	Part 1: Gen	eral Information		
YODA Project (Protocol) ID:	2018-3556			
Date:	13 September 2018			
Product Name:	Infliximab/ Ustekinumab/ Golimumab			
Therapeutic Area:	Immunology			
Product Class:	-			
	Antirheumatic Agents - Biologic Response Modifiers			
Condition(s) Studied:	Crohn's Disease, Ulcerative Colitis			
Protocol Number(s) and	Infliximab: NCT00036439	C0160T27		
Title(s):	NCT00036439			
	NCT00098633			
	NCT00207675			
	NCT00034438			
	NCT00336492			
	NCT00207662			
	NCT00207788			
	NCT00004941			
	NCT00537316 NCT01551290			
		C0168T16 (initial treatment ph	ace and repeated	
	treatment phas		ase and repeated	
	li catillent phas	se studies _j		
	Ustekinumab:			
	NCT02407236	CNTO1275UCO3001		
	NCT00771667	C0743T26		
	NCT01369329	CNTO1275CRD3001		
	NCT01369342	CNTO1275CRD3002		
	NCT01369355	CNTO1275CRD3003		
	NCT00265122	C0379T07		
	Golimumab:			
	NCT00487539	C0524T17		
	NCT00487533			
	NCT00488774			
		CNTO148UCO3001		
		CNTO148UCO2001		
	Part 2: Da	ata Availability		
Data Holder has authority to pr		l data or development partner	Yes	
has agreed to share clinical tria	l data.			
Comments:				
Data Holder has sharable electi	onic clinical trial	data or data can be converted	Yes	
to electronic format.				
Comments:				
De-identification and redaction	of clinical trial d	ata in accordance with current	Yes	
HIPAA and EU criteria allows pr	otection of parti	cipant privacy and		
confidentiality.	-			
Comments:				

The product and relevant indication studied has either been approved by	Yes			
regulators in the US and EU, or terminated from development.				
Comments:				
Data Holder has completed the clinical trial and trial has been completed for a	Yes			
period of at least 18 months (or results published in peer-reviewed				
biomedical literature).				
Comments:				
Part 3: Data Availability Summary				
Based on the responses to the above Data Availability questions, the				
requested clinical trial data are available for a data sharing request.				

Part 4: Proposal Review					
Question:			Response:		
Summary-level CSR data is appropriate for the proposed analysis.			No		
Participant-level data is appropriate for the proposed analysis.			Yes		
A similar analysis is underway or completed/pending disclosure by Janssen.			No		
Comments:					
	Part 5: Cost				
Estimated cost to share the requested clinical trial data.			Tier 2		
Tier 1: < \$5,000	Tier 2: \$5,000 - \$25,000	Tier 3: > \$25,00	00		