

# British Society for Rheumatology Biologics Register for Rheumatoid Arthritis (BSRBR-RA)




Monitoring the long-term safety of biologic, biosimilar, and other new targeted therapies in the UK.

Over 30,000 patients registered in the study since 2001.


**NEW REGISTRATIONS AND FOLLOW-UP DATA IS SUBMITTED ONLINE, DIRECTLY IN TO THE SECURE STUDY DATABASE.**

To sign up please visit <https://bsrbr.org/database/>

## Eligibility for registration

	Diagnosis of Rheumatoid Arthritis		Aged 16 Years or over	Starting eligible biologic treatment	
---	-----------------------------------	---	-----------------------	--------------------------------------	---

ANTI-TNF originators	BIOSIMILARS	OTHER TARGETED THERAPIES				
<p>adalimumab originator* <b>Humira</b></p> <p>etanercept originator* <b>Enbrel</b></p> <p>infliximab originator* <b>Remicade</b></p> <p><small>*patients must be biologic, biosimilar &amp; targeted therapy naive to be eligible</small></p>	<table border="0"> <tr> <td style="vertical-align: top;"> <p>adalimumab biosimilars <b>Amgevita</b> <b>Hulio</b> <b>Hyrimoz</b> <b>Idacio</b> <b>Imraldi</b> <b>Yuflyma</b></p> <p>infliximab biosimilars <b>Flixabi</b> <b>Inflectra</b> <b>Remsima IV</b> <b>Remsima SC</b></p> </td> <td style="vertical-align: top;"> <p>etanercept biosimilars <b>Benepali</b> <b>Erelzi</b></p> <p>rituximab biosimilars <b>Rixathon</b> <b>Ruxience</b></p> <p>tocilizumab biosimilars <b>Tyenne</b></p> </td> </tr> </table>	<p>adalimumab biosimilars <b>Amgevita</b> <b>Hulio</b> <b>Hyrimoz</b> <b>Idacio</b> <b>Imraldi</b> <b>Yuflyma</b></p> <p>infliximab biosimilars <b>Flixabi</b> <b>Inflectra</b> <b>Remsima IV</b> <b>Remsima SC</b></p>	<p>etanercept biosimilars <b>Benepali</b> <b>Erelzi</b></p> <p>rituximab biosimilars <b>Rixathon</b> <b>Ruxience</b></p> <p>tocilizumab biosimilars <b>Tyenne</b></p>	<table border="0"> <tr> <td style="vertical-align: top;"> <p>baricitinib <b>Olumiant</b></p> <p>certolizumab <b>Cimzia</b></p> <p>tofacitinib <b>Xeljanz</b></p> <p>filgotinib <b>Jyseleca</b></p> </td> <td style="vertical-align: top;"> <p>tocilizumab <b>RoActemra</b></p> <p>sarilumab <b>Kevzara</b></p> <p>upadacitinib <b>Rinvoq</b></p> </td> </tr> </table>	<p>baricitinib <b>Olumiant</b></p> <p>certolizumab <b>Cimzia</b></p> <p>tofacitinib <b>Xeljanz</b></p> <p>filgotinib <b>Jyseleca</b></p>	<p>tocilizumab <b>RoActemra</b></p> <p>sarilumab <b>Kevzara</b></p> <p>upadacitinib <b>Rinvoq</b></p>
<p>adalimumab biosimilars <b>Amgevita</b> <b>Hulio</b> <b>Hyrimoz</b> <b>Idacio</b> <b>Imraldi</b> <b>Yuflyma</b></p> <p>infliximab biosimilars <b>Flixabi</b> <b>Inflectra</b> <b>Remsima IV</b> <b>Remsima SC</b></p>	<p>etanercept biosimilars <b>Benepali</b> <b>Erelzi</b></p> <p>rituximab biosimilars <b>Rixathon</b> <b>Ruxience</b></p> <p>tocilizumab biosimilars <b>Tyenne</b></p>					
<p>baricitinib <b>Olumiant</b></p> <p>certolizumab <b>Cimzia</b></p> <p>tofacitinib <b>Xeljanz</b></p> <p>filgotinib <b>Jyseleca</b></p>	<p>tocilizumab <b>RoActemra</b></p> <p>sarilumab <b>Kevzara</b></p> <p>upadacitinib <b>Rinvoq</b></p>					

Registration within 6 months of the therapy start date\* 

\*For patients already registered who are starting a new therapy, cohort switch request needs to be made within 24 months of therapy start.

Please contact the BSRBR-RA team with any questions

<https://bsrbr.org/> 0161 275 1652 [biologics.register@manchester.ac.uk](mailto:biologics.register@manchester.ac.uk)

[https://twitter.com/BSRBR\\_RA](https://twitter.com/BSRBR_RA)