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1 BACKGROUND / OBJECTIVE



To evaluate long-term outcomes, relapse patterns and clinicopathological factors associated with recurrence in patients with clinical stage I seminoma managed in routine practice over three decades.

2 METHODS



- Retrospective single-centre study.
- Clinical stage I seminoma, diagnosed 1993–2025.
- Variables: baseline features, post-orchietomy strategy, relapse events/timing.
- Management: active surveillance or adjuvant carboplatin.
- Associations assessed via descriptive statistics and Cramér's V.

3 PATIENT CHARACTERISTICS

n = 113 patients



Median age at diagnosis **36 years**



Stage IB disease **21.2%**



Tumour size >4 cm **56.0%**



Rete testis invasion **26.5%**



Lymphovascular invasion **16.8%**



Diagnosis period **1993–2025**

4 MANAGEMENT AND RELAPSE PATTERNS



Active surveillance: 59.3%



Adjuvant carboplatin: 39.8%



Overall relapses: 6 patients (5.3%)



All relapses occurred after surveillance



Mean time to relapse: 17.2 months



Low overall relapse rate



No relapses after adjuvant carboplatin



Recurrences concentrated in surveillance group



Structured follow-up is essential

5 KEY OUTCOMES AND RISK FACTORS



Patients included

113



Relapse rate

5.3%



Relapses after surveillance

6 / 6



Mean time to relapse

17.2 months



Strongest association with relapse

Lymphovascular invasion



Cramér's V / p-value

0.30 / 0.006

6 CONCLUSIONS



In this real-world cohort of clinical stage I seminoma, relapse rates were low and all recurrences occurred after surveillance, supporting the overall safety of a de-escalated post-orchietomy strategy when combined with structured follow-up. Lymphovascular invasion emerged as a potential marker of relapse risk in this dataset, although the small number of events precludes definitive conclusions. These findings support individualized counselling and highlight the value of long-term real-world series to refine risk-adapted management in stage I seminoma.