

Grapefruit–medication interactions: Forbidden fruit or avoidable consequences?

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Our research group discovered the interaction between grapefruit and certain medications more than 20 years ago.^{1–3} Currently, more than 85 drugs, most of which are available in Canada, are known or predicted to interact with grapefruit. This interaction enhances systemic drug concentration through impaired drug metabolism.

Many of the drugs that interact with grapefruit are highly prescribed and are essential for the treatment of important or common medical conditions. Recently, however, a disturbing trend has been seen. Between 2008 and 2012, the number of medications with the potential to interact with grapefruit and cause serious adverse effects (i.e., torsade de pointes, rhabdomyolysis, myelotoxicity, respiratory depression, gastrointestinal bleeding, nephrotoxicity) has increased from 17 to 43, representing an average rate of increase exceeding 6 drugs per year. This increase is a result of the introduction of new chemical entities and formulations.

This review identifies the key scientific concepts and clinical implications of grapefruit–drug interactions relevant to medical practice. We focus on grapefruit because it is the most widely examined, but other citrus fruits may have similar consequences. It was recently found that grapefruit and certain other citrus juices act by an additional mechanism to cause diminished systemic concentration of certain drugs by inhibiting drug transporters.⁴ However, we will focus on the most well-known and well-studied interaction: that of interference in the activity of the cytochrome P450 3A4 (CYP3A4) enzyme. A summary of the evidence used in this review, which comes mainly from randomized controlled trials, is found in Box 1.

What are the key scientific concepts of grapefruit–drug interactions?

The actions of drugs are terminated through several biological mechanisms. The most important

is drug metabolism involving oxidation by enzymes belonging to the cytochrome P450 superfamily. Cytochrome P450 3A4 is particularly essential, because it is involved in the bioinactivation of about 50% of all drugs.⁵ CYP3A4 is located in epithelial cells (enterocytes) lining the small intestines and colon, and in the parenchymal cells of the liver (hepatocytes) (Figure 1). Consequently, orally administered drugs can be metabolized twice before reaching the systemic circulation. Thus, the percentage of drug absorbed unchanged (oral bioavailability) can be markedly attenuated. For example, the oral bioavailability of the antihypertensive drug felodipine is normally reduced to 15% of the oral dose.⁶ In other words, felodipine has low innate bioavailability. For this reason, it is subject to a potentially dramatic increase in systemic exposure and associated higher risk of overdose with grapefruit as a result of diminished CYP3A4 activity, primarily in the small intestine rather than in the liver.

The chemicals in grapefruit involved in this interaction are the furanocoumarins.⁷ Furanocoumarins are metabolized by CYP3A4 to reactive intermediates that bond covalently to the active site of the enzyme, causing irreversible inactivation (mechanism-based inhibition).⁸ Con-

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KEY POINTS

- Currently, more than 85 drugs have the possibility of interacting with grapefruit; of these drugs, 43 have interactions that can result in serious adverse effects.
- Drugs that interact with grapefruit have all of the following characteristics: they are administered orally, they have very low to intermediate absolute bioavailability, and they are metabolized by the cytochrome P450 3A4 enzyme (CYP3A4).
- All sources of grapefruit and certain related citrus fruits can irreversibly inhibit CYP3A4 in the gastrointestinal tract; to prevent this interaction, affected drugs should not be consumed with any of these fruits during the treatment period, or noninteracting alternative medications should be prescribed.
- Older patients have the greatest possibility of ingesting grapefruit and interacting medications and are the most vulnerable to the adverse clinical consequences.

sequently, CYP3A4 activity in the small intestine is impaired until de novo synthesis returns the enzyme to its previous level. This mechanism explains the important clinical effects on drug pharmacokinetics, specifically the peak plasma drug concentration (C_{max}) and the area under the drug concentration–time curve (AUC). These key parameters of oral bioavailability are increased, whereas systemic elimination half-life is unaltered. The pharmacokinetics of intravenously administered drugs are unchanged.^{9,10}

Because these chemicals are innate to grapefruit, all forms of the fruit (freshly squeezed juice, frozen concentrate and whole fruit) have the potential to reduce the activity of CYP3A4. One whole grapefruit or 200 mL of grapefruit juice is sufficient to cause clinically relevant increased systemic drug concentration and subsequent adverse effects.^{11,12} Seville oranges, (often used in marmalades), limes and pomelos also produce this interaction.^{13–15} Varieties of sweet orange, such as navel or valencia, do not contain furanocoumarins and do not produce this interaction.²

What determines which drugs are affected?

The interaction between medications and grapefruit is drug-specific and is not a class effect. Medications currently documented or predicted to have augmented oral bioavailability with grapefruit are shown in Table 1 and Appendix 1 (available at www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.120951/-/DC1). Affected drugs possess 3 essential characteristics: they have an oral route of administration, they have very low (< 10%) to intermediate (> 30%–70%) intrinsic oral bioavailability, and they are metabolized by CYP3A4. These criteria can often be found in the product monograph or prescribing information (under “clinical pharmacology”) for a drug,

particularly for recently marketed drugs, and enable the prediction of whether an interaction might occur. In principle, this would assist practitioners in formulating appropriate management strategies without exposing patients to potentially harmful combinations.

What determines the clinical significance of the interaction?

The clinical significance of any particular interaction depends on the seriousness of the dose-related drug toxicity and the extent to which the systemic drug concentration increases. The latter relies on multiple factors that include the innate oral bioavailability of the interacting drug, the circumstances under which the grapefruit or other citrus fruit is consumed and the vulnerability of the patient to the interaction.

Oral bioavailability

The lower innate oral bioavailability of the drug, the greater the possible increase in systemic drug concentration. Grapefruit-interacting drugs can be separated into the 4 categories of very low (< 10%), low (10%–30%), intermediate (> 30%–70%) and high (> 70%) absolute bioavailability. Drugs with very low bioavailability are the most likely to interact with grapefruit in a way that substantially alters their pharmacokinetics (i.e., analogous to consuming many doses of the drug alone). Conversely, drugs with high bioavailability have a marginally clinically relevant increase in systemic drug concentration.

Circumstances of grapefruit consumption

Although some pharmacokinetic studies have tested a higher than usual amount of grapefruit juice to determine the maximum effect, this should not be interpreted to mean that a relevant pharmacokinetic interaction will only occur with high levels of consumption. Indeed, a single usual amount (i.e., 200–250 mL juice or a whole grapefruit) has sufficient potency to cause a pertinent pharmacokinetic interaction.^{8,11,12} For example, felodipine combined with such a quantity of grapefruit had an average systemic drug concentration that was 3-fold that seen with water.^{8,11} With twice the amount of grapefruit, there was only a modestly greater increase in the systemic concentration of felodipine, showing that a near-maximal pharmacokinetic interaction had already occurred with the consumption of the single quantity.¹¹ With repeated ingestion of grapefruit (250 mL of juice, 3 times/d for 6 d), a single dose of felodipine increased to 5 times the systemic concentration seen with water, suggesting that frequent consumption of a usual

Box 1: Evidence used in this review

We conducted a comprehensive search of the PubMed database for all available scientifically valid evidence using the keyword “grapefruit” and the following additional terms: “drug,” “drug interaction,” “pharmacokinetics,” “cytochrome P450,” “CYP3A4,” “case report” or “review.” In addition, we obtained product monographs and prescribing information for drugs recently introduced (i.e., in the last 4 yr) to the Canadian market. We assessed the following sections of these documents for relevant information: “Summary Product Information,” “Warnings and Precautions,” “Contraindications,” “Adverse Reactions,” “Drug Interactions” and “Action and Clinical Pharmacology.” We identified 190 relevant publications (161 articles from PubMed; 29 product monographs or prescribing information sheets). Most of the information was from randomized controlled clinical trials ($n = 102$). The measured outcome from these studies was change in drug pharmacokinetics, and this was used to assess the potential for adverse clinical consequences.

quantity daily augmented the pharmacokinetic effect moreso than the lone quantity.⁸

The interval between the ingestion of grapefruit and the administration of the interacting drug has some effect on pharmacokinetics. For example, a single glass (200 mL) of grapefruit juice ingested within 4 hours before felodipine produced the maximal pharmacokinetic interaction.¹⁶ Thereafter, an increased interval between ingesting the 2 substances slowly decreased the size of the effect — an interval of 10 hours produced an effect that was 50% of the maximum, and an interval of 24 hours produced an effect that was 25% of the maximum.¹⁶ Thus, a modest solitary quantity of grapefruit can have sufficient duration of action to affect interacting drugs that are administered once daily at any time during the dosing interval. Furthermore, repeated ingestion of grapefruit (200 mL of juice, 3 times/d for 7 d) doubled the size of the interaction for 24 hours, consistent with a cumulative inhibitory action.¹⁷

Theoretically, the batch, type (i.e., white or pink) and storage conditions of grapefruit could possibly influence the size of the interaction. However, to our knowledge, these aspects have not been systematically studied.

Patient vulnerability

Patient vulnerability to this pharmacokinetic interaction varies markedly. For example, individual systemic felodipine concentrations with a single serving of grapefruit juice (250 mL) ranged from 0 to 8-fold that seen with water.⁸ Individual biopsies of the small intestine showed that higher CYP3A4 levels before ingesting grapefruit juice resulted in greater decrease in enzyme levels and greater increase in oral drug bioavailability after consumption of the juice. Accordingly, patients with elevated levels of CYP3A4 in the small intestine appear to be at increased risk for this interaction. It is impractical to routinely determine enterocyte CYP3A4 content in clinical practice. However, patients with substantial intestinal levels of CYP3A4 could require a higher dose of a grapefruit-interactive drug to achieve adequate systemic concentration. Thus, this is a possible means of identifying patients at greater risk before exposure to an interacting combination for medications routinely titrated to therapeutic effect.

Despite current science from well-conducted clinical studies, there remains the key practical issue of the frequency of adverse outcomes from this interaction occurring in routine clinical prac-

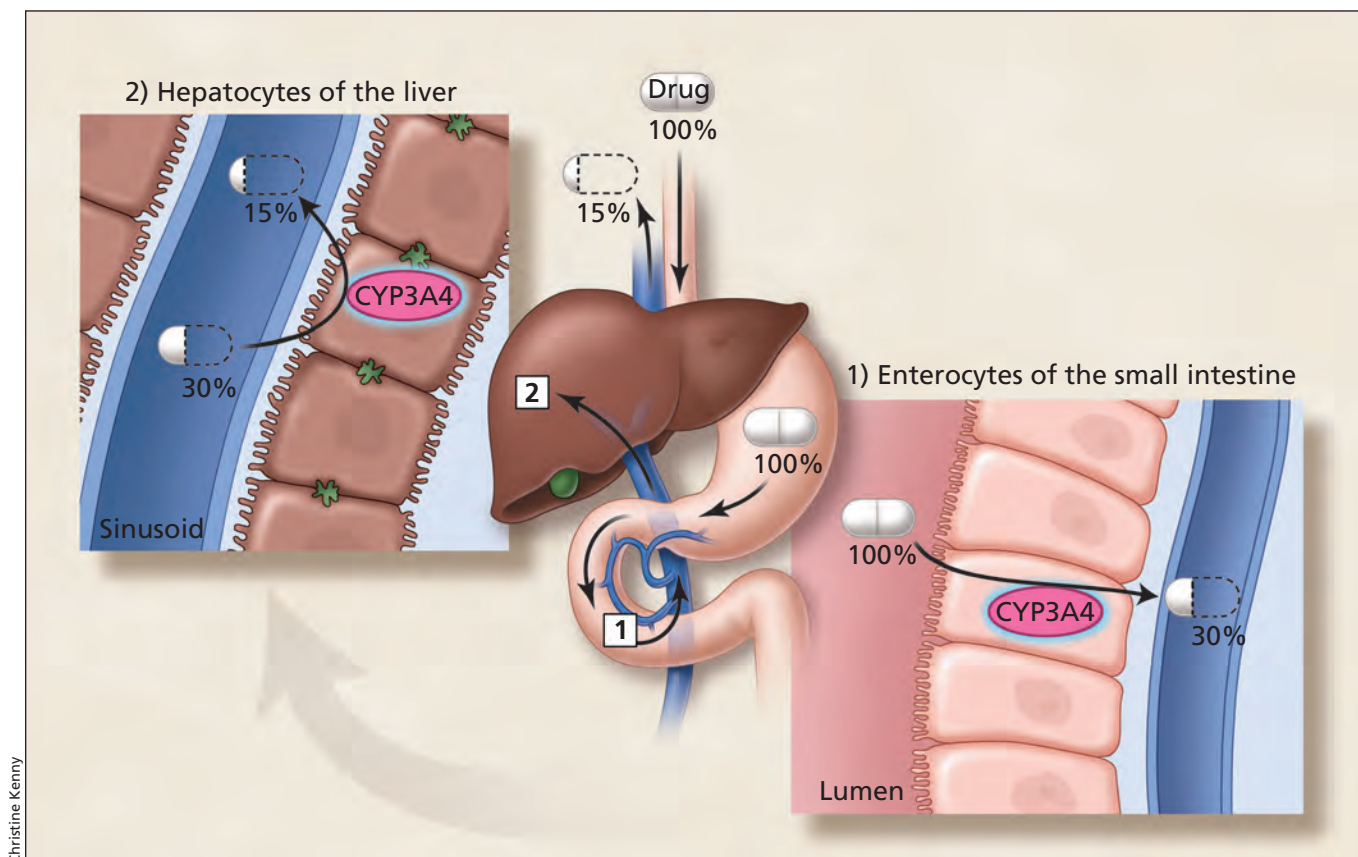


Figure 1: Sequential first-pass elimination of a drug, such as felodipine, through metabolism in enterocytes of the small intestine, then hepatocytes of the liver. The percentages of the initial dose that are available before and after passage through the gut wall and liver are shown. Although felodipine is 100% absorbed from the gastrointestinal tract, its bioavailability is only 15% after oral administration. CYP3A4 = cytochrome P450 enzyme 3A4.

tice. Because multiple factors likely need to combine to achieve a marked increase in systemic drug concentration, it is reasonable to state that just exposure to any interacting combination would not be sufficient to elicit a clinically important change in drug response in all, if not most, cases. However, important toxic events have been docu-

mented for grapefruit–drug interactions (Table 2).^{18–26} These case reports uniformly cited the circumstance of a patient whose therapeutic dose of a susceptible drug was stabilized, who subsequently showed serious toxicity that occurred after several days of simultaneous intake of the drug and grapefruit in a normal or high quantity.

Table 1 (part 1 of 2): Selected drugs that interact with grapefruit, associated oral bioavailability, adverse event(s), predicted risk and possible alternative agents

Interacting drugs	Innate oral bioavailability*	Dose-related adverse event(s)	Predicted interaction risk†	Potential alternative agent(s)‡
Anticancer agents				
Crizotinib	Intermediate	Torsade de pointes, myelotoxicity	High	
Dasatinib	Not known	Torsade de pointes, myelotoxicity	High	Imatinib
Erlotinib	Intermediate	Myelotoxicity	High	
Everolimus	Low	Myelotoxicity, nephrotoxicity	High	
Lapatinib	Incomplete	Torsade de pointes, myelotoxicity	High	
Nilotinib	Intermediate	Torsade de pointes, myelotoxicity	High	Imatinib
Pazopanib	Incomplete	Torsade de pointes, myelotoxicity	High	Sorafenib
Sunitinib	Not known	Torsade de pointes, myelotoxicity	High	Sorafenib
Vandetanib	Not known	Torsade de pointes, myelotoxicity	High	
Venurafenib	Not known	Torsade de pointes, myelotoxicity	High	
Anti-infective agents				
Erythromycin	Intermediate	Torsade de pointes	High	Clarithromycin
Halofantrine	Low	Torsade de pointes	Very high	Doxycycline
Maraviroc	Low	Postural hypotension, syncope	Very high	Enfuvirtide
Primaquine	Intermediate	Myelotoxicity	High	Doxycycline
Quinine	Intermediate	Torsade de pointes	High	Doxycycline
Rilpivirine	Not known	Torsade de pointes	High	Nevirapine
Antilipemic agents				
Atorvastatin	Low	Rhabdomyolysis	High	Pravastatin, rosuvastatin, fluvastatin
Lovastatin	Very low	Rhabdomyolysis	Very high	Pravastatin, rosuvastatin, fluvastatin
Simvastatin	Very low	Rhabdomyolysis	Very high	Pravastatin, rosuvastatin, fluvastatin
Cardiovascular agents				
Amiodarone	Intermediate	Torsade de pointes	High	Sotalol
Apixaban	Intermediate	GI bleeding	High	Warfarin
Clopidogrel	Very low	Loss of efficacy	High	Acetylsalicylic acid
Dronedarone	Low	Torsade de pointes	Very high	Sotalol
Eplerenone	Intermediate	Hyperkalemia, serious arrhythmias	High	Spironolactone
Felodipine	Low	Hypotension, peripheral edema	Intermediate	Amlodipine
Nifedipine	Intermediate	Hypotension, peripheral edema	Intermediate	Amlodipine

But how big a problem are such interactions? Unless health care professionals are aware of the possibility that the adverse event they are seeing might have an origin in the recent addition of grapefruit to the patient's diet, it is very unlikely that they will investigate it. In addition, the

patient may not volunteer this information. Thus, we contend that there remains a lack of knowledge about this interaction in the general health care community. Consequently, current data are not available to provide an absolute or even approximate number representing the true inci-

Table 1 (part 2 of 2): Selected drugs that interact with grapefruit, associated oral bioavailability, adverse event(s), predicted risk and possible alternative agents

Interacting drugs	Innate oral bioavailability*	Dose-related adverse event(s)	Predicted interaction risk†	Potential alternative agent(s)‡
Cardiovascular agents (continued)				
Quinidine	High	Torsade de pointes	Intermediate	
Rivaroxaban	High	GI bleeding	Intermediate	Warfarin
Ticagrelor	Intermediate	GI or kidney bleeding	High	Acetylsalicylic acid
CNS agents				
Alfentanil (oral)	Intermediate	Respiratory depression	High	Hydromorphone, morphine
Bupirone	Very low	Dizziness, sedation	High	Oxazepam, tamazepam
Dextromethorphan	Very low	Hallucinations, somnolence	High	
Fentanyl (oral)	Intermediate	Respiratory depression	High	Hydromorphone, morphine
Ketamine (oral)	Low	Respiratory depression	Very high	Hydromorphone, morphine
Lurasidone	Low	Torsade de pointes, orthostatic hypotension, syncope	Very high	Haloperidol, risperidone, olanzapine
Oxycodone	Intermediate	Respiratory depression	High	Hydromorphone, morphine
Pimozide	Intermediate	Torsade de pointes	High	Haloperidol, risperidone, olanzapine
Quetiapine	Very low	Dizziness, somnolence	High	Haloperidol, risperidone, olanzapine
Triazolam	Intermediate	Sedation	Intermediate	Alprazolam, lorazepam
Ziprasidone	Intermediate	Torsade de pointes	High	Haloperidol, risperidone, olanzapine
Gastrointestinal agents				
Domperidone	Low	Torsade de pointes	Very high	Metoclopramide
Immunosuppressants				
Cyclosporine	Low	Nephrotoxicity	High	
Everolimus	Low	Myelotoxicity, nephrotoxicity	High	
Sirolimus	Low	Myelotoxicity, nephrotoxicity	High	
Tacrolimus	Low	Nephrotoxicity	High	
Urinary tract agents				
Darifenacin	Low	Urinary retention, constipation	Intermediate	
Fesoterodine	Intermediate	Urinary retention, constipation	Intermediate	
Solifenacin	High	Torsade de pointes	Intermediate	
Silodosin	Intermediate	Postural hypotension, dizziness	Intermediate	
Tamsulosin	Intermediate	Postural hypotension, dizziness	Intermediate	

Note: CNS = central nervous system, GI = gastrointestinal.

*Population average: very low < 10%, low >10%–30%, intermediate >30%–70%, high > 70%.

†Based on the seriousness of the adverse effect and adjusted for the innate oral bioavailability of the drug, which is used to determine the potential increase in systemic drug concentration. For older patients, particularly the elderly, it is recommended that grapefruit or other citrus fruits be contraindicated for drugs in the very high or high category, or that a suggested alternative noninteracting drug be employed when available.

‡Commonly prescribed drugs (Top 100 prescribed in 2011 in Canada) with the same therapeutic indication that have no or minor pharmacokinetic interaction with grapefruit or other citrus fruits.

dence of grapefruit–drug interactions in routine practice. Nonetheless, there are certain situations in which this interaction has a predictably greater likelihood of producing particularly adverse clinical results.

Who is at higher risk of grapefruit–drug interactions?

Although patient vulnerability is largely unknown, people older than 45 years are the prime purchasers of grapefruit and receive the most prescriptions for drugs.^{27,28} Because of the size of this population, substantial exposure to this interaction is likely. In addition, a pronounced pharmacokinetic interaction has been shown to occur in patients older than 70 years.²⁹ Furthermore, older adults can have decreased capacity to compensate for excessive systemic drug concentrations. For example, felodipine (which normally lowers blood pressure) does not cause a compensating increase in heart rate in older adults when ingested with grapefruit, but does in young and middle-aged people, likely because of attenuated baroreceptor sensitivity associated with age.²⁹ Consequently, older people appear to be an especially vulnerable population for grapefruit–drug interactions.

The predicted interaction risk for grapefruit–interacting drugs (Table 1, Appendix 1) (i.e., very high, high, intermediate, low) can assist clinicians in prescribing drugs for vulnerable patients and determining whether grapefruit or other citrus fruits should be contraindicated during pharmacotherapy, or whether an alternative therapy can be used.

What are examples of important grapefruit–drug interactions?

Examples of grapefruit–drug interactions were selected to illustrate documented pharmacokinetic changes for which clinical outcomes are considered serious. These effects include torsade de pointes, rhabdomyolysis, nephrotoxicity and breast cancer.

Torsade de pointes

Torsade de pointes and risk of sudden death can occur with excessive prolongation of the corrected QT interval. The antiarrhythmic agent amiodarone had a mean C_{max} with grapefruit juice (300 mL at 0, 3 and 9 h relative to drug administration) corresponding to 180% of that with water; the AUC was 150% of that with water.³⁰ The combination was also reported to markedly prolong the corrected QT interval and to cause ventricular arrhythmias, including torsade de pointes, in clinical practice.¹⁸ Dronedarone, the chemical analog of amiodarone, was associated with reports of ventricular arrhythmia, cardiac arrest and torsade de pointes in clinical practice.³¹ Dronedarone with grapefruit juice (300 mL, 3 times/d) resulted in a systemic drug concentration that was 300% of that of a control.³²

Torsade de pointes can also occur with certain anticancer agents. The tyrosine kinase inhibitor nilotinib had a mean C_{max} with a single glass of grapefruit juice (480 mL) that was 160% of that with water, and an AUC that was 129% of that with water.³³ Sunitinib, another tyrosine kinase inhibitor, had a mean bioavailability with grapefruit consumption (200 mL, 3 times/d for 3 d) that was 111% of that compared with the level seen when grapefruit is not consumed.³⁴ Although the pharmacokinetic interaction was weaker with sunitinib, the potential seriousness of the adverse effect and concern for interpatient variability would still warrant the avoidance of grapefruit.

Rhabdomyolysis

Rhabdomyolysis is the consequence of profound damage to skeletal muscle tissue, the release of large quantities of proteins, such as myoglobin, into the blood and acute renal failure. The active forms of all statins can produce this toxicity at excessive systemic concentrations.³⁵

Simvastatin with grapefruit juice at high volume (400 mL, 3 times/d for 3 d) had an AUC that was 700% of that for water; at a more usual amount of juice (200 mL, once daily for 3 d), the AUC was 330% of that with water.^{36,37} Rhabdomyolysis was also reported after 10 days concomitant consumption of fresh grapefruit.²³ Lovastatin with grapefruit juice at a high level of

Table 2: Case reports of serious adverse events related to grapefruit–drug interaction^{18–26}

Serious adverse event	Drug	Amount of grapefruit consumed
Torsade de pointes	Amiodarone ¹⁸	Juice, 1–1.5 L/d on a regular basis
	Quinine in tonic water ¹⁹	Juice, high volume during preceding days
Complete heart block	Verapamil ²⁰	Juice, high volume during preceding days
Rhabdomyolysis	Atorvastatin ^{21,22}	Juice, 1–2 glasses/d for 5 d; juice from fresh grapefruit daily for 2 mo
	Simvastatin ²³	Whole fruit, 1 fruit/d for 2 wk
Nephrotoxicity	Tacrolimus ²⁴	Marmalade, 1.5 kg eaten during preceding 1 wk
Myelotoxicity	Colchicine ²⁵	Juice, 1 L/d for preceding 2 mo
Venous thrombosis	Ethinylestradiol ²⁶	Whole fruit, 1 fruit/d for breakfast for preceding 3 d

consumption (400 mL, 3 times/d for 3 d) caused an AUC that was 500% of that with water.³⁸ Atorvastatin, the most frequently prescribed drug in Canada in 2011, had an AUC with grapefruit juice (250–400 mL, 3 times/d for 2–4 d) that ranged from 180% to 250% of that with water.^{39–41}

Rhabdomyolysis has also been reported with grapefruit ingested at usual amounts (Table 2).^{21–23} Thus, this adverse outcome with certain statins can occur with the ingestion of much less grapefruit than was previously expressed by the US Food and Drug Administration.⁴²

However, taking atorvastatin in the evening and drinking grapefruit juice in the morning (300 mL/d from a specific lot prepared by the Florida Department of Citrus) resulted in drug serum concentrations that were 119%–126% of those seen with no consumption of grapefruit, with no evidence of skeletal muscle toxicity (e.g., elevated creatine phosphokinase, myalgia).⁴³ In addition, pravastatin does not produce a pharmacokinetic interaction with grapefruit,^{39,40} rosuvastatin is eliminated unchanged,³⁵ and fluvastatin is metabolized by an enzyme (cytochrome P450 2C9) that is not affected by grapefruit.³⁵ Although staggering the ingestion of atorvastatin and grapefruit may reduce risk, substituting pravastatin, rosuvastatin or fluvastatin, or eliminating grapefruit juice from the diet, appears more preferable.

Nephrotoxicity

Nephrotoxicity can occur with the calcineurin inhibitors cyclosporine and tacrolimus, which are vital in preventing organ rejection after transplantation. Both of these drugs have a narrow range of therapeutic blood concentrations (i.e., below which they lack sufficient efficacy and above which they cause toxicity). Cyclosporine with grapefruit juice (single serving of 250 mL) produced a mean oral bioavailability that was 162% of that with water.¹⁰ One of the 9 patients involved in this study had systemic drug availability increased to 670%. In addition, a case report showed cyclosporine concentration to be increased to 600% with grapefruit.⁴⁴ Taking tacrolimus after ingesting grapefruit juice (250 mL, 4 times/d for 3 d) resulted in a 1000% higher trough blood concentration, causing profound inhibition of calcineurin phosphatase in a recipient of a liver transplant.⁴⁵ Moreover, taking tacrolimus after consuming a large amount of grapefruit marmalade during the previous week caused a 500% greater blood concentration of the drug and acute renal dysfunction.²⁴

Breast cancer

Based on increased oral bioavailability of estrogens (ethinylestradiol and 17- β -estradiol) with

grapefruit juice, 2 large epidemiologic studies assessed the risk of breast cancer.^{46,47} The initial investigation found higher risk (relative risk 1.30, 95% confidence interval 1.06–1.58) in women postmenopause who were taking estrogens and consuming one-quarter grapefruit or more per day compared with women not eating grapefruit.⁴⁸ However, a follow-up study involving the same population found no such association.⁴⁹ Thus, there is controversy over the risk of breast cancer in women postmenopause receiving estrogen therapy and consuming grapefruit.

Gaps in knowledge

Although much effort has been expended to provide a complete list of currently known and predicted drugs that will interact with grapefruit, the absence of a drug from Table 1 and Appendix 1 should not be interpreted to mean that it lacks this interaction. The few case reports presented in this review should not be considered as a useful index of the frequency of occurrence of serious grapefruit–drug interactions in general practice, as these are likely underreported.

Conclusion

Grapefruit and certain other citrus fruits represent examples of foods generally considered to be healthful, but with the potential for a pharmacokinetic interaction causing greatly enhanced oral drug bioavailability. The current trend of increasing numbers of newly marketed grapefruit-affected drugs possessing substantial adverse clinical effects necessitates an understanding of this interaction and the application of this knowledge for the safe and effective use of drugs in general practice.

References

- Bailey DG, Spence JD, Edgar B, et al. Ethanol enhances the hemodynamic effects of felodipine. *Clin Invest Med* 1989;12:357–62.
- Bailey DG, Arnold JMO, Spence JD. Grapefruit juice–drug interactions. *Br J Clin Pharmacol* 1998;46:101–10.
- Bailey DG, Spence JD, Munoz C, et al. Interaction of citrus juices with felodipine and nifedipine. *Lancet* 1991;337:268–9.
- Bailey DG. Fruit juice inhibition of uptake transport: a new type of food–drug interaction. *Br J Clin Pharmacol* 2010;70:645–55.
- Wilkinson GR. Drug metabolism and variability among patients in drug response. *N Engl J Med* 2005;352:2211–21.
- Edgar B, Regardh CG, Johnsson G, et al. Felodipine kinetics in healthy man. *Clin Pharmacol Ther* 1985;38:205–11.
- Guo LQ, Yamazoe Y. Inhibition of cytochrome P450 by furanocoumarins in grapefruit juice and herbal medicines. *Acta Pharmacol Sin* 2004;25:129–36.
- Lown KS, Bailey DG, Fontana RJ, et al. Grapefruit juice increases felodipine oral availability in humans by decreasing intestinal CYP3A protein expression. *J Clin Invest* 1997;99:2545–53.
- Lundahl J, Regardh CG, Edgar B, et al. Effects of grapefruit juice ingestion — pharmacokinetics and haemodynamics of intravenously and orally administered felodipine in healthy men. *Eur J Clin Pharmacol* 1997;52:139–45.
- Ducharme MP, Warbasse LH, Edwards DJ. Disposition of intravenous and oral cyclosporine after administration with grapefruit juice. *Clin Pharmacol Ther* 1995;57:485–91.

11. Edgar B, Bailey D, Bergstrand R, et al. Acute effects of drinking grapefruit juice on the pharmacokinetics and dynamics of felodipine and its potential clinical relevance. *Eur J Clin Pharmacol* 1992;42:313-7.
12. Bailey DG, Dresser GK, Kreeft JH, et al. Grapefruit–felodipine interaction: effect of unprocessed fruit and probable active ingredients. *Clin Pharmacol Ther* 2000;68:468-77.
13. Malhotra S, Bailey DG, Paine MF, et al. Seville orange juice–felodipine interaction: comparison with dilute grapefruit juice and involvement of the furanocoumarins. *Clin Pharmacol Ther* 2001;69:14-23.
14. Bailey DG, Dresser GK, Bend JR. Bergamottin, lime juice and red wine as inhibitors of CYP3A4 activity: comparison with grapefruit juice. *Clin Pharmacol Ther* 2003;73:529-37.
15. Guo LQ, Chen QY, Wang X, et al. Different roles of pummelo furanocoumarin and cytochrome P450 3A5*3 polymorphism in the fate and action of felodipine. *Curr Drug Metab* 2007;8:623-30.
16. Lundahl J, Regardh CG, Edgar B, et al. Relationship between time of intake of grapefruit juice and its effect on pharmacokinetics and pharmacodynamics of felodipine in healthy subjects. *Eur J Clin Pharmacol* 1995;49:61-7.
17. Takanaga H, Ohnishi A, Murakami H, et al. Relationship between time after intake of grapefruit juice and the effect on the pharmacokinetics and pharmacodynamics of nisoldipine in healthy subjects. *Clin Pharmacol Ther* 2000;67:201-14.
18. Agosti S, Casalino L, Bertero G, et al. A dangerous fruit juice. *Am J Emerg Med* 2012;30:248.e5-248.e8.
19. Hermans K, Stockman D, Van den Branden F. Grapefruit and tonic: a deadly combination in a patient with the long QT syndrome. *Am J Med* 2003;114:511-2.
20. Pillai U, Muzaffar J, Sen S, et al. Grapefruit juice and verapamil: a toxic cocktail. *South Med J* 2009;102:308-9.
21. Mazokopakis EE. Unusual causes of rhabdomyolysis. *Intern Med J* 2008;38:364-7.
22. Karch AM. The grapefruit challenge: the juice inhibits a crucial enzyme, with possibly fatal consequences. *Am J Nurs* 2004;104:33-5.
23. Dreier JP, Endres M. Statin-associated rhabdomyolysis triggered by grapefruit consumption. *Neurology* 2004;62:670.
24. Peynaud D, Charpiat B, Vial T, et al. Tacrolimus severe overdose after intake of masked grapefruit in orange marmalade. *Eur J Clin Pharmacol* 2007;63:721-2.
25. Goldbart A, Press J, Sofer S, et al. Near fatal acute colchicine intoxication in a child. A case report. *Eur J Pediatr* 2000;159:895-7.
26. Grande LA, Mendez RD, Krug RT, et al. Attention grapefruit! *Lancet* 2009;373:1222.
27. Johannot L, Somerset SM. Age-related variations in flavonoid intake and sources in the Australian population. *Public Health Nutr* 2006;9:1045-54.
28. Genser D. Food and drug interaction: consequences for the nutrition/health status. *Ann Nutr Metab* 2008;52(Suppl 1):29-32.
29. Dresser GK, Bailey DG, Carruthers SG. Grapefruit juice–felodipine interaction in the elderly. *Clin Pharmacol Ther* 2000;68:28-34.
30. Libersa CC, Brique SA, Motte KB, et al. Dramatic inhibition of amiodarone metabolism induced by grapefruit juice. *Br J Clin Pharmacol* 2000;49:373-8.
31. Kao DP, Hiatt WR, Krantz MJ. Proarrhythmic potential of dronedarone: emerging evidence from spontaneous adverse event reporting. *Pharmacotherapy* 2012 June 28 [Epub ahead of print].
32. Multaq (dronedarone HCl) [product monograph]. Laval (QC); Sanofi-Aventis; 2009.
33. Yin OQ, Gallagher N, Li A, et al. Effect of grapefruit juice on the pharmacokinetics of nilotinib in healthy participants. *J Clin Pharmacol* 2010;50:188-94.
34. van Erp NP, Baker SD, Zandvliet AS, et al. Marginal increase of sunitinib exposure by grapefruit juice. *Cancer Chemother Pharmacol* 2011;67:695-703.
35. Neuvonen PJ, Niemi M, Backman JT. Drug interactions with lipid-lowering drugs: mechanisms and clinical relevance. *Clin Pharmacol Ther* 2006;80:565-81.
36. Lilja JJ, Kivisto KT, Neuvonen PJ. Grapefruit juice–simvastatin interaction: effect on serum concentrations of simvastatin, simvastatin acid, and HMG-CoA reductase inhibitors. *Clin Pharmacol Ther* 1998;64:477-83.
37. Lilja JJ, Neuvonen M, Neuvonen PJ. Effects of regular consumption of grapefruit juice on the pharmacokinetics of simvastatin. *Br J Clin Pharmacol* 2004;58:56-60.
38. Kantola T, Kivisto KT, Neuvonen PJ. Grapefruit juice greatly increases serum concentrations of lovastatin and lovastatin acid. *Clin Pharmacol Ther* 1998;63:397-402.
39. Lilja JJ, Kivisto KT, Neuvonen PJ. Grapefruit juice increases serum concentrations of atorvastatin and has no effect on pravastatin. *Clin Pharmacol Ther* 1999;66:118-27.
40. Ando H, Tsuruoka S, Yanagihara H, et al. Effects of grapefruit juice on the pharmacokinetics of pitavastatin and atorvastatin. *Br J Clin Pharmacol* 2005;60:494-7.
41. Fukazawa I, Uchida N, Uchida E, et al. Effects of grapefruit juice pharmacokinetics of atorvastatin and pravastatin in Japanese. *Br J Clin Pharmacol* 2004;57:448-55.
42. Lipid-altering agents (also called statins). *Avoid food–drug interactions: a guide from the National Consumers League and U.S. Food and Drug Administration*. Publication no. (FDA) CDER 10-1933, p 13. Available: www.fda.gov/downloads/Drugs/ResourcesForYou/Consumers/BuyingUsingMedicineSafely/EnsuringSafeUseofMedicine/GeneralUseofMedicine/UCM229033.pdf (accessed 2012 Oct. 25).
43. Reddy P, Ellington D, Zhu Y, et al. Serum concentrations and clinical effects of atorvastatin in patients taking grapefruit juice daily. *Br J Clin Pharmacol* 2011;72:434-41.
44. Jürgen Roth H, Wintergalen M. Unusual high levels of ciclosporin in a female patient — the impact of lifestyle? *Clin Lab* 2005;51:425-7.
45. Fukatsu S, Fukudo M, Masuda S, et al. Delayed effect of grapefruit juice on pharmacokinetics and pharmacodynamics of tacrolimus in a living-donor liver transplant recipient. *Drug Metab Pharmacokinet* 2006;21:122-5.
46. Schubert W, Cullberg G, Edgar B, et al. Inhibition of 17 β -estradiol metabolism by grapefruit juice in ovariectomized women. *Maturitas* 1994;20:155-63.
47. Weber A, Jäger R, Börner A, et al. Can grapefruit juice influence ethinylestradiol bioavailability? *Contraception* 1996;53:41-7.
48. Monroe KR, Murphy SP, Kolonel LN, et al. Prospective study of grapefruit intake and risk of breast cancer in postmenopausal women: the Multiethnic Cohort Study. *Br J Cancer* 2007;97:440-5.
49. Spencer EA, Key TJ, Appleby PN, et al. Prospective study of the association between grapefruit intake and risk of breast cancer in the European Prospective Investigation into Cancer and Nutrition (EPIC). *Cancer Causes Control* 2009;20:803-9.

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