Additional file 6

Time	**Visit 1** **Pre-Screening**	**Visit 2** **Baseline visit** **Confirmation of eligibility & Randomisation**			**Telephone call**	**Visit 3** **Week 4 Treatment Visit**	**Telephone call**	**Visit 4** **Week 12 EoT**	**Telephone call**
	Week -1	**Week 0**			**Week 2**	**Post Week 4**	**Week 6**	**Post week 12**	**4 weeks post visit 4**
Trial Discussed / PIS given[1]	X								
Informed Consent[2]		X							
Photography				X					
Medical History and demographics			X						
EQ5D[3]			X						
DLQI[3]			X						
Patient treatment questionnaire								X	
Compliance assessment					X		X		
Volume measurement				X		X		X	
Tumour Pain Assessment				X		X		X	
Pregnancy Test[4]			X			X		X	
Blood sample								X	
Trial medication dispensed				X		X			
Trial medication returned						X		X	
Patient Diary				X		X		X	

Lesions biopsied								X	
Adverse events						X		X	X
Concomitant medications			X			X		X	
CRF completion				X		X		X	X

[1] Patient information sheet can be post to avoid any unnecessary journey.

[2] A minimum of 24hr for review of patient information sheet before patient can sign informed consent form.

[3] QoL only to be completed if patient was not part of the cohort 1 trial.

[4]Pregnancy test should be urine dipstick and for all women of childbearing potential.