<b>85mm</b> ← → →		<b>85mm</b> →
Erelzi (Etanercept) Patient Card	$\uparrow$	Congestive Heart Failure
This card contains important safety information that you need to be aware of before you are given Erelzi and during treatment with Erelzi. If you do not understand this information, please ask your doctor to explain it to you  Show this card to any doctor involved in your treatment. See the Erelzi package leaflet for more information. Keep this card with you for 2 months after the Erelzi dose, since side effects may occur after your last dose of Erelzi.	25mm	If you develop symptoms suggestive of congestive heart failure or worsening of existing congestive heart failure, such as shortness of breath, swelling of ankles, persistent cough or fatigue, seek medical attention immediately.
Infections		Other Information (please complete):
Erelzi may increase your risk of getting infections, which		• Patient's name:
could be serious.  • You should not use Erelzi if you have an infection. If you		• Doctor's name:
<ul> <li>are not sure, ask your doctor.</li> <li>If you develop symptoms suggestive of infections, such as fever, persistent cough, weight loss, or listlessness, seek medical attention immediately.</li> <li>You should be evaluated for tuberculosis (TB). Ask your doctor to record the dates and the results of your last screening for TB below:</li> </ul>		• Doctor's phone:
Test:		It is important that you and your doctor record the brand name and batch number of your medication.
Date:		Brand Name:
Results:		Batch Number:
Test:		<b>Further Information</b> Please read the Erelzi Package Leaflet for further information.
Date:  Results:		MHRA approval: June 2022
Please ask your doctor to list your other medications that may increase your risk of infection.		
		UK 212890 0000000000

**FRONT PAGE** 

## **BACK PAGE**

Artwork P	roof Box:				
Variation:	N004: RMP update v 5.1 and alignment to originator + approval date			Technical Colours:	
Proof no:	003.1			Legend:	
Date prepared:	09/06/2022			Cutting:	
Font size:	10pt				
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SKUs:	00000000				
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Novartis data	if applicable:				
Version:	0000000				
Print Proof date:	0000000				
AQWA#:	0000000				
Production site:	0000000				
Technical num:	0000000			A 65555	
Old Comp num:	00000000			& SANDOZ	
Signature	e Panel:				