**Review Paper: Drug Interactions of Psychiatric and COVID-19 Medications**

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**Introduction:** Coronavirus disease 2019 (COVID-19) has become a pandemic with 1771514 cases identified in the world and 70029 cases in Iran until April 12, 2020. The co-prescription of psychotropics with COVID-19 medication is not uncommon. Healthcare providers should be familiar with many Potential Drug-Drug Interactions (DDIs) between COVID-19 therapeutic agents and psychotropic drugs based on cytochrome P450 metabolism. This review comprehensively summarizes the current literature on DDIs between antiretroviral drugs and chloroquine/hydroxychloroquine, and psychotropics, including antidepressants, antipsychotics, mood stabilizers, and anxiolytics.

**Methods:** Medical databases, including Google Scholar, PubMed, Web of Science, and Scopus were searched to identify studies in English with keywords related to psychiatric disorders, medications used in the treatment of psychiatric disorders and COVID-19 medications.

**Results:** There is a great potential for DDIs between psychiatric and COVID-19 medications ranging from interactions that are not clinically apparent (minor) to those that produce life-threatening adverse drug reactions, or loss of treatment efficacy. The majority of interactions are pharmacokinetic interactions via the cytochrome P450 enzyme system.

**Conclusion:** DDIs are a major concern in the comorbidity of psychiatric disorders and COVID-19 infection resulting in the alteration of expected therapeutic outcomes. The risk of toxicity or lack of efficacy may occur due to a higher or lower plasma concentration of medications. However, psychiatric medication can be safely used in combination with COVID-19 pharmacotherapy with either a wise selection of medication with the least possibility of interaction or careful patient monitoring and management.
1. Introduction

In December 2019, the reports of Coronavirus Disease 2019 (COVID-19) were documented in Wuhan, China. This condition was caused by a novel beta-coronavirus, called Severe Acute Respiratory Syndrome coronavirus 2 (SARS-CoV-2). Nowadays, the Coronavirus Disease 2019 (COVID-19) has become a global pandemic; with more than hundred thousand cases have been identified in Iran. COVID-19 affects people’s mental health and quality of life. The COVID-19 causes panic, anxiety, depression and insomnia, especially in patients and health care professionals. There is a possibility of starting psychiatric drugs in patients with COVID-19. Also, some patients with COVID-19 have been taking psychiatric drugs due to various problems. As a result of this, the likelihood of medication interaction (DDI) between psychiatric drugs and drugs used to treat COVID-19 can be a concern. Many psychiatric drugs are metabolized via the cytochrome P450 system. Therefore, their hepatic metabolism and ability to alter activity of cytochrome P450 can lead to pharmacokinetic interactions when they are used with medications in COVID-19 treatment. Risk of toxicity or lack of efficacy may occur due to higher or lower plasma concentration of medications. Moreover intensification of similar adverse effects might lead to patients harm. Psychiatric medication can be safe in combination with COVID-19 pharmacotherapy with close attention to selection of medication with the least possibility of interaction and also careful monitoring.

Highlights

- Drug interactions are major challenge in comorbidity of psychiatric disorders and COVID-19 infection
- QTc prolongation is a major concern while using antiviral medications and/or Chloroquine/hydroxychloroquine in combination with many psychotropics. Cardiac monitoring especially in high risk patients is highly recommended.
- Concomitant use of SSRIs with antiviral medications and/or Chloroquine/hydroxychloroquine increase the risk of hypoglycemia.
- Concomitant use of pimozide or midazolam with antiviral medications is contraindicated.

Plain Language Summary

In December 2019, the reports of Coronavirus Disease 2019 (COVID-19) were documented in Wuhan, China. This condition was caused by a novel beta-coronavirus, called as Severe Acute Respiratory Syndrome coronavirus 2 (SARS-CoV-2). Nowadays, the Coronavirus Disease 2019 (COVID-19) has become a global pandemic; with more than hundred thousand cases have been identified in Iran. COVID-19 affects people’s mental health and quality of life. The COVID-19 causes panic, anxiety, depression and insomnia, especially in patients and health care professionals. There is a possibility of starting psychiatric drugs in patients with COVID-19. Also, some patients with COVID-19 have been taking psychiatric drugs due to various problems. As a result of this, the likelihood of drug-drug interaction (DDI) between psychiatric drugs and drugs used to treat COVID-19 can be a concern. Many psychiatric drugs are metabolized via the cytochrome P450 system. Therefore, their hepatic metabolism and ability to alter activity of cytochrome P450 can lead to pharmacokinetic interactions when they are used with medications in COVID-19 treatment. Risk of toxicity or lack of efficacy may occur due to higher or lower plasma concentration of medications. Moreover intensification of similar adverse effects might lead to patients harm. Psychiatric medication can be safe in combination with COVID-19 pharmacotherapy with close attention to selection of medication with the least possibility of interaction and also careful monitoring.
plays a key role in the metabolism of medications and their interactions (Ogu & Maxa, 2000). For example, the hepatic CYP isoenzymes, which are responsible for metabolizing the tricyclic antidepressants consisting of 2D6, 1A2, 3A4, 2C19 (Gillman, 2007). Most first- and second-generation antipsychotics are metabolized by the cytochrome P450 system mainly include 2D6, 1A2, and 3A4 (Wijesinghe, 2016). On the other hand, some Selective Serotonin Reuptake Inhibitors (SSRIs) are cytochrome P450 inhibitors and can result in DDIs by changing the blood concentration of the drugs activated or metabolized by these enzymes (Preskorn, 1997). Therefore, their hepatic metabolism and ability to alter the activity of cytochrome P450 can make pharmacokinetic interactions when they are used with medications in COVID-19 treatment.

The purpose of this review is to comprehensively explore DDIs, especially pharmacokinetic DDIs between psychotropic medications and COVID-19 medications. Also, we try to find the adverse effects and clinical issues that must be considered for combined medication regimens in psychiatry.

2. Methods

2.1. Search strategy

For this narrative review, the relevant articles published from 2000 to April 2020 were searched in Google Scholar, PubMed, Web of Science, and Scopus databases. The articles were limited to the English language. The keywords used in titles and abstracts were depression or anxiety, psychosis or psychiatric disorders and specific names of medications used in the treatment of psychiatric disorders, antidepressants, antipsychotics, anxiolytics, or mood stabilizers, as well as psychotropic medication in combination with pharmacological treatments of COVID-19 i.e. chloroquine, hydroxychloroquine, ritonavir, lopinavir, or Kaletra. Article references were also checked to find additional relevant studies.

The inclusion criteria were any type of study, including reviews and systematic reviews, clinical trials and case reports with the subject of drug-drug interactions of any psychotropic drug with ritonavir, lopinavir, or Kaletra and or chloroquine/hydroxychloroquine with a study population of either healthy volunteers or patients with HIV/AIDS, hepatitis, or malaria.

Articles’ titles and abstracts were reviewed for relevant information by two reviewers and the discrepancies were checked with the first author.

In total, 30 studies met the inclusion criteria. All publications were reviewed in full text by the researchers.

2.2. Antidepressants

2.2.1 Selective serotonin reuptake inhibitors

The Selective Serotonin Reuptake Inhibitors (SSRIs) are used to treat major depressive disorder, anxiety, and some other psychiatric disorders. All SSRIs are metabolized by cytochrome P450 enzymes and some of them (such as fluoxetine) can inhibit these enzymes. Therefore, SSRIs’ interactions with other cytochrome P450 substrates should be considered (DEPRESSION).

Fluoxetine is one of the most recommended SSRIs. The utmost amount of fluoxetine N-demethylation is mediated by CYP2D6, CYP2C9, CYP2C19, and CYP3A4. Fluoxetine and its active metabolite (nor-fluoxetine) inhibit CYP2D6 (strong), CYP2C9 (moderate), CYP2C19 (weak to moderate), CYP3A4 (weak to moderate), and CYP1A2 (weak) (Jin et al., 2016; Margolis, O’Donnell, Mankowski, Ekins, & Obach, 2000). Fluoxetine and its metabolites increase the Area Under The Curve (AUC) of ritonavir by up to 19% through inhibiting CYP2D6 and CYP3A4 (especially CYP2D6) without significant change in other kinetic parameters of ritonavir. This relatively low drug-drug interaction does not need dose modification (Ouellet et al., 1998). On the other hand, there is a possibility of serotonin syndrome in concomitant use of ritonavir with SSRIs. Desilova et al. reported four cases of serotonin syndrome as a result of the co-administration of ritonavir and fluoxetine (DeSilva, Le Flore, Marston, & Rimland, 2001). Fluoxetine as a CYP2D6 inhibitor (strong) may decrease the metabolism of CYP2D6 substrates like chloroquine and hydroxychloroquine (Bjornsson et al., 2003). Consequently, the side effects of chloroquine and hydroxychloroquine should be considered.

Paroxetine is metabolized mainly by CYP2D6 (high affinity) and CYP3A4 (low affinity) and partially by CYP1A2, CYP2C19, and CYP3A5. Its protein binding is 95% and can inhibit the CYP2D6 enzyme (Jornil, Jensen, Larsen, & Linnet, 2010). In Manon.et al. study, 26 healthy participants were given paroxetine (20 mg/d) and fosamprenavir/ritonavir (700 mg/100 mg twice daily) for 10 days. They found that paroxetine levels dropped by an average of 55% compared with when it was used alone because of protein binding displacement (Van Der Lee et al., 2007).
Citalopram and S-enantiomer of citalopram (escitalopram) are metabolized by CYP3A4, CYP2C19, and CYP2D6. They both have a weak inhibitory effect on CYP2D6 (von Molk et al., 2001). Ritonavir inhibits CYP3A4 but the interaction with citalopram and its active S-enantiomer is not clinically significant. In a study by Marcelo on 21 healthy volunteers who received daily escitalopram (20 mg) and ritonavir (600 mg), there was no significant change in the pharmacokinetic profile of escitalopram except for the volume of distribution, which decreased by 10% during concomitant use with ritonavir (Gutierrez, Rosenberg, & Abramowitz, 2003). Nonetheless, ketoconazole as a potent CYP3A4 inhibitor did not alter the pharmacokinetics of citalopram, indicating that ketoconazole and other CYP3A4 inhibitors may potentially be safe with citalopram and no dose adjustment is necessary, although this issue has not been studied explicitly (Gutierrez & Abramowitz, 2001). Citalopram and escitalopram inhibit CYP2D6 (weak), so it does not appear to cause significant changes in ritonavir levels (Hill & Lee, 2013).

Medications like citalopram/escitalopram and chloroquine/hydroxychloroquine are known for causing cardiac complications (such as QTc prolongation). Possibility of this complication seemed additive and can be aggravated in the presence of further risk factors, such as old age, cardiac disease, electrolyte disturbances ( hypokalemia, hypomagnesemia), and bradycardia. If there is no alternative medicine, co-administration of these drugs must be done with monitoring the patient’s clinical symptoms and Electrocardiogram (ECG) (Meid et al., 2017). Citalopram may enhance the hypoglycemic effect of hydroxychloroquine. Results of two case-control studies show that consuming SSRIs, especially for more than 3 years in people treated with blood sugar lowering drugs, can increase the risk of hypoglycemia (Derijks et al., 2008; Ghaeli et al., 2015; Han et al., 2016; Santra, Banerjee, Munshi, & Chaudhuri, 2015). Hydroxychloroquine can induce symptomatic or asymptomatic hypoglycemia in diabetic and nondiabetic patients (Unübol, Ayhan, Gucen, & Guncu, 2011). Also, chloroquine can lead to hypoglycemia by increasing glucose uptake in muscle cells and stimulating glycogen synthesis (Halaby, Kaslein, & Yang, 2013). Thus, one should contemplate further monitoring of glycemic control in patients receiving an agent with blood glucose-lowering effects concurrently with an SSRI and chloroquine or hydroxychloroquine.

2.2.2. Serotonin-norepinephrine reuptake inhibitors

Serotonin-Norepinephrine Reuptake Inhibitors (SNRIs) are used to treat depressive disorders (e.g. dysthymia), anxiety disorders (e.g. panic disorder, general anxiety disorder, and social anxiety disorder), and chronic pain syndromes, such as diabetic peripheral neuropathy (Nelson, 2019).

Venlafaxine is metabolized mainly by CYP2D6 and CYP3A4 and partially by CYP2C19. It does not affect (inhibit or induce) CYP450 microsomes. Venlafaxine
may cause arrhythmia associated with QTc prolongation in high-risk people (e.g., those who use drugs that prolong QT intervals like chloroquine and hydroxychloroquine, preexisting cardiac disease, old age, female sex, electrolyte imbalance, etc.). In the case of concurrent use of several medications associated with QT interval prolongation, close monitoring, and careful clinical assessment should be done (Howell, Wilson, & Waring, 2007). In a study investigating 21 healthy participants on venlafaxine and potent CYP3A4 inhibitor ketoconazole, it was found that levels of venlafaxine increased with the concomitant use of ketoconazole (Lindh, Annas, Meurling, Dahl, & Ayman, 2003). Also, ritonavir has a strong inhibitory effect on CYP3A4.

Duloxetine is metabolized by CYP1A2 and CYP2D6. It has a moderate inhibitory effect on CYP2D6. In a study ion the effect of duloxetine on CYP2D6, healthy volunteers used desipramine (CYP2D6 substrate) and paroxetine (a CYP2D6 inhibitor) along with duloxetine. The results indicated that desipramine levels were 2.9 times higher than when administered alone. On the other hand, concomitant use of duloxetine with paroxetine lead to increased levels of duloxetine. Side effects of duloxetine alone or in combination with desipramine or paroxetine were similar. Chloroquine may affect the concentration of duloxetine by inhibiting CYP2D6 ( Skinner et al., 2003). Despite the inhibitory effect on CYP2D6 and induced CYP1A2 by ritonavir, concomitant use of duloxetine with ritonavir is allowed (Smolders et al., 2016).

2.2.3. Serotonin modulators

Trazodone is used for the treatment of major depressive disorder, anxiety disorder, and obsessive-compulsive disorder (Bossini et al., 2015). The metabolism of trazodone in humans is mediated through CYP3A4 and CYP2D6. Trazodone does not affect the activity of CYP isozymes (Zalma et al., 2000). Trazodone peak plasma concentration and the AUC significantly increase 34% and 137%, respectively in concomitant use with ritonavir (a CYP3A4 inhibitor). Trazodone elimination half-life was prolonged 122% and oral clearance decreased 52% by ritonavir. Drowsiness, lethargy, and performance impairment also increased in co-administration with ritonavir, also three subjects experienced nausea, dizziness, and hypotension (Zalma et al., 2000). In a case report, a patient was treated with fluoxetine, diazepam, and trazodone went to the emergency room with sedation, dizziness, and nausea after taking 60 units trazodone, which showed a QTc prolongation in her ECG (Levenson, 1999). Thus, additive effects (e.g. some potent CYP450 3A4 inhibitors like ritonavir, chloroquine, hydroxychloroquine) may result in QTc prolongation when used with trazodone (Tönnesmann et al., 2013).

2.2.4. Norepinephrine-serotonin modulator

Mirtazapine is metabolized by CYP1A2, CYP3A4, and CYP2D6 (Gillman, 2007) and ritonavir boosted lopinavir is a CYP3A4 inhibitor (Lim et al., 2004). Clinically significant interactions between mirtazapine and ritonavir-boosted lopinavir have not been reported. However, it is suggested that the lowest efficient dose of mirtazapine be used when it is co-administered with CYP3A4 inhibitors (Goodlet, Zmarlicka, & Peckham, 2019). No interaction has been found between mirtazapine and hydroxychloroquine, chloroquine, ribavirin, and interferons.

2.2.5. Norepinephrine-dopamine reuptake inhibitor

Bupropion is classified as a dopamine norepinephrine reuptake inhibitor and used to treat major depression and other mood disorders (Horsït & Preskorn, 1998). Bupropion is metabolized by CYP2B6 isoenzyme to hydroxybupropion (Spina, Pisani, & de Leon, 2016) and CYP2B6 inducers such as ritonavir may decrease its serum concentration (Hesse, von Moltke, Shader, & Greenblatt, 2001). In one study on 12 healthy volunteers, the ritonavir boosted lopinavir at the standard dose (400 mg/100 mg twice daily) and reduced the bupropion AUC by 57% (Hogeland et al., 2007). Another study showed the ritonavir administration at high (600 mg twice daily) and low (100 mg twice daily) doses resulted in 67% and 22% reductions in bupropion concentrations, respectively (Park et al., 2010). This interaction can occur only 3 days after starting ritonavir (Kharasch, Mitchell, Coles, & Blanco, 2008). In contrast, one study indicated that short-term use of ritonavir had minimal impact on bupropion pharmacokinetics (Hesse, Greenblatt, von Moltke, & Court, 2006). We suggest that the clinical effectiveness of bupropion is monitored and a higher dose may be required.

No study or case report has documented the DDI between bupropion and chloroquine but theoretically, there is a pharmacokinetic interaction. Bupropion is a strong CYP2D6 inhibitor (Kotlyar et al., 2005; Xue, Zhang, & Cai, 2017) and chloroquine is a CYP2D6 substrate. Thus bupropion may decrease the metabolism of chloroquine. We think that this interaction may be clinically unimportant because chloroquine can be metabolized by other isoenzymes like CYP2C8 and CYP3A4 with low-affinity, high-capacity systems (Projean et al., 2003). There is no study and case report about bupropion interaction with other drugs that are used for the treatment of COVID-19.
2.2.6. Tricyclic antidepressants

In 1958, Tricyclic Antidepressants (TCAs) were introduced to the world by using imipramine for melancholic depression (Kuhn, 1958). This class was the first-line treatment for depression until the SSRIs were introduced. Nevertheless, tricyclic antidepressants continue to be used to treat many other psychiatric disorders there with depression (Gelenberg et al., 2010). The principle site for metabolism and elimination of TCAs is the liver. These drugs are metabolized through the hepatic CYP isoenzymes, mainly CYP2D6, CYP1A2, CYP3A4, and CYP1C19 (Gillman, 2007). Since protease inhibitors such as ritonavir can inhibit CYP2D6 and CYP3A4, the serum concentration of TCAs may increase. In one study on the effect of low-dose ritonavir on the activity of cytochrome P450, the CYP2D6 dose of 100 mg twice daily of ritonavir increased 26% in the AUC of desipramine. Low-dose ritonavir which is present in combination product of lopinavir/ritonavir has a medium inhibitory effect on the function of CYP2D6. Therefore, it seems that dose reduction is not necessary and only the patient should be monitored for toxic effects of TCAs (Aarnoutse et al., 2005).

No interaction has been reported between TCAs and hydroxychloroquine, although it is possible due to the risk of QT interval prolongation with both hydroxychloroquine and TCAs. So, there is a potential drug-drug interaction between them (Norman, Dunn, Bakker, Allen, & De Albuquerque, 2013). The same is true for chloroquine.

2.2.7. Monoamine Oxidase Inhibitors (MAOIs)

Monoamine Oxidase Inhibitors (MAOIs) are not the first-line drugs for the treatment of depression because of their side effect profile and drug-drug and food-drug interactions. Nevertheless, this class may be used in resistant depression (Yamada & Yasuhara, 2004). No interaction has been reported between MAOIs and hydroxychloroquine, but several articles reported the hypoglycemic effect of MAOIs (Cooper & Ashcroft, 1966; Wickström & Pettersson, 1964) and hydroxychloroquine (Cansu & Korkmaz, 2008; De-Heer & Doherty, 2018; Shojania, Koehler, & Elliott, 1999; Unübol et al., 2011). Hypothetically, their co-administration may lead to hypoglycemia. Therefore, it is recommended that susceptible patients be monitored for signs and symptoms of hypoglycemia. Similar to hydroxychloroquine, the effect of lowering blood glucose has been reported for chloroquine (Goyal & Borda, 1995; Mostafavi, 1998; Phillips et al., 1986). Thus, concomitant use of chloroquine with MAOIs may cause hypoglycemia like hydroxychloroquine. So far no study has not been published about the interaction between MAOIs and protease inhibitors, ribavirin, and interferons.

Among antidepressants, fluoxetine and paroxetine have the greatest inhibitory effect on CYP2D6; fluvoxamine strikingly inhibits CYP1A2 and CYP2C19. Hence, when these medications are prescribed with substrates of the relevant isozymes together, clinically interactions may be anticipated, especially those with a narrow therapeutic range. Duloxetine, sertraline, and bupropion inhibit CYP2D6 moderately, but sertraline at a high dose may have more potency to inhibit CYP2D6. Citalopram, escitalopram, venlafaxine, mirtazapine have a weak or negligible inhibitory effect on CYP isozymes in vitro, so significant interactions are less likely to occur with concomitant use of this cytochrome’s substrates (Spina, Santoro, & D’Arrigo, 2008).

2.2.8. Newer antidepressants

Lopinavir and ritonavir can potentially increase the level of vortioxetine by affecting CYP2D6 (Spina, Pisani, & de Leon, 2016). The clinical significance of this interaction is unknown as there is no published report about their co-administration.

Similarly, both ritonavir and lopinavir can increase the level of vilazodone by affecting the CYP3A4 (Spina, Pisani, & de Leon, 2016). The interaction may be serious and 50% vilazodone dose reduction or using an alternative is suggested (Boinpally, Gad, Gupta, & Pericelou, 2014). There is no interaction between the above-mentioned antidepressants and chloroquine or hydroxychloroquine.

2.2.9. Mood stabilizers

Mood stabilizers, including lithium, lamotrigine, valproic acid, and carbamazepine are mainstay drugs for the treatment of patients with bipolar disorders in the acute and maintenance phase (Gould, Chen, & Manji, 2002).

2.2.10. Lamotrigine

Lopinavir may decrease the serum concentration of lamotrigine because lopinavir/ritonavir induces Uridine Diphosphate Glucuronosyltransferases (UGTs), enzymes responsible for lamotrigine metabolism. Lamotrigine efficacy and toxicity symptoms, including skin rash and hematologic toxicities should be closely monitored in their co-administration. One study on 24 healthy participants who received lamotrigine (100 mg BD) in combination with
Kaletra (400 mg/100 mg BD) showed a 50% reduction in half-life, AUC, and Cmin. Also, lamotrigine clearance increased by approximately 85% (van der Lee et al., 2006). Accordingly, concomitant administration of lamotrigine with Kaletra that is a glucuronidation inducer may decrease lamotrigine concentrations so dose increase is recommended for dose titration.

2.2.11. Valproic acid

Valproic acid is metabolized by three pathways of extensive UGT glucuronidation (UGT1A6, UGT1A9, UGT2B7), mitochondrial β-oxidation, and minimal cytochrome P450-mediated oxidation. UGT enzyme inhibitors or inducers can increase or reduce serum concentrations of valproic acid, respectively. Sheehan NL et al. reported a patient with bipolar disorder in maintenance treatment with valproic acid who experienced mania symptoms a few days after initiating antivirals regimen ritonavir and lopinavir, potentially due to decrease in the serum concentration of valproate product (Ethell, Anderson, & Burchell, 2003; Sheehan, Brouillette, Delisle, & Allan, 2006). Also according to the NIH guidelines, lopinavir AUC increases by approximately 75% with concomitant valproic acid use. Therefore, patients under combination therapy should be monitored for lopinavir toxicity and virologic response (Dybul, Fauci, Bartlett, Kaplan, & Pau, 2002). In two studies, the reduced valproate concentrations were observed in patients treated with valproic acid and lopinavir/ritonavir (DiCenzo et al., 2004; Sheehan et al., 2006).

2.2.12. Carbamazepine

Carbamazepine is both a substrate and a potent inducer of CYP3A4. Higher doses of lopinavir as a substrate for CYP450-3A4 may be necessary in combination with carbamazepine and close monitoring should be provided (Hill & Lee, 2013).

Chloroquine is metabolized via CYP3A4 (Ducharme & Farinotti, 1996). Therefore, concomitant use of chloroquine with a CYP3A4 inducer, such as carbamazepine appears to reduce chloroquine levels and dose adjustment is offered.

2.2.13. Lithium

Lithium is excreted unchanged in the urine and has no known effects on CYP enzymes. Hence it is the least likely mood stabilizer that has DDIs with antiretroviral drugs (Thompson, Silverman, Dzeng, & Treisman, 2006). However, it is rational to inspect patients for interactions. No interaction has been reported between mood stabilizers with ribavirin and hydroxychloroquine.

2.2.14. Antipsychotics

Antipsychotics are the main drugs for the treatment of schizophrenia. The principle site of metabolism of most antipsychotics (the first and second-generation) is CYP isoenzymes, in particular CYP3A4 and CYP2D6. Therefore, DDIs in combined use with drugs that have a strong effect on CYP pathway metabolism such as CYP450, CYP3A4, CYP2D6 inducer, and inhibitors are possible (Hill & Lee, 2013; Lally & MacCabe, 2015). Protease inhibitors such as Kaletra (ritonavir/lopinavir) are strong CYP3A4 inhibitors and affect the activity of cytochrome resulting in antipsychotic accumulation and toxicity (Corbett, Lim, & Kashuba, 2002; Jernigan, Kipp, Rather, Jenkins, & Chung, 2013; Wyen et al., 2008).

2.2.15. Aripiprazole

Aripiprazole is a substrate of CYP2D6 and CYP3A4, and ritonavir is a strong inhibitor of these two enzymes (Azuma et al., 2012; Hendset, Hermann, Lunde, Refsum, & Molden, 2007; Kirby et al., 2011; Preskorn, 2012; Sheehan, Sliwa, Amatniek, Grinspan, & Canuso, 2010; Suzuki et al., 2011; Uritchuk, Prior, Dursun, & Baker, 2008; Wyen et al., 2008). Concomitant use of aripiprazole and ritonavir has the potential additive effect on metabolic disturbances and increases the adverse/toxic effects of ritonavir that should be monitored. Aripiprazole dose adjustment (reduced dose) is often appropriate in patients with CYP2D6 poor metabolizers or when combined with a CYP2D6 inhibitor or a CYP3A4 inhibitor (Hill & Lee, 2013).

2.2.16. Olanzapine

The serum concentration of olanzapine may be decreased by ritonavir (dose depending) because of ritonavir induction of CYP1A2 and or glucuronosyltransferase-mediated olanzapine metabolism. In a study on 14 healthy volunteers, ritonavir dose tapered up gradually and reduced the olanzapine Cmax, AUC, and half-life by 40%, 53%, and 50%, respectively (Hill & Lee, 2013; Jacobs, Colbers, Velthoven-Graafland, Schouwenberg, & Burger, 2014; Penzak et al., 2002).

2.2.17. Quetiapine

Ritonavir is a strong CYP3A4 inhibitor that has an inhibitory effect on quetiapine metabolism and may increase its serum concentration and adverse reactions.
(Hill & Lee, 2013; Pollack, McCoy, & Stead, 2009). Therefore, it is best to reduce the dose of quetiapine and monitor it closely.

2.2.18. Ziprasidone

Protease inhibitors (e.g. ritonavir) with CYP3A4 inhibitory property can increase the level of ziprasidone. Therefore, it needs more monitoring (Hill & Lee, 2013; Manini, Raspberry, Hoffman, & Nelson, 2007).

2.2.19. Clozapine

There are overlap metabolic toxicities (e.g. hyperglycemia, weight gain, hyperlipidemia) with clozapine and all of the protease inhibitors. So their administration should be monitored or alternative agents be considered (Hill & Lee, 2013)

2.2.20. Chlorpromazine

Chlorpromazine is a typical antipsychotic agent that was first developed in 1951. Chlorpromazine is a high-risk drug for QT-prolongation and chloroquine increased its serum concentrations 3 hours after an oral dose in patients under treatment (Makanjuola, Dixon, & Oforah, 1988).

2.2.21. Pimozide

Pimozide is an antipsychotic and CYP3A4 substrate. Its co-administration with ritonavir should be avoided because of increasing pimozide levels and the risk of QTc prolongation (Hill & Lee, 2013). QT prolongation is a cardiovascular side effect of pimozide (Krähenbühl et al., 1995). Therefore, we should avoid concomitant use with medications that can affect QT interval (e.g. chloroquine).

2.2.22. Risperidone

CYP2D6 and CYP3A4 are the main sites for risperidone metabolism. Ritonavir is a strong inhibitor for both enzymes. Studies have shown that co-administration of these drugs increases the side effects of risperidone (Jover, Cuadrado, Andreu, & Merino, 2002; Kelly, Béique, & Bowmer, 2002). Theoretically, concomitant use of antipsychotics like ziprasidone and quetiapine with chloroquine or hydroxychloroquine can increase the risk of QTc prolongation. No interaction has been reported between antipsychotic drugs and ribaviirin.

2.3. Anxiolytics

2.3.1. Benzodiazepines

Benzodiazepines (BZDs) are a group of psychotropic medications with sedative, hypnotic, and anxiolytic effects that are metabolized via CYP450 enzymes particularly subtype CYP3A4 in the liver (Griffin, Kaye, Bueno, & Kaye, 2013; Riss, Cloyd, Gates, & Collins, 2008). Therefore, medications with CYP3A4 inhibitory effect like Kaletra (ritonavir/lopinavir) that is a strong CYP3A4 inhibitor may decrease the metabolism of a CYP3A4 substrate such as chlorzoxazone, diazepam, clonazepam, flurazepam, triazolam, midazolam, and alprazolam. The patient should be monitored closely and even using an alternative agent with lower interaction (e.g. lorazepam, oxazepam, and temazepam) or dose adjustments should be considered (Dybul et al., 2002).

2.3.2. Midazolam

Midazolam is a CYP3A4 substrate (Leiri et al., 2013; Kronbach, Mathys, Umeno, Gonzalez, & Meyer, 1989). Some studies have reported AUC increase (5 to 28 fold) or reduction in oral and intravenous midazolam clearance (65%-96%) after co-administration with ritonavir or Kaletra (ritonavir/lopinavir) (Greenblatt et al., 2000; Greenblatt et al., 2003; Knox et al., 2008; Mathias, West, Hui, & Kearney, 2009; Schmitt, Hofmann, Riek, Zwanziger, & Patel, 2009; Yeh et al., 2006).

2.3.3. Diazepam

Diazepam is metabolized via CYP3A4 (Griffin, Kaye, Bueno, & Kaye, 2013; Manini, Raspberry, Hoffman, & Nelson, 2007). Some studies have reported increased serum concentrations of diazepam after co-administration with ritonavir or Kaletra (ritonavir/lopinavir) (Greenblatt et al., 2000; Greenblatt et al., 2003; Knox et al., 2008; Mathias, West, Hui, & Kearney, 2009; Schmitt, Hofmann, Riek, Zwanziger, & Patel, 2009; Yeh et al., 2006).

2.3.4. Alprazolam

Alprazolam is a benzodiazepine metabolized via CYP3A4, and to a smaller extent via CYP2C19 (Kakuda, Schöller-Gyüre, & Hoetelmans, 2011). Therefore co-administration of diazepam with ritonavir can decrease diazepam metabolism and in combination therapy, the lower dose of diazepam is recommended (Dybul et al., 2002).

2.3.5. Triazolam

Triazolam is metabolized via CYP3A4 (Griffin, Kaye, Bueno, & Kaye, 2013; Manini, Raspberry, Hoffman, & Nelson, 2007). Some studies have reported increased serum concentrations of triazolam after co-administration with ritonavir or Kaletra (ritonavir/lopinavir) (Greenblatt et al., 2000; Greenblatt et al., 2003; Knox et al., 2008; Mathias, West, Hui, & Kearney, 2009; Schmitt, Hofmann, Riek, Zwanziger, & Patel, 2009; Yeh et al., 2006).

Hill & Lee, 2013; Pollack, McCoy, & Stead, 2009). Therefore, it is best to reduce the dose of quetiapine and monitor it closely.
Table 1. Summary of drug-drug interaction between psychotropics and pharmacological treatments used for COVID-19

| Psychotropic Medication | Lopinavir/Ritonavir | Chloroquine/hydroxychloroquine | Implication |
|-------------------------|---------------------|-------------------------------|-------------|
| Antidepressants         |                     |                               |             |
| Fluoxetine              | ↑ Ritonavir via CYP2D6 and CYP3A4 | Risk of hypoglycemia | Caution in patients with diabetes |
| Sertraline              | ↑ Sertraline via CYP3A4 | Risk of hypoglycemia |             |
| SSRIs                   | Citalopram          | ↑QTc                          | Cardiac monitoring especially in high-risk patients |
|                         | Escitalopram        | ↑QTc                          | Cardiac monitoring especially in high-risk patients |
|                         | Paroxetine          | ↑ Paroxetine via CYP2D6 and protein binding displacement | Risk of hypoglycemia |
|                         | Fluvoxamine         | ↑ Ritonavir via CYP3A4 | Risk of hypoglycemia |
| SNRIs                   |                     |                               |             |
| TCAs                    | ↑ TCAs via CYP2D6   | ↑QTc                          |    |
| MAOIs                   |                     |                               | Cardiac monitoring especially in high-risk patients dose reduction is not necessary, monitor TCA side effects |
| Bupropion               | ↓ Bupropion via CYP2B6 |                               | Monitor clinical effectiveness of bupropion |
| Trazodone               | ↑ Trazodone via CYP3A4 | ↑QTc                          | Cardiac monitoring especially in high-risk patients Sedation may also occur |
| Mirtazapine             | ↑ Mirtazapine via CYP3A4 |                               | Use the lowest efficient dose |
| Vortioxetine            | ↑ Vortioxetine via CYP2D6 |                               | Not clinically significant |
| Vilazodone              | ↑ Vilazodone via CYP3A4 |                               | Dose reduction by 50% or using and alternative is suggested |
| Mood stabilizers        | Valproic acid       | ↓ Valproate ↑ Lopinavir       | Monitor for lopinavir toxicity and virologic response |
|                         | Lamotrigine         | ↓ Lam via UGTs                | A dose increase is recommended |
|                         | Carbamazepine       | ↓ Lopinavir ↑ Chloroquine     | |
|                         | Lithium             |                               | |

Mohebbi, N., et al. (2020). Drug Interactions of Psychiatric and COVID-19. BCN, 11(2). Covid19, 185-200.
### Table: Psychotropic Medication Interactions with Lopinavir/Ritonavir and Chloroquine/Hydroxychloroquine

| Psychotropic Medication          | Lopinavir/Ritonavir | Chloroquine/Hydroxychloroquine | Implication                  |
|---------------------------------|---------------------|--------------------------------|------------------------------|
| **Antipsychotics**              |                     |                                |                              |
| Risperidone                     | ↑ Risperidone via CYP3A4 and 2D6 |                                | Dose reduction               |
| Aripiprazole                    | ↑ Aripiprazole via CYP3A4 and 2D6 |                                |                              |
| Olanzapine                      | ↓ Olanzapine via CYP1A2 | ↑QTc                         |                              |
| Quetiapine                      | ↑ Quetiapine via CYP3A4 | ↑QTc                         | Cardiac monitoring           |
| Ziprasidone                     | ↑ Que via CYP3A4     | ↑QTc                         |                              |
| Clozapine                       | Additive metabolic toxicities | ↑QTc                         | Consider alternative agents  |
| Chlorpromazine                  | ↑ Pimozide via CYP3A4 | ↑QTc                         | Contraindicated              |
| **Anxiolytics**                 |                     |                                |                              |
| Midazolam                       | ↑ Midazolam via CYP3A4 |                                | Contraindicated              |
| Diazepam                        | ↑ Diazepam via 3A4   |                                | Dose reduction               |
| Alprazolam                      | ↑ Alprazolam via 3A4 |                                | Monitor sedation and dose reduction |
| Triazolam                       | ↓ Triazolam via 3A4  |                                | Not clinically significant    |
| Zolpidem                        | ↑ Zolpidem via 3A4   |                                | Dose adjustment is not usually necessary |
| Buspirone                       | ↑ Buspirone via 3A4  |                                |                              |

**SSRIs**: Selective Serotonin Reuptake Inhibitors; **SNRIs**: Serotonin-Norepinephrine Reuptake Inhibitors; **TCAs**: Tricyclic Antidepressants; **MAOIs**: Monoamine Oxidase Inhibitors

with ritonavir (Greenblatt, von Moltke, Daily, Harmatz, & Shader, 1999).

#### 2.3.6. Non-benzodiazepine

Non-benzodiazepines are another sedative, hypnotic, and anxiolytic drugs, including zolpidem, eszopiclone, zaleplon, and ramelteon that is extensively metabolized via the CYP450 system. So, there is a potential for drug-drug interaction (Mandrioli, Mercolini, & Augusta Raggi, 2010).

#### 2.3.7. Zolpidem

Pathways for zolpidem metabolism are CYP3A4, CYP2C9, CYP1A2, CYP2D6, and CYP2C19 (Greenblatt et al., 2000; Wyen et al., 2008). The concomitant use of strong CYP3A4 inhibitors such as ritonavir may increase the serum concentration of zolpidem because CYP3A4 is an isoenzyme partially responsible for zolpidem metabolism (Valuck, 2004). In a pharmacokinetic study of healthy participants, ritonavir (200 mg twice daily for 2 days) increased the zolpidem (5 mg single dose bedtime) AUC, and Cmax 28% and 22%, respectively and no pharmacodynamic changes related to zolpidem effects were found (Greenblatt et al., 2000).

#### 2.3.8. Buspirone

Buspirone is an anxiolytic drug that its main pathway for metabolism is CYP3A4 isoenzyme. So one should be cautious and carefully monitor it in concomitant use with CYP3A4 inhibitors such as ritonavir. However, dose adjustment is not necessary (Clay & Adams, 2003).

No interaction has been reported between anxiolytics and ribavirin, hydroxychloroquine, and chloroquine.

We should consider that the inhibitory effect of ritonavir on CYP2D6 is dose-dependent. Low-dose ritona-
vir (100 mg two times a day) had only a slight effect on CYP2D6, but the curative dose of ritonavir (600 mg two times a day) had a greater effect (Smolders et al., 2016).

Azithromycin use, as one of the alternatives used in COVID-19 patients, needs caution because it may enhance the QTc-prolonging effect of QT-prolonging agents like citalopram, especially in patients at high risk (preexisting cardiac disease, old age, female sex, electrolyte imbalance, etc.) (Huang, Wu, Hsia, & Yin Chen, 2007; Kezerashvili, Khattak, Barsky, Nazari, & Fisher, 2007). No interaction has been reported between amantadine and chloroquine, hydroxychloroquine, ritonavir, and ribavirin. Table 1 summarizes the drug-drug interactions mentioned in this article.

3. Conclusion

Drug interactions are major challenges in the comorbidity of psychiatric disorders and COVID-19 infection resulting in the alteration of expected therapeutic outcomes. The risk of toxicity or lack of efficacy may occur due to a higher or lower plasma concentration of medications. Moreover, intensification of similar adverse effects might harm the patients. However, psychiatric medication could be used safely in combination with COVID-19 pharmacotherapy if medications were selected with the least possibility of interaction and also careful monitoring. Psychiatrists should be aware of psychotropic drug interactions for optimal pharmacotherapy in the management of mental problems in COVID-19 patients.

Bjernsson, T. D., Callaghan, J. T., Einolf, H. J., Fischer, V., Gan, L., Grimm, S., et al. (2003). The conduct of in vitro and in vivo drug-drug interaction studies: A Pharmaceutical Research and Manufacturers of America (PhRMA) perspective. Drug Metabolism and Disposition, 31(7), 815-32. [DOI:10.1124/dmd3.17.815] [PMID]

Boinpally, R., Gad, N., Gupta, S., & Periclou, A. (2014). Influence of CYP3A4 induction/inhibition on the pharmacokinetics of vilazodone in healthy subjects. Clinical Therapeutics, 36(11), 1638-49. [DOI:10.1016/j.clinthera.2014.08.003] [PMID]

Bossini, L., Colaccia, A., Casolaro, L., Benbow, J., Amodeo, G., De Giorgi, R., et al. (2015). Off-label trazodone prescription: Evidence, benefits and risks. Current Pharmaceutical Design, 21(23), 3343-51. [DOI:10.2174/1381612821650161902236] [PMID]

Cansu, D. U., & Korkmaz, C. (2008). Hypoglycaemia induced by hydroxychloroquine in a non-diabetic patient treated for RA. Rheumatology [Oxford, England], 47(3), 378-9. [DOI:10.1093/rheumatology/kem378] [PMID]

Chen, C. J., Sugiyama, K., Kubo, H., Huang, C., & Makino, S. (2004). Murine coronavirus nonstructural protein p28 arrests cell cycle in G0/G1 phase. Journal of Virology, 78(19), 10410-9. [DOI:10.1128/JVI.18.19.10410-10419.2004] [PMID] [PMCID]

Clay, P. G., & Adams, M. M. (2003). Pseudo-Parkinson disease secondary to ritonavir-buspirone interaction. Annals of Pharmacotherapy, 37(2), 202-5. [DOI:10.1177/106002800303700207] [PMID]

Cooper, A. J., & Ashcroft, G. (1966). Potentiation of insulin hypoglycaemia by MA 0.1. Antidepressant drugs. The Lancet, 287(7434), 407-9. [DOI:10.1016/S0140-6736(66)91399-7]

Corbett, A. H., Lim, M. L., & Kashuba, A. D. (2002). Kaletra (lopinavir/ritonavir). Annals of Pharmacotherapy, 36(7-8), 1193-203. [DOI:10.1345/aph.1A363] [PMID]

De-Heer, R., & Doherty, T. (2018). A case of hydroxychloroquine induced hypoglycaemia in a non-diabetic patient. Journal of Rheumatic Diseases and Treatment, 4(3), 066. [DOI:10.23937/2469-5726-1510066]

Valuck, R. (2004). Selective serotonin reuptake inhibitors: A class review. P&T Community, 29(4), 234-43. https://www.ptcommunity.com/journal/article/archives/2004/4/234/

References

Aarnoutse, R. E., Kleinnijenhuis, J., Koopmans, P. P., Touw, D. J., Wieling, J., Helster, Y. A., et al. (2005). Effect of low-dose ritonavir (100 mg twice daily) on the activity of cytochrome P450 3D6 in healthy volunteers. Clinical Pharmacology & Therapeutics, 78(6), 664-74. [DOI:10.1016/j.clpt.2005.09.001] [PMID]

Axelson, D. A., Perel, J. M., Birmaher, B., Rudolph, G. R., Nuss, S., Bridge, J., et al. (2002). Sertraline pharmacokinetics and dynamics in adolescents. Journal of the American Academy of Child & Adolescent Psychiatry, 41(9), 1037-44. [DOI:10.1097/00004583-200209000-00003] [PMID]

Azuma, J., Hasunuma, T., Kubo, M., Miyatake, M., Koue, T., Higashi, K., et al. (2012). The relationship between clinical pharmacokinetics of aripiprazole and CYP2D6 genetic polymorphism: Effects of CYP enzyme inhibition by coadministration of paroxetine or fluvoxamine. European Journal of Clinical Pharmacology, 68(1), 29-37. [DOI:10.1007/s00228-011-1094-4] [PMID] [PMCID]

De-Heer, R., & Doherty, T. (2018). A case of hydroxychloroquine induced hypoglycaemia in a non-diabetic patient. Journal of Rheumatic Diseases and Treatment, 4(3), 066. [DOI:10.23937/2469-5726-1510066]
DeVane, C. L., & Gill, H. S. (1997). Clinical pharmacokinetics of fluvoxamine: Applications to dosing regimen design. The Journal of Clinical Psychiatry, 58 Suppl 5, 7-14. [PMID]

Di Paola, L., & Giuliani, A. (2020). Mapping active allosteric loci SARS-CoV spike proteins by means of protein contact networks. arXiv:2003.05200. https://arxiv.org/abs/2003.05200

DiCenzo, R., Peterson, D., Crutenden, K., Morse, G., Riggs, G., Gelbard, H., et al. (2004). Effects of valproic acid coadministration on plasma efavirenz and lopinavir concentrations in human immunodeficiency virus-infected adults. Antimicrobial Agents and Chemotherapy, 48(11), 4328-31. [DOI:10.1128/AAC.48.11.4328-4331.2004] [PMID] [PMCID]

Ducharme, J., & Farinotti, R. (1996). Clinical pharmacokinetics and metabolism of chloroxazone. Clinical Pharmacokinetics, 31(4), 257-74. [DOI:10.2165/00023210-199631040-00003] [PMID]

Dubyl, M., Fauci, A. S., Bartlett, J. G., Kaplan, J. E., Pau, A. K., & Panel on Clinical Practices for Treatment of HIV. (2002). Guidelines for using antiretroviral agents among HIV-infected adults and adolescents. Annals of Internal Medicine, 137(5 Pt 2), 381-433. [DOI:10.7326/0003-4819-137-5_Part_2-20020903010001] [PMID]

Ehbell, B. T., Anderson, G. D., & Burchell, B. (2003). The effect of valproic acid on drug and steroid glucuronidation by expressed human UDP-glucuronosyltransferases. Biochemical Pharmacology, 65(9), 1441-9. [DOI:10.1016/j.bcp.2003.07.067] [PMID]

Frye, R., Bertz, R., Granneman, G. R., Qian, J., Lamm, J., Dennis, S., & Valdes, J. (1997). Effect of ritonavir on the pharmacokinetics and pharmacoodynamics of alprazolam. Paper presented at InterScience Conference on Antimicrobial Agents and Chemotherapy, Metro Toronto, Canada, 28 September-1 October 1997.

American Psychiatric Association. (2000). Practice guideline for the treatment of patients with major depressive disorder (revision). The American journal of psychiatry, 157(4 Suppl), 1-45.

Ghaeli, P., Shahsavand, E., Mesbah, M., Kamkar, M. Z., Sadeghi, M., & Dashi-Khavasadi, S. (2004). Comparing the effects of 8-week treatment with fluoxetine and imipramine on fasting blood glucose of patients with major depressive disorder. Journal of Clinical Psychopharmacology, 24(4), 386-8. [DOI:10.1097/00023210-200406000-00013] [PMID]

Gillman, P. K. (2007). Tricyclic antidepressant pharmacology and therapeutic drug interactions updated. British Journal of Pharmacology, 151(6), 737-48. [DOI:10.1038/sj.bjp.0707253] [PMID] [PMCID]

Goodyer, K. J., Zmarlicka, M. T., & Peckham, A. M. (2019). Drug-drug interactions and clinical considerations with co-administration of antiretrovirals and psychotropic drugs. CNS Spectrums, 24(3), 287-312. [DOI:10.1017/S109285981900113X] [PMID]

Gould, T. D., Chen, G., & Manji, H. K. (2002). Mood stabilizer psychopharmacology. Clinical Neuroscience Research, 2(3-4), 193-212. [DOI:10.1016/S1566-2772(02)00044-0]

Goyal, K., Chauhan, P., Chikara, K., Gupta, P., & Singh, M. P. (2020). Fear of COVID 19: First suicidal case in India. Asian Journal of Psychiatry, 49, 101989. [DOI:10.1016/j.ajp.2020.101989] [PMID] [PMCID]

Greenblatt, D. J., von Moltke, L. L., Harmatz, J. S., Durol, A. L., Daily, J. P., Graf, J. A., et al. (2000). Differential impairment of triazolam and zolpidem clearance by ritonavir. Journal of Acquired Immune Deficiency Syndromes, 24(2), 129-36. [DOI:10.1097/00126334-200006010-00007]

Greenblatt, D. J., von Moltke, L. L., Daily, J. P., Harmatz, J. S., & Shader, R. I. (1999). Extensive impairment of triazolam and alprazolam clearance by short-term low-dose ritonavir: The clinical dilemma of concurrent inhibition and induction. Journal of Clinical Psychopharmacology, 19(4), 293-6. [DOI:10.1097/00004714-199908000-00001] [PMID]

Greenblatt, D. J., von Moltke, L. L., Harmatz, J. S., Fogelman, S. M., Chen, G., Graf, J. A., et al. (2003). Short-term exposure to low-dose ritonavir impairs clearance and enhances adverse effects of trazodone. The Journal of Clinical Pharmacology, 43(4), 414-22. [DOI:10.1177/009127003251864] [PMID]

Griffin, C. E., Kaye, A. M., Bueno, F. R., & Kaye, A. D. (2013). Benzodiazepine pharmacology and central nervous system-mediated effects. The Ochsner Journal, 13(2), 214-23. [PMID] [PMCID]

Goyal, V., & Borda, A. (1995). The hypoglycemic effect of chloroquine. The Journal of the Association of Physicians of India, 43(1), 17-8. [PMID]

Heser, L. M., Greenblatt, D. J., von Moltke, L. L., & Court, M. H. (2006). Ritonavir has minimal impact on the pharmacokinetic disposition of a single dose of bupropion administered to human volunteers. The Journal of Clinical Pharmacology, 46(5), 567-76. [DOI:10.1177/0091270060626981] [PMID]

Heser, L. M., von Moltke, L. L., & Greenblatt, D. J. (2003). Clinically important drug interactions with zopiclone, zolpidem and zaleplon. CNS Drugs, 17(7), 513-32. [DOI:10.2165/00032310-20031707-00004] [PMID]
Hesse, L. M., von Moltke, L. L., Shador, R. L., & Greenblatt, D. J. (2003). Ritonavir, efavirenz, and nelfinavir inhibit CYP2B6 activity in vitro: Potential drug interactions with bupropion. Drug Metabolism and Disposition, 31(2), 100-2. [PMID]

Hill, L., & Lee, K. C. (2013). Pharmacotherapy considerations in patients with HIV and psychiatric disorders: Focus on antidepressants and antipsychotics. Annals of Pharmacotherapy, 47(1), 75-89. [DOI:10.1345/aph.1R343] [PMID]

Hogeland, G. W., Swindells, S., McNabb, J. C., Kashuba, A. D., Yee, G. C., & Lindley, C. M. (2007). Lopinavir/ritonavir reduces bupropion plasma concentrations in healthy subjects. Clinical Pharmacology & Therapeutics, 81(1), 69-75. [DOI:10.1038/sj.clpt.6100272] [PMID]

Horst, W. D., & Preskorn, S. H. (1998). Mechanisms of action and clinical characteristics of three atypical antidepressants: Venlafaxine, nefazodone, bupropion. Journal of Affective Disorders, 51(3), 257-54. [DOI:10.1016/S0165-0327(98)00222-5]

Hou, X. J., Xu, J. H., Wang, J., & Yu, Y. Y. (2013). Can antidepressants prevent pegylated interferon-alpha/ribavirin-associated depression in patients with chronic hepatitis C? Meta-analysis of randomized, double-blind, placebo-controlled trials? PLoS One, 8(10), e76799. [PMID]

Hou, X. J., Xu, J. H., Wang, J., & Yu, Y. Y. (2013). Can antidepressants prevent pegylated interferon-alpha/ribavirin-associated depression in patients with chronic hepatitis C? Meta-analysis of randomized, double-blind, placebo-controlled trials? PLoS One, 8(10), e76799. [PMID] [PMCID]

Howell, C., Wilson, A. D., & Waring, W. (2007). Cardiovascular toxicity due to venlafaxine poisoning in adults: A review of 235 consecutive cases. British Journal of Clinical Pharmacology, 64(2), 192-7. [DOI:10.1111/j.1365-2125.2007.02849.x] [PMID] [PMCID]

Huang, B. H., Wu, C. H., Hsia, C. P., & Chen, C. Y. (2007). Azithromycin-induced torsades de pointes. Pacing and Clinical Electrophysiology, 30(12), 1579-82. [DOI:10.1111/j.1540-8159.2007.00912.x] [PMID]

Ieiri, I., Tsunemitsu, S., Maeda, K., Ando, Y., Izumi, N., Kimura, M., et al. (2013). Mechanisms of pharmacokinetic enhancement between ritonavir and saquinavir: micro/small dosing tests using midazolam (CYP3A4), fexofenadine (P-glycoprotein), and pravastatin (OATP1B1) as probe drugs. The Journal of Clinical Pharmacology, 53(6), 654-61. [DOI:10.1002/jcph.62] [PMID]

Jacobs, B. S., Colbers, A. P., Veltkoven-Graafland, K., Schouwenberg, B. J., & Burger, D. M. (2014). Effect of fosamprenavir/ritonavir on the pharmacokinetics of single-dose olanzapine in healthy volunteers. International Journal of Antimicrobial Agents, 44(2), 173-7. [DOI:10.1016/j.ijantimicag.2014.03.014] [PMID]

Jernigan, M. G., Kipp, G. M., Rather, A., Jenkins, M. T., & Chung, A. M. (2013). Clinical implications and management of drug-drug interactions between antiretroviral agents and psychotropic medications. Mental Health Clinician, 2(9), 274-85. [DOI:10.9740/mhcclin139874]

Jin, X., Potter, B., Luong, T. L., Nelson, J., Vuong, C., Potter, C., et al. (2016). Pre-clinical evaluation of CYP2D6 dependent drug-drug interactions between primaquine and SSR1/SNRI antidepressants. Malaria Journal, 15(1), 280. [DOI:10.1186/s12936-016-1329-z] [PMID] [PMCID]

Jornil, J., Jensen, K. G., Lansen, F., & Linnet, K. (2010). Identification of cytochrome P450 isoforms involved in the metabolism of paroxetine and estimation of their importance for human paroxetine metabolism using a population-based simulator. Drug Metabolism and Disposition, 38(3), 376-85. [DOI:10.1124/dmd.109.030551] [PMID]

Jover, F., Cuadrado, J. M., Andreu, L., & Merino, J. (2002). Reversible coma caused by risperidone-ritonavir interaction. Clinical Neuropharmacology, 25(5), 251-3. [DOI:10.1097/00002826-200209000-00004]

Kakuda, T. N., Schiller-Gyure, M., & Hoetelmans, R. M. W. (2011). Pharmacokinetic interactions between etravirine and non-antiretroviral drugs. Clinical Pharmacokinetics, 50(1), 25-39. [DOI:10.2165/11534740-000000000-00000] [PMID]

Kelly, D. V., Beique, L. C., & Bowmer, M. I. (2002). Extrapyramidal symptoms with ritonavir/indinavir plus risperidone. Annals of Pharmacotherapy, 36(5), 827-30. [DOI:10.1345/aph.1A335] [PMID]

Kezerashvili, A., Khattak, H., Barsky, A., Nazari, R., & Fisher, J. D. (2007). Azithromycin as a cause of QT-interval prolongation and torsade de pointes in the absence of other known precipitating factors. Journal of Interventional Cardiac Electrophysiology, 18(3), 243-6. [DOI:10.1007/s10840-007-9124-y] [PMID]

Kharasch, E. D., Mitchell, D., Coles, R., & Blanco, R. (2008). Rapid clinical induction of hepatic cytochrome P4502B6 activity by ritonavir. Antimicrobial Agents and Chemotherapy, 52(6), 1663-9. [DOI:10.1124/ AAC.107.115067] [PMID] [PMCID]

Khoza, S., & Barner, J. C. (2011). Glucose dysregulation associated with antidepressant agents: An analysis of 17 published case reports. International Journal of Clinical Pharmacy, 33(3), 484-92. [DOI:10.1007/s11096-011-9507-0] [PMID]

Kim, J. A. A., Chang, J. S., & Abela, G. S. (2013). An Association of Takotsubo Cardiomyopathy and Use of Antidepressants. Paper presented at 18th World Congress on Heart Disease, Vancouver, Canada, 26-29 July 2013.

Kirby, B. J., Collier, A. C., Kharasch, E. D., Whittington, D., Thummel, K. E., & Unadkat, J. D. (2011). Complex drug interactions of HIV protease inhibitors 1: Inactivation, induction, and inhibition of cytochrome P450 3A4 by ritonavir or nelfinavir. Drug Metabolism and Disposition, 39(6), 1070-8. [DOI:10.1124/dmd.110.037522] [PMID] [PMCID]

Knox, T. A., Olesen, L., von Moltke, L. L., Kaufman, R., C. W., Wanke, C. A., & Greenblatt, D. J. (2008). Ritonavir greatly impairs CYP3A activity in HIV infection with chronic viral hepatitis. JAIDS Journal of Acquired Immune Deficiency Syndromes, 49(4), 358-68. [DOI:10.1097/QAI.0b013e31818f7674] [PMID]

Kotlyar, M., Brauer, L. H., Tracy, T. S., Hutsukami, D. K., Harris, J., Bronars, C. A., et al. (2005). Inhibition of CYP2D6 activity by bupropion. Journal of Clinical Psychopharmacology, 25(3), 226-9. [DOI:10.1097/01.jcp.0000162805.46453.e3] [PMID]

Krähenbühl, S., Sauter, B., Kupferschmid, H., Krause, M., Wyss, P. A., & Meier, P. J. (1995). Reversible QT prolongation with torsades de pointes in a patient with pimozone intoxication. The American Journal of the Medical Sciences, 309(6), 315-6. [DOI:10.1097/00002826-199506000-00003] [PMID]

Kronbach, T., Mathys, D., Umeno, M., Gonzalez, F. J., & Meyer, U. A. (1989). Oxidation of midazolam and triazolam by hu-
man liver cytochrome P450IIA4. Molecular Pharmacology, 36(1), 89-96. http://molpharm.aspetjournals.org/content/36/1/89

Kuhn, R. (1958). The treatment of depressive states with G 22355 (imipramine hydrochloride). American Journal of Psychiatry, 115(5), 459-64. [DOI:10.1176/ajp.115.5.459] [PMID]

Lai, C. C., Shih, T. P., Ko, W. C., Tang, H. J., & Hsueh, P. R. (2020). Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) and corona virus disease-2019 (COVID-19): The epidemic and the challenges. International Journal of Antimicrobial Agents, 55(3), 105924. [DOI:10.1016/j.ijantimicag.2020.105924] [PMID] [PMCID]

Lally, J., & MacCabe, J. H. (2015). Antipsychotic medication in hypnotic drugs: An analytical point of view. Current Drug Metabolism, 16(10), 1732-8. [DOI:10.2174/138920010794328887] [PMID] [PMCID]

Manini, A. F., Raspberry, D., Hoffman, R. S., & Nelson, L. S. (2007). QT prolongation and torsades de pointes following overdose of ziprasidone and amantadine. Journal of Medical Toxicology, 3(4), 178-81. [DOI:10.1007/BF03160936] [PMID] [PMCID]

Margolis, J. M., O’Donnell, J. P., Mankowski, D. C., Ekins, S., & Obach, R. S. (2000). (R)-, (S)-, and racemic fluoxetine N-demethylation by human cytochrome P450 enzymes. Drug Metabolism and Disposition, 28(10), 1187-91. [PMID]

Mathias, A. A., West, S., Hui, J., & Kearney, B. P. (2009). Dose-response of ritonavir on hepatic CYP3A activity and elvitra-grav oral exposure. Clinical Pharmacology & Therapeutics, 85(1), 64-70. [DOI:10.1038/clpt.2008.168] [PMID]

Meid, A. D., Bighelli, I., Mächler, S., Mikus, G., Carrà, G., Castellazzi, M., et al. (2017). Combinations of QTc-prolonging drugs: Towards disentangling pharmacokinetic and pharmacodynamic effects in their potentially additive nature. Therapeutic Advances in Psychopharmacology, 7(12), 251-64. [DOI:10.1177/204512317216622] [PMID] [PMCID]

Mostafavi, H. (1998). Effect of chloroquine on blood glucose levels in patients with non insulin dependent diabetes mellitus. Acta Medica Iranica, 36(2), 109-12. http://acta.turns.ac.ir/index.php/acta/article/view/1753

Nelson, C. (2020). Serotonin-Norepinephrine Reuptake Inhibitors (SNRIs): Pharmacology, administration, and side effects. Retrieved from https://www.uptodate.com/contents-serotonin-norepinephrine-reuptake-inhibitors-suris-pharmacology-administration-and-side-effects

Nguyen, H. C., Nguyen, M. H., Do, B. N., Tran, C. Q., Nguyen, T. T. P., Pham, K. M., et al. (2020). People with suspected COVID-19 symptoms were more likely depressed and had lower health-related quality of life: The potential benefit of health literacy. Journal of Clinical Medicine, 9(4), E965. [DOI:10.3390/jcm9040965] [PMID]

Norman, E. S., Dunn, G., Bakker, K., Allen, D. M., & De Albuquerque, R. C. (2013). Water security assessment: Integrating governance and freshwater indicators. Water Resources Management, 27(2), 335-51. [DOI:10.1007/s11269-012-0200-4]

Ogu, C. C., & Maxa, J. L. (2000). Drug interactions due to cytochrome P450. Proceedings (Baylor University. Medical Center), 13(4), 421-3. [DOI:10.1080/08998280.2000.11927719] [PMID] [PMCID]

Ouellet, D., Hsu, A., Qian, J., Lamm, J. E., Cavanaugh, J. H., Leonard, J. M., et al. (1998). Effect of fluoxetine on pharmacokinetics of ritonavir. Antimicrobial Agents and Chemotherapy, 42(12), 3107-12. [DOI:10.1128/AAC.42.12.3107] [PMID] [PMCID]

Park, J., Vousden, M., Brittain, C., McConnell, D. J., Lavarone, L., Ascher, J., et al. (2010). Dose-related reduction in bupropion plasma concentrations by ritonavir. The Journal of Clinical Pharmacology, 50(10), 1180-7. [DOI:10.1177/0091270090935924] [PMID]

Penzak, S. R., Hon, Y. Y., Lawhorn, W. D., Shirley, K. L., Spratlin, V., & Jann, M. W. (2002). Influence of ritonavir on olanzapine pharmacokinetics in healthy volunteers. Journal of Clinical Psychopharmacology, 22(4), 366-70. [DOI:10.1097/00004714-200208000-00006] [PMID]

Phillips, R. E., Looareesuwan, S., White, N. J., Chanthavanich, P., Karbwang, J., Supanaranond, W., et al. (1986). Hypoglycaemia and antimalarial drugs: Quinidine and release of insulin. British Medical Journal (Clinical research ed.), 292(6531), 1319-21. [DOI:10.1136/bmj.292.6531.1319] [PMID] [PMCID]

Pollack, T. M., McCoy, C., & Stead, W. (2009). Clinically significant adverse events from a drug interaction between queti-a
pine at azanavir-ritonavir in two patients. Pharmacotherapy, 29(11), 1386-91. [DOI:10.1592/phco.29.11.1386] [PMID]

Preskorn, S. H. (1997). Clinically relevant pharmacology of selective serotonin reuptake inhibitors: An overview with emphasis on pharmacokinetics and effects on oxidative drug metabolism. Clinical Pharmacokinetics, 32 Suppl 1, 1-21. [DOI:10.2165/00003495-199732012-00003] [PMID]

Preskorn, S. H. (2012). Clinically important differences in the pharmacokinetics of the ten newer “atypical” antidepressants: Part 2. Metabolism and elimination. Journal of Psychiatric Practice, 18(3), 361-8. [DOI:10.1097/JPP.0b00008820268128. ad] [PMID]

Projean, D., Baune, B., Farinotti, R., Flinois, J. P., Beuneu, P., Taburet, A. M., et al. (2003). In vitro metabolism of chloroquine: Identification of CYP2C8, CYP3A4, and CYP2D6 as the main isoforms catalyzing N-desethylichloroquine formation. Atazanavir and Disposition, 31(6), 748-54. [DOI:10.1124/dndi.31.6.748] [PMID]

Riss, J., Cloyd, J., Gates, J., & Collins, S. (2008). Benzodiazepines and a substrate of cytochrome P4503A4 activity in healthy volunteers using midazolam as a probe. Pharmacotherapy, 29(10), 1175-81. [DOI:10.1592/phco.29.10.1175] [PMID]

Schmitt, C., Hofmann, C., Riek, M., Patel, A., & Zwanziger, E. (2009). Effect of saquinavir-ritonavir on cytochrome P450 3A4 activity in healthy volunteers using midazolam as a probe. Pharmacotherapy, 29(10), 1175-81. [DOI:10.1592/phco.29.10.1175] [PMID]

Sheehan, J., Sliva, J., Amatniek, J., Grinspan, A., & Canuso, C. (2010). Atypical antidepressant metabolism and excretion. Current Drug Metabolism, 11(6), 516-25. [DOI:10.2174/138920010971636202] [PMID]

Sheehan, N. L., Brouillette, M. J., Delisle, M. S., & Allan, J. (2006). Hypoglycemic episodes in a case of premenstrual dysphoric disorder. The Journal of Rheumatology, 26(1), 195-6. [DOI:10.1097/RHU.0b013e3182098e1f] [PMID]

Thompson, A., Silverman, B., Dzeng, L., & Treisman, G. (2006). Psychotropic medications and HIV. Clinical Infectious Diseases, 42(9), 1305-10. [DOI:10.1086/501454] [PMID]

Tönnesmann, E., Kandolf, R., & Lewalter, T. (2013). Chloroquine cardiomyopathy—a review of the literature. Immunopharmacology and Immunotoxicology, 35(3), 434-42. [DOI:10.3109/01492918.2013.780078] [PMID]

Tuite, A. R., Bogoch, I., Sherbo, R., Watts, A., Fisman, D. N., & Khan, K. (2020). Estimation of COVID-2019 burden and potential for international dissemination of infection from Iran. medRxiv. [DOI:10.1101/2020.02.24.20027357]

Untúbal, M., Ayhan, M., & Guney, E. (2011). Hypoglycemia induced by hydroxychloroquine in a patient treated for rheumatoid arthritis. Journal of Clinical Rheumatology, 17(1), 46-7. [DOI:10.1097/RHU.0b013e3182098e1f] [PMID]

Urichuk, L., Prior, T. I., Dursun, S., & Baker, G. (2008). Metabolism of atypical antidepressants: Involvement of cytochrome P450 enzymes and relevance for drug-drug interactions. Current Drug Metabolism, 9(5), 410-8. [DOI:10.2174/13892000878474673] [PMID]

van der Lee, M. J., Blenke, A. A. M., van der Lee, M. J., Blenke, A. A. M., Rongen, G. A., Verwey, A., van der Lee, M. J., Dawood, L., ter Hofstede, H. J. M., de Graaff, E. A. M., et al. (2007). Interaction study of the combined use of paroxetine and fosamprenavir-ritonavir in healthy subjects. An open-label, randomized, controlled, parallel-group study. Pharmacological Research, 55(11), 1094-109. [DOI:10.1016/j.phrs.2016.02.014] [PMID]

van der Lee, M. J., Dawood, L., ter Hofstede, H. J. M., van Ewijk, W., Wickström, L., & Pettersson, K. (1964). Treatment of diabetics with chloroquine and atazanavir-ritonavir in two patients. Pharmacotherapy, 29(11), 1386-91. [DOI:10.1592/phco.29.11.1386] [PMID]
Wijesinghe, R. (2016). A review of pharmacokinetic and pharmacodynamic interactions with antipsychotics. Mental Health Clinician, 6(1), 21-7. [DOI:10.9740/mhc.2016.01.021] [PMID] [PMCID]

Wu, Z., & McGowan, J. M. (2020). Characteristics of and important lessons from the coronavirus disease 2019 (COVID-19) outbreak in China: Summary of a report of 72,314 cases from the Chinese Center for Disease Control and Prevention. JAMA, 323(13), 1239-42. [DOI:10.1001/jama.2020.2648]

Wyen, C., Fuhr, U., Frank, D., Aarnoutse, R. E., Klaassen, T., Lazar, A., et al. (2008). Effect of an antiretroviral regimen containing ritonavir boosted lopinavir on intestinal and hepatic CYP3A, CYP2D6 and P-glycoprotein in HIV-infected patients. Clinical Pharmacology & Therapeutics, 84(1), 75-82. [DOI:10.1038/sj.clpt.6100452] [PMID]

Xue, C., Zhang, X., & Cai, W. (2017). Prediction of drug-drug interactions with bupropion and its metabolites as CYP2D6 inhibitors using a physiologically-based pharmacokinetic model. Pharmaceutics, 10(1), 1. [DOI:10.3390/pharmaceutics10010001] [PMID] [PMCID]

Yamada, M., & Yasuhara, H. (2004). Clinical pharmacology of MAO inhibitors: Safety and future. Neurotoxicology, 25(1-2), 215-21. [DOI:10.1016/S0161-813X(03)00097-4]

Yeh, R. F., Gaver, V. E., Patterson, K. B., Rezk, N. L., Baxter-Meheux, F., Blake, M. J., et al. (2006). Lopinavir/ritonavir induces the hepatic activity of cytochrome P450 enzymes CYP2C9, CYP2C19, and CYP1A2 but inhibits the hepatic and intestinal activity of CYP3A as measured by a phenotyping drug cocktail in healthy volunteers. Journal of Acquired Immune Deficiency Syndromes, 42(1), 52-60. [DOI:10.1097/01.qai.0000219774.20174.64] [PMID]

Zalma, A., von Moltke, L. L., Granda, B. W., Harmatz, J. S., Shader, R. I., & Greenblatt, D. J. (2000). In vitro metabolism of trazodone by CYP3A: Inhibition by ketoconazole and human immunodeficiency viral protease inhibitors. Biological Psychiatry, 47(7), 655-61. [DOI:10.1016/S0006-3223(99)00176-4]