							MRC Population Health Research Unit				RSITY OF FORI)
8.		Participant ID:	9]
		Site ID:	9]
C	Consent form											•••
E	SSENTIAL parts of the EMPA-KIDNE	EY study (please initial <u>ea</u>	<u>ch</u> box)							ease	
be	confirm that I have read and understoodelow). I have had the opportunity to contaitsfactorily.		sk que	stions	The	se ha	ve be			er 🔽	<u>(on e</u>	a
	understand that my participation in EM ithout giving any reason, and without m					to wit	hdraw	ı at a	ny tim	e,		
	agree to blood/urine samples being tak y GP (if necessary).	en for EMPA-KIDNEY-rela	ted tes	ts and	for r	esult	s to b	e pro	vided	to		
lo Kl	give permission for relevant sections of my medical notes, and information collected during the study, to be boked at, in confidence, by authorised individuals from my local study site, the University of Oxford, EMPA- KIDNEY Regional Coordinating Centres, Boehringer Ingelheim, and regulatory authorities (UK or foreign) to sheck that the study is being carried out correctly.											-
in re	understand that my identifiable data wi formation about my health (including gistration and end-stage renal disease ay be provided to EMPA-KIDNEY both) but not necessarily limit) held by NHS Digital, othe	ed to r NHS	cance bodie	r, ho s and	ospita d the	l adn UK R	nissio	ns, civ	/il		
6 I.	understand that my GP will be informed	of my participation in the E	MPA-ł		Y stu	dy.				L		
	agree to the collection, storage, processing, transfer and use of my personal data including blood/urine imples as explained in the EMPA-KIDNEY Participant Information Leaflet (version number above).											
3 Ia	I agree to take part in the EMPA-KIDNEY study.										Pleas Yes (_
0	<u>OPTIONAL</u> parts of the EMPA-KIDNEY study (please initial the box indicating yes <u>or</u> no)											,
	erstand that my participation in this optional part is voluntary and that I am free to withdraw at any time, but giving any reason, and without my medical care or legal rights being affected.										Yes	
m	agree that the left over samples of my b ay be stored, processed and used for articipant Information Leaflet (version n	future scientific research	(other	than g	enet	ics) a	is exp	laine	d in th		Yes Yes	
st	agree that left over samples of my bloc ored, processed and used for future <u>g</u> eaflet (version number above). I unders	enetic scientific research a	s expla	ained i	n the	Part	icipar				Yes	
	I agree to be contacted by the study coordinators to provide updated information about my health or be invited to new studies after the scheduled follow-up period of the EMPA-KIDNEY study.										163	
						D	D/M	MM	/ Y Y	ΥY		
PF	RINTED name of consenting patient	Signati					То	day's	date			
						D			/ Y Y	ΥY		
PF	RINTED name of consent taker	Signati	ıre				То	day's	date			

Top copy (yellow): participant's trial file - Middle copy (pink): co-ordinating centre (where relevant) - Bottom copy (white): participantEMPA-KIDNEY Informed Consent Form V1.2. IRAS no.: 23621126-APR-2018