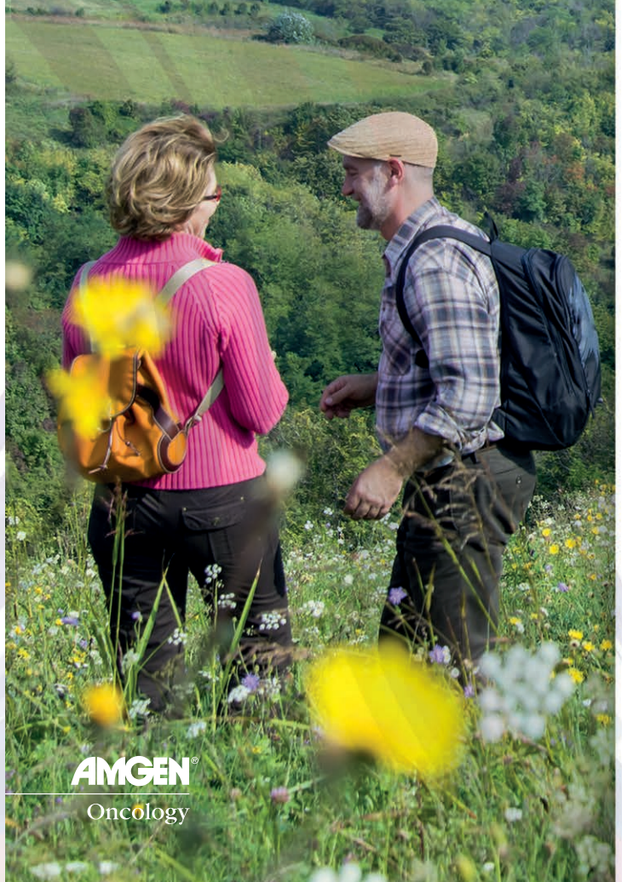
If you require additional copies of this Treatment Diary, please speak to your healthcare professional.

AMGEN® Oncology

Amgen Ireland Limited, 21 Northwood Court, Santry, Dublin 9.
UKIE-NPC-CARF-1118-069857
September 2019



KYPROLIS®▼ (CARFILZOMIB)

Treatment Diary

UKIE-NPC-CARF-1118-069857
September 2019

CONTACTS

DOCTOR'S NAME

Phone Number

HOSPITAL TELEPHONE NUMBER

CLINICAL NURSE SPECIALIST'S NAME

Phone Number

OUT OF HOURS TELEPHONE NUMBER

PHARMACIST'S NAME

OTHER CONTACTS

CARFILZOMIB TREATMENT DIARY

THIS TREATMENT DIARY HAS BEEN GIVEN TO YOU TO HELP YOU RECORD DETAILS OF YOUR TREATMENT:

- A reminder of when to take your medicines for myeloma
- A record of other medication you are taking
- A diary of your appointments

You may also wish to use the diary as a place to record any advice or suggestions from your healthcare professional (i.e. doctor or nurse) and the results of blood tests.

Please bring the diary with you when you come to your appointment. It will help your chemotherapy team see how you are getting on with treatment.

Using the diary
Your healthcare professional will explain how to use the diary. They will mark the days you will receive carfilzomib and add in the details of your appointments. Your healthcare professional will also add in other medicines you are taking in combination with carfilzomib and the days on which these should be taken.

They will also let you know if a medicine should be taken at a specific time of day.

Put a cross in the box to keep track of when you have received or taken each medicine in the diary.

Each double page includes one cycle of treatment. The first dose of carfilzomib is cycle one, day one. Each cycle is 28 days long. Cycle 2 begins immediately after the end of cycle 1.

Reporting side effects
Kyprolis®▼(carfilzomib) is subject to additional monitoring. This will allow quick identification of new safety information. If you get any side effects, talk to your doctor, pharmacist or nurse straightaway. This includes any possible side effects not listed in the package leaflet.

Adverse reactions/events should be reported to the Health Products Regulatory Authority (HPRA) using the available methods via www.hpra.ie. Adverse events should also be reported to Amgen Limited on +44 (0) 1223 436441.

By reporting side effects, you can help provide more information on the safety of this medicine.

For more information about carfilzomib, and a full list of side effects, refer to the patient information leaflet, which you can request from your healthcare professional or view online at: https://www.medicines.ie/medicines/kyprolis-32623/

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Please complete the 'other medications' table later in this diary with details of any additional medication you are taking

ADD THE TREATMENT COMBINATION YOU HAVE BEEN PRESCRIBED HERE:

DD / MM / YY

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Please use this box to note any comments from your healthcare professional during your appointments

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OTHER MEDICATIONS

OTHER MEDICATION YOU ARE TAKING		
NAME (Please add the name of the medication you are taking)	**DOSE** (Please add the dose of the medication prescribed)	**NOTES** (Please add any notes about the medication and any advice you have been given e.g. where to take the medicine, time of day to take, with or without food etc.)

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