Patient Alert Card: CAPRELSA® (vandetanib)

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Information for the patient

- This card contains important safety information that you should know before you are given vandetanib and during treatment with vandetanib.
- Show this card to any doctor involved in your treatment. If you get any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in the package leaflet.
- You can report side effects directly to HPRA Pharmacovigilance, website **www.hpra.ie**. Side effects should also be reported to Esteve Pharmaceuticals Ltd, Tel: **+44 (0)330 1359 437** or by Email: **medinfo.uk@esteve.com**
- By reporting side effects you can help provide more information on the safety of this medicine.

Vandetanib can cause a change in the electrical activity of your heart called QTc prolongation, which can cause irregular heartbeats and life-threatening changes in heart rhythm.

A condition called **posterior reversible encephalopathy syndrome (PRES; also known as reversible posterior leukoencephalopathy syndrome [RPLS])** can occur while taking vandetanib.

During vandetanib treatment, telephone your doctor or tell your caregiver immediately if you:

- Feel faint, dizzy or feel your heart beating irregularly, as these may be symptoms related to QTc prolongation.
- Experience headaches, seizures, convulsions, confusion, problems seeing or have problems concentrating, as these may be symptoms of PRES.

Do not stop taking vandetanib, or change your dose, unless told to by your doctor.
If you take too many vandetanib tablets, telephone your doctor immediately.

Ver.1 Date of preparation September 2026

Read the package leaflet that came with your medicine for more information.

Please make sure that you have a list of all your other medicines with you at any visit to your doctor.

Patient's name: _____
Caregiver's name: _____
Caregiver's telephone number: _____
Doctor's name: _____
Doctor's telephone number: _____
Start date of vandetanib treatment: _____