Ovarian cancer symptoms, routes to diagnosis and survival – Population cohort study in the ‘no screen’ arm of the UK Collaborative Trial of Ovarian Cancer Screening (UKCTOCS)

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Objective. There are widespread efforts to increase symptom awareness of ‘pelvic/abdominal pain, increase abdominal size/bloating and difficulty eating/feeling full’ in an attempt to diagnose ovarian cancer earlier. Long-term survival of women with these symptoms adjusted for known prognostic factors is yet to be determined. This study explored the association of symptoms, routes and interval to diagnosis and long-term survival in a population-based cohort of postmenopausal women diagnosed with invasive epithelial tubo-ovarian cancer (iEOC) in the ‘no screen’ (control) UKCTOCS arm.

Methods. Of 101,299 women in the control arm, 574 were confirmed on outcome review to have iEOC between randomisation (2001–2005) and 31 December 2014. Data was extracted from medical notes and electronic records. A multivariable model was fitted for individual symptoms, time interval from symptom onset to diagnosis, route to diagnosis, speciality, morphological Type, age at diagnosis, year of diagnosis (period effect), stage, primary treatment, and residual disease.

Results. Women presenting with symptoms listed in the NICE guidelines (HR1.48, 95%CI1.16–1.89, p = 0.001) or the modified Goff Index (HR1.68, 95%CI1.32–2.13, p < 0.0001) had significantly worse survival than those who did not. Each additional presenting symptom decreased survival (HR1.20, 95%CI1.12–1.28, p < 0.0001). In multivariable analysis, in addition to advanced stage, increasing residual disease and inadequate primary treatment, abdominal pain and loss of appetite/feeling full were significantly associated with increased mortality.

HIGHLIGHTS

• This study explored the association of symptoms of ovarian cancer, interval and route to diagnosis with survival.
• Focus on ‘high alert’ symptoms: pelvic/abdominal pain, increase abdominal size/bloating and difficulty eating/feeling full
• The ovarian cancer ‘high alert’ symptom complexes identify postmenopausal women with a significantly poorer prognosis.
• The study could not however exclude the possibility of better outcomes in those who are aware and acted on these symptoms.

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1. Introduction

Ovarian cancer continues to be diagnosed at advanced stage with high fatality rates. A significant contributing factor is lack of clear alarm symptoms. To address this, substantial work in exploring symptoms has been undertaken. Although symptoms in women diagnosed with early and late stage disease were described as early as 1985 [1], it was Goff et al. [2,3] in 2004 who gave impetus to this effort by describing a symptom triad (pelvic/abdominal pain, increased abdominal size/bloating and difficulty eating/feeling full) frequently associated with ovarian cancer. This Goff Index [3] with some modifications [4,5] has since been widely used across the globe to drive awareness campaigns [6] among the public and primary care physicians.

Testing symptomatic women was studied in the DOvE randomised controlled trial [7]. Although there was no stage shift, there was a trend to higher complete tumour resection rates in women diagnosed with ovarian cancer in the intervention compared to the standard care arm (73% vs 44%; p = 0.075). Clinical implementation of symptom triggered testing in the UK found that while more women with ovarian cancer accessed expedited care pathways (two-week urgent referral), there was no stage shift [8]. This was in keeping with retrospective studies that reported no difference in stage or survival associated with time to diagnosis in a population cohort of ovarian cancer patients diagnosed in Australia [9] or with two-week urgent referral in an English hospital-based cohort [10].

We are not aware of any studies in women with ovarian cancer that explore the association of symptoms and routes to diagnosis with long term survival adjusted for prognostic factors. Data from the population cohort of women presenting clinically with invasive epithelial tubo-ovarian cancer [11] (IEOC) in the ‘no screening’ (control) arm of the UK Collaborative Trial of Ovarian Cancer Screening (UKTOCOS) provides an opportunity to explore this issue in more depth. We report on association of symptoms, intervals and routes to diagnosis with survival adjusted for prognostic factors.

2. Materials and methods

UKTOCOS is a randomised controlled trial assessing the impact of population screening on ovarian cancer mortality [12,13]. In brief, following random invitation from population registers between 2001 and 2005, 202,638 postmenopausal women aged ≥50–74 years, were recruited through 13 regional centres in England, Wales and Northern Ireland. They were randomised to no screening (Control group – 101,359) or annual screening using a multimodal (MMS 50,640) or ultrasound (USS, 50,639) strategy. The cohort only included women with at least one ovary, no significant family history of ovarian cancer and no personal history of ovarian cancer or an active non-ovarian malignancy.

2.1. Follow-up and confirmation of diagnosis

Participants were followed up via electronic health record linkage to national cancer and death registrations (England and Wales - NHS digital; Northern Ireland-Health and Social Care Business Services Organisation and Northern Ireland Cancer Registry). Additional sources included two rounds of postal questionnaires (3–5 years after randomisation and in 2014) and direct communication from participants. For women based in England, data was also obtained from the National Cancer Intelligence Network (April 2001–March 2010) and Hospital Episode Statistics administrative records. Censorship date for this analysis was 31st Dec 2014.

As previously detailed [13,14], medical notes were retrieved for all women with notifications of a possible ovarian cancer diagnosis. An independent outcomes review committee assigned the final diagnosis [15], date of diagnosis, FIGO 2003 stage, morphological Type [16] (Type I- low-grade serous, low-grade endometrioid, clear cell, mucinous, Type II - mainly high-grade serous carcinoma eTable 1), and cause of death (where applicable).

2.2. Subjects

All women with confirmed diagnosis as per WH0 2014 of primary invasive epithelial tubo-ovarian cancer (includes tubal/peritoneal - ICD-10 C57·0, C48·1, C48·2 in addition to C56.0) on outcomes review were included in the analysis [17]. Women with borderline and non-epithelial ovarian cancer were excluded.

2.3. Symptom ascertainment

Symptom data was retrieved by hand searching of medical records, which included copies of GP and hospital letters, multidisciplinary team (MDT) summaries, hospital notes as well as trial records. All symptoms were captured unless notes review confirmed they were longstanding which was defined as ≥12 months. Any accompanying diagnoses (e.g. joint pain with diagnosis of osteoarthritis) were noted. No limit was placed on the number of symptoms that could be recorded for each woman.

Women were classified as ‘symptomatic’ if they had presented with any symptoms or ‘asymptomatic’ if this was documented or no symptoms were mentioned despite the availability of comprehensive documentation. Women with ‘insufficient’ documentation were classified as having missing data. The symptoms were grouped as National Institute for Health and Care Excellence (NICE) UK guideline [4] - positive (abdominal or pelvic pain, increased abdominal size or bloating, loss of appetite/feeling full and increased urinary urgency or frequency) or modified Goff Symptom index positive (abdominal or pelvic pain, increased abdominal size or bloating and loss of appetite/feeling full). The Goff symptom index includes duration and frequency of symptoms. The latter was not included in our analysis as frequency was often not captured in the hospital notes. Symptoms were also grouped according to system (gynaecological, abdominal, gastrointestinal, urinary, systemic, other) as described previously [18]. Symptoms not previously described were allocated to the most appropriate system upon agreement of two clinical researchers (JD and UM) (eTable 2).

Symptom interval was calculated from onset to date of diagnosis. Date of onset was derived using GP and secondary care records. Where women had multiple episodes rather than a persistent symptom, the start date of the first episode was recorded as date of onset. When only the month was available, the midpoint (15th) was used. Diagnosis was based on histological confirmation with the date used that of primary surgery or biopsy. In cases with no surgery or biopsy, the date cytology was taken was used. Where none of these were available the date confirmatory imaging was used.
The date the woman was first seen in hospital, the speciality involved and the route to that appointment was recorded. For those residents in England, this was supplemented where missing, with HES data. The routes were classified as

1) emergency presentation (via accident and emergency department)
2) two-week urgent ‘cancer’ referral to rapid access outpatient diagnostic clinic
3) routine referral to secondary care (e.g. gynaecology or colorectal outpatient clinic). The patients who were seen privately (outside the NHS) were included in this group.

In women who had multiple appointments prior to diagnosis, the first was taken as their route to diagnosis.

2.5. Primary treatment and surgical outcome

The medical notes especially the surgical records and details of chemotherapy received were used to classify women into primary treatment categories (eTable 1). In patients who underwent surgery whether primary or interval debulking, the amount of residual disease as recorded by the surgeon in the operation record was extracted.

2.6. Statistical analysis

Standard survival methods were used to analyse the data, with entry fixed at date of diagnosis, exit at date of death from ovarian cancer, or date of censorship (loss to follow-up or 31st December 2014). Cox models were used to obtain hazard ratio estimates of variables. Formal tests of the proportional hazards assumption were performed using Schoenfeld residuals.

A multivariable model was fitted for individual symptoms, time interval from symptom onset to diagnosis, route to diagnosis, speciality, Type, age at diagnosis, year of diagnosis (period effect), BMI, stage, primary treatment, and residual disease. To look at period effects, the year of diagnosis was categorised with 2010 as the reference, as this was the year prior to the introduction of NICE guidelines in UK. To address the issue of missing data, we performed a multiple imputation (MI) using chained equations (MICE) to preserve as much information as possible and provide correct inference for the multivariable Cox model. The three modified Goff symptom variables were imputed using a logit model, time from symptom to diagnosis using a linear regression model after first log transforming, stage using an ordered logit model, and route to diagnosis and speciality using a multinomial logit model. 20 fully imputed datasets were created using MICE and following model fitting on all 20 sets, parameter estimates with standard errors were calculated using Rubin’s rules [19].
Table 1

Presenting symptoms/symptom complexes (only symptoms with ≥10% women) Number of women who had the symptom n (%) Histological

| Presenting symptoms/symptom complexes | Type I (n = 88) | Type II (n = 424) | Type uncertain (n = 62) | p-Value between all Types | p-Value between Type I and II |
|---------------------------------------|----------------|-----------------|------------------------|--------------------------|-----------------------------|
| Abdominal or pelvic discomfort/pain    | 227 (39.5%)    | 26 (29.5%)      | 170 (40.1%)            | 30 (48.4%)               | 0.055                       | 0.058                       |
| Increased abdominal size/bloating     | 225 (39.2%)    | 22 (25%)        | 167 (39.4%)            | 35 (56.5%)               | 0.001                       | 0.023                       |
| Change in bowel habit                 | 115 (20%)      | 11 (12.5%)      | 88 (20.8%)             | 15 (24.2%)               | 0.133                       | 0.068                       |
| Loss of appetite/feeling full         | 84 (14.6%)     | 7 (8%)          | 62 (14.6%)             | 15 (24.2%)               | 0.022                       | 0.096                       |
| Weight loss                           | 67 (11.7%)     | 6 (6.8%)        | 53 (12.5%)             | 8 (12.9%)                | 0.304                       | 0.129                       |
| Lump or mass felt by woman            | 45 (7.8%)      | 12 (13.6%)      | 29 (6.8%)              | 4 (6.5%)                 | 0.089                       | 0.033                       |
| Respiratory symptoms                  | 44 (7.7%)      | 3 (3.4%)        | 33 (7.8%)              | 8 (12.9%)                | 0.097                       | 0.144                       |
| Vaginal bleeding                      | 44 (7.7%)      | 11 (12.5%)      | 30 (7.1%)              | 3 (4.8%)                 | 0.167                       | 0.106                       |
| Urinary frequency                     | 39 (6.8%)      | 8 (9.1%)        | 31 (7.3%)              | 0 (0%)                   | 0.066                       | 0.567                       |
| Nausea                                | 29 (5.1%)      | 2 (2.3%)        | 23 (5.4%)              | 4 (6.5%)                 | 0.408                       | 0.212                       |
| Other symptom                         | 171 (29.8%)    | 24 (28.4%)      | 126 (29.7%)            | 20 (32.3%)               |                            |                            |
| Asymptomatic                          | 13 (2.3%)      | 2 (2.3%)        | 16 (3.7%)              | 0 (0%)                   |                            |                            |
| Abdominal                             | 40 (6.9%)      | 9 (10.2%)       | 29 (6.8%)              | 2 (3.2%)                 |                            |                            |
| Gastrointestinal                      | 375 (65%)      | 51 (58%)        | 277 (65.3%)            | 47 (76.6%)               |                            |                            |
| Systemic                              | 156 (27%)      | 18 (20.5%)      | 118 (27.8%)            | 20 (32.3%)               |                            |                            |
| Urinary                               | 133 (23%)      | 11 (12.5%)      | 101 (23.8%)            | 21 (33.9%)               |                            |                            |
| Respiratory                           | 76 (13%)       | 13 (14.8%)      | 61 (14.4%)             | 2 (3.2%)                 |                            |                            |
| Gynaecological                        | 60 (10%)       | 12 (13.6%)      | 45 (10.6%)             | 3 (4.8%)                 |                            |                            |
| Other                                 | 65 (11%)       | 7 (8%)          | 49 (11.6%)             | 9 (14.5%)                |                            |                            |
| Modified Goff                         | 362 (63%)      | 47 (53.4%)      | 268 (63.2%)            | 47 (75.8%)               |                            |                            |
| NICE                                  | 381 (66%)      | 53 (60.2%)      | 281 (66.3%)            | 47 (75.8%)               |                            |                            |

Interval from symptom onset to diagnosis in days- median (inter quartile)

| Interval from symptom onset to diagnosis in days- median (inter quartile) | Type I | Type II | Type uncertain |
|--------------------------------------------------------------------------|--------|---------|----------------|
| No. of patients with symptoms                                           | 521    | 77      | 384            |
| No. of patients with data on interval                                   | 316    | 46      | 232            |
| Overall                                                                  | 80 (83) | 93 (122) | 78 (82.5)     |
| Early Stage patients                                                     | 91 (88.5) | 92 (79) | 88 (71)       |
| Latent Stage patients                                                    | 75 (84) | 98 (144) | 76 (82)       |
| One and five year survival                                               | 574    | 88      | 424            |
| One year survival (IQR)                                                  | 76%    | 93%     | 78%            |
| Five year survival (IQR)                                                 | 33%    | 76%     | 29%            |
| One year survival early stage patients                                   | 94%    | 97%     | 91%            |
| Five year survival early stage patients                                  | 79%    | 88%     | 70%            |
| One year survival late stage patients                                    | 71%    | 78%     | 75%            |
| Five year survival late stage patients                                   | 19%    | 28%     | 21%            |

Fig. 1. Survival curves of women with invasive epithelial tubo-ovarian cancer in the ‘no screening’ arm of UKCTOCS, with presenting symptoms included in (A) NICE guidelines vs not included (B) modified Goff symptom index vs not included.

Fig. 2. Survival curves of women with invasive epithelial tubo-ovarian cancer in the ‘no screening’ arm of UKCTOCS. (A) With the three individual symptoms that constitute the modified Goff index (B) With symptoms affecting different systems*. *Abdominal, GI, gynae, urinary, systemic, other and none.
presenting as an emergency had significantly worse survival (eFig. 1D, eTable 6). Irrespective of route to diagnosis, compared to women initially managed by a gynaecologist, those initially managed by the emergency physicians or gastrointestinal physicians/surgeons had worse survival (eFig. 1E).

The multivariable analysis involves 573 patients – one woman had no treatment data and was excluded. In addition to advanced stage, inadequate primary treatment and increasing residual disease, abdominal pain, loss of appetite/feeling full, were significantly associated with increased mortality (Table 2).

4. Discussion

In this population cohort study of postmenopausal women with invasive epithelial ovarian/tubal/peritoneal cancer, those with symptoms listed on the NICE guidelines or modified Goff Index had significantly worse survival compared to those who did not. Abdominal pain and loss of appetite/feeling full was significantly associated with increased mortality on multivariate analysis alongside advanced stage, increasing residual disease and inadequate primary treatment. It needs to be noted however, that this study cannot exclude the possibility that within the group of symptomatic women who collectively have a poor outcome, those who act early on their symptoms may have a better outcome than those who do not.

4.1. Strength and weakness

The focus is on invasive epithelial cancer, which is the major contributor to mortality in this disease. The key strength is that selection bias is minimised as significant efforts were made via linkage to multiple national registries and postal follow-up to ensure all those with the disease were identified. Site, stage and Type were confirmed by independent outcome review. Similar incidence and survival of ovarian cancer in this group compared to the UK age matched population [14] ensures that the cohort is representative and the findings therefore generalisable. Symptom data were collected using both primary and secondary care records. To allow comparison, similar rules as reported in the literature [18], were used to group symptoms by system. Although symptom data was collected prior to any diagnosis of ovarian cancer, the retrospective review of records is a weakness. Consequently the collection of symptom data is different from the Goff index that uses prospective questionnaire to screen women for symptoms of ovarian cancer, the retrospective review of records is a weakness. Hence, previous work [20] has suggested questionnaires and medical records have comparable specificity. The NICE symptom index relies on women presenting to healthcare professionals and volunteering symptoms, a method, which is more closely aligned to that used in this study. The Goff symptoms index uses both duration and frequency in the evaluation process. Though data was collected for duration of symptoms, it was missing in 45% (258/574). Multiple imputation was undertaken using MICE to preserve as much information as possible and provide correct inference. It was not possible to determine the frequency or severity of symptoms. A further potential weakness is that the model does not include performance status or medical comorbidity, both of which could be confounders when interpreting certain symptoms and in survival.

4.2. Findings in the context of literature

Since the publication of the Goff symptom index [3], symptom awareness has been promoted as an approach that would enable earlier diagnosis of ovarian cancer and has been incorporated into national guidelines such as NICE [4]. Our findings of worse survival in women with NICE or modified Goff symptoms are in keeping with reports that these symptoms are more likely in women with advanced disease [21].

The incidence of common symptoms reported in our study (abdominal/pelvic discomfort or pain, increased abdominal size or bloating and

| Variable | HR   | 95% CI | p value | se  | t     |
|----------|------|--------|---------|-----|-------|
| Asymptomatic (compared to symptomatic) | 0.78 | 0.28–2.21 | 0.643 | 0.415 | −0.46 |
| Loss of appetite/Feeing full | 1.43 | 1.04–1.96 | 0.026 | 0.229 | 2.23 |
| Abdominal/Pelvic pain | 1.34 | 1.05–1.72 | 0.018 | 0.168 | 2.37 |
| Increase abdominal size/Bloating | 1.12 | 0.85–1.48 | 0.411 | 0.158 | 0.82 |

| Initial specialist seen in secondary care (compared to Emergency department) | Overall p value for variable | 0.648 |
|---|---|---|
| Emergency department | 1.00 |
| Gynaecology | 0.80 | 0.48–1.33 | 0.386 | 0.208 | −0.87 |
| Gastrointestinal | 0.87 | 0.55–1.40 | 0.588 | 0.209 | −0.57 |
| Other | 0.73 | 0.42–1.29 | 0.279 | 0.211 | −1.08 |

| Route to diagnosis (compared to two-week urgent cancer) | Overall p value for variable | 0.780 |
|---|---|---|
| Two-week urgent cancer | 1.00 |
| Routine | 1.02 | 0.76–1.37 | 0.892 | 0.152 | 0.14 |
| Emergency | 1.12 | 0.79–1.59 | 0.506 | 0.199 | 0.66 |
| Time to symptoms to diagnosis | 0.95 | 0.79–1.14 | 0.589 | 0.087 | −0.54 |

| Stage (compared to Stage I) | Overall p value for variable | 0.001 |
|---|---|---|
| 2 | 1.00 |
| 3 | 1.36 | 0.63–2.92 | 0.422 | 0.531 | 0.79 |
| 4 | 3.27 | 1.75–6.11 | −0.001 | 1.043 | 3.71 |
| 5 | 2.66 | 1.37–5.15 | 0.004 | 0.897 | 2.89 |

| Type (compared to Type I) | Overall p value for variable | 0.326 |
|---|---|---|
| II | 1.00 |
| III | 1.18 | 0.69–2.03 | 0.53 | 0.327 | 0.6 |
| Uncertain | 1.50 | 0.80–2.81 | 0.209 | 0.482 | 1.26 |

| Year (compared to 2010) | Overall p value for variable | 0.063 |
|---|---|---|
| 2010 | 1.00 |
| 2002 | 1.52 | 0.51–4.51 | 0.453 | 0.843 | 0.75 |
| 2004 | 1.57 | 0.66–3.75 | 0.312 | 0.697 | 0.10 |
| 2005 | 1.68 | 0.93–3.03 | 0.087 | 0.505 | 1.71 |
| 2006 | 1.49 | 0.87–2.57 | 0.149 | 0.414 | 1.44 |
| 2007 | 1.51 | 0.92–2.49 | 0.105 | 0.384 | 1.62 |
| 2009 | 0.73 | 0.41–1.22 | 0.209 | 0.196 | −1.26 |
| 2010 | 0.10 | 0.06–1.63 | 0.948 | 0.244 | 0.07 |
| 2011 | 0.00 | 0.61–1.63 | 0.997 | 0.250 | 0.0 |
| 2012 | 1.05 | 0.62–1.78 | 0.846 | 0.281 | 0.19 |
| 2013 | 1.26 | 0.75–2.13 | 0.383 | 0.336 | 0.87 |
| 2014 | 1.27 | 0.72–2.25 | 0.413 | 0.370 | 0.82 |

| Surgery- residual disease (compared to 0 mm) | Overall p value for variable | 0.001 |
|---|---|---|
| Zero | 1.00 |
| >0 mm and ≤10 mm | 1.58 | 1.05–2.37 | 0.027 | 0.328 | 2.21 |
| >10 mm and ≤20 mm | 2.67 | 1.71–4.16 | −0.001 | 0.604 | 4.34 |
| >20 mm or no surgery | 2.03 | 0.91–4.50 | 0.083 | 0.825 | 1.73 |

| Primary treatment (compared to Neo adjuvant chemotherapy and interval debulking surgery) | Overall p value for variable | 0.001 |
|---|---|---|
| Neo adjuvant chemotherapy and interval debulking surgery | 1.00 |
| Primary surgery plus adjuvant chemotherapy where appropriate | 0.66 | 0.47–0.91 | 0.012 | 0.110 | −2.5 |
| Primary chemotherapy | 1.91 | 0.89–4.12 | 0.097 | 0.749 | 1.66 |
| No treatment with curative intent | 5.99 | 2.69–13.35 | −0.001 | 2.448 | 4.38 |
| Primary surgery and no adjuvant chemotherapy although recommended | 12.41 | 6.33–24.34 | −0.001 | 4.265 | 7.33 |

* Includes Stage I patients where chemotherapy was not recommended.
change in bowel habit) closely matches that in similar studies [22–24]. As previously reported [22], increasing number of symptoms led to a worsening of survival. This is a reflection of increasing disease burden, which increases the clinical complexity, adding to the diagnostic challenge.

Our findings that emergency presentation confers worse survival compared with two-week urgent ‘cancer’ referral are in line with Barclay et al. [25] and Altman et al. [26]. It highlights the need to ensure for fast tracking of referrals to gynaecological services. However, the time interval between initial onset of symptoms and diagnosis did not independently influence survival once other factors such as age, stage and Type were included in the analysis. This aligns with Nagle et al. [9] who found that in symptomatic ovarian cancer reducing time to diagnosis does not alter stage or survival. Although reducing the time to diagnosis is associated with more favourable outcomes in breast, colo-rectal, head and neck, melanoma and testicular cancers [27], the benefit varies between cancers. The lack of benefit in iEOC is probably a reflection of disease biology. Across all stages, larger volumes of residual disease led to a significantly worse survival, which is in keeping with current surgical philosophy of maximal surgical effort to removal all disease [26,28,29].

The introduction of the NICE guidelines did not improve survival in our study. A recent UK based study [8] found that while the introduction of the NICE guidelines did increase in number of case of ovarian cancer being diagnosed, it did not result in a shift in stage at diagnosis.

4.3. Implications

There are two important implications of the findings of this population-based study. The first that the symptom complexes identify women with a poorer prognosis. While it is necessary to continue measures to ensure prompt referral and reduce emergency presentations, it is difficult to assess the impact of these measures on overall survival. To decrease deaths from ovarian cancer, it is critical we remain focussed on understanding disease biology, exploring preventative strategies, refining the current screening strategies by incorporating novel tests and optimising surgical and adjuvant treatment.

The second that the evidence cannot exclude the possibility of better outcomes in those who are aware and act on these symptoms compared to those who do not. A worse survival in women with more symptoms would support this. Definitive proof of this requires further randomised controlled trials like DoVe. Meanwhile, women should be encouraged to seek help as longer intervals to diagnosis and treatment are associated with reduced overall quality of life and decreased patient satisfaction [30].

5. Conclusions

Symptoms awareness using the NICE Guidance 2011 and modified Goff Index and other similar [5,31] guidance has been widely adopted as a method to identify ovarian cancer earlier. Women with these symptom complexes are likely to have advanced disease and poorer survival. The lack of significant impact on survival of healthcare interventions such as route or interval to diagnosis or secondary care team involved in initial management suggests that in invasive epithelial tubo-ovarian cancer, tumour biology is the overriding driver of survival.

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CRedIT authorship contribution statement

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Declaration of competing interest

UM has stocks in Abcodia awarded to her by UCL. IJ and SJS are co-inventors of the Risk of Ovarian Cancer Algorithm (ROCA) which has been licensed to Abcodia Ltd. by Massachusetts General Hospital (MGH) and Queen Mary University of London (QMUL). IJ has a financial interest in Abcodia Ltd. as a shareholder and director and is entitled to a royalty payments via MGH and QMUL from any commercial use of the ROCA. SJS co-developed the ROCA with all rights assigned to MGH and QMUL. SJS receives personal fees from the LUNGevity Foundation and SISCAPA-Technologies as member of their Scientific Advisory Boards and from Abcodia as a consultant. All other authors declare no competing interests.

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Disclaimer

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Paper presentation

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