The YODA Project Research Proposal Due Diligence Assessment

DMARD Therapy NCT01606761 CNTO136ARA3003 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite Anti-TNF-Alpha Therapy Part 2: Data Availability Question: Response: Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A	Part 1: General Information			
Product Name: Sirukumab Therapeutic Area: Immunology Product Class: Antirheumatic Agents - Biologic Response Modifiers Condition(s) Studied: Arthritis, Rheumatoid Protocol Number(s) and Title(s): NCT01604343 CNT0136ARA3002 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-III-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite DMARD Therapy NCT01606761 CNT0136ARA3003 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-III-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite Anti-TNF-Alpha Therapy Part 2: Data Availability Question: Response: Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to Yes electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in Yes the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature).	YODA Project (Protocol) ID:	2019-4015		
Therapeutic Area: Product Class: Antirheumatic Agents - Biologic Response Modifiers Condition(s) Studied: Arthritis, Rheumatoid Protocol Number(s) and Title(s): NCT01604343 CNT0136ARA3002 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite DMARD Therapy NCT01606761 CNT0136ARA3003 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite Anti-TNF-Alpha Therapy Part 2: Data Availability Question: Response: Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to Yes electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA Yes and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in Yes the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature).	Date:	18 December 2019		
Product Class: Antirheumatic Agents - Biologic Response Modifiers Condition(s) Studied: Arthritis, Rheumatoid Protocol Number(s) and Title(s): NCT01604343 CNT0136ARA3002 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite DMARD Therapy NCT01606761 CNT0136ARA3003 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite Anti-TNF-Alpha Therapy Part 2: Data Availability Question: Response: Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to Yes electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in Yes the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature).	Product Name:	Sirukumab		
Condition(s) Studied: Arthritis, Rheumatoid NCT01604343 CNTO136ARA3002 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite DMARD Therapy NCT01606761 CNTO136ARA3003 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite Anti-TNF-Alpha Therapy Part 2: Data Availability Question: Response: Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in Yes the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A	Therapeutic Area:	Immunology		
Protocol Number(s) and Title(s): NCT01604343 CNT0136ARA3002 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite DMARD Therapy NCT01606761 CNT0136ARA3003 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite Anti-TNF-Alpha Therapy Part 2: Data Availability Question: Response:	Product Class:	Antirheumatic Agents - Biologic Response Modifiers		
Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite DMARD Therapy NCT01606761 CNTO136ARA3003 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite Anti-TNF-Alpha Therapy Part 2: Data Availability Question: Response: Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to Yes electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in Yes the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A	Condition(s) Studied:	Arthritis, Rheumatoid		
Question: Response: Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Yes Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to electronic format. Yes Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality. Yes Comments: N/A The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development. Yes Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Yes Comments: N/A	1	Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite DMARD Therapy NCT01606761 CNTO136ARA3003 A Multicenter, Randomized, Double-blind, Placebo-controlled, Parallel Group Study of CNTO 136 (Sirukumab), a Human Anti-IL-6 Monoclonal Antibody, Administered Subcutaneously, in Subjects With Active Rheumatoid Arthritis Despite		
Data Holder has authority to provide clinical trial data or development partner has agreed to share clinical trial data. Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A				
agreed to share clinical trial data. Comments: N/A Data Holder has sharable electronic clinical trial data or data can be converted to electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A		Question:	Response:	
Data Holder has sharable electronic clinical trial data or data can be converted to electronic format. Comments: N/A De-identification and redaction of clinical trial data in accordance with current HIPAA Yes and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in Yes the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A	agreed to share clinical trial data.			
De-identification and redaction of clinical trial data in accordance with current HIPAA Yes and EU criteria allows protection of participant privacy and confidentiality. Comments: N/A The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A	Data Holder has sharable electronic clinical trial data or data can be converted to electronic format.			
The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development. Comments: N/A Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A	De-identification and redaction of clinical trial data in accordance with current HIPAA Yes and EU criteria allows protection of participant privacy and confidentiality.			
Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature). Comments: N/A	The product and relevant indication studied has either been approved by regulators in the US and EU, or terminated from development.			
	Data Holder has completed the clinical trial and trial has been completed for a period of at least 18 months (or results published in peer-reviewed biomedical literature).			
Part 3: Data Availability Summary				
Based on the responses to the above Data Availability questions, the requested clinical trial data can be made available for data sharing.	, , , , , , , , , , , , , , , , , , , ,			
Part 4: Proposal Review				
Question: Response:	Question:			
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:	No			