



**British Society for
Rheumatology**

Rheumatoid Arthritis Register

Add a New Follow Up: Biologic/Targeted Therapy

In this section please record any changes to the participant's biologic / targeted therapy including temporary breaks in treatment and the reasons for any breaks.

Once a follow up is open you will have 14 days to enter data before the edit window is closed.

Patient Status: Current [\[Click here to reveal Patient Identifiable Data\]](#)

Study ID: **Cohort:** Erelzi **Follow-Up:** 1 **FUP Status:** **Due Date:** 01/05/2021 **Last FUP Date Entered:** 15/11/2021

Patient

Patient summary

Clinician FUP

Biologic Targeted Therapy

Other Current Therapy

Adverse Events

Disease Activity

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Biologic Targeted Therapy

Therapy changes

Have there been any changes to the patient's Rheumatoid Arthritis target therapy details in this follow-up period? (Excluding existing drug's additional dose infusions)

No

Yes (use 'Add New' to enter details after clicking 'Update')

[Update](#)

	Drug	Start Date	Stop Date	Dose	Frequency	Stop Reason	Doses & Batch	
Edit	Erelzi (etanercept biosimilar)	01/11/2021		50 mg	Once a week		01/11/2021	Delete

[Add New](#) [Tutorial](#) [Advance to Next Section](#)

If there are no changes to record select **No** and **Update**.

Patient

Patient summary

Clinician FUP

Biologic Targeted Therapy

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Biologic Targeted Therapy

Therapy changes

Have there been any changes to the patient's Rheumatoid Arthritis target therapy details in this follow-up period? (Excluding existing drug's additional dose infusions)

No **1 Select No**

Yes (use 'Add New' to enter details after clicking 'Update')

Update **2 Click Update to save answer**

	Drug	Start Date	Stop Date	Dose	Frequency	Stop Reason	Doses & Batch	
Edit	Erelzi (etanercept biosimilar)	01/11/2021		50 mg	Once a week		01/11/2021	Delete

Add New **Tutorial** **Advance to Next Section** **3 Click Advance to Next Section**

If there has not been a change in biologic / targeted therapy but the participant has had further IV infusions of a prescribed drug, select **Edit** to add dose details. See Slide 7 for further details.

If there are changes to make to an existing biologic / targeted therapy record select **Yes** and **Update**.
Select **Edit** to edit the record.

Patient

Patient summary

Clinician FUP

Biologic Targeted Therapy

Other Current Therapy

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Biologic Targeted Therapy

Therapy changes

Have there been any changes to the patient's Rheumatoid Arthritis target therapy details in this follow-up period? (Excluding existing drug's additional dose infusions)

No

Yes (use 'Add New' to enter details after clicking 'Update')

1 Select Yes

2 Click Update to save answer

3 Select Edit

	Drug	Start Date	Stop Date	Dose	Frequency	Stop Reason	Doses & Batch	
Edit	Erelzi (etanercept biosimilar)	01/11/2021					01/11/2021	Delete

Add New **Tutorial** **Advance to Next Section**

Rheumatoid Arthritis Target Therapy

Drug

Biologic: Erelzi (etanercept biosimilar)

Commenced date: 1/11/2021

Dose: 50 mg

Route: SC

Frequency: Once a week

Dose Delay Reason:

Stop date: 15/11/2021 

4 Add stop date

Stop reason: Adverse Event 

5 Add stop reason, please provide details of Adverse Advent in 'other stop reason' box

Adverse Event/Other stop reason:

Adverse Event Details

Please remember to give details of all adverse events in the Adverse Events section of this follow up.

Save page

Cancel

6 Select Save Page

Add the biologic stop date and stop reason (e.g. Adverse Event)

Select **Save page**.

The biologic therapy record is now updated.

If a patient has restarted the same drug or a new biologic please add a new record. A new entry for the same biologic drug can be added (e.g. if the biologic was stopped temporarily due to an adverse event).

If the patient has started a new biologic, please enter details of this in the Biologic Targeted Therapy section. A stop date for the current biologic / targeted therapy must be added before a new biologic / targeted therapy start date can be added.

Patient

Patient summary

Clinician FUP

Biologic Targeted Therapy

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Biologic Targeted Therapy

Therapy changes

Have there been any changes to the patient's Rheumatoid Arthritis target therapy details in this follow-up period? (Excluding existing drug's additional dose infusions)

No

Yes (use 'Add New' to enter details after clicking 'Update')

Update

	Drug	Start Date	Stop Date	Dose	Frequency	Stop Reason	Doses & Batch	
Edit	Erelzi (etanercept biosimilar)	01/11/2021	15/11/2021	50 mg	Once a week	Adverse Event	01/11/2021	Delete
Edit	Erelzi (etanercept biosimilar)	25/11/2021		50 mg	Once a week		25/11/2021	Delete

Add New  **Click Add New to add Biologic Therapy**

If there has not been a change in biologic / targeted therapy but the participant has had further doses of an IV drug select **Edit** to add dose details.


	Drug	Start Date	Stop Date	Dose	Frequency	Stop Reason	Doses & Batch	
Edit	Remsima (im	01/02					01/02/2019	Delete

[Add New](#) [Tutorial](#)

Doses & Batch
You can only delete the doses entered in the current follow up.

Dose date	Dose	Batch
01/02/2019	180	

Add drug dose or batch details

Dose date:  **1** Enter dose date

Dose: mg **2** Enter dose

Batch:

[Save drug dose or batch details](#) **3** Click Save drug dose or batch details to save information

The Biologic / targeted therapy section is now completed.
The next section is **Other Current Therapy**.

Study ID **Cohort:** Anti-TNF **Follow-Up:** 5 **FUP Status:** In Edit Window **Due Date:** 04/02/2022 **Last FUP Date Entered:**

Patient

Patient summary

Clinician FUP

Biologic Targeted Therapy

Other Current Therapy

Adverse Events

Disease Activity

Additional Info

Administrative

Preview Queries

Close Edit Window

Feedback / Comments

Biologic Targeted Therapy

Therapy changes

Have there been any changes to the patient's Rheumatoid Arthritis target therapy details in this follow-up period? (Excluding existing drug's additional dose infusions)

No

Yes (use 'Add New' to enter details after clicking 'Update')

Update

	Drug	Start Date	Stop Date	Dose	Frequency	Stop Reason	Doses & Batch	
Edit	Humira (adalimumab originator)	15/08/2019		40 mg	Fortnightly			Delete

Add New **Tutorial** **Advance to Next Section**

Click Advance to Next Section