<i>Please com</i> Name: Hosp. No.: NHS/CHI: DoB: Gender:	nplete or c	e E Female	cker: Address:	BADBIR II	BIR Biologics and Immunomodulators Register
BAD Bi	ologics	and Immuno	omodulato	rs Register Baseline Cli	nical Questionnaire
Today's Da	ate:			Date of Consent:	Sent to BADBIR?
Date Enter	ed on to	Database:			
oriasis 1. Does th	e patient h	ave a <u>past history</u> o	f the following? No		Νο
Eryth	rodermic ps	oriasis		Generalised pustular psoriasis	
2. What t	ype of psori	asis does the patier	nt <u>currently</u> have	2	
			Yes No		
	Chroni	c plaque psoriasis	-	Small (≤3cm diam) Large	e (>3cm diam)
	Flexura	ll/intertriginous			
	Seborr	hoeic psoriasis			
	Scalp				
	Palms/	soles (non pustular)			
	Nails		-	 Indicate number of nails affected 	
	Guttate	e psoriasis			
	Unstab	le psoriasis			
	Erythro	odermic			
	Genera	lised pustular psori	asis	Yes	No
	Localis	ed pustular psoriasi	5	Acrodermatitis Hallopeau	
	Other	(please specify belo	w)	Palmoplantar pustulosis	
2 Plassa		e following details:			
	•	-		1	
Year of o	diagnosis (be	est approximation)		Year first seen by a dermato	ologist
4. Does th sibling or sease Severity	-	ave a family history	of psoriasis? (i.e	. first-degree relative such as parent	t, Yes No Don't know
5. Does th	-	ave diagnosis by a r	-	· –	Year of Diagnosis
6. Please	document a	III recent PASIs & PO	GAs including the	pre-BADBIR registration treatment	PASI:
PASI	Date	Psoriasis Global	Patient	nl	patient has pustular psoriasis ease document BSA:
		Assessment	Completed PGA		BSA Date
				Moderate	
				Mild Almost clear	I
				• Clear	

Current Drug Therapy 7. Is the patient currently on any of the following topical treatments?												
Topical pimecrolimus Yes No Topical tacrolimus Yes No												
8. Please list all the patient's curre	-	py for ar	y indica	tion (I	Please note topi	ical treatme	ents apo	art fro				
the two listed above are not requi	-	ato Star	tod		DRU	G		r	Date St	artad		
d	<mark>Date Started</mark> d m m y y			у		<u>u</u>	d	d L	m	m	у у	/
]							
] ī							
Psoriasis Treatment		· · · ·			<u> </u>			·	. <u> </u>			
9. Is the patient currently receiving Amgevita (adalimumab)	g <u>biologi</u>	<u>treatm</u>	<u>ent</u> for t	heir p	soriasis?	Yes	No					
								b t	l m	m	у	У
Benepali (etanercept) Cimzia (certolizumab pegol)	Comm	encemei	nt date o	of this	episode of biolo	gic therapy:						
Cosentyx (secukinumab)	Is this	the patie	ent's first	t expo	sure to a biologi	c agent:	Yes		N	10		
Erelzi (etanercept)	n	ose:				le: Was the rec	_				lowed?	
Hyrimoz (adalimumab)					Yes	No*			ly unkno			
Ilumetri (tildrakizumab)	Frequ	ency:				ease provide de		eviation	ii oiii sch	euule.		
Imraldi (adalimumab)												
Kyntheum (brodalumab)			ZESSLY O	NLY: Pr	rovide	RECOMMENDED OPENING SCHEDULES: Amgevita: 80mg week 0, 40mg fortnightly from week 1						
Skyrizi (risankizumab)	administration dates d d m m y y				Cimzia: 400 mg at weeks 0, 2 and 4 Batch number Cosentyx: 300mg at weeks 0, 1, 2, 3 & 4							
Taltz (ixekizumab)					Hyrimoz: 80mg week 0, 40mg fortnightly from week Ilumetri: 100mg at weeks 0 & 4. 12 weekly thereafte			fter				
Tremfya (guselkumab)					Imraldi: 80mg week 0, 40mg fortnightly from week 1 Kyntheum: 210 mg at weeks 0, 1 and 2							
Zessly (infliximab)					Skyrizi: 150mg at weeks 0 & 4. 12 weekly thereafter Taltz: 160mg at week0, 80mg at weeks 2, 4, 6, 8, 10, and 12 Tremfya: 100mg at week 0, 100mg at week 4							
						Treninya. 1001		K 0, 100h				
10. Is the patient currently receiv	ing a <u>sm</u> a			nunon			psorias	is?	Yes	s	No	
<u>(Please</u> <u>DRUG Tick)</u> <u>Dos</u>	se (mg)	<u>Freque</u>	<u>ncy</u>		Date Starte	<u>d</u>						
Skilarance				Γ								
(dimethyl fumarate)		Average I	Daily Dose									
11. Is the patient currently receiving <u>conventional therapy</u> for their psoriasis? Yes No												
	-				-							
DRUG		<u>ease</u> ck)	J/cm ² o mg	<u>or</u>	<u>Frequency</u>		Date St			.,		
Oral PUVA MTX Only:							only:					
		1			1						Oral	Sub-Cut
Methotrexate					<u> </u>							
Ciclosporin					Average Daily Dose	<u> </u>						
Acitretin												
Fumaderm					Average Daily Dose							
Hydroxycarbamide												
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12. Please list <u>all previous</u> systemic anti-psoriatic therapy:

If none please tick

Drug	Start date	Stop date	Stop reason*				
*Stop reasons: Adverse Events, Clinical Trial, Contraindication, Death, Financial Consideration, Inefficacy, Inefficacy and							
Adverse Events, Other (please provide details), Patient Choice, Patient Non-Compliance, Remission, Titration							
morbidities							
13. Has the patient <u>ever</u> had (i.e. <u>requ</u>	ired treatment for) any of	the following illnesses?					

Hypertension	Yes	Year of Onset
Hypertension		
Cardiovascular Disease	Yes	Year of Onset
Angina		
Myocardial Infarction		
Stroke / Cerebrovascular		
Disease		
Peripheral Vascular		
Disease		
Dyslipidaemia		
Diabetes	Yes	Year of Onset
Type 1		

(please tick <u>all</u> that apply)

Autoimmune Disorders	Yes	Year of	f Onset
Туре 2			
Type 1			

Thyroid Disease			
Alopecia Areata			
Vitiligo			
Psoriatic Arthritis			

Thrombosis	Yes	Year of Onset
Deep vein thrombosis		
Pulmonary embolism		
Asthma		
COPD (including chronic bronchitis, emphysema)		

Liver Disease	Yes	Year of Onset
NAFLD (non-alcoholic fatty		
liver disease, including fatty		
liver and NASH)		
Alcoholic Liver Disease		
Viral Hepatitis		
Autoimmune Hepatitis		
Inherited Liver Disease		
(inc. haemochromatosis)		

If none please tick

Kidney Disease	Yes	Year of Onset
Chronic Kidney Disease		
Glomerular Disease		
Renovascular Kidney		
Disease		
Inherited Renal Disease		
(polycystic kidney disease)		
•		N (0)
Peptic Ulcer	Yes	Year of Onset
Peptic Ulcer		
Demyelination	Yes	Year of Onset
Optic Neuritis	103	
Multiple Sclerosis		
Transverse Myelitis		
Chronic Inflammatory De-		
myelinating Polyneuropathy		
Guillain-Barre Syndrome		
Guillant-Barre Synuronne		
Epilepsy	Yes	Year of Onset
Epilepsy		
Peptic Ulcer	Yes	Year of Onset
Peptic Ulcer		
Non-Skin Cancer	Yes	Year of Onset
Please specify type / site:		
Douchistria	Vee	Veen of Oreest
Psychiatric Depression	Yes	Year of Onset
· · ·		
Anxiety		
Inflammatory Bowel	Yes	Year of Onset
Crohns		
Ulcerative Colitis		
Other (please specify)	Yes	Year of Onset

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Skin

Skin Cancer risk factors:

Burns easily, never tans Burns easily, tans minimally

Burns minimally, tans well Rarely burns, tans profusely Never burns, deeply pigmented

Burns moderately, tans gradually

Description

14a) Please indicate Fitzpatrick skin type in box below

(Please indicate number) and site below)

Fitzpatrick Skin Type	Please tick	Туре	Site	Number
1		SCC		
2		BCC		
3		Melanoma		
4		Melanoma in situ		
5		Actinic keratosis		
6		Bowen's disease		
	<u> </u>	Keratoacanthoma		

Additional Information

Date:

UV Therapy

15. Has the patient ever had	UV thera	py? Yes	No If <u>YES</u> , p	please complete the fo	ollowing:
UV Therapy Details	Yes	No. of Courses	No. of Treatments	Cumulative Dose (J/cm ²)	Data Known to be Accurate?
Broadband UVB					
Narrowband LIVB					

Narrowband UVB			
TOTAL BODY PUVA			
Oral PUVA			
Topical PUVA			
HAND AND FOOT PUVA			
Oral PUVA			
Topical PUVA			

Lab Values

16. Please complete the following laboratory 17. What is the patient's current values (recent i.e. within last 6 months): (i.e. at the time that the biologic/systemic agent was started) blood pressure? LABORATORY VALUES Result Date Systolic mm Haemoglobin count (g/dL) Diastolic White cell count $(x10^{9}/L)$ mm Platelet count (x10⁹/L) 15. What is the patient's <u>current</u> (i.e. at the time that the biologic/systemic agent was started) Creatinine (µmol/L) height, weight and waist circumference? Transaminase ALT (U/L) Height cm Cholesterol (mmol/L) Weight kg Triglyceride (mmol/L) Waist cm HDL (mmol/L) circumference PBQ & QoL Questionnaires If paediatric patient: The following patient CAGE PBQ ⁽²⁾HAQ questionnaires PBQ ⁽¹⁾DLQI cDLQI should also be completed: ⁽²⁾cHAQ HADS EQ-5D-y EuroQol (1) It is not essential but a DLQI taken prior to drug commencement is preferred (2) (Only if patient has a rheumatologist's diagnosis of inflammatory arthritis) Signature

Please sign and date below:

Signature:

Name:

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1	.4b)	History of prior neoplastic or pre-cancerous lesions?	Yes
			• •

No