and others).

