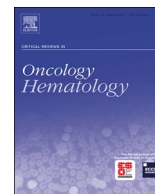




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COVID 19 therapies and anti-cancer drugs: A systematic review of recent literature



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ABSTRACT

Background: It is reasonable to think that cancer patients undergoing chemotherapy, targeted therapy or immunotherapy could have a more aggressive course if positive for Coronavirus disease CoV-2 (COVID-19).

Methods: We conducted a literature review on <https://www.ncbi.nlm.nih.gov/pubmed/>, <https://scholar.google.com>, www.arxiv.org, www.biorxiv.org, of all articles published using the keywords COVID-19 therapy or treatment and cancer until May 2, 2020. A total of 205 articles were identified and 53 were included in this review.

Results: We describe the ongoing COVID-19 therapies that should be known by oncologists and highlight the potential interactions with antineoplastic drugs, commonly used in clinical practice. The main drug interactions were found with tocilizumab, ruxolitinib and colchicine. **Conclusions.** The literature provides an inconclusive picture on potential preferred treatments for COVID-19 and their interactions with antineoplastic agents. Future clinical trials are needed to better understand the interactions between different drugs in the context of COVID-19 pandemic.

1. Introduction

In our recent history there have been three epidemics related to coronavirus infections: the SARS-CoV (severe acute respiratory syndrome), 2002-03; the MERS-CoV (Middle-East-Respiratory-Syndrome), 2012; and currently the SARS-CoV-2, (2019-2020) (Author 1, 2020). Until May 2, 2020, there were approximately 209000 confirmed cases, with 28 000 deaths due to coronavirus disease CoV-2 (COVID-19) in Italy, according to the Italian Civil Protection bulletin (Author 2, 2020). According to the World Health Organization (WHO) as of May 2 2020 due to COVID-19, in Spain there were 215000 confirmed cases and 25000 deaths; in the United States, 1 million confirmed cases and 57000 deaths; France has 128000 confirmed cases with 24000 deaths while United Kingdom has 177000 cases and 27000 deaths (Author 3, 2020).

From the available scientific literature it is evident that about 19.4% of the deaths with the coronavirus had an oncological pathology as comorbidity (Remuzzi et al., 2020; Landmann et al., 2020). Thus, during this COVID-19 crisis, cancer patients are regarded as a highly vulnerable group. It was found that within 14 days, anti-cancer treatments were significantly associated with occurrence of severe clinical events in COVID-19 infection (Zhang et al., 2020). The cancer population subjected to chemotherapy and/or radiotherapy is more exposed to infections in general and, therefore, also to that from Coronavirus primarily due to the effect of the cytotoxic action on the hematopoietic and immune systems with a reduction in the number of neutrophils, the first bulwark of infections, and decreased immune capacity (Remuzzi et al., 2020). Although there is no data yet on the risks of contracting coronavirus infection or on the clinical course of the infection during

Abbreviations: COVID-19, corona virus disease; AIFA, Italian Medicines Agency (Agenzia Italiana del Farmaco); IL6, interleukin 6; CPI, checkpoint inhibitors; CYP, cytochrome P450; ARSI, androgen receptor signaling inhibitors; DMARDs, disease-modifying antirheumatic drugs; Ace2, angiotensin-converting enzyme 2; DPP4, Dipeptidyl peptidase 4; HIV-1, human immunodeficiency virus

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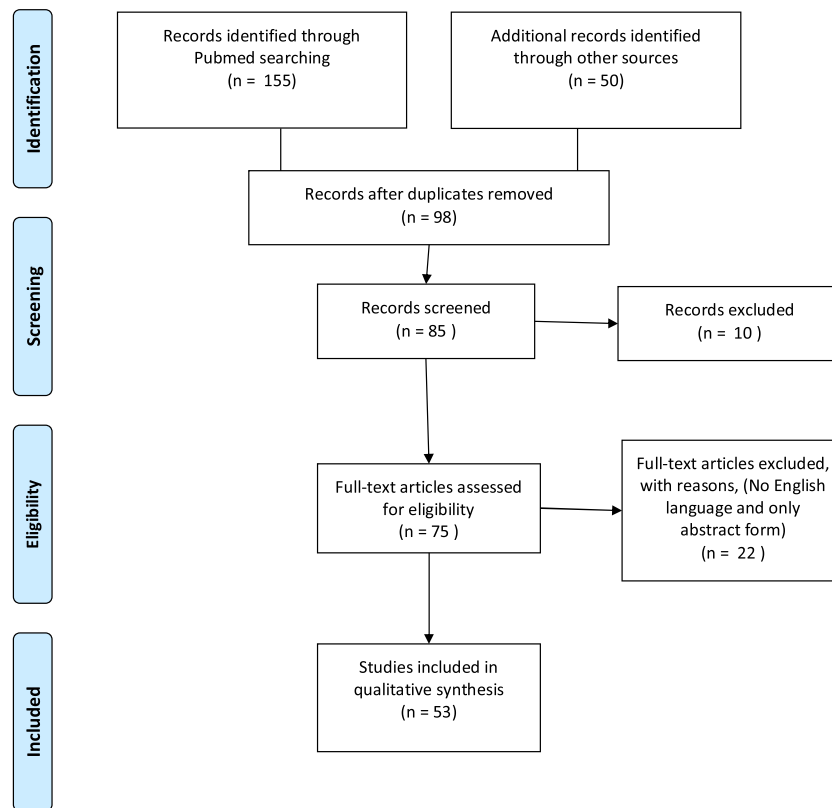


Fig. 1. Flow-chart of articles selection.

immunotherapy and/or immunosuppressive treatment with chemotherapy. It is reasonable to think, by analogy of what happens in the case of seasonal flu, due to the presence of immunosuppression, that in treated cancer patients, there may be a greater number of complications and the clinical course to be more serious (Author 1, 2020).

Therefore there remains an urgent need to answer whether COVID-19-positive cancer patients will have worse outcomes, such as death, from the coronavirus-induced pneumonia for example, and whether cancer patients should receive anti-cancer treatments. Additionally, oncologists are required to know the toxic effects of the drugs used in the experimental therapy of COVID-19 and the possible interactions of these drugs with the commonly used antineoplastic drugs.

2. Methods

2.1. Data sources and literature search strategy

The systematic review followed the PRISMA guidelines (Fig. 1) (Stewart et al., 2015). Two investigators (EL and RDT) independently conducted literature search using as combined keywords COVID-19 therapy or treatment and cancer on <https://www.ncbi.nlm.nih.gov/pubmed/>, www.arxiv.org (Author 4, 2020), www.biorxiv.org (Author 5, 2020) and <https://scholar.google.com> (Author 6, 2020). The database search was run of all the published articles from database inception until May 2, 2020. In Pubmed the following strategy was used: (COVID-19 OR Novel Coronavirus-Infected Pneumonia OR 2019 novel coronavirus OR 2019-nCoV or SARS-CoV-2 therapy or treatment) AND cancer.

The strategy was then adapted for the other databases, including website of Italian Medicines Agency (AIFA) for ongoing trials (<https://www.aifa.gov.it/emergenza-covid-19>) (Author 7, 2020).

2.2. Study selection and data synthesis

All studies reporting information on both COVID-19 therapy/treatment and cancer were included. 205 articles were identified and reviewed independently by two authors (EL AND RDT) and 53 articles were considered relevant to the scope of the current review, as described in Fig. 1 (Zhang et al., 2000; Mohile et al., 2020; AminJafari and Ghasemi, 2020; Salako et al., 2020; Russell et al., 2020; De Felice et al., 2020; Akladios et al. 2020, Banna et al., 2020; Al-Shamsi et al., 2020; Zhang et al., 2020; Hanna et al., 2020; Ying et al., 2015; Coles et al., 2020; Debureau et al., 2020; Di Saverio et al., 2020; Givi et al., 2020; Jin et al., 2020; Bersanelli, 2020; Ueda et al., 2020; Deng et al., 2020; Liang et al., 2020; Combs et al., 2020; Wang et al., 2020a; Zaorsky et al., 2020; Lim et al., 2020; Kalil, 2020; Wang et al., 2020b; Stebbin et al., 2020; Yang et al., 2020; Yu et al., 2020; Williams et al., 2020; Sheahan et al., 2020; Gautret et al., 2020; Chatre et al., 2018; Nhean et al., 2018; Gerson et al., 2018; Naqash et al., 2018; La Regina et al., 2019; Zheng et al., 2020; Author 8, 2020; Author 9, 2020; Author 10, 2020; Author 11, 2020; Author 12, 2020; Author 13, 2020; Deisseroth et al., 2015; Author 14, 2020; Author 15, 2020; JAK Inhibitors: Prospects in Connective Tissue Diseases et al., 2020; Iacobellis, 2020; Thachil, 2020; Author 16, 2020; de Bono et al. 2011; Author 17, 2020). Any inconsistencies were resolved by consensus with a third author (GDL). All health outcomes were included, due to the anticipated scarcity of data.

3. Results

The lists of the drugs being tested and their side-effects are listed in Table 1. They are presented below in more detail.

3.1. Antiviral drugs

Considering the etiological therapy of SARS-CoV-2, antiviral agents,

Table 1
Undesirable effects of the drugs tested in COVID19.

Drugs tested in COVID-19	Side effects
Favipiravir	Most common: nausea, vomiting, diarrhea, abdominal pain, anorexia, headache, anemia, muscle and joint pain.
Remdesivir	Common: leukopenia and thrombocytopenia, abnormal liver function; confusion, aggressive behaviour, agitation, migraine, tremors, paresthesia, dysgeusia.
Lopinavir	Uncommon: muscle weakness, pancreas dysfunction, lethargy, convulsions, coma, hallucinations, ataxia, kidney failure
Ritonavir	
Ribavirin	
Umifenovir	
Chloroquine / hydroxychloroquine	Common: anorexia, weight loss, nausea, abdominal pain, diarrhea, vomiting, corneal changes; uncommon: affective lability, nervousness, headache, dizziness, accommodation disorders Rare: retinopathy with changes in pigmentation and visual field defects, edema and opacity dizziness, tinnitus, bundle branch block, atrioventricular block, abnormal liver function tests, hypoglycemia skin rash, graying of the hair, alopecia, rashes, tiredness, psoriasis not sensitive to light
Azithromycin	Most common: diarrhea. Common: nausea, vomiting, abdominal pain, headache. Uncommon: asthenia, irritability, dermatological disorders as rash skin, edema, urticaria Rare: abnormal liver function
Anakinra	Most common: itching, erythema, pain, bruising, bleeding in the inoculation site, headache; common : abdominal pain, nausea, diarrhea Most common: upper airway infections, hypercholesterolemia
Tocilizumab	
Ivermectin	Common: pneumonia, cellulite, herpes zoster, abdominal pain, neutropenia, thrombocytopenia, abnormal liver function, gastrointestinal perforation, episodes of hypertension during drug infusion, headache, skin reactions Uncommon: hypothyroidism, stomach ulcers
Ecuzumab	Common: dizziness, nausea, diarrhea, swelling of the joints, rapid heart beat Uncommon: loss of appetite, vomiting, constipation, weakness, drowsiness, swelling of the eyes, face, arms, hands, feet
Emapalumab	Most common: headache Common: dizziness, nausea and diarrhea, abdominal pain, arthralgias, asthenia Uncommon: back pain, abnormal liver function, kidney failure.
Sarilumab	
Ruxolitinib	
	Most common: infections, hypertension Most common: neutropenia; common: ALT elevation, injection site erythema Most common: Myelosuppression, anemia, urinary tract infections, bleeding and increased systolic blood pressure. Common: neutropenia

*Adverse events frequency:

Most common $\geq 1/10$; Common $< 1/10 - \geq 1/100$.

Uncommon $< 1/100 - > 1/1000$; Rare $< 1/1000 - > 1/10000$.

already on the market and in use for other viral pathologies, in monotherapy or in combination are currently being tested.

3.2. Favipiravir (Avigan)

It has a mechanism of action related to the selective inhibition of the Viral polymerase-RNA-dependent RNA and is used as a backup drug when other therapies do not work (Sheahan et al., 2020).

3.3. Ribavirin

Commonly used for the treatment of hepatitis C and for inflammatory lung diseases, such as bronchiolitis, this antiviral drug is always used in combination with other medicines, such as interferon alfa and peginterferon alfa and can be used both in the treatment of adult patients and in the treatment of pediatric patients (Zheng et al., 2020).

3.4. Remdesivir

It is an antiviral drug in the class of nucleotide analogues. It was developed as a treatment for Ebola virus disease and Marburg virus infections. It has also been shown to have antiviral activity against RNA viruses such as human respiratory syncytial virus and coronaviruses, including viruses that cause the MERS and SARS (Sheahan et al., 2020).

On 29 April, the director of the US National Institute of Allergy and Infectious Diseases (NIAID), announced the results of a study in which patients taking remdesivir recovered in 11 days compared with 15 days for those on a placebo. The shortened recovery time was so significant that investigators decided to stop the trial (Author 8, 2020).

This news contradicts previous mixed results on the drug. Gilead Sciences announced that in a non-randomized trial, more than half of

participants with severe COVID-19 had recovered from their disease within two weeks of receiving treatment. Another smaller trial in China announced that it had found no benefits from remdesivir when compared with a placebo. The NIAID did not release information on safety data (Author 8, 2020).

3.5. Lopinavir-Ritonavir

Lopinavir and Ritonavir is indicated, in combination with other antiretroviral medicines, for the treatment of adults, and children over the age of 2 years with human immunodeficiency virus (HIV-1) infection (Sheahan et al., 2020). Lopinavir-Ritonavir in association with Ribavirin is being evaluated.

3.6. Anticytokines

3.6.1. Tocilizumab

It is a drug that counteracts the over-reaction of the immune system, at the origin of some of the most serious complications of COVID-19 and which in these days seems to offer hope to patients with coronavirus-induced pneumonia. In addition to rheumatoid arthritis therapy, the medicine is also used to treat cytokine release syndrome, a side effect of CAR-T anticancer therapy which consists of a massive release of inflammatory molecules in response to the immune cells used in the treatment (Author 9, 2020).

3.6.2. Emapalumab

It is an anti-interferon gamma. This drug keeps children with hemophagocytic lymphohistiocytosis (HLH) alive, awaiting transplantation. The administration of emapalumab is able to "turn off" the abnormal and excessive inflammatory response in patients with HLH, neutralizing the effects deriving from the excessive production of

interferon-gamma (Author 10, 2020).

3.6.3. Sarilumab

Sarilumab is an interleukin-6 (IL-6) receptor antagonist indicated for the treatment of adult patients with moderately to severely active rheumatoid arthritis who have had an inadequate response or intolerance to one or more disease-modifying antirheumatic drugs (DMARDs) (Author 11, 2020).

3.6.4. Eculizumab

It is a member of the drug class selective immunosuppressants and is used to treat Hemolytic Uremic Syndrome, Myasthenia Gravis, Neuromyelitis Optica and Paroxysmal Nocturnal Hemoglobinuria (Author 12, 2020).

3.6.5. Anakinra

It inhibits the activity of interleukin-1 (IL-1), and has already been approved for the treatment of a group of rare and potentially fatal auto-inflammatory diseases. The US Food and Drugs Administration (FDA) has granted a study to test the safety and effectiveness of individually or simultaneously blocking IL-6 and IL-1 versus standard of care on blood oxygenation and systemic cytokine release syndrome in patients with COVID-19 coronavirus infection and acute hypoxic respiratory failure and systemic cytokine release syndrome (Author 13, 2020).

3.6.6. Siltuximab

Siltuximab like tocilizumab, binds to interleukin-6 and is indicated for the treatment of adult patients with Castleman disease (Deisseroth et al., 2015). The SISCO study (Siltuximab In Serious COVID-19) is configured as an observational study conducted both on hospitalized patients and those already in intensive care (Author 18, 2020).

Table 2 shows Italian Medicines Agency (AIFA) approved

Table 2

AIFA approved experimental studies in COVID-19 therapy.

Study	Promoter	CE Single Opinion Date	Documents
A randomized, double-blind, placebo-controlled, multicenter study to evaluate the safety and efficacy of tocilizumab in patients with severe covid-19 pneumonia (Tocilizumab 2020-001154-22).	F.Hoffmann-La Roche Ltd	30/03/2020	https://www.aifa.gov.it/documents/20142/1131319/TocilizumabDocumenti.zip
An open-label randomized multicenter study to evaluate the efficacy of early administration of Tocilizumab (TCZ) in patients with COVID-19 pneumonia (RCT-TCZ-COVID-19).	Local Health Unit-IRCCS of Reggio Emilia	27/03/2020	https://www.aifa.gov.it/documents/20142/1131319/RCT-TCZ-COVID19documenti.zip
Multicenter study on the efficacy and tolerability of tocilizumab in the treatment of patients with COVID-19 pneumonia (TOCIVID-19).	National Cancer Institute, IRCCS, G. Pascale Foundation	22/03/2020	https://www.aifa.gov.it/documents/20142/1131319/TOCIVID-19documenti.zip
A randomized, double-blind, placebo-controlled adaptive phase 2/3 study evaluating the efficacy and safety of sarilumab for hospitalized patients with COVID-19 (Sarilumab COVID-19).	Sanofi-aventis Recherche & Développement	26/03/2020	https://www.aifa.gov.it/documents/20142/1131319/Sarilumabdocumenti.zip
A phase 2/3, randomized, open-label, parallel, 3-arm, multicenter study investigating the efficacy and safety of intravenous doses of emapalumab, a monoclonal anti-interferon gamma (anti-IFN γ) and anakinra antibody, a interleukin -1 receptor antagonist (IL-1), compared to the standard of care, in reducing hyperinflammation and respiratory difficulties in patients with SARSCoV-2 infection (Sobi.IMMUNO-101).	SOBI	25/03/2020	https://www.aifa.gov.it/documents/20142/1131319/Sobi.IMMUNO-101documenti.zip
A randomized phase 3 study to evaluate the safety and antiviral activity of Remdesivir (GS-5734 [™]) in participants with moderate COVID-19 treatment compared to the standard of therapy. (Study GS-US-540-5774)	Gilead Sciences, Inc	11/03/2020	https://www.aifa.gov.it/documents/20142/1131319/GS-US-540-5774documenti.zip
A randomized phase 3 study to evaluate the safety and antiviral activity of Remdesivir (GS-5734 [™]) in participants with severe COVID-19. (Study GS-US-540-5773)	Gilead Sciences, Inc	11/03/2020	https://www.aifa.gov.it/documents/20142/1131319/GS-US-540-5773_documenti.zip
A randomized controlled two-phase phase 2 trial to evaluate the efficacy, safety and tolerability of baricitinib in addition to the usual treatment in patients with pneumonia in COVID19	Pisan University Hospital	23/04/2020	https://www.aifa.gov.it/documents/20142/1131319/BARCHID_documenti.zip
Colchicine to counteract the inflammatory response in COVID pneumonia 19	University Hospital of Parma	23/04/2020	https://www.aifa.gov.it/documents/20142/1131319/ColCoviddocumenti.zip
Intermediate dose enoxaparin in hospitalized patients with moderate-severe COVID19: a pilot phase II single-arm study, INHIXACOV19	Bologna University	23/04/2020	https://www.aifa.gov.it/documents/20142/1131319/INHIXACOV19_documenti.zip

AIFA: Italian Medicines Agency.

experimental studies in COVID-19 therapy (Author 19, 2020; Author 20, 2020; Author 21, 2020; Author 22, 2020; Author 23, 2020; Author 24, 2020; Author 25, 2020; Author 26, 2020; Author 27, 2020; Author 28, 2020; Author 29, 2020).

The main currently ongoing studies are:

- 1) **TOCIVID-19**, multicenter study on the efficacy and tolerability of Tocilizumab in the treatment of patients with COVID-19 pneumonia;
- 2) **Sobi study, IMMUNO-101**, a randomized, open, 3 parallel, multicenter, phase 2/3 groups, that evaluates the efficacy and safety of intravenous doses of Emapalumab, anti-interferon gamma monoclonal antibody (anti-IFN γ), and Anakinra, interleukin receptor antagonist-1 (IL-1), compared to standard therapy;
- 3) **Sarilumab COVID-19 study**: a randomized, double-blind, placebo-controlled phase 2/3 study evaluating the efficacy and safety of intravenous administration of Sarilumab, an interleukin receptor antagonist- 6 (IL- 6).

3.7. Other potential therapeutic agents

The required urgency for the identification of potential treatments has necessitated the further investigation of a wide-range of potential therapeutic agents. The ones identified in this review are listed below in some detail.

3.7.1. Ivermectine

The pesticide Avermectina (otherwise known as Abamectina and Ivermectina) is registered by the US Environmental Protection Agency (EMA) as a pesticide. The study with the pesticide Ivermectin, approved by the FDA as a pesticide against scabies and already used against the HIV, Zika, Dengue, West Nile and Influenza viruses. This study, carried

Table 3
Pharmacological interactions of the drugs tested in COVID-19 with the most common antineoplastic drugs.

Anti-covid19	Interaction	Side effects
Azithromycin	Regorafenib	Therapeutic effect reduction
Azithromycin	Vinblastina	Increased serum levels of P-glycoprotein; Increased toxicity (neutropenia, myalgias, constipation)
Chloroquine	Doxorubicina	cardiomyopathy or conduction system abnormalities
Chloroquine	Taxanes	Increase in Plasma concentrations
Chloroquine	Trastuzumab	cardiomyopathy or conduction system abnormalities
Anakinra	Fluorouracile	Increased immunosuppressive action
Anakinra	Durvalumab	Therapeutic effect reduction
Emapalumab, Sarilumab	no interactions with antineoplastic drugs	
Favipiravir	Enzalutamide	Interference with cytochromes: reduce efficacy of antivirals
Lopinavir		
Ritonavir		
Ribavirin		
Umifenovir		

out at the Monash University's Biomedicine Discovery Institute of the Peter Doherty Institute of Infection and Immunity in Australia, has shown that a single dose of the medicine would be able to block the growth of the pathogen in culture, eliminating all the viral genetic material within 48 hours (Author 14, 2020).

3.7.2. Association of chloroquine or hydroxychloroquine in combination with a macrolide (azithromycin)

Two studies have been published in France on the combination of hydroxychloroquine and azithromycin in patients with COVID-19 by the same research team as The Méditerranée Infection University Hospital Institute in Marseille. In summary, the two studies have documented a significant reduction in viral load and a clinical improvement compared to the natural progression of the disease triggered by SARS-CoV-2 infection. A death was recorded, while three patients were hospitalized in the Intensive Care wards (pooled data of the two studies) (Gautret et al., 2020; Author 15, 2020).

3.7.3. Ruxolitinib

Ruxolitinib is an inhibitor of the signal transmission pathway mediated by Janus Kinase (Jak), with anti-inflammatory effects related to the inhibition of the release of cytokines. Normally ruxolitinib is used in the hematology field and is indicated for the treatment of splenomegaly or disease-related symptoms in adult patients with primary myelofibrosis. It is also approved for the treatment of adult patients with polycythemia vera who are resistant or intolerant to hydroxyurea (JAK Inhibitors: Prospects in Connective Tissue Diseases et al., 2020). A compassionate use of Ruxolitinib has also been approved by AIFA in Covid-19 patients with respiratory failure who do not require invasive assisted ventilation.

3.7.4. Baricitinib

Baricitinib is another JAK inhibitor indicated for the treatment of active rheumatoid arthritis (JAK Inhibitors: Prospects in Connective Tissue Diseases et al., 2020). AIFA has licensed a randomized phase 2 trial to evaluate the efficacy, safety and tolerability of baricitinib in addition to the usual treatment in patients with pneumonia in COVID-19 (Barcivid study) (Author 27, 2020).

3.7.5. Anti-diabetes

A new line of research comes from diabetologists. According to a study, in addition to the main entrance door of virus, the angiotensin-converting enzyme 2 (ACE2) receptor, the DPP4 receptor must be evaluated. The Dipeptidyl peptidase 4 (Dpp4) receptor is present on all human cell types (bronchi and heart) and is the same on which many medicines for diabetes work (Iacobellis, 2020).

3.7.6. Colchicine

Colchicine is a molecule, capable of interfering with the inflammatory immune response observed in subjects with COVID-19. Colchicine is a drug that has been used for a long time and is effective for the treatment of acute attack of gouty arthritis. Specifically, it manages to reduce the release of cytokines, molecules that, like IL-6, are responsible for the inflammatory (La Regina et al., 2019; Author 9, 2020). A study has recently started in Italy (Author 28, 2020).

3.7.7. Heparin

A reduction of up to 20% in mortality among COVID-19 patients with a marked increase in an indicator of the presence of blood clots, was achieved due to the use of an old drug, heparin (Thachil, 2020). This was reported by a study according to which the use of heparin in COVID-19 patients could have anticoagulant effects, as well as anti-inflammatory and potentially even antiviral ones. It has been shown that new coronavirus infections are associated with a high mortality in the presence of high value of D-dimer, a particularly important marker for coagulopathy (Thachil, 2020). However, it will be necessary to study in depth at what dosage the drug could have such potential antiviral properties. A recent study has been approved by AIFA with enoxeparin (INHIXACOVID) (Author 29, 2020).

3.8. Anti cancer drugs and anti COVID-19

Table 3 highlights the pharmacological interactions of some molecules tested against COVID-19 with the most common antineoplastic drugs. We have not found an overall high rate of interactions. The main drug-interactions reported are with Chloroquine/Hydroxychloroquine and chemotherapy or trastuzumab. Conduction disorders, ventricular hypertrophy and valvular dysfunction were the main side effect reported (Gautret et al., 2020; Author 16, 2020).

Enzalutamide, androgen receptor signaling inhibitors (ARSI), commonly used in prostate cancer could potentially interfere with antiviral drugs due to interference on cytochromes (Nhean et al., 2018).

Table 4 shows interferences with tocilizumab, ruxolitinib and colchicine. Tocilizumab increases immunosuppressive action and inhibits Fluorouracil-resistance (Ying et al., 2015); it interferes with pharmacodynamic activity and the therapeutic efficacy of durvalumab, nivolumab, atezolizumab (Ying et al., 2015; Jin et al., 2020; Bersanelli, 2020). Moreover it is important to emphasize that tocilizumab decreases the concentration of several medications as a cytochrome P450 (CYP), isoenzyme CYP3A4 inducer (De Felice et al., 2020, Akladios et al., 2020, Banna et al., 2020; Al-Shamsi et al., 2020; Zhang et al., 2020; Hanna et al., 2020; Ying et al., 2015; Author 17, 2020). Therefore if drugs in lung cancer are used, such as ceritinib, crizotinib, brigatinib, gefitinib, docetaxel, a reduction in the anticancer drug may become noticeable, as well as paying attention in the case of concomitant use of

Table 4
Pharmacological interactions of the Tocilizumab, Ruxolitinib and colchicine with the most common antineoplastic drugs.

Antineoplastic agents	Drug interaction with antiCOVID-19 agents	Mechanism of interaction
Abiraterone	Tocilizumab Colchicine	Tocilizumab reduces abiraterone as a CYP3A4 inducer. Increase abiraterone as CYP3A4 inhibitor
Alectinib	Ruxolitinib	Increased bradycardic effect.
Axitinib	Tocilizumab	Tocilizumab reduces axitinib as a CYP3A4 inducer.
Bosutinib	Tocilizumab	Tocilizumab reduces bosutinib as a CYP3A4 inducer.
Brigatinib	Ruxolitinib	Increased bradycardic effect.
Cabozantinib	Tocilizumab	Tocilizumab reduces brigatinib serum as a CYP3A4 inducer.
Ceritinib	Tocilizumab Colchicine Ruxolitinib	Tocilizumab reduces cabozantinib as a CYP3A4 inducer. Increased colchicine serum concentration. Increased ruxolitinib serum concentration.
Cisplatin	Tocilizumab	Tocilizumab reduces ceritinib as a CYP3A4 inducer.
Cobimetinib	Colchicine	adjust doses because uricemia increases
CPI	Tocilizumab Ruxolitinib Baricitinib	Tocilizumab reduces cobimetinib, as a CYP3A4 inducer. Therapeutic effect reduction or synergism (nivolumab) Interference with pharmacodynamic activity and therapeutic efficacy of these molecules
Crizotinib	Tocilizumab Colchicine Ruxolitinib	Increased colchicine and increased bradycardic effect as CYP3A4 inhibitor
Dasatinib	Tocilizumab	Tocilizumab reduces crizotinib as a CYP3A4 inducer
Docetaxel	Tocilizumab Colchicine	Tocilizumab reduces docetaxel as a CYP3A4 inducer. Increase docetaxel as a CYP 3A4 inhibitor.
Enzalutamide	Tocilizumab	Tocilizumab reduces enzalutamide as a CYP3A4 inducer.
Erlotinib	Tocilizumab	Tocilizumab reduces erlotinib as a CYP3A4 inducer.
Everolimus	Tocilizumab	Tocilizumab reduces serum concentration as a CYP3A4 inducer.
Fluorouracil	Tocilizumab (eculizumab)	Increased immunosuppressive action; Fluorouracil-resistance inhibition
Exemestane	Tocilizumab	Tocilizumab reduces exemestane, as a CYP3A4 inducer.
Gefitinib	Tocilizumab	Tocilizumab reduces gefitinib as a CYP3A4 inducer.
Ibrutinib	Tocilizumab	Tocilizumab reduces ibrutinib as a CYP3A4 inducer.
Idelalisib	Tocilizumab	Tocilizumab reduces idelalisib as a CYP3A4 inducer.
Ixazomib	Tocilizumab	Tocilizumab reduces ixazomib as a CYP3A4 inducer.
Lapatinib	Colchicine Tocilizumab	Lapatinib increases colchicine as a glycoprotein-P, ABCB1 inhibitor. Tocilizumab reduces lapatinib as a CYP3A4 inducer.
Neratinib	Colchicine Tocilizumab	Neratinib increases colchicine as glycoprotein-P ABCB1 inhibitor. Tocilizumab reduces neratinib as a CYP3A4 inducer.
Nilotinib	Colchicine Ruxolitinib Tocilizumab	Nilotinib increases colchicine as a CYP 3A4 inhibitor. Nilotinib increases ruxolitinib as a CYP 3A4 inhibitor. Tocilizumab reduces nilotinib as a CYP3A4 inducer.
Olaparib	Tocilizumab	Tocilizumab reduces olaparib as a CYP3A4 inducer.
Paclitaxel	Tocilizumab	Tocilizumab reduces paclitaxel serum as a CYP3A4 inducer
Palbociclib	Tocilizumab	Tocilizumab reduces palbociclib as a CYP3A4 inducer.
Panobinostat	Tocilizumab	Tocilizumab reduces panobinostat as a CYP3A4 inducer.
Pazopanib	Tocilizumab	Tocilizumab reduces pazopanib as a CYP3A4 inducer.
Pomalidomide	Tocilizumab	Increased immunosuppressive effect.
Regorafenib	Tocilizumab	Tocilizumab reduces regorafenib as a CYP3A4 inducer.
Ribociclib	Colchicine Ruxolitinib Tocilizumab	Ribociclib increases colchicine as a CYP 3A4 inhibitor. Ribociclib increases colchicine as a CYP 3A4 inhibitor. Tocilizumab reduces ribociclib, as a CYP3A4 inducer.
Ruxolitinib	Tocilizumab	Tocilizumab reduces ruxolitinib as a CYP3A4 inducer.
Sunitinib	Tocilizumab	Tocilizumab reduces sunitinib as a CYP3A4 inducer.
Tamoxifene	Colchicine Tocilizumab	Tamoxifene increases colchicine as a glycoprotein-P, ABCB1 inhibitor. Tocilizumab reduces tamoxifene as a CYP3A4 inducer.
Vandetanib	Tocilizumab	Tocilizumab reduces vandetanib as a CYP3A4 inducer.
Venetoclax	Tocilizumab	Tocilizumab reduces venetoclax as a CYP3A4 inducer.
Alcaloids of the vinca	Colchicine	Synergy of action and enhancement of toxicity.

CPI: Check Point inhibitors.

tyrosine kinase inhibitors.

Anakinra and ruxolitinib could interfere with checkpoint inhibitors (CPI) in terms of antagonisms or synergy but this is not yet entirely clear (Ying et al., 2015; Bersanelli, 2020; Author 13, 2020). Several drugs, such as CYP3A4 inhibitors, increase concentration of colchicine and attention should be paid when the latter is used with cisplatin or vinca alkaloids (La Regina et al., 2019). Furthermore, one should not forget a concomitant use of antidiabetic drugs with antineoplastics that require cortisone (de Bono et al. 2011) or cytostatic drugs that reduce the value of platelets with the concomitant use of heparin (Thachil, 2020).

4. Discussion

Based on the increased risk of disease aggression of COVID-19 in patients with cancer it is important to describe which anti COVID-19 drugs could be administered and which not in cancer patients. The current review attempts to provide a first systematic glance at this issue. There are certain emergent points of attention that have already been described.

Specifically, particular attention to the potential cardiotoxicity of chloroquine with anthracyclines is necessary. Hydroxychloroquine is an antimalarial that has become a mainstay in the management of systemic lupus erythematosus and rheumatoid arthritis. Similar to chloroquine

in structure, hydroxychloroquine is used more frequently because it confers lower toxicity. Cardiotoxicity manifests as restrictive or dilated cardiomyopathy or with conduction system abnormalities such as atrioventricular and bundle-branch block (Gautret et al., 2020). Recently, some doubts related to cardiovascular health have emerged in COVID-19 patients. In fact cardiac rhythm disturbances and cardiac arrest in some French hospitals are currently under consideration by the French National Agency for the Safety of Medicines (Author 16, 2020).

If one considers that the metabolism of most drugs is catalyzed by the enzyme group of CYP, composed of a large number of isoenzymes, the most important of which in the metabolism of drugs are CYP1A2, CYP2C9, CYP2C19, CYP2D6 and CYP3A4. Enzalutamide, recognized as a strong CYP3A4 and a moderate CYP2C9 and CYP2C19 inducer, significantly reduces concentrations of several anti-HIV agents such as darunavir, ritonavir, etravirine, and raltegravir. The latter drugs are the subject of a series of trials in progress and could be soon active in the fight against COVID-19 (Nhean et al., 2018).

It is important also to note that some chemotherapeutic agents or hormonal therapies such as abiraterone require concomitant steroids that are immunosuppressive agents and could weaken anticitokines drugs such as tocilizumab (de Bono et al. 2011). However in some cases, the use of steroids could be reduced without interfering with the efficacy and toxicity profile. Tocilizumab decreases concentrations of many anticancer drugs due to interference with CYP (Author 17, 2020).

Deserving to underline that today many patients carry out CPI, currently used in daily practice for the treatment of solid tumors. CPI, activating T-lymphocytes, increase the risk of autoimmune disease such as pneumoniae (Gerson et al., 2018). Few case-reports have described significant reduction in IL-6 and CPI immunorelated toxicity after tocilizumab administration (Naqash et al., 2018). To date the interaction between CPI and anti COVID-19 agents is not well-described. Consideration could be given to either extending the timing of the doses or delaying pretreated patients.

The present review should be interpreted in light of its limitations. Firstly, there are no studies with COVID-19 therapies in cancer populations being treated with antineoplastic drugs and for this reason the informations collected are indirect and from studies in which these antiviral experimental agents have been used in cancer patients for other indications. Thus more research on this topic is needed before concrete recommendations can be made.

Secondly, the type and dosage of therapeutic agents varied between studies.

Thirdly the data are based on retrospective findings, while the results of prospective cohorts are being awaited and expected to shed more light.

Moreover considering the rapid evolution of information related to the pandemic we conducted a search on 2 arxiv and biorxiv databases indentifying 50 articles of interest; these are preliminary reports that have not been peer-reviewed. Among the 50 articles only 2 were considered as relevant (Yu et al., 2020; Williams et al., 2020) and subsequently 1 has been published in peer-reviewed journal (Yu et al., 2020), testifying to the current urgent need of relevant, well-designed studies.

5. Conclusions

The literature so far available provides an inconclusive picture on potential preferred treatments for COVID-19 and their interactions with antineoplastic agents.

In order to continue treating cancer patients likely to be infected with COVID-19, we believe that it is appropriate to modify available therapeutic choices and arrive at a new therapeutic protocol that provides:

- 1) where there are no interactions to continue or delay antineoplastic therapy by 1-2 weeks;
- 2) where there are interactions and the patient has already done many

- cycles to stop the administration of the anticancer drug;
- 3) for newly diagnosed cancer patients to prefer antineoplastic drugs without interactions;
- 4) where further attention is given to CPI, the choice of which may vary from case to case.

This new therapeutic proposal awaits the constructive suggestions of the oncological community in order to make improvements, such as to better protect our patients, already fragile because of cancer. While the information flow has been largely uninhibited during the pandemic, findings from current and future clinical trials are still needed to better understand the interactions between different pharmaceutical agents in the context of COVID-19 infections.

Declaration of conflicting interests

All authors declare no conflict of interest.

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