- 1 An umbrella review of systematic reviews on the impact of the COVID-19 pandemic on cancer
- 2 prevention and management, and patient needs

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## ABSTRACT

24	The COVID-19 pandemic led to relocation and reconstruction of health care resources and systems,
25	and to a decrease in healthcare utilization, and this may have affected the treatment, diagnosis,
26	prognosis, and psychosocial well-being of patients with cancer. We aimed to summarize and quantify
27	the evidence on the impact of the COVID-19 pandemic on the full spectrum of cancer care. An
28	umbrella review was undertaken to summarize and quantify the findings from systematic reviews on
29	impact of the COVID-19 pandemic on cancer treatment modification, delays, and cancellations;
30	delays or cancellations in screening and diagnosis; psychosocial well-being, financial distress, and use
31	of telemedicine as well as on other aspects of cancer care. PubMed and WHO COVID-19 Database
32	was searched for relevant systematic reviews with or without meta-analysis published before
33	November 29th, 2022. Abstract, full text screening and data extraction were performed by two
34	independent reviewers. AMSTAR-2 was used for critical appraisal of included systematic reviews. 51
35	systematic reviews evaluating different aspects of cancer care were included in our analysis. Most
36	reviews were based on observational studies judged to be at medium and high risk of bias. Only 2 of
37	the included reviews had high or moderate scores based on AMSTAR-2. Findings suggest treatment
38	modifications in cancer care during the pandemic versus the pre-pandemic period were based on low
39	level of evidence. Different degrees of delays and cancellations in cancer treatment, screening and
40	diagnosis were observed, with low-and-middle income countries and countries that implemented
41	lockdowns being disproportionally affected. A shift from in-person appointments to telemedicine use
42	was observed, but utility of telemedicine, challenges in implementation and cost-effectiveness in
43	different areas of cancer care were little explored. Evidence was consistent in suggesting psychosocial
44	well-being (e.g., depression, anxiety, and social activities) of patients with cancer deteriorated, and
45	cancer patients experienced financial distress, albeit results were in general not compared to pre-
46	pandemic levels. Impact of cancer care disruption during the pandemic on cancer prognosis was little
47	explored. In conclusion, Substantial but heterogenous impact of COVID-19 pandemic on cancer care
48	has been observed. Evidence gaps exist on this topic, with mid- and long-term impact on cancer care
49	being most uncertain.

## INTRODUCTION

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51	The coronavirus disease 2019 (COVID-19) pandemic and the mitigation measures that were
52	undertaken posed major challenges to cancer care. The rapid spread of COVID-19 and early data
53	showing patients with cancer were at increased risk of morbidity and mortality after Severe Acute
54	Respiratory Syndrome Coronavirus 2 (SARS-CoV-2) infection, prompted changes in healthcare
55	delivery <sup>1</sup> . These changes included reduction of medical activities, reallocation of healthcare workers,
56	shifting in-person appointments to remote consultations, and limiting access of patients to care
57	facilities <sup>2</sup> .
58	Concerns have been raised that disruption of health care services might have had multidimensional
59	impact in cancer care. Indeed, several studies have described delays and cancellation in treatment,
50	screening, and diagnosis <sup>3-5</sup> . For example, two meta-analyses showed that during the pandemic there
51	was a $\sim 50\%$ reduction in breast and cervical cancer screening, and that there was 18.7% reduction for
52	all cancer treatments, with surgical treatment showing the highest reduction <sup>3 4</sup> . In addition, several
53	studies have highlighted deterioration of psychological well-being of patients with cancer, and
64	psychological, ethical, spiritual, and financial needs of patients with cancer were also affected <sup>67</sup> .
65	While several systematic reviews have examined the impact of COVID-19 on cancer care, they
66	evaluated different outcomes and periods of the pandemic, and thus the available review findings are
67	rather fragmented <sup>3 4 8-14</sup> . A comprehensive review of impact of COVID-19 on several aspects of
68	cancer would be essential to understand gaps and scale-up evidence-based interventions, including
69	learning lessons for future pandemics. In addition, although systematic reviews are important for
70	public health and policy decision-making during the pandemic, the level of methodological rigor they
71	implemented is unclear.
72	In the current study we performed an umbrella review of systematic reviews to summarize the impact
73	of COVID-19 on several aspects of cancer care, including treatment, diagnosis, financial,
74	psychological and social dimensions. We assessed the amount and geographical breadth of the
75	available evidence and methodological rigor of the primary studies included in each review (as
76	assessed by the reviewers) and of the systematic reviews themselves; and summarized the conclusions
77	from different reviews on COVID-19 impact.
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79	RESULTS

Our search strategy identified 1172 citations. Based on title and abstract screening, we retrieved full texts of 96 articles for further screening. Of those, 45 articles did not meet our eligibility criteria, thus leaving 51 articles to be included in our final analysis. **Figure 1** summarizes our screening procedure.

No additional study was found from screening of references of the included studies.

84	Characteristics of the included systematic reviews
85	Of the 51 included systematic reviews, 14 articles also included a quantitative analysis/meta-analysis
86	with one being individual participant meta-analysis. 2-47 Other key characteristics of the 51 systematic
87	reviews are shown in Table 1 (more extensive details appear in Supplementary File 1a and
88	Supplementary File 2). The median number of bibliographic databases/data sources that were
89	searched was 3; the most searched databases were PubMed (n=35), Medline (n=25), Embase (n=22),
90	Scopus (n=19), Web of Science (n=13) and The Cumulative Index to Nursing and Allied Health
91	Literature- CINAHL database (n=10). One review searched for mobile applications using the iOS
92	App Store and Android Google Play <sup>20</sup> . The median number of studies included in the systematic
93	reviews was 31 (interquartile range, 15; 51). The type of study designs included across reviews
94	varied, but most reviews included data from observational study designs of cross-sectional and
95	retrospective nature. Twenty-one reviews focused/reported exclusively on studies that include pre-
96	pandemic controls. Twenty reviews provided data only on site-specific cancers, while the rest for any
97	cancer-site with or without data on site-specific cancers. Nineteen reviews assessed only one aspect of
98	cancer care, while the rest examined two or more of our pre-defined outcomes. The date of last search
99	varied from April 2020 to May 2022, with 16 reviews ending searches during 2020, 25 during 2021
100	and 5 during 2022; 4 reviews did not provide information on date of last search.
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102	Geographical distribution
103	Out of 51 reviews, 46 provided some information on geographical distribution of the included primary
104	studies. Of those, most reviews provided data from different countries, while only two studies (3.9%)
105	focused on data from India <sup>26</sup> and Italy <sup>32</sup> exclusively. Also the majority of the evidence was derived
106	from high- and middle-income countries.
107	Risk of bias of primary studies included in the systematic reviews and GRADE assessments
108	Of the 51 reviews, 32 assessed risk of bias of the included studies (Table 2 and details in
109	Supplementary File 1b). Thirteen different risks of bias checklists were used, and the most common
110	checklists used to assess methodological rigor were Newcastle-Ottawa Scale (NOS) (n=10) and
111	Joanna Briggs Institute tools (n=7). Of the systematic reviews that assess methodological rigor of the
112	individual studies, 8 concluded strong evidence, 19 mixed evidence, 3 weak evidence and 2 did not
<ul><li>112</li><li>113</li></ul>	individual studies, 8 concluded strong evidence, 19 mixed evidence, 3 weak evidence and 2 did not provide any results. Excluding the NOS assessments (since NOS has been criticized to not provide
113	provide any results. Excluding the NOS assessments (since NOS has been criticized to not provide
113 114	provide any results. Excluding the NOS assessments (since NOS has been criticized to not provide accurate assessment of methodological rigor <sup>48</sup> ), the respective numbers were 3, 14, 3, and 2. Only two

Methodological rigor of included systematic reviews

11	8 <b>Table 3</b> shows the AMSTAR-2 evaluations for the included systematic reviews. Only two reviews
11	9 scored moderate to high quality, while the rest were evaluated as low or critically low quality due to
12	not meeting one or more of the seven domains considered critical. Most of the studies did not provide
12	the list of excluded studies during the full text screening, and did not account for methodological rigor
12	2 of included studies when interpreting/discussing the results of the reviews.
12	3 Results and conclusions of systematic reviews and of meta-analyses
12	The main results and conclusions of the eligible systematic reviews are presented in <b>Supplementary</b>
12	files 1c-1j for various aspects of cancer care. Table 4 lists the effect sizes and confidence intervals for
12	the systematic reviews that used formal meta-analysis as well as heterogeneity metrics. Figure 2
12	7 provides a summary of main findings of this umbrella review. Here, we present some key findings for
12	8 each type of outcome:
12	9 Modification of treatment
13	There were 15 reviews assessing modification of treatment <sup>5 9 10 15 16 18 23 28 31 34 37 49-52</sup> . Main findings for
13	each individual review are outlined in <b>Supplementary File 1c and Table 4</b> . All reviews were
13	consistent reporting changes in treatment, with downscaling treatments plans in patients with cancer
13	being a significant intervention. Di Cosimo S et al. 2022 reported changes in treatment plans in 65%
13	4 (95%CI, 53%-75%; I <sup>2</sup> , 98%) of centers <sup>31</sup> . Guidelines recommended use of non-surgical treatment over
13	surgical treatments, as it was seen in head and neck cancer management. However, reviews suggested
13	patients being assessed in a case-by-case basis and that individual factors should be considered for
13	7 individualized treatment ( <b>Supplementary File 1c</b> ). Garg PK et al. 2020 found that available
13	guidelines were based on low level of evidence and had significant discordance for the role and
13	9 timing of surgery, especially in early tumors <sup>18</sup> .
14	0 Delayed and/or cancelled treatment
14	Supplementary File 1d and Table 4 summarize the main findings from the 15 reviewes <sup>2 4 5 13 14 19 25 29</sup>
14	2 31 33 35 37 41 47 52 that assessed and reported on treatment delays and cancellations of cancer treatment.
14	3 Most reviews mentioned that cancellations of treatment were observed, although to what extend this
14	4 happened was not consistently provided <sup>19 25 29 31 33 37 41</sup> . However, reviews reported that these
14	reductions were more pronounced during a lockdown. In the meta-analysis by Teglia F et al., 2022, it
14	6 was found an overall reduction of -18.7% (95% CI, -13.3 to -24.1) in the total number of cancer
14	7 treatments administered during January-October 2020 compared to the previous periods, with surgical
14	8 treatment having a larger decrease compared to medical treatment (-33.9% versus -12.6%); among
14	9 cancers, the largest decrease was observed for skin cancer (-34.7% [95% CI, -22.5 to -46.8]) <sup>4</sup> . This
15	0 difference would depend on the period, with the review reporting a U-shape for the period January-
15	October 2020. Lignou S et al. 2022 <sup>35</sup> reported that between 18 <sup>th</sup> to 31 <sup>st</sup> of January 2021, pediatric and

- noncancer surgical activities were occurring at less than a third of the rate of the previous year, while
- Di Cosimo SD et al. 2022<sup>31</sup> reported cancellation/delays of treatment in 58% (95%CI, 48%-67%; 1<sup>2</sup>,
- 154 98%) of centers. Majeed A et al., 2022<sup>14</sup> showed that shortage of treatment and delays and
- interruptions to cancer therapies in general were more common in low- and middle-income countries.
- 156 Delayed and/or cancelled screening
- The results of 11 reviews <sup>3 30 32-34 36 38 39 43 46 53</sup> reporting on cancer screening are summarized in
- Supplementary File 1e and Table 4. Of these, 5 included a meta-analysis. Overall, reviews showed
- a decline in screening rates across all cancer types, and that differences by demographic area and time
- periods were observed; for instance, countries that implemented lockdowns showed a higher decline
- in screening rates. Within colorectal and gastric cancers, most reviews reported a reduction of at least
- 50% in number of endoscopies and gastroscopies compared to previous years. In the meta-analysis by
- Teglia F et al<sup>3</sup>., while colorectal screening on average was reduced by 44.9% (95% CI, -53.8% to -
- 164 36.1%) during January-October 2020, a U-shape association was observed. Within women-specific
- cancers, the meta-analyses showed a decrease in breast and cervical cancers screening rates of at least
- 40-50%. A meta-analysis focused on cytopathology practice showed that on average there was a
- sample volume reduction of 45.3% (range, 0.1%-98.0%), although the results would depend on the
- tissue sampled<sup>46</sup>. Similar findings were reported by Alkatoul et al. 2021<sup>30</sup>.
- 169 Reduced cancer diagnosis
- Main findings of the 11 reviews 5 14 30 32-35 37 39 46 51 providing data on reduction in cancer diagnosis are
- 171 provided in **Supplementary File 1f** and **Table 4**. Reviews were consistent in reporting decreased
- diagnosis of new cancer cases during the pandemic, although the reduction depended on the
- geographical area, the period being investigated and type of cancer. For example, there was a 73.4%
- decrease in cervical cancer diagnoses in Portugal during 2020, and in Italy, while there was up to 62%
- reduced diagnosis of colorectal cancer in 2020 compared to pre-pandemic years, the reduction was
- more pronounced in Northern Italy where strict lockdowns were implemented. Indeed, reviews
- 177 showed that countries that implemented lockdowns measures showed the highest reduction in number
- of new cancer cases being diagnosed. Breast cancer diagnosis rates dropped by an estimate between
- 179 18-29% between 2019 and 2021<sup>39</sup>.
- 180 Reduced uptake of HPV vaccination
- 181 There was only one review to summarize data on HPV vaccination, showing up to 96% reduction in
- number of vaccine doses administered in March-May 2020 among adolescents and young girls aged
- 9-26 years; the one- year period reduction reported was much smaller (13%)<sup>33</sup>.
- 184 Psychological needs/distress

185	Thirteen reviews covered topics related to psychological needs and distress that patients with cancer
186	experienced during the pandemic <sup>2 5-7 11 17 19 21 24 26 29 34 52</sup> ; the findings are summarized in
187	Supplementary File 1f and Table 4. Reviews reported that the pandemic negatively impacted the
188	psychosocial and physical wellbeing of cancer survivors and patients with cancer experienced
189	different levels of anxiety, depression, and insomnia. In a meta-analysis, Ayubi E et al. 2021 reported
190	an overall prevalence of depression and anxiety of 37% (95%CI, 27-47, I <sup>2</sup> , 99.05) and 38% (95%CI,
191	31-46%, I <sup>2</sup> , 99.08) in patients with cancer, respectively <sup>17</sup> . Similar findings were reported by Zhang et
192	al. 2022 <sup>6</sup> . Compared to controls, patients with cancer had higher anxiety level [standard mean
193	difference (SMD 0.25 (95% CI, 0.08, 0.42)] <sup>17</sup> .
194	Telemedicine
195	Telehealth was investigated and reported in 12 of the included reviews <sup>2 10 12 16 20 22 27 29 31 35 51 54</sup> ; a
196	summary of main findings is provided in <b>Supplementary File 1h</b> . Salehi F et al. 2022 <sup>27</sup> reported that
197	telemedicine use in breast cancer patients was the most common investigated in studies exploring
198	cancer-specific use of telemedicine. Telemedicine was used for various reasons, with provision of
199	virtual visit services and consultation being the most common <sup>27</sup> . One study explored various symptom
200	tracking apps for patients with cancer, available in the mobile health market, and found that only a
201	limited number of apps exist for cancer-specific symptom tracking (27%) <sup>20</sup> . In addition, of the 41
202	apps found, only one was tested in a clinical trial for usability among patients with cancer <sup>20</sup> . While
203	little research exists on how patients perceived telemedicine during the COVID-19 pandemic, early
204	data showed that majority of patients found telemedicine service helpful and that obtaining a
205	telemedicine service helped solve their health problem. Nevertheless, there were concerns that use of
206	telehealth for people with cancer suggests a greater proportion of missed diagnoses <sup>35</sup> , and that
207	telemedicine cannot be a substitute for face-to-face appointments <sup>22</sup> .
208	Financial distress and Social isolation
209	Five reviews reported the economic impact of COVID-19 and social isolation of patients with cancer
210	during the pandemic (Supplementary File 1i) <sup>27111952</sup> . While there is little research on this topic,
211	overall, the reviews suggested financial distress with direct and indirect costs burden and social
212	isolation being a common issue for patients with cancer. Reviews also were consistent in reporting
213	social isolation and loneliness among patients with cancer. Several factors contributed to social
214	isolation, including fear of infection, social distancing measures, not having visitors and lack of social
215	interaction during treatment.
216	Tobacco use and cessation
217	There was only one systematic review and meta-analysis to explore tobacco use and cessation during
218	the pandemic <sup>42</sup> . Based on data from 31 studies, Sarich P et al. 2022 found that, compared to pre-

pandemic period, the proportion of people smoking during the pandemic was lower (pooled prevalence ratio of 0·87 (95%CI:0·79-0·97). In addition, there was similar proportions among smokers before pandemic who smoked more or smoked less during the pandemic, and on average 4% (95%CI: 1-9%) reported stopping smoking. 2% reported starting smoking during the pandemic. High heterogeneity was observed across the meta-analyses results.

Other aspects of cancer care

Eighteen reviews<sup>8-10 13-16 23 25 26 31 35 40 44 45 47 51</sup> reported on mitigations strategies and cancer service restructuring, impact of measures on cancer prognosis, and on quality of recommendations provided during COVID-19 for cancer care; findings are summarized in Supplementary File 1j. In the metaanalysis by Di Cosimo S et al., routine use of PPE by patient and healthcare personnel was reported by 81% and 80% of centers, respectively; systematic SARS-CoV-2 screening by nasopharyngeal swabs was reported by only 41% of centers<sup>31</sup>. Five reviews also reported on potential impact of mitigation strategies on cancer outcomes/prognosis<sup>30 35 40 44 47</sup>. It was estimated that 59,204–63,229 years of life lost might be attributable to delays in cancer diagnosis alone because of the first COVID-19 lockdown in the UK, albeit the findings were based on single study. Delayed cancer screening was estimated to cause globally the following additional numbers of cancer deaths secondary to breast, esophageal, lung, and colorectal cancer, respectively: 54,112-65,756, 31,556-32,644, 86,214-95,195, and 143,081–155,238<sup>30</sup>. Tang et al. 2022<sup>44</sup> de Bock et al. 2022<sup>47</sup> found no deterioration in the surgical outcomes of all types of cancer or colorectal cancer surgery: also no reduction in the quality of cancer removal was observed. Similar findings were also reported by Pararas N et al. 2022<sup>40</sup>, despite the number of patients presenting with metastases during the pandemic was significantly increased. Thomson JD et al. 2020<sup>45</sup>, by exploring recommendations for hypofractionated radiation therapy, found that in general the recommendations during the pandemic were based on lower quality of evidence than the highest quality routinely used dose fractionation schedules.

DISCUSSION

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The current umbrella review summarized and appraised systematically the evidence on the extent to which several aspects of cancer care were disrupted during the COVID-19 pandemic. The summary message provided by 51 systematic reviews is that there have been modifications, delays and cancellation of treatment, delays and cancellation in cancer screening and diagnosis, and patients with cancer may have experienced additional psychological, social, and financial distress. Nevertheless, appraisal of the impact of COVID-19 on cancer care is mainly based on limited and low-quality evidence, and that data mainly derive from high-income countries, with little understanding of consequences of COVID-19 on cancer care in low- and- middle income countries. In addition, limited

evidence exists on whether disruptions in cancer care during the pandemic had adverse impact in prognosis of patients with cancer and mortality.

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Several guidelines were provided for cancer care during the pandemic, including recommendations on mitigation strategies to prevent SARS-CoV-2 infection and cancer treatment modalities. Nevertheless, most recommendations were based on expert opinions, and little quantitative evidence was provided to support them. This aspect was highlighted also in the systematic review by Thomson JD et al. 2020<sup>45</sup>. The authors explored recommendations for hypofranctionated radiation therapy before and during pandemic and found that during the pandemic there was a significant shift from established higher-quality evidence to lower-quality evidence and expert opinions for the recommended hypofractionated radiation schedules. Similar findings were reported also by Garg PK et al. 2020<sup>18</sup>, suggesting not only guidelines were based on low level of evidence, but also there was significant discordance for the role and timing of surgery, especially in early tumors.

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Specific recommendations established from the guidelines such as prioritization of high-grade malignancy, as well as other aspects such as lockdowns, social restrictions, restructure of cancer care with prioritization of high-risk malignancies and use of telemedicine, fear of infection, financial distress and shortage in medications could explain the delays and cancellation in cancer treatment, screening and diagnosis reported in several studies. For example, Lignou S et al. 202<sup>35</sup> raised concerns that use of telehealth for people with cancer suggests a greater proportion of missed diagnoses. Most of examined systematic reviews reported a substantial reduction in treatment, screening, and diagnosis of several cancers during the pandemic, which was more pronounced for countries that implemented a lockdown. In addition, differences were observed by geographical area, suggesting that the impact on cancer treatment, screening and diagnosis could depend on mitigation strategies countries implemented as well as on country-specific health care organization and resources. For example, shortage of treatment and delays and interruptions to cancer therapies in general were more pronounced in low- and middle-income countries<sup>14</sup>. The findings on disruption of cancer treatment, screening and diagnosis are in line with findings reported for other chronic diseases, such as cardiovascular disease<sup>55</sup>, suggesting the adverse impact might not be cancer specific. Future research should explore and compare how different chronic diseases were impacted. Evidence is limited on evaluating how disruption of cancer care during COVID-19 affected prognosis of patients with cancer. Limited evidence showed that the number of patients presenting with metastases during the pandemic was significantly increased, and emergency presentations and palliative surgeries were more frequent during the pandemic<sup>40</sup>. No deterioration in the surgical

outcomes of colorectal cancer surgery including mortality or reduction in the quality of cancer

removal was observed<sup>40 44</sup>. A study<sup>56</sup> in UK estimated that 59,204–63,229 years of life lost might be attributable to delays in cancer diagnosis alone because of the first COVID-19 lockdown, but estimates were based on modelling. Several studies<sup>57 58</sup> have shown a decline in elective cancer such as colorectal cancer, despite findings showing that gastrointestinal cancer surgery during pandemic is safe with appropriate isolation measures and no delays should be implemented for both early and advanced cancer<sup>59</sup>. A recent meta-analysis<sup>60</sup> showed that delaying colorectal cancer longer than 4 weeks could be associated with poorer outcomes.

Several studies and systematic reviews thereof have investigated the impact of the pandemic on psychological wellbeing, financial distress, and social isolation of patients with cancer, as well as the role of telemedicine in cancer care. While studies suggested depression, anxiety, post traumatic disorder, insomnia and fear of cancer progression being highly reported by cancer patients with estimates reaching beyond 50%, high heterogeneity was observed, and in general systemic analysis comparing the findings with pre-pandemic period rates was lacking. The pandemic was reported to have financial burden on cancer patients with direct and indirect costs. Social isolation was commonly reported and mainly driven by fear of infection, social distancing measures and lack of social interaction during treatment. Nevertheless, there was limited effort to quantify social isolation and economic impact on cancer care. Telemedicine and remote consultations were sharply increased in use for different aspects of cancer care, including treatment, screening, and rehabilitation.

However, evidence is limited in evaluating and quantifying the positive and negative impact, as well as cost-effectiveness of telemedicine. While limited evidence suggested telemedicine reduced costs of cancer care for both patients and health care provider, there were concerns especially from patients that telemedicine could not have similar benefits to on-site consultations.

Our study has certain limitations. Although our search was based on recent recommendations on optimal databases needed to be searched for umbrella reviews<sup>61</sup>, we cannot rule out missing some other relevant systematic reviews. Most systematic reviews included in this umbrella review were based on intermediate and high risk of bias studies, and the findings were mainly based on case-series, cross-sectional and retrospective observational study designs which are prone to residual confounding and poor in determining temporal associations. Prevalence and incidence estimates are also subject to selection biases. In some instances, data were derived from one study or from studies with small sample sizes and limited number of events, leading to large uncertainty. Many studies did not include any pre-pandemic controls. Furthermore, some of the evidence overlapped among the systematic reviews that were included in this umbrella review, but this allows comparing notes on results and conclusions for the overlapping efforts. Some systematic reviews were published early (in 2020), and

323 thus they had even more limited evidence and the impact of the disruptions may have differed across 324 different pandemic waves. Most findings were derived from high-income and/or western countries, 325 limiting the generalizability of the findings to low- and middle-income countries. Lastly, concreate 326 conclusions on intermediate, and long-term impact remain unclear. Finally, the suboptimal 327 methodological rigor of many included reviews is notable. 328 In summary, evidence shows a diverse and substantial impact of the COVID-19 pandemic on cancer 329 care, including delays in treatment, screening and diagnosis. Also, patients with cancer had been 330 affected psychologically, socially, and financially during the COVID-19 crisis. However, large 331 uncertainty and gaps exist in the literature on this topic. Most of the evidence on the topic is derived 332 mainly from high and middle-income countries, and low-quality studies, and thus, future high-quality 333 studies with larger geographical capture and properly performed, rigorous systematic reviews with 334 careful meta-analyses will continue to have value in this field. 335 **MATERIALS and METHODS** We performed an umbrella review following the recent published guideline<sup>62</sup>, and for reporting we 336 adhered to the Preferred Reporting Items for Overviews of Reviews- PRIOR checklist<sup>63</sup> 337 338 (Supplementary File 1k). The protocol has been registered with the Open Science Framework 339 (https://osf.io/qjgxv) 340 341 **Search Strategy** 342 Literature search was performed in PubMed and WHO COVID-19 Database using the search strategy 343 in Supplementary File 11. No language restriction was applied. We searched for studies published until November 3<sup>rd</sup>, 2022; an update of the search was performed until November 29<sup>th</sup>, 2022. 344 345 References cited in the final included studies for analysis were further screened to identify other 346 relevant publications. 347 Screening, Study selection and Eligibility criteria 348 Retrieved items were first screened based on the title and abstract and potentially eligible references 349 were then screened in full text. Screening was performed by two reviewers and in case of 350 discrepancies, a final decision to include or exclude was settled with discussion. We included studies 351 if they fulfilled all the following criteria: (i) were systematic reviews with our without meta-analysis 352 or individual participant meta-analysis; (ii) included individuals diagnosed with any type of cancer 353 and at any cancer stages (early to advanced), or individuals targeted for cancer screening; (iii) 354 assessed the impact of the COVID-19 pandemic, and thus had data collected during the pandemic 355 period (2020-2022) (the included studies may nevertheless have used also control pre-pandemic

periods in order to assess the magnitude of change during the pandemic); and assessed any of the following outcomes: delay/cancellation of treatment (overall, and per specific treatment); modification of treatment (overall, and per specific treatment); delayed/cancelled screening (overall and per specific type of screening); reduced diagnoses (overall and per specific diagnosis); psychological needs; ethical needs; social needs; financial burden and distress; social impact/ isolation; psychological distress; use of telehealth/virtual visits and other aspects of cancer care such as impact of the COVID-19 pandemic on prognosis. In addition, irrespective of including patients with cancer, we included reviews that looked at impact of COVID-19 on uptake of HPV vaccination and tobacco use and cessation.

## **Data extraction and Critical appraisal**

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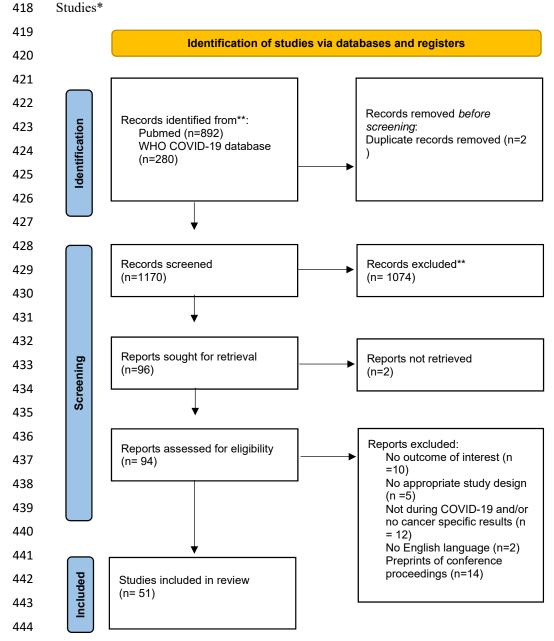
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The data extraction was performed by one of the authors and the extracted data were further checked by two other authors; differences were settled by discussion. In case an eligible article included data from several diseases, when feasible, we extracted information only on cancer-related outcomes of our interest. First, we extracted general information from the eligible reviews, including information on authors, year of publication, type of studies considered (design), number of eligible studies, COVID-19 period covered (until when), whether it has considered studies with pre-pandemic controls (yes exclusively/yes for some/not at all), the outcomes examined and for which cancers each outcome was examined, and methods of analysis and heterogeneity (if provided). To provide the geographical breadth of the evidence, we extracted information on location(s) of the individual studies included in the eligible reviews; for example, retrieving information on countries and areas or whether the studies were done in multiple countries. Concerning the methodological rigor, for each systematic review we extracted information on whether the authors used any previously validated tool or any other set of extracted items to assess the methodological rigor of the included studies. If yes, we recorded the tool used and the main conclusions of the assessment were grouped in the broad categories: most studies were weak in methodological rigor, most studies were strong in methodological rigor, or mixed/ intermediate pattern between the other two categories. Two reviewers assessed methodological rigor of the included systematic reviews using the AMSTAR-2 tool<sup>64</sup>; any discrepancies were settled with the help of a third reviewer. AMSTAR-2 is based on a 16 item or domain checklist, with seven of these items considered critical for the overall validity of a review. The domains considered critical are: (i) protocol registration before starting the review; (ii) adequate and comprehensive search of the literature; (iii) providing justification for the exclusion of individual studies; (iv) risk of bias assessment of the studies included in the review; (v) use of appropriate statistical methods in performing a meta-analysis; (vi) accounting for risk of bias when interpreting the results; (vii) and evaluation of the presence and impact of publication bias. Last, based on abstract and full text reading, we extracted information on main conclusions derived from each of the included reviews. When the

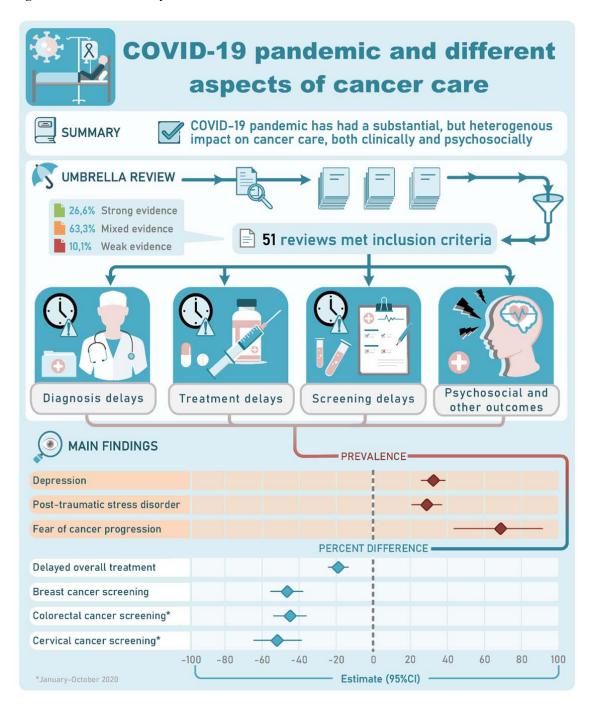
391 review included several disease areas, we extracted information on main findings of the included 392 individual studies within the review that were relevant to cancer. 393 Statistical analysis 394 Due to high heterogeneity in the designs, study questions, outcomes, and metrics, a descriptive 395 analysis was performed. We calculated the proportion of reviews that provided information on single 396 countries and multiple countries. Median and interquartile range were calculated for some of the 397 characteristics of the eligible reviews (e.g., number of databases searched). Separate tables were 398 created for the methodological appraisal of the systematic reviews, the methodological appraisal of 399 the studies in each systematic review, for the characteristics and subject matter information of each 400 systematic review, and for the final conclusions of each systematic review. In addition, we created a 401 separate table for reviews that implemented meta-analysis, providing the summary estimates, 95% 402 confidence intervals and heterogeneity estimates. Limitations and areas of limited evidence were 403 noted. 404 405 Acknowledgement 406 We would like to thank Beatrice Minder for helping with search strategy and Dr. Erand Llanai 407 (Department of Molecular Epidemiology, German Institute of Human Nutrition Potsdam-Rehbruecke, 408 Nuthetal, Germany) for designing and illustrating the graphical abstract. 409 Availability of data and materials: All relevant data are included in the manuscript and 410 supplemental material. Competing interests: TM has found Epistudia, an online platform on evidence synthesis. All other 411 412 authors have no disclosures to report. 413 Funding: No funding was provided for this project 414 Supplementary Files 1a-1L: Table characteristics, main findings, PRISMA and search strategy 415 Supplementary File 2: Bibliographic databases used from each review (see excel file)

**Figure 1**. Flowchart of Identification, Screening, Eligibility, Inclusion, and Exclusion of Retrieved Studies\*



<sup>\*</sup>In the search, we did not include any language restriction filter. However, during full text screening we included only studies that were in English.

\*\*WHO COVID-19 database does not allow to specify the search by both date and month, and the search for this specific database is up to end-December 2022. Any full text (n=0) that was eligible and published after November 29<sup>th</sup>, 2022, was excluded.



Author, year of	Meta-	Number of		Pre-pandemic			Last
publication	analysis	included studies	Countries*	controls	Cancer types	Aspects assessed	search
							15-Jul-
Adham, 2022 <sup>15</sup>	No	5	Globally	No	H&N	MT, O	20
			Multiple countries, including				
			US, TW, BE, NL, JP, IT, UK, AS,				28-Dec-
Alkatoul, 2021 <sup>30</sup>	No	16	CA	Yes	ALL	DCS, RD	20
							1-Sep-
Alom, 2021 <sup>16</sup>	No	72	Multiple counties	No	All	MT, TL, O	20
							3-Jan-
Ayubi, 2021 <sup>17</sup>	Yes	34	Multiple counties	No	All	PSND, O	21
							End of
Azad. 2021 <sup>50</sup>	No	51	Multiple counties	No	Glioma	MT	2020
							01-Apr-
Beterra et al. 2022	No	8	NP	No	ALL	TL	2021
40							08-Apr-
Crosby, 2022 <sup>49</sup>	No	45	NP	No/NS	H&N	MT	2020
						Delayed and/or	
						cancelled treatment	
De Bock et al.							21-Mar-
2022 <sup>47</sup>	Yes	24	Multiple counties	Yes	ALL, BC	Other aspects	2021
			Multiple counties, including IT, US,			DCT, DCS, PSND,	1-Dec-
Dhada, 2021 <sup>2</sup>	No	19	UK, NL	No	ALL	TL, FBD, SIA	20
Di Cosimo,							11-Dec-
2022 <sup>31</sup>	Yes	56	Multiple counties	Yes	ALL	MT, DCT, TL, O	20
			Multiple counties, including CN, IR,				3-Aug-
Donkor, 2021 <sup>8</sup>	No	11	BR, ZA	No	ALL	0	20
							31-Jan-
Fancellu, 2022 <sup>32</sup>	No	7	IT	Yes	CRC	DCS, RD	22

Ferrar, 2022 <sup>33</sup>	No	33	Multiple counties	Yes	CV	DCT, DCS, RD, RHPV	8-Feb- 22
1 011011, 2022	110		Multiple counties, including IN, SL,				15-Dec-
Gadsden, 2022 <sup>13</sup>	No	17	BA	Yes	ALL	DCT, O	21
Garg, 2020 <sup>18</sup>	No	212	Multiple counties	No	ALL	MT	2-May- 20
Gascon, 2020 <sup>9</sup>	No	23	Multiple counties	No	H&N	MT, O	1-May- 20
Hesary, 2022 <sup>34</sup>	No	22	Multiple counties, including IT, UK, PG, NL, CN, IN, JP, TU, IR, SN	Yes	GA	MT, DCS, RD, PSND	31-Dec- 21
Hojaij, 2020 <sup>10</sup>	No	35	Multiple counties	No	H&N,OTO	MT, TL, O	31-Dec- 20
Jammu, 2021 <sup>19</sup>	No		Multiple counties	No	ALL	DCT, PSND, FBD	27-Aug- 20
Kirby, 2022 <sup>7</sup>	No	56	Multiple counties	No	ALL	PSND, FBD, SIA	31-Mar- 21
Legge, 2022 <sup>11</sup>	No	18	Multiple counties	No	ALL	PSND, FBD, SIA	25- May-22
Lignou, 2022 <sup>35</sup>	No	32	Multiple counties	Yes	PC	DCT, RD, TL	1-Aug- 21
Lu, 2021 <sup>20</sup>	No	41**	NP	No	ALL	TL	1-May- 20
Majeed, 2021 <sup>14</sup>	No	60	Multiple counties	Yes, but NS	PC	DCT, RD, TL	3-Nov- 21
Mayo, 2021 <sup>36</sup>	Yes	13	Multiple counties, including IT, AU, TW, US, FR, NL	Yes	ALL	DCT, DCS	10-Feb- 21
Mazidimoradi, 2021 <sup>37</sup>	No	43	Multiple counties	Yes	CRC	MT, DCT, RD	1-Jun- 21
Mazidimoradi, 2022 <sup>38</sup>	No	25	Multiple counties	Yes	CRC	DCS	1-Jun- 21
Moemenimovahe d, 2021 <sup>21</sup>	No	55	Multiple counties	No	ALL	PSND	30-Jun- 21
Mostafaei, 2022 <sup>22</sup>	No	22	Multiple counties	No	ALL	TL	1-Jun- 21

							15-Apr-
Moujaess, 2020 <sup>23</sup>	No	88	Multiple counties	No	ALL	DCT, O	20
Muls, 2022 <sup>24</sup>	No	51	Multiple counties	No	ALL	PSND	1-Oct- 21
							31-Mar-
Murphy A, 2022 <sup>12</sup>	No	37	Multiple counties	No	ALL	TL	21
Ng, 2022 <sup>39</sup>	Yes	31	Multiple counties	Yes	BC	DCS, RD	1-Oct- 20
Nikolopoulos,							10-Feb-
20225	No	15	Multiple counties	Yes, but NS	GC	MT, DCT, RD, PSND	21
Pacheco, 2021 <sup>25</sup>	No	9	Multiple counties, including US, IT, CN, SP, UK, IR	No	ALL	DCT, O	NP
Pararas, 2022 <sup>40</sup>	Yes	10	Multiple counties	Yes	CRC	0	NP
Pascual et al.			Multiple counties from Low- and		Surgical Neuro-		01-Sep-
2021 <sup>51</sup>	No	12	Middle-income countries	Yes, but NS	Oncology	MD, RD, TL, O	2021
Piras et al. 2022 <sup>52</sup>	No	281	Multiple counties	No	ALL	MT, DCT, SIA, PSND	31-Dec- 2021
Riera, 2021 <sup>41</sup>	No	62	Multiple counties	Yes	ALL	DCT	NP
			•				3-Feb-
Rohilla, 2021 <sup>26</sup>	No	6	IN	No	ALL	PSND, O	21
27							1-Apr-
Salehi, 2022 <sup>27</sup>	No	16	Multiple counties	No	ALL	TL	21
Sarich, 2021 <sup>42</sup>	Yes	44	Multiple counties	Yes	NA	RF	5-Nov- 20
Sasidharanpillai,			Multiple counties, including				1-Sep-
2022 <sup>43</sup>	Yes	7	SL, IT, CA, SC, BE, US	Yes	CV	DCT, RD	21
							1-Feb-
Sun P, 2021 <sup>28</sup>	No	6	IT, AM, UK	No	BC	MT	21
44							12-Jan-
Tang, 2022 <sup>44</sup>	Yes	14	TU, CN, UK, IT, DN, AS, AU	Yes	CRC	0	22
T 1: 20223	**	20	Ar to 1	**	DG GDG GU	DOT DD	12-Dec-
Teglia, 2022 <sup>3</sup>	Yes	39	Multiple counties	Yes	BC, CRC, CV	DCT, RD	21
Teglia, 2022 <sup>4</sup>	Yes	47	Multiple counties	Yes	ALL	DCT	12-Dec- 21

								1-Jun-
ď	Thomson, 2020 <sup>45</sup>	Yes	54	NP	Yes	ALL	0	21
								30-Apr-
	Vigliar, 2020 <sup>46</sup>	Yes	41***	Multiple counties	Yes	ALL	DCS, RD	20
	Zapala, 2022 <sup>29</sup>	No	160	NP	No	ALL	DCT, PSND, TL	NP
								31-Jan-
L.	Zhang, 2022 <sup>6</sup>	Yes	40	Multiple counties	No	ALL	PSND	22

AM, America; BC; AS, Austria; AU, Australia; BA, Bangladesh; BC, breast cancer; BE, Belgium; BR, Brazil; CA, Canada; China; CRC, colorectal cancer; CV, cervical cancer; DN, Denmark; FR, France; GA, gastric cancer; GC, gynecological cancer; H&N, head and neck cancer; IN, India; IR, Iran; IT, Italy; JP, Japan; NA, not applicable; NL, Netherlands; NP, not provided; OTO, otorhinolaryngology cancer; PC, pediatric cancer; PG, Portugal; SC, Scotland; SL, Slovenia or Sri Lanka; SN, Singapore; SP, Spain; TU, Turkey; TW, Taiwan; UK, United Kingdom; United States; ZA, Zambia;

MT, modification of treatment; DCT, delayed and/or cancelled treatment; DCS, delayed and cancelled screening; RD, reduced diagnosis: RHPV, reduced uptake of HPV vaccination; TL, telemedicine; PSND, psychological needs/distress; FBD, Financial burden/ distress; SIA, social isolation; O, other aspects \*Multiple countries refer to inclusion of studies for final analysis that used data from more than one country. If complete information on location from all primary studies were provided, then specific countries were listed.

\*\*apps; \*\*\*respondents

Author	Checklist use	Methodological rigor conclusion category	GRADE
Adham M et al. 2022	CEBM	Not provided	Not provided
Alkatoul et al. 2021	NOS	Strong evidence	Not provided
Alom S et al. 2021	NHLBI, NIH	Not provided	Not provided
Ayubi E et al. 2021	Not applied	Not provided	Not provided
Azad MA et al. 2021	Not applied	Not provided	Not provided
Beterra GMF et al. 2022	Not applied	Not provided	Not provided
Cosimo SD et al. 2022	CLARITY	Mixed/Intermediate	Not provided
Crosby DL et al. 2022	Not applied	Not provided	Not provided
De Bock E et al. 2022	ROBINS-I	Strong evidence	Not provided
Dhada S et al. 2021	CASP, NHLBI, NIH	Mixed/Intermediate	Not provided
Donkor et al.	JBI	Weak	Not provided
Fancellu A et al	Not applied	Not provided	Not provided
Ferrar P et al. 2022	NOS	Strong evidence	Not provided
Gadsden T et al. 2022	JBI, ROBINS-I	Mixed/Intermediate	Not provided
Garg PK et al. 2020	Not applied	Not provided	Not provided
Gascon L et al. 2020	Agree II	Mixed/Intermediate	Not provided
Hesary FB et al. 2022	NOS	Mixed/Intermediate	Not provided
Hojaij FC et al.2020	Not applied	Not provided	Not provided
Jammu As et al	Not applied	Not provided	Not provided
Kirby A et al. 2022	JBI, CHEC	Mixed/Intermediate	Not provided
Legge H et al. 2022	MMAT	Strong evidence	Not provided
Lignou S et al. 2022	Not applied	Not provided	Not provided
Lu DJ et al. 2021	MARS	Mixed/Intermediate	Not provided
Majeed A et al. 2021	Not applied	Not provided	Low to moderate certainty
Mayo M et al. 2021	NOS	Mixed/Intermediate	Moderate to high

Mazidimoradi A et al.2021	NOS	Mixed/Intermediate	Not provided
Mazidimoradi A et al.2022	NOS	Strong evidence	Not provided
Moemenimovahed Z et al. 2021	Not applied	Not provided	Not provided
Mostafaei A et al. 2022	JBI	Mixed/Intermediate	Not provided
Moujaess E et al. 2020	Not applied	Not provided	Not provided
Muls A et al. 2022	MMAT	Mixed/Intermediate	Not provided
Murphy A et al. 2022	JBI, CHEC	Mixed/Intermediate	Not provided
Ng JS et al. 2022	NOS	Mixed/Intermediate	Not provided
Nikolopoulos M et al. 2022	NOS	Mixed/Intermediate	Not provided
Pacheco RF et al. 2021	JBI, ROBINS-I	Weak	Not provided
Pararas N et al. 2022	NOS	Strong evidence	Not provided
Pascual JSG et al. 2021	Not applied	Not provided	Not provided
Piras A et al. 2022	Not applied	Not provided	Not provided
Riera R et al. 2021	ROBINS-I	Mixed/Intermediate	Not provided
Rohilla KK et al. 2021	Not applied	Not provided	Not provided
Salehi F et al. 2022	Not applied	Not provided	Not provided
Sarich P et al. 2021	ROBINS-I	Weak evidence	Not provided
Sasidharanpillai S et al. 2022	NHLBI, NIH	Strong evidence	Not provided
Sun P et al. 2021	Not applied	Not provided	Not provided
Tang G et al. 2022	NOS	Strong evidence	Not provided
Teglia F et al. 2022	CASP	Mixed/Intermediate	Not provided
Teglia F et al. 2022	CASP	Mixed/Intermediate	Not provided
Thomson JD et al. 2020	ASTRO	Mixed/Intermediate	Not provided
Vigliar E et al. 2020	Not applicable	Not provided	Not provided
Zapala J et al. 2022	Not applied	Not provided	Not provided
Zhang L et al. 2022	JBI	Mixed/Intermediate	Not provided

CEBM, Critical appraisal tool of qualitative studies from Centre of Evidence-based Medicine (CEBM), University of Oxford; ASTRO, The American Society of Radiation Oncology; CASP, https://casp-uk.net/casp-tools-checklists/; CHEC, Consensus on Health Economic Criteria: CLARITY, "Risk of bias

instrument for cross-sectional surveys of attitudes and practices" from the CLARITY Group at McMaster University"; JBI, Joanna Briggs Institute; MARS, Mobile Apps Rating Scale; MMAT, Mixed Methods Appraisal Tool; NHLBI, NHI, National Institute of Health Checklist; NOS, Newcastle-Ottawa Quality Assessment: RBC, Risk of Bias Checklist for Prevalence Studies by Hoy Damian et al. 2012

476 Table 3. Methodological assessment of the included reviews- AMSTAR 2 evaluation (16 questions)\*

Authors, year of publication	q1	q2	q3	q4	q5	q6	<b>q</b> 7	q8	q9**	q10	q11	q12	q13	q14	q15	q16	Overall Assessment
Adham M et al. 2022	n	n	n	ру	n	n	n	n	у	n	na	na	na	n	na	n	Critical low
AlkatouI et al. 2021	n	ру	у	ру	n	n	n	ру	у	n	na	na	n	n	na	У	Critical Low
Alom S et al., 2021	n	n	n	ру	n	у	n	ру	у	n	na	na	у	n	na	У	Critical Low
Ayubi E et al. 2021	у	n	n	ру	n	n	n	у	n	n	у	n	n	n	у	У	Critical low
Azad MA et al., 2021	n	n	n	ру	у	у	n	у	py	n	у	n	n	n	у	У	Critical low
Beterra GMF et al., 2021	у	n	n	n	n	n	n	у	n	n	na	na	n	n	na	у	Critical low
Crosby DL et al., 2020	n	n	n	n	n	n	n	n	n	n	na	na	na	n	na	у	Critical low
de Bock E et al, 2022	У	n	у	ру	у	у	n	у	у	n	у	n	n	у	n	у	Critical low
Dhada S et al. 2021	n	ру	n	ру	n	n	n	у	у	n	na	na	n	n	na	у	Critical Low
Di Cosimo et al. 2022	n	n	n	ру	у	n	n	у	у	n	у	у	у	у	у	у	Critical low
Donkor et al. 2021	n	n	n	ру	у	у	n	у	у	n	na	na	na	n	na	У	Critical low
Fancellu A et al. 2022	у	n	n	n	n	n	n	n	n	n	na	na	n	n	n	n	Critical low
Ferrara P et al. 2022	n	py	n	ру	у	у	n	n	у	n	na	na	y	n	na	У	Low
Gadsden T et al. 2022	у	ру	n	ру	у	n	n	у	у	n	na	na	y	n	na	У	Low
Garg PK et al. 2020	n	n	n	ру	у	у	n	n	n	n	na	na	n	У	na	У	Critical low
Gascon L et al. 2020	у	у	n	у	у	у	n	na	у	у	na	na	na	n	na	У	Low
Hesary FB et al. 2022	n	ру	n	ру	n	n	n	n	у	n	na	na	n	n	na	У	Critical Low
Hojaij FC et al. 2020	n	n	n	n	n	n	n	n	n	n	na	na	na	n	na	У	Critical low

Jammu AS et al. 2021	n	n	n	ру	у	у	n	n	n	n	na	na	n	n	na	У	Critical low
Kirby A et al. 2022	У	ру	n	у	n	у	n	ру	У	n	na	na	n	n	na	У	Critical Low
Legge H et al. 2022	У	ру	у	ру	у	у	n	у	у	n	na	na	n	n	na	у	Critical Low
Lignou S et al. 2022	У	n	n	n	у	у	n	у	n	n	na	na	n	n	na	У	Critical low
Lu DJ et al. 2021	У	n	na	ру	n	n	n	у	na	n	na	na	na	n	na	У	Critical Low
Majeed A et al. 2022	n	у	n	ру	n	у	n	n	ру	n	na	na	n	n	na	у	Critical Low
Mayo M et al. 2021	n	у	n	ру	у	у	n	n	ру	n	n	у	у	n	n	у	Critical low
Mazidimoradi A et al. 2022	n	ру	n	ру	n	n	n	ру	у	n	na	na	n	n	na	у	Critical Low
Mazidimoradi A et al.2021	n	ру	n	ру	n	n	n	у	у	n	na	na	n	n	na	у	Critical Low
Momenimovahed Z et al. 2021	n	n	n	ру	n	n	n	n	n	n	na	na	n	n	na	у	Critical low
Mostafaei A et al. 2022	n	ру	n	n	n	n	у	ру	у	n	na	na	n	n	na	у	Critical low
Muls A et al. 2022	У	ру	у	ру	n	у	n	у	у	n	na	na	n	n	na	у	Critical Low
Murphy A et al. 2022	n	n	n	у	n	n	n	у	у	n	na	na	n	n	na	у	Critical low
Ng JS et al. 2022	n	ру	n	ру	n	n	n	ру	у	n	у	n	у	у	у	у	Low
Nikolopoulos M et al. 2022	n	ру	n	ру	n	n	n	n	у	n	na	na	n	n	na	у	Critical Low
Pacheco RF et al. 2021	У	у	у	ру	у	у	у	ру	у	у	na	na	у	n	na	у	High quality
Pararas N et al. 2022	n	у	n	у	y	n	n	n	у	n	n	n	n	у	у	у	Critical low
Pascual JSG et al., 2022	У	n	у	ру	у	у	n	у	n	n	na	na	n	у	na	n	Critical low
Piras A et al., 2025	n	n	n	ру	n	n	n	ру	n	n	na	na	n	n	na	у	Critical low
Riera R et al. 2021	n	ру	у	ру	у	у	у	у	у	у	na	na	n	у	na	у	Moderate quality

Rohilla KK et al. 2021	n	n	n	ру	n	у	n	n	n	n	na	na	n	n	na	у	Critical low
Salehi F et ak. 2022	n	n	n	ру	у	n	n	n	n	n	na	na	n	n	na	У	Critical low
Sarich P et al. 2022	у	у	у	ру	у	у	n	у	у	n	У	у	n	У	n	У	Critical low
Sasidharanpillai et al. 2022	n	ру	n	ру	n	n	n	у	У	n	У	У	У	у	у	у	Low
Sun P et al. 2021	n	n	n	ру	n	n	n	n	n	n	na	na	na	n	na	n	Critical low
Tang G et al. 2022	у	n	n	n	n	n	n	n	У	ру	n	n	n	У	n	У	Critical low
Teglia F et al. 2022	у	ру	у	ру	У	У	n	n	У	n	n	n	n	n	у	У	Critical low
Teglia F et al. 2022	у	ру	у	ру	У	У	n	ру	У	n	n	n	n	У	n	У	Critical low
Thomson JD et al. 2020	n	n	n	n	n	n	n	n	У	n	У	n	n	n	na	У	Critical low
Vigliar E et al., 2020**	na																
Zapala J et al. 2022	n	n	n	n	n	n	n	n	n	n	na	na	n	n	na	У	Critical low
Zhang L et al. 2022	У	У	У	ру	n	у	n	ру	у	n	У	У	У	у	y	У	Low

n, no; NA, not applicable; py, partially yes; y, yes 477

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\*The review scored yes if study used a checklist to evaluate methodological rigor, and partial yes if only GRADE assessment was provided without applying 478 a checklist for assessing methodological rigor. \*Individual participant meta-analysis and thus not applicable the AMSTAR evaluation 479

AMSTAR-2 overall assessment rating: high —the review provides an accurate and comprehensive summary of the results of the available studies that addresses the question of interest; moderate—the review has more than one weakness, but no critical flaws. It may provide an accurate summary of the results of the available studies; low—the review has a critical flaw and may not provide an accurate and comprehensive summary of the available studies that address the question of interest; or critically low—the review has more than one critical flaw and should not be relied on to provide an accurate and comprehensive summary of the available studies

O1; Did the research questions and inclusion criteria for the review include the components of PICO?

Q2; Did the report of the review contain an explicit statement that the review methods were established prior to the conduct of the review and did the report 486 justify any significant deviations from the protocol?

- 488 Q3; Did the review authors explain their selection of the study designs for inclusion in the review?
- 489 Q4; Did the review authors use a comprehensive literature search strategy?
- 490 Q5; Did the review authors perform study selection in duplicate?
- 491 Q6; Did the review authors perform data extraction in duplicate?
- 492 Q7; Did the review authors provide a list of excluded studies and justify the exclusions?
- 493 Q8; Did the review authors describe the included studies in adequate detail?
- 494 Q9; Did the review authors use a satisfactory technique for assessing the risk of bias (RoB) in individual studies that were included in the review?
- 495 Q10; Did the review authors report on the sources of funding for the studies included in the review?
- 496 Q11; If meta-analysis was performed did the review authors use appropriate methods for statistical combination of results?
- 497 Q12; If meta-analysis was performed, did the review authors assess the potential impact of RoB in individual studies on the results of the meta-analysis or
- 498 other evidence synthesis?
- 499 Q13; Did the review authors account for RoB in individual studies when interpreting/ discussing the results of the review?
- Q14; Did the review authors provide a satisfactory explanation for, and discussion of, any heterogeneity observed in the results of the review?
- Q15; If they performed quantitative synthesis did the review authors carry out an adequate investigation of publication bias (small study bias) and discuss its
- likely impact on the results of the review?
- Q16; Did the review authors report any potential sources of conflict of interest, including any funding they received for conducting the review?

Author	No. of studies	Outcome	Estima te	LCI	UCI	$\mathbf{I}^2$	P- heterogeniet y	Metri c
Ayubi et al. 2021	15	Depression	0.37	0.27	0.47	99	< 0.001	Prev*
	17	Anxiety	0.38	0.31	0.46	99	< 0.001	Prev*
	4	Anxiety	0.25	0.08	0.42	68	0.02	SMD *
Zhang et al.2022	28	Depression	0.325	0.26	0.39	99	<0.001	Prev*
		Anxiety	0.313	0.25	0.37 5	99		Prev*
	8	PTSD	0.288	0.20 7	8	99	< 0.001	Prev*
	5	Distress	0.539		9	67	0.016	Prev*
	5	Insomia	0.232		0.29	91	<0.001	Prev*
	3	Fear of cancer progression	0.674	0.43	0.91	93	<0.001	Prev*
Cosimo et al. 2022	28	Cancellation/delay of treatment	0.58	0.48	0.67	98	<0.01	Prop*
	14	Modification of treatment	0.65	0.53	0.75	98	< 0.01	Prop* a
	10	Delay of clinic visits	0.75	0.49	0.95	99	< 0.01	Prop* a
	14	Reduction in activity	0.58	0.47	0.68	93	< 0.01	Prop* a
	25	Use of remote consultation	0.72	0.59	0.84	99	< 0.01	Prop* a
	7	Routine use of PPE (patients)	0.81	0.75	0.95	96	< 0.01	Prop* a
	16	Routine use of PPE (workers)	0.8	0.61	0.94	99	< 0.01	Prop* a

	18	Routine screening SARA-CoV-2 swab	0.41	0.3	0.53	96	< 0.01		Prop* a
	5	≥T2 stage during the COVID-19 pandemic compared to the prepandemic control group	1.00	0.72	1.38	58	0.05		OR**
	4	≥T3 stage during the COVID-19 pandemic compared to the prepandemic control group	0.95	0.69	1.32	39	0.18		OR**
De Beck et al. 2022	5	≥N1 stage during the COVID-19 pandemic compared to the prepandemic control group	1.55	0.87	2.74	3	0.39		OR**
Mayo et al. 2021	6	Screening breast cancer	0.63	0.53	0.77	10 0	< 0.001		IRR**
	5	Screening conlonc cancer	0.11	0.05	0.24	10 0	< 0.001		IRR**
	3	Screening cervical cancer	0.1	0.04	0.24	10 0	< 0.001		IRR**
Ng et al. 2022	3	Screening breast cancer rigistry-based study	0.59	0.46	0.7	10 0	< 0.001		RR**
	10	Screening breast cancer non rigistry-based study	0.47	0.38	0.58	10 0	<0.001		RR**
	4	Diagnosis breast cancer registry-based study	0.82	0.63	1.06	99	< 0.001		RR**
	18	Diagnosis breast cancer non-registry-based study	0.71	0.63	0.8	92	< 0.001		RR**
Praras et al. 2022	5	Tis-T1 stage	1.14	0.87	1.48	41		0.15	OR**
	5	T2 stage	0.91	0.78	1.06	0		0.6	OR**
	5	T3 stage	1.18	0.82	1.7	88	< 0.001		OR**
	6	T4 stage	1.19	0.79	1.8	80	< 0.001		OR**
	6	N+ stage	1	0.89	1.11	0		0.54	OR**
	6	M+ stage	1.65	1.02	2.67	91	< 0.001		OR**
	7	Right-sided tumors	0.88	0.51	1.52	99	< 0.001		OR**
	7	Left-sided tumors	0.91	0.56	1.5	96	< 0.001		OR**

	8	Rectal tumors	0.93	0.63	1.37	95	< 0.001	OR**
	3	Emergency presantations	1.74	1.07	2.84	95	< 0.001	OR**
	3	Complicated tumor	1.72	0.78	3.78	82	0.004	OR**
	3	Neoadjuvant therapy	1.22	1.09	1.37	0	0.4	OR**
	4	Palliative internt surgery	1.95	1.13	3.36	54	0.09	OR**
	6	Minimally invasive surgery	0.68	0.37	1.24	98	< 0.001	OR**
	5	Stoma formation	0.91	0.51	1.62	94	< 0.001	OR**
	2	Morbidity	0.92	0.55	1.55	25	0.25	OR**
	3	Leng of hospital stay	0.51	-0.93	1.94	79	0.008	
	3	Lymph node harvest	1.57	-1.99	5.13	64	0.06	WMD **
Sarich et al. 2022	12	Smoking prevalence	0.87	0.79	0.97	99	< 0.001	PR**
	17	Among smokers, smoking less prevalence	0.21	0.14	0.3	99	< 0.001	Prev*
	22	Among smokers, smoking more	0.27	0.22	0.32	98	< 0.001	Prev*
	17	Among smokers, smoking unchanged	0.5	0.41	0.58	99	< 0.001	Prev*
	6	Among smokers, quit smoking	0.04	0.01	0.09	95	< 0.001	Prev*
	4	Among non-smokers, started smoking	0.02	0.01	0.03	92	< 0.001	Prev*
Sasidharanpillai et al. 2022	7	Women screened before the COVID-19 pandemic	0.0979	0.06	0.13 59	10 0	<0.001	Prop
	7	Women screened during the COVID-19 pandemic	0.0424	0.02 77	0.05 71	10	<0.001	Prop
Tang et al. 2022	10	Postoperative morbidity	0.9	0.8	1.01	26	0.22	OR**
	8	Postoperative mortality	1.27	0.92	1.75	0	0.57	OR**
	4	Converion rate	1.07	0.75	1.52	31	0.23	OR**
	5	Incidence of anastomotic leakage	0.71	0.07	19.2 2	0	0.74	OR**

	2	Intensive care unit demand rate	0.73	0.29	1.85	0	0.5	OR**
	4	R1 resections rate	0.46	0.11	1.9	0	0.48	OR**
	5	Mean lymph node yield	0.16	-2.26	2.59	54	0.07	MD**
	7	Length of hospital stay	-0.05	-2.28	2.19	98	< 0.001	MD**
				0.37	0.37	NT		PRED
Teglia et al. 2022	21	Breast cancer screening January-October 2020	0.467	8	8	P	NP	**
	21	Breast cancer screening April 2020	0.74	0.56 7		N P	NP	PRED **
	21	Breast cancer screening June-October 2020	0.13	-0.07	0.33	N P	NP	PRED **
	22		0.449	0.36	0.53		NP	PRED **
		Colonoscopy screening January-October 2020	0.525	0.38	0.66	N	NP	PRED **
	21	Fecal occult blood test or fecal immunochemical test January-October	0.378	0.25	0.49		NP	PRED **
	21		0.693	0.36		N P	NP	PRED **
		Colorectal cancer screening June-October 2020	0.234	0.02	0.44		NP	PRED **
		Cervical cancer screening January-October 2020	0.518	0.38	0.64		NP	PRED **
	21		0.788	0.58	0.99		NP	PRED **
								PRED **
Teglia et al. 2022	NP	Overall treatment January-October 2020	0.187	0.13	0.24	N P	NP	PRED **
	NP	Overall treatment January-February 2020	0.027	0.04	0.1	N P	NP	PRED **
	NP	Overall treatment March 2020	0.156	0.07 6	0.23		NP	PRED **

NP	Overall treatment April 2020	0.283	0.19 4	0.37		NP	PRED
INP	Overan treatment April 2020	0.283	0.17	$\frac{2}{0.04}$		NP	PRED
NP	Overall treatment May 2020	0.262	6	1	P	NP	**
	· ·		0.04	0.27	N		PRED
NP	Overall treatment June-October 2020	0.16	1		P	NP	**
			0.27	0.39			PRED
NP	Overall surgical treatment January-October 2020	0.339	9	9	P	NP	**
			- 0.00	0.22	N.T		PRED
NID	Overall sympical treatment January Eshavery 2020	0.072	0.09	0.23	N P	NP	**
NP	Overall surgical treatment January-February 2020	0.072	0.21	0.39		NP	PRED
NP	Overall surgical treatment March 2020	0.307	9	6.39		NP	**
INI	Overail surgical iteatment materi 2020	0.307	0.23	0.44		INI	PRED
NP	Overall surgical treatment April 2020	0.342	9	5		NP	**
		7.0	0.31	0.51			PRED
NP	Overall surgical treatment May 2020	0.416	8		P	NP	**
			0.18	0.51			PRED
NP	Overall surgical treatment June-October 2020	0.351	6		P	NP	**
			0.04	0.20			PRED
NP	Overall medical treatment January-October 2020	0.126	8	4	P	NP	**
			-				PRED
NID		0.015	0.05	0.08		NID	**
NP	Overall medical treatment January-February 2020	0.015	5	4	P	NP	DDED
			0.01	0.23	N		PRED
NP	Overall medical treatment March 2020	0.116	2	3	P	NP	
111	O TOTALI III GALONI (TOMINONI IVIMIONI 2020	0.110		0.40		111	PRED
NP	Overall medical treatment April 2020	0.248	0.09	7		NP	**
			0.08	0.30			PRED
NP	Overall medical treatment May 2020	0.196	5	6	P	NP	**
			-				PRED
			0.07	0.23			**
NP	Overall medical treatment June-October 2020	0.079	8	6	P	NP	

								PRED
				0.00		N		PRED
Vigliar et al. 2020	41	Cytological samples over 4 weeks of the COVID-19 pandemic	0.453	1	0.98	P	NP	**
	41	Ratio of exfoliative to fine needle aspiration samples	0.89	0.74	1.08	95	< 0.01	OR**
				0.03	0.07			
	27	Malignant diagnosis	0.0556	77	35	81	< 0.01	RD**

LCI, lower confidence interval; IRR, incidence rate ratio; MD, mean difference; OR, odds ratio; PRED, percent reduction; PR, prevalence ratio; Prev, prevalence: Prop, proportion; RD, risk difference; RR, rate ratio; PPE, personal protective equipment; NP, not provided; UCI, upper confidence interval; SMD, standardized mean difference; WMD, weighted mean difference

<sup>a</sup>, surveyed centers/operators; \*, estimates are during pandemic; \*\*. estimates are pandemic vs. pre-pandemic

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