Please complete or attach patient sticker: Name: Hosp. No.: NHS/CHI: DoB: Badbir ID: Badbir ID:							
BAD Biologic Interventions Register Clinical Follow-Up Questionnaire							
BIOLOGIC COHORT ONLY:							
Since the patient's last follow up have their been any changes to their biologic therapy? No If we place record all changes for biologic							
If yes, please record all changes for biologic							
Drug Batch Number Unit Frequency Date started ddmmyy Date of final dose ddmmyy *Stop reason							
*Stop reasons (may have than one reason) (1) Inefficacy (2) Remission (3) Adverse Events (4) Other (please provide detail)							
If <u>infliximab or ustekinumab</u> please provide the administration dates d d m m y y mg/kg							
CONVENTIONAL COHORT ONLY:							
Conventional Therapy Since the patient's last follow up have there been any changes to their systemic therapy? Yes If yes, please record all changes for systemic agents:							
Drug Dose / Date started ddmmyy Date of final dose ddmmyy *Stop reason							
*Stop reasons (may have than one reason) (1) Inefficacy (2) Remission (3) Adverse Events (4) Other (please provide detail)							
Since the patient's last follow up have they started biologic therapy? No If yes please enter details of their biologic therapy in the Biologics Only box at the top of this page. Version 8 01/07/2015 p.1 of 4							

UV Therapy			
IIV Iherany	4 10 4 10 1		
	UV Inerany		

Since the patients last follow-up have they had any UV therapy? If yes please complete the following:

Yes		
No		

UV Therapy Details	Yes	No. of Courses	No. of Treatments	Cumulative Dose (J/cm²)	Data Known to be Accurate?
Broadband UVB					
Narrowband UVB					
TOTAL BODY PUVA					
Oral PUVA					
Topical PUVA					
HAND AND FOOT PUVA					
Oral PUVA					
Topical PUVA					
UVA 1					

Concomitant Therapy

Since the patient's last follow up have they had any changes to their concomitant therapy? Please list all changes below: (please note we do not need details of topical therapy

for psoriasis except for tacrolimus and pimecrolimus)

Drug	Start date	Stop date	Are these dates estimated?

Lab Values

Please complete the following laboratory values (recent i.e. within last 6 months):

LABORATORY VALUES	Result	Date
Haemoglobin count (g/dL)		
White cell count (x10 ⁹ /L)		
Platelet count (x10 ⁹ /L)		
Creatinine (μmol/L)		
Transaminase ALT (U/L)		
Cholesterol (mmol/L)		
Triglyceride (mmol/L)		
HDL (mmol/L)		

FUP7 +: Lab Values not required

	Tuberculosis (Not Latent)	on/Failure •	Serious Hepatic Dysfunction/Failure	ious Hepat	• Ser	vens Disease)	Melanoma / Skin Cancer (inc. Bowens Disease)		of 4	Version 8 01/07/2015 p.3 of 4	ion 8 01,	Ven
disation Only)	Surgery (Overnight Hospitalisation Only)	otic Neuritis	Serious Demyelination/Optic Neuritis	ious Demy	• Ser		• Malignancy (not inc. skin)	vent	Medically Important Event	Medicall	7	
	Serious Skin Reaction	Failure	Serious Congestive Heart Failure	ious Conge	• Ser		Lymphoproliferative Disease	reatening	Was in any way life threatening	Was in a	6	
vernight Hospitalisation	Serious Psoriasis Flare (Overnight Hospitalisation Only)		tion House					ion	Congenital malformation	Congenit	5	
Illness	serious Lupus/Lupus like iliness		holism	Pulmonary Embolism	• Pul			Significant loss of function or disability	nt loss of func	Significar	4	
	Serious linection (excl. 18)			Preanancy	• Pre		Cerebrovascular Accident (CVA)	ungals	IV Anti-biotics/virals/fungals	IV Anti-b	ω	
R)	Serious Infection (avel T		zse	Coronary Disease	Cor		serious neutropenia		isation	Hospitalisation	2	
Reaction	Serious Hypersensitivity Reaction	. tte	Mvocardial Infarction/Acute	ocardial In	• 8	lor	Aplastic angemia, pancytopaenia or			Death	_	
eeds to be completed:	the following, an Event of Special Interest (ESI) form needs to be	Event of Special	owing, an		de any o	listed includ	If any of the events you have listed include any of	fication	SAE Classification	TO .	Code	
Resolved Resolved	If 'Hospitalisation' Admission Date: Discharge Date:	If 'Yes' Name of										
Resolved Resolved W/ Sequelae Not Resolved M/ Sequelae Not Resolved Dunknown Death	If 'Hospitalisation' Admission Date: Discharge Date:	If 'Yes' Name of										
Resolved Resolved Resolved w/ Sequelae	If 'Hospitalisation' Admission Date: Discharge Date:	If 'Yes' Name of										
Resolved Aresolved w/Sequelae Not Resolved Unknown Death	If 'Hospitalisation' Admission Date: Discharge Date:	If 'Yes' Name of Biologic:										
Resolved	If 'Hospitalisation' Admission Date:	If 'Yes' Name of										
Is the event an ESI? If yes please select from list (see below) Outcome of the event?	Is the event a SAE ? If yes please select code (see below)	Is the event related to biologic therapy? Yes, No or Possibly Not required for conventional cohort patients Yellow Card	Estimated? Is the event ongoing?	od atte	Start Date Estimated? Stop	Start date	Description of event	Desc	gription please gnosis if ole)	Short description of event (please record diagnosis if available)	Event No.	
		ous) from this follow-up period	m this fo	rious) fro	non-se	h serious and	Please enter details of $\overline{\sf ALL}$ adverse events (both serious and non-seri	Please enter de	No	7		
			¥	box below	າs in the	classification	A <u>serious adverse event</u> (SAE) is defined by the classifications in the b	A <u>serious adver</u>	experienced any adverse events)? γ_{es}	experienced a events)?	eve	
itment or medic	ot related to any trea	nt whether or n	in a patie	ccurring	l event o	lly untoward	An <u>adverse event (</u> AE) is defined as any medically untoward event occurring in a patient whether or not related to any treatment or medication	An <u>adverse eve</u>	last data r patient	Since date of last data entry has your patient	Sin eni	
										nts	Adverse Events	Ad

Current Disease Severity									
Please indicate the current diseas	se severit	y (i.e.	at the time	the patie	ent started the new o	drug)			
BSA			Only if the	natient ha	s pustular psoriasis,				
Date o	of BSA .	/	/	Julient nu	s pusturur psorrusis,				
Please details of all PASI's that	have bee	en com	pleted sinc	e the pat	ients last follow-up.				
PASI			te of PASI		Psoriasis Global As				
Pagriasia Clahal As				(1) Causa					
Psoriasis Global As	sessmen	. score	: :	(1) Seve	erate to severe				
				(3) Mod					
				(4) Mild					
(4) Mild (5) Almost clear									
				(6) Clear	r				
Has the patient been diagnosed w						es No			
*if this is a new diagnosis please ren	nember to	add th	nis as an advi	erse event	*				
Additional Information									
What is the patient's <u>cu</u>	urrent we	ight a	nd waist cir	cumferer	nce?				
		W	eight		kg		JP9 +:		
	Wa		umference				eight / W ot require		
	vva	ist circ	difference		cm				
Detical Fall control of the control									
Patient Follow-up Questionnaire Occupation	on Qus		Lifestyle C	us	If paediat	ric patient:			
The patient questionnaire should also be completed containing:			CAGE		cDLQ				
	oQol _		*HAQ		EQ-5D-y		*cHAQ		
FUP 7+: Patient Questionnaire is not required									
<u></u>					t has a rheumatologist's lammatory arthritis)				
Signature	Ple	ease si	gn and date	below					
	<u></u>	<u> </u>	, aa aate						
Clinician's signature:			D	ate:					
			•						

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