VI.2 Elements for a public summary

VI.2.1 Overview of disease epidemiology

This medicine is used to lower levels of total cholesterol, LDL cholesterol ("bad" cholesterol), and fatty substances called triglycerides in the blood. In addition, this medicine raises levels of HDL cholesterol ("good" cholesterol). Abnormal cholesterol and triglycerides levels are called dyslipidemia.

Dyslipidemia is a major, modifiable risk factor for cardiovascular disease. Cardiovascular disease is a leading cause of morbidity and mortality in the world; there are an estimated 16.7 million deaths each year from cardiovascular disease worldwide. Dyslipidemia is usually asymptomatic and is not fatal; however, if improperly managed or left untreated, it can eventually contribute to coronary artery disease or peripheral artery disease, both of which can be fatal.

VI.2.2 Summary of treatment benefits

The active substances (simvastatin and ezetimibe) work in different ways and their action have a complementary effect.

Simvastatin belongs to the group called 'statins'. It reduces total blood cholesterol by blocking the action of HMG-CoA reductase, an enzyme in the liver involved in the production of cholesterol. As the liver needs cholesterol to produce bile, the reduced blood cholesterol level causes the liver cells to produce receptors that draw cholesterol from the blood, reducing its level even further. The cholesterol drawn out of the blood in this way is the LDL, or 'bad' cholesterol.

Ezetimibe inhibits intestinal uptake of dietary and biliary cholesterol without affecting the absorption of fat-soluble nutrients. By inhibiting cholesterol absorption at the level of the brush border of the intestine, ezetimibe reduces the amount of lipoprotein cholesterol circulated to the liver. In response to reduced cholesterol delivery, the liver reacts by up-regulating LDLR, which in turn leads to increased clearance of LDL from the blood.

Study which analyzed 27 previously published studies compared effectiveness of combination of simvastatin and ezetimibe and simvastatin alone. Combination of simvastatin and ezetimibe was more effective than simvastatin in lowering total cholesterol, LDL cholesterol and triglycerides levels and in increasing HDL cholesterol levels

VI.2.3 Unknowns relating to treatment benefits

There was only limited clinical trial experience in children 10 - 17 years of age. Also no clinical trials in children less than 10 years of age were performed.

VI.2.4 Summary of safety concerns

Important identified risks

Risk	What is known	Preventability
Muscle disorder	On rare occasions, muscle	Contact your doctor
	problems can be serious, including	immediately if you
(rhabdomyolysis/myopathy)	muscle breakdown resulting in	experience unexplained
	kidney damage; and very rare	muscle pain, tenderness,
	deaths have occurred.	or weakness.
	The risk of muscle breakdown is	
	greater at higher doses of	Talk with your doctor if
	ezetimibe/simvastatin,	any of the following
	particularly the 10/80-mg dose.	applies:
	The risk of muscle breakdown is	• you have kidney
	also greater in certain patients.	problems
		• you have thyroid
		problems
	Like all medicines,	• you are 65 years or
	ezetimibe/simvastatin can cause	older
	side effects, although not	• you are female
	everybody gets them.	• you have ever had
	- muscle aches	muscle problems during
	- elevations in laboratory blood	treatment with
	muscle (CK) function	cholesterol lowering
		medicines called
		"statins" (like
		simvastatin,
		atorvastatin, and
		rosuvastatin) or fibrates
		(like gemfibrozil and
		bezafibrate)
		• you or close family
		members have a
		hereditary muscle
		disorder.
		Taking
		ezetimibe/simvastatin
		with any of these drugs
		can increase the risk of
		muscle problems

	• • • • •
	• ciclosporin (a
	medicine often used in
	organ transplant
	patients)
	• danazol (a man-made
	hormone used to treat
	endometriosis)
	• medicines like
	itraconazole,
	ketoconazole,
	fluconazole or
	posaconazole
	(medicines for fungal
	infections)
	• fibrates like
	gemfibrozil and
	bezafibrate (medicines
	for lowering
	cholesterol)
	• erythromycin,
	• •
	clarithromycin,
	telithromycin, or fusidic
	acid (medicines for
	bacterial infections)
	• HIV protease
	inhibitors such as
	indinavir, nelfinavir,
	ritonavir, and
	saquinavir (medicines
	for AIDS)
	• boceprevir or
	telaprevir (medicines
	for hepatitis C virus
	infection)
	• nefazodone (a
Grapefruit juice contains one or	medicine for
more components that alter the	depression)
metabolism of some medications,	• amiodarone (a
including ezetimibe/simvastatin.	medicine for an
G G G G G G G G G G	irregular heartbeat)
	• verapamil, diltiazem,
	or amlodipine
	annourphie

		(medicines for high
		blood pressure, chest
		pain associated with
		heart disease, or other
		heart conditions)
		,
		• large amounts (1 gram
		or more each day) of
		niacin or nicotinic acid
		(medicines for lowering
		cholesterol)
		• colchicine (a medicine
		used to treat gout).
		If any of the side effects
		gets serious, or if you
		notice any side effects
		not listed in this leaflet,
		please tell your doctor
		or pharmacist.
		Consuming grapefruit
		juice should be avoided
		as it may increase your
		risk of muscle
		problems.
		If any of the side effects
		gets serious, or if you
		notice any side effects
		not listed in this leaflet,
		please tell your doctor
		or pharmacist.
Liver disease or unexplained	Like all medicines,	Your doctor should do a
persistant elevation in some	ezetimibe/simvastatin can cause	blood test before you
enzymes	side effects, although not	start taking
	everybody gets them.	ezetimibe/simvastatin
(Abnormal liver function)	The following common side	and if you have any
(1.01011111111111111111111)	effects were reported	symptoms of liver
	• elevations in laboratory blood	problems while you
	tests of liver (transaminases)	take
	• inflammation of the liver with the	ezetimibe/simvastatin.
	following symptoms: yellowing of	
	Tonowing symptoms. yenowing of	THIS IS TO CHECK HOW

	the skin and avec itaking dark	wall wave liver in
	the skin and eyes, itching, dark coloured urine or pale coloured stool, feeling tired or weak, loss of appetite; liver failure; gallstones or inflammation of the gallbladder (which may cause abdominal pain, nausea, vomiting)	 well your liver is working. Do not take ezetimibe/simvastatin if: you currently have liver problems Your doctor may also want you to have blood tests to check how well your liver is working after you start taking ezetimibe/simvastatin.
		If any of the side effects gets serious, or if you notice any side effects not listed in leaflet, please tell your doctor or pharmacist.
Hypersensitivity	These medications may cause allergic reactions as all medicines can.	Yes, by monitoring for early symptoms and avoiding the drugs you are already known to be allergic of.
		Please tell your doctor if you are undergoing or are planning to undergo desensitisation therapy.
		Be careful and tell your doctor if you have experienced symptoms such as sudden wheeziness, chest pain, shortness of breath or difficulty in breathing, swelling of eyelids, face or lips, severe skin reactions or other
		reactions or other allergic reactions.

Drug interaction with	Both active components in ezetimibe/simvastatin can effect	Yes, by monitoring for early symptoms.
warfarin, another coumarin anticoagulant, or fluindione	warfarin action but very rare cases of elevated INR (prothrombin time) have been reported. So it is important that doctor monitor INR before starting therapy and more frequently during early therapy to ensure that no significant alteration of prothrombin time occurs.	Tell your doctor if you are taking any other drugs. You should take your medicine at the same time each day with a drink of water.
Concomitant administration of ciclosporin	Taking ezetimibe/simvastatin with other drugs can increase the risk of some side effects.	Donottakeezetimibe/simvastatinif:you are taking one ormore than one of thefollowing drugs at thesame time:•gemfibrozil(amedicine for loweringcholesterol)•ciclosporin(amedicine often used inorgantransplantpatients)•danazol (a man-madehormone used to treatendometriosis).Ask your doctor if youare not sure if yourmedicineislisted

Important potential risks:

Risk	What is known (including reason why it is considered a	
	potential risk)	
Inflammation of the pancreas	Simvastatin can cause inflammation of the pancreas, which is	
(Pancreatitis)	presented as pain in the upper abdomen.	

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Inflammation of the	Ezetimibe is an active component of ezetimibe/simvastatin.
gallbladder/gallstones	Concomitant fenofibrate or gemfibrozil administration
(Cholecystitis/cholelithiasis)	modestly increased ezetimibe concentrations (ezetimibe is
	In patients receiving fenofibrate and Ezoleta, physicians
	should be aware of the possible risk of cholelithiasis and
	gallbladder disease.
	If cholelithiasis is suspected in a patient receiving ezetimibe
	and fenofibrate, gallbladder investigations are indicated and
	this therapy should be discontinued.
	Coadministration of ezetimibe with other fibrates has not been
	studied.
	Fibrates may increase cholesterol excretion into the bile,
	leading to cholelithiasis. In animal studies, ezetimibe
	sometimes increased cholesterol in the gallbladder bile, but
	not in all species.
Different lung condition	Simvastatin, active component of ezetimibe/simvastatin,
(Interstitial lung disease)	belongs to the group called 'statins'. Exceptional cases of
(Interstituti lung uiseuse)	interstitial lung disease have been reported with some statins,
	especially with long term therapy. Symptoms of interstitial
	lung disease include breathing problems including persistent
	cough and/or shortness of breath or fever. Interstitial lung
	disease describes a large group of disorders, most of which
	cause progressive scarring of lung tissue and can have severe
	outcome.
	Tall your destor if you have severe lung disease
Streets at a time	Tell your doctor if you have severe lung disease.
Simvastatin	These medications may cause allergic reactions as all
hypersensitivity syndrome	medicines can.
	The doctor or pharmacist should be informed if you have
	allergic reactions to other simvastatin containing medicine.
New onset	Simvastatin, active component of ezetimibe/simvastatin,
diabetes/impaired glucose	belongs to the group called 'statins'.
metabolism	Some evidence suggests that statins as a group raise blood
	glucose (sugar) and in some patients, at high risk of
	developing future diabetes, may produce a level of
	hyperglycaemia (high levels of sugars) where formal diabetes
	care is appropriate. This risk, however, is outweighed by the
	reduction in vascular risk with statins and therefore should not
	be a reason for stopping treatment with this medicine.
	While you are on this medicine your doctor will monitor you
	closely if you have diabetes or are at risk of developing

	diabetes. You are likely to be at risk of developing diabetes if
	you have high levels of sugars and fats in your blood, are
	overweight and have high blood pressure.
Haemorrhagic stroke	Talk to your doctor or pharmacist before taking this medicinal
	product if you have had a previous stroke with bleeding into
	the brain, or have small pockets of fluid in the brain from
	previous strokes.
	All types of stroke are dangerous, but a few of them are
	notorious for causing severe disability and/or a rapid
	progression to death.
	Take special care with if you have had a previous stroke with
	bleeding into the brain, or have small pockets of fluid in the
	brain from previous strokes.

Missing information

Risk	What is known	
Pregnancy and breastfeeding	Some drugs used during pregnancy and breastfeeding	
	can have temporary or permanent effects on the fetus.	
(Exposure during pregnancy and	The safety of simvastatin in pregnant women has not	
lactation)	been established. No clinical data are available on the	
	use of ezetimibe during pregnancy.	
	Do not take ezetimibe/simvastatin if you are pregnant,	
	are trying to get pregnant or think you may be pregnant	
	as there is not enough information on the safety of the	
	drug.	
	If you get pregnant while taking ezetimibe/simvastatin,	
	stop taking it immediately and tell your doctor.	
	Do not take ezetimibe/simvastatin if you are breast-	
	feeding, because it is not known if the medicine is	
	passed into breast milk.	
Use in children (limited clinical	As insufficient data on the safety and efficacy of the	
trial experience in children 10 –	drug is not available, the drug is not recommended for	
17 years of age. No clinical	al children under age 10.	
experience in children < 10 years		
of age)		

VI.2.5 Summary of risk minimisation measures by safety concern

All medicines have a Summary of Product Characteristics (SmPC) which provides physicians, pharmacists and other health care professionals with details on how to use the medicine, the risks and recommendations for minimising them. An abbreviated version of this in lay language is provided in the form of the package leaflet (PL). The measures in these documents are known as routine risk minimisation measures.

The Summary of Product Characteristics and the Package leaflet for this product can be found at the agency's EPAR page.

This medicine has no additional risk minimisation measures.

VI.2.6 Planned post authorisation development plan

Not applicable. No postauthorisation studies are planned.

VI.2.7 Summary of changes to the Risk Management Plan over time

Not applicable.

ⁱ Leiter LA, Betteridge DJ, Farnier M, Guyton JR, Lin J, Shah A, et al. Lipid-altering efficacy and safety profile of combination therapy with ezetimibe/statin vs. statin monotherapy in patients with and without diabetes: an analysis of pooled data from 27 clinical trials. Diabetes Obes Metab 2011; 13(7): 615-28.