CAR-T treatment for cancer patients

Funding request form



This form is to be used to request Bupa funding for your patient with acute lymphoblastic leukaemia (ALL), diffuse large B-cell lymphoma (DLBCL), mantle cell lymphoma (MCL) or primary mediastinal B-cell lymphoma (PMBCL) to receive CAR-T treatment.

For us to accurately assess your funding request, you need to:

- 1. Complete this form in full including all relevant elements of your patient's current medical condition and medical history
- 2. Attach multidisciplinary team (MDT) and/or NCCP meeting notes
- 3. Attach your patient's full medical notes

We're unable to agree funding based on incomplete forms or evidence. If we need to ask you for more information, this is likely to delay our funding decision and risks delaying your patient's treatment.

. About the patient			
Title (please tick) Miss Mrs Mrs M	s Mr Dr Other (please state	e)	
Name			
Date of birth	YYY		
Bupa membership number			
2. Clinician's details			
Name of requesting consultant	Specialty		
Bupa provider number	Hospital name		
Phone number	Email address		
S. Questions about the patie			
The patient has a creatinine clearance (as estimate	ent's general fitness	Last records	
The patient has a creatinine clearance (as estimated) The patient has a serum ALT/AST ≤ 2.5 ULN	ent's general fitness	Last recorde	ed value
The patient has a creatinine clearance (as estimated) The patient has a serum ALT/AST ≤ 2.5 ULN	ent's general fitness		ed value
The patient has a creatinine clearance (as estimated as a serum ALT/AST ≤ 2.5 ULN) The patient has a total bilirubin ≤1.5 mg/dl The patient has a cardiac ejection fraction ≥ 50%	ent's general fitness ted by Cockcroft Gault) ≥ 60 mL/min with no evidence of pericardial effusion	Last recorde	ed value
The patient has a creatinine clearance (as estimated as a serum ALT/AST ≤ 2.5 ULN) The patient has a total bilirubin ≤1.5 mg/dl The patient has a cardiac ejection fraction ≥ 50% as determined by an ECHO, and no clinically sign	ent's general fitness ted by Cockcroft Gault) ≥ 60 mL/min with no evidence of pericardial effusion	Last recorde	ed value ed value No
The patient has a creatinine clearance (as estimated as a serum ALT/AST ≤ 2.5 ULN) The patient has a total bilirubin ≤1.5 mg/dl The patient has a cardiac ejection fraction ≥ 50%	ent's general fitness ted by Cockcroft Gault) ≥ 60 mL/min with no evidence of pericardial effusion hificant ECG findings	Last recorde Last recorde	ed value ed value No
The patient has a creatinine clearance (as estimated as a serum ALT/AST ≤ 2.5 ULN) The patient has a total bilirubin ≤1.5 mg/dl The patient has a cardiac ejection fraction ≥ 50% as determined by an ECHO, and no clinically sign	ent's general fitness ted by Cockcroft Gault) ≥ 60 mL/min with no evidence of pericardial effusion ifficant ECG findings	Last recorde Last recorde Yes Last recorde	ed value No ed value
The patient has a creatinine clearance (as estimated as a serum ALT/AST ≤ 2.5 ULN) The patient has a total bilirubin ≤1.5 mg/dl The patient has a cardiac ejection fraction ≥ 50% as determined by an ECHO, and no clinically sign The patient has a no clinically significant pleural expressions.	ent's general fitness ted by Cockcroft Gault) ≥ 60 mL/min with no evidence of pericardial effusion afficant ECG findings effusion % on room air	Last recorde Last recorde Yes Last recorde	ed value One No ed value No

4. The patient's treatment to date

Past cand	er treatment					
Line of treatment	Drug regimen	Treatment response	Number of cycles	Treatment start date		Date of relapse
First				D D M M Y	Υ	D D M M Y Y
Second				D D M M Y	Υ	D D M M Y Y
Other				D D M M Y	Y	D D M M Y Y
Other				D D M M Y	Y	D D M M Y Y
Other				D D M M Y	Y	D D M M Y Y
All current	medication		-			
5. Que	estions about	the proposed tro	eatment	:		
Please ar	nswer the following	questions for any indicati	on			
Proposed	treatment start date		D D	M M Y Y	Υ	
Is the lead	consultant a haematolo	ogist accredited in the use of t	his drug?		Yes	s No
hospital/cl	inic and have the consu	mitting rights to a Bupa accred Itant and the hospital/clinic sig Bliver CAR-T therapy to Bupa p	gned a CAR-T		Yes	s No
	consultant a member o	f the Bupa accredited CAR-T	treatment hos	pital/clinic	Yes	s No
Has the histological diagnosis been either made by or reviewed and confirmed by a designated stem cell transplant centre?						s No
		ore lines of systemic therapy (d after the last line of systemic		cified therapies	Yes	s No
Does the p	patient have sufficient e	nd organ function to tolerate t	reatment?		Yes	s No
	tient had previous treat unotherapy?	ment with any genetically mo	dified autolog	ous or allogeneic	Yes	s No
	oses of tocilizumab be a release syndrome?	available for use in this patient	in the event c	of the development	Yes	s No
Will the CA (SPC)?	AR-T cells otherwise be	used as set out in its Summary	y of Product C	haracteristics	Yes	s No
1. Local M					Yes	
2. Nationa	al CAR-T Clinical Panel ((NCCP) or equivalent body?			Yes	s No
When was	this patient's case revie	ewed by the NCCP or	D D	M M Y Y	Υ	

Which CAR-T branded product is intended for this patient? Please explain why

equivalent body?

5. Questions about the proposed treatment (continued)

Has the patient been offered NHS CAR-T treatment?	Yes	No			
Have the patient's treatment funding options (Bupa and local NHS facility) been explained to them?	Yes	No			
Has the patient been made aware:					
That there are no clinical differences in the delivery of, and priority of access to, CAR-T treatment based on whether the treatment is funded privately or by the NHS?	Yes	No			
 Of any operational differences (such as access to a private room or other facilities)? 	Yes	No			
Please answer these further questions if the patient has acute lymphoblastic leukae	mia (ALL)				
Does the patient have ALL and CD19 ALL positivity in the bone marrow which is detectable using flow cytometry?	Yes	☐ No			
Does the patient have an isolated extramedullary ALL relapse following a stem cell transplant or two lines of systemic therapy?	Yes	☐ No			
Does the patient have an isolated extramedullary ALL relapse?	Yes	No			
Does the patient have known active central nervous system involvement in relation to their lymphoma?	Yes	☐ No			
Has the patient been treated with blinatumomab?	Yes	No			
Is the patient aged under 26 years on the date of the treatment request?	Yes	No			
Does the patient have a Karnofsky (age ≥16 years) or a Lansky (<16 years) performance status of at least 50%?	Yes	☐ No			
Please answer these questions if the patient has diffuse large B-cell lymphoma, mantle cell lymphoma or primary mediastinal B-cell lymphoma					
Has progressive disease been defined radiologically as per RECIST version 1.1 and is based on CT or MR scans? (not an increased SUV or PET scan)	Yes	☐ No			
Does the patient have primary central nervous system lymphoma?	Yes	No			
Does the patient have known active central nervous system involvement in relation to their lymphoma?	Yes	☐ No			
Is the patient aged 18 years or older on the date of request?	Yes	No			
Does the patient have an ECOG performance score of 0 or 1?	Yes	No			
Supporting evidence					
Please tick the boxes below to confirm that you've shared the following reports containing medic CAR-T funding request	al notes with B	Supa as part of this			
Patient's MDT and/or NCCP meeting notes Medical report including: Diagnosis and stage of cancer Previous and current cancer treatments Co-morbiditie Allergies Performance status Family history or previous genetic testing Palliative care information if relevant					
Pulmonary function test (necessary where the patient has poor lung or cardiac function) Pathology					
Imaging reports					
Electrocardiogram (ECG)					
Echocardiogram (ECHO)					
PET scan of whole body					
Brain MRI (necessary where the patient has CNS involvement)					

5. Questions about the proposed treatment (continued) Lab reports Full blood count with differential Chemistry panel Liver function test (LFT) Lactate dehydrogenase (LDH) test Ferritin test

6. Consultant's declaration

Please complete this section to confirm that the information on this form is accurate, that you've obtained informed consent from the patient and have explained all the risks and alternatives associated with this treatment.

I understand that the clinical information I've supplied may be considered to be a medical report for insurance purposes. I confirm

that my patient (or their legal representative) has given their permission for me to shar review this information, they've been given an opportunity before I submitted this form	
Consultant's name	Date D M M Y Y Y
General Medical Council number	
Further information	
Please send your completed form and supporting information to us within 24 hours of specialistnursesupport@bupa.com	receipt by secure email
Information you send to this email address may not be secure unless you send us your Egress account, go to https://switch.egress.com	email through Egress. To sign up for a free
We'll let you know by phone or secure email within seven working days of receiving yo patient's treatment is clinically eligible and covered by their policy.	ur completed form whether your Bupa
Please let us know how you'd prefer us to contact you about this Phone or secure email	
What's the best phone number or email address for us to use?	

If you've any questions, please call us on 0845 850 0465. We're here between 8am and 6pm Monday to Friday. We may record or monitor our calls.

7. Appendix

MDT requirements for Bupa CAR-T network hospitals

Before we can let you know whether we'll fund your patient's proposed CAR-T treatment, their case needs to be reviewed by an MDT based at a Bupa accredited CAR-T treatment hospital/clinic and, if applicable, by the NCCP or an equivalent body, to endorse clinical eligibility and prioritisation. Bupa CAR-T network hospital MDTs need to be quorate and include sufficient representation from the following:

- At least two haemato-oncologists (either haematologists or some medical oncologists) who specialise in your patient's tumour type being discussed,
- one from each hospital contributing to the MDT
- At least one haematopathologist from the Specialist Integrated Haematological Malignancy Diagnostic Services to provide the diagnostic information.
- Input from the clinical oncologist, as required, when radiotherapy is delivered for your patient if they have acute lymphoblastic leukaemia (ALL) or lymphoma
- At least one radiologist specialising in haematology/lymphoma and input from neuro-radiology as required
- At least one clinical nurse specialist acting as your patient's advocate and accountable for ensuring that frailty is taken into account and documented.
- At least one specialist palliative care doctor on the specialist register or a nurse experienced in palliative care to liaise with specialists from other sites
- At least one geriatrician if your patient is over the age of 80.
- A neurologist if your patient has significant neurological comorbidities.
- A cardiologist if your patient has significant cardiology comorbidities
- Support staff to organise team meetings, provide secretarial support and submit to Bupa all required documents

In line with our contract, we may need a peer review process to take place to confirm the patient's clinical suitability for CAR-T treatment.