# Primary Surgery vs. Neoadjuvant Chemotherapy in Advanced Ovarian, Tubal, and Peritoneal Carcinoma: Real-World **Evidence from a Cancer Registry**

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## Background

Standard: Primary debulking surgery (PDS) + chemotherapy (CTx)

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• Alternative: Neoadjuvant chemotherapy (NACT) + interval debulking surgery (IDS) - efficacy uncertain

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- Evidence:
  - EORTC-GCG & CHORUS → NACT+IDS non-inferior

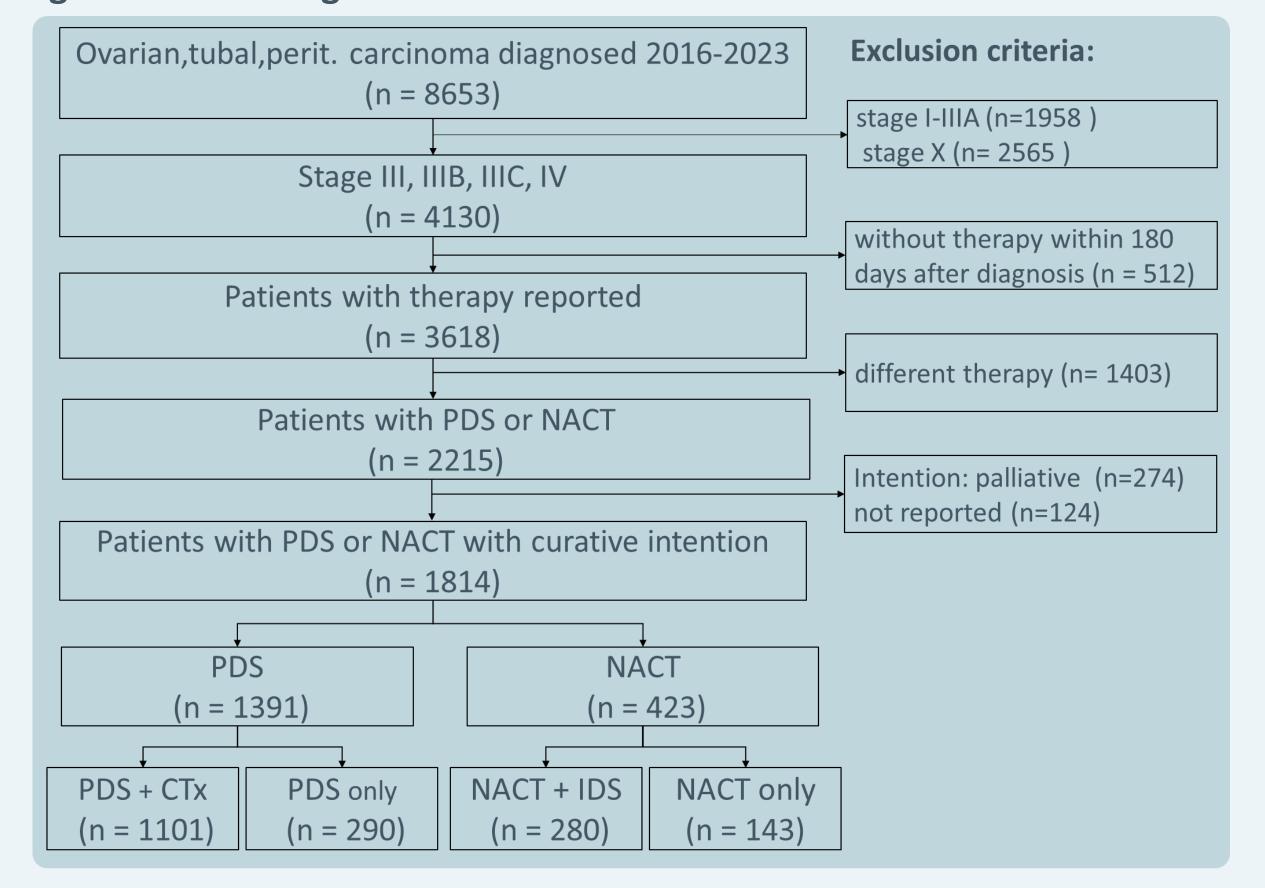
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- TRUST → PDS improved PFS (esp. <65 yrs, stage III), no OS benefit
- Aim: Compare oncologic outcomes of NACT vs. PDS in advanced ovarian, tubal & peritoneal carcinoma using real-world data of Baden-Wuerttemberg Cancer Registry (BWCR), Germany

### Methods

- Study type: Cancer registry study
- Data source: Baden-Wuerttemberg Cancer Registry (BWCR), Germany
- Patients: FIGO IIIB-IV ovarian, tubal, or peritoneal carcinoma (diagnosed 2016-2023)
- **Treatment**: PDS or NACT with paclitaxel-carboplatin (+/- bevacizumab).
- Statistics: Fisher's exact/chi-square test; Kaplan-Meier and Cox models.
- Endpoints:
  - Primary Overall survival (OS), progressive-free survival (PFS)
  - Secondary surgical outcomes (macroscopic complete resection (CR) vs. residual disease) and 90-day postoperative mortality.

Fig. 1: Consort Diagram



- No surgery after NACT: 143 (33.8%) patients.
- No adjuvant CTx after PDS: 290 (20.8%) patients

#### Results

**Table 1: Patient characteristics** 

	Overall	NACT	PDS	P - value
Total – no. (%)	1814	423 (23.3)	1391 (76.7)	
Age – mean (SD)	64.0	66.0	64.0	0.036
stage – no. (%)				0.38
<b>=</b> []]	1243 (68.5)	282 ( 66.7)	961 (69.1)	
	571 (31.5)	141 (33.3)	430 ( 30.9)	
grade – no. (%)				0.176
<b>1</b> -2	147 (8.4)	27 ( 6.6)	120 ( 8.9)	
<b>3</b>	1610 (91.6)	381 (93.4)	1229 (91.1)	
Localisation – no. (%)				<0.001
<ul><li>Ovar</li></ul>	1512 (83.4)	352 (83.2)	1160 (83.4)	
<ul><li>Fallopian tube</li></ul>	218 (12.0)	34 ( 8.0)	184 ( 13.2)	
<ul><li>Peritoneum</li></ul>	84 ( 4.6)	37 ( 8.7)	47 ( 3.4)	

#### Baseline characteristics:

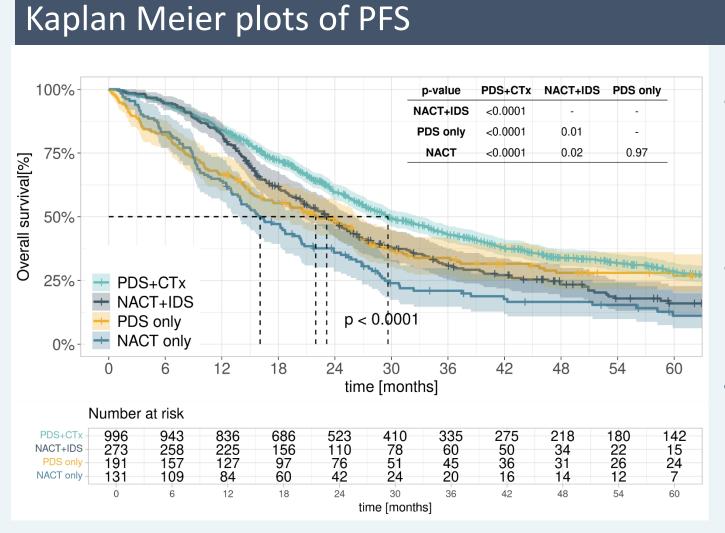
- Well balanced between treatment groups
- Stage IV disease: 28.8% of the cohort.

**Table 2: Oncological Outcomes** 

	Overall	NAT	PDS	P - value
90d mortality – no. (%)	54 ( 3.0)	4 ( 0.9)	50 ( 3.6)	0.008
Residual disease – no.(%)				0.018
CR	927 (66.2)	183 (73.8)	744 ( 64.6)	
<ul> <li>Microscopic residual</li> </ul>	243 (17.4)	31 ( 12.5)	212 ( 18.4)	
<ul> <li>Macroscopic residual</li> </ul>	230 (16.4)	34 ( 13.7)	196 ( 17.0)	

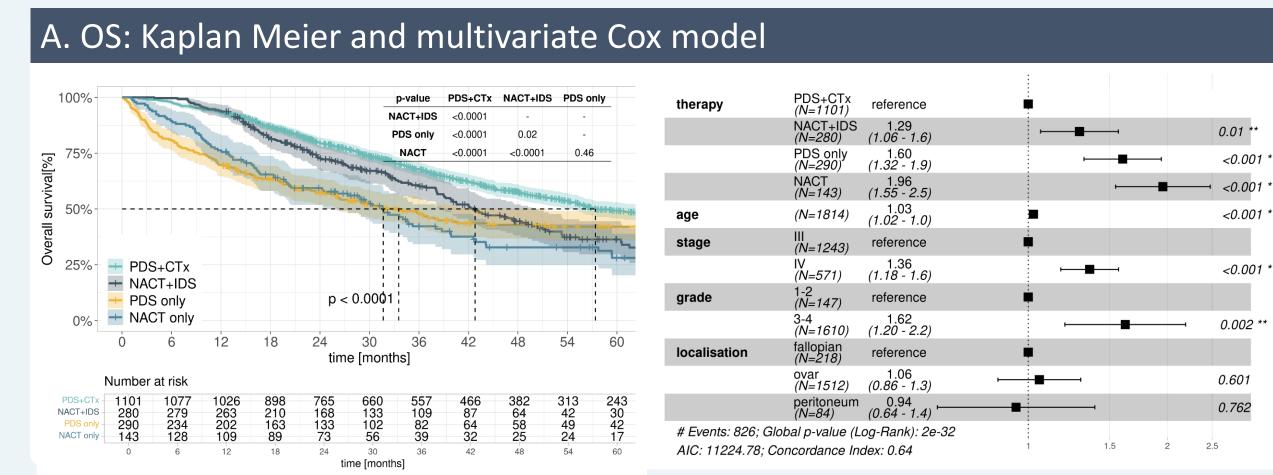
- Complete resection rates: Higher in NACT+IDS (73.8% vs. 64.6% (p = 0.018))
- 90-day postoperative mortality: Significantly higher after PDS:
  - 3.6% (PDS) vs. 0.9% (NACT+IDS) (p = 0.008)

Fig. 2: Progressive free survival

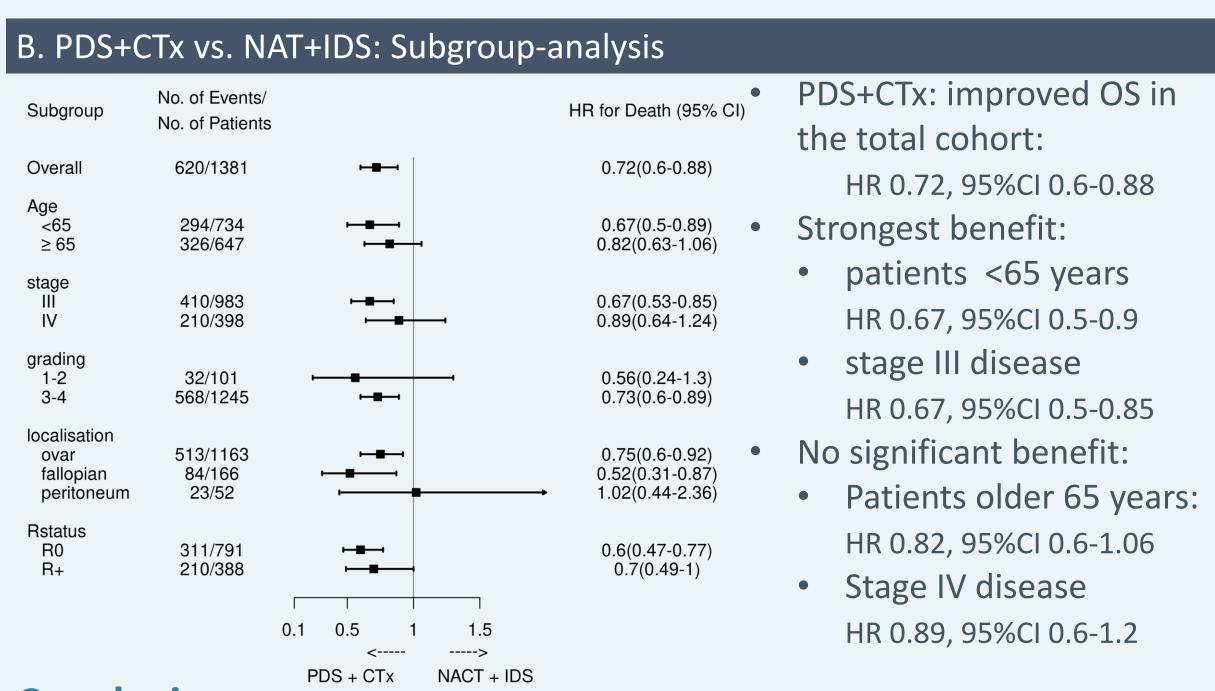


- longer PFS than PDS+CTx NACT+IDS
- mPFS: 29.6 (95%CI: 27.7-33.0) vs 23.1 (95% CI: 20.6-26.1) months
- NACT without IDS → poorest PFS: **mPFS:** 16.1 months (95% CI: 13.9.-
- PDS only / NACT+IDS → similar outcomes • **mPFS**: 22.0 (95% CI: 16.5-26.4) vs
- 23.1 (95% CI: 20.6-26.1) months

Fig. 3: Overall Survival



- median follow-up: 53.6 months
- PDS+CTx was associated with longer OS compared to NACT+IDS:
  - **mOS:** 57.4 vs 42.8 months (p=0.01)
- Poorer outcomes were observed in patients receiving NACT without subsequent IDS and patients with PDS only:
- **mOS**: 32 (33.6) months (p<0.0001)
- Independent prognostic factors:
  - age (HR 1.03, 95%Cl 1.02-1.04), tumor grade (HR 1.6, 95%Cl 1.2-2.2), and stage (HR 1.36, 95% CI 1.2-1.6)



#### Conclusion

- Key determinant of outcome is treatment pathway completion; failure to complete multimodal therapy predicts the poorest prognosis.
- NACT+IDS achieved higher complete resection rates and lower 90-day mortality, but patients without subsequent surgery had poor outcomes.
- PDS provided a survival benefit in patients under 65 years and with stage III disease, while outcomes were similar in older patients and those with stage IV disease.
- Real-world data from state-run cancer registries help bridge the gap between clinical trials and daily oncology practice, supporting treatment decisions in real-world settings.



**Conflict of interest:** No conflict of interest



