Division	:	Worldwide Development
Information Type	:	Reporting and Analysis Plan (RAP)

Title	:	Reporting and Analysis Plan for An Open-Label Two-way Interaction Clinical Trial to Evaluate the Pharmacokinetic Interactions Between GSK3640254 and Dolutegravir in Healthy Subjects
Compound Number	:	GSK3640254
Effective Date	:	01-APR-2019

Description:

- The purpose of this RAP is to describe the planned analyses and output to be included in the Clinical Study Report for Protocol 209712.
- This RAP is intended to describe the full analyses required for the study.
- This RAP will be provided to the study team members to convey the content of the Statistical Analysis Complete (SAC) deliverable.

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1. INTRODUCTION

The purpose of this reporting and analysis plan (RAP) is to describe the analyses to be included in the Clinical Study Report for Protocol: 209712

2. SUMMARY OF KEY PROTOCOL INFORMATION

This is an open-label, single-sequence, two-way drug interaction study to investigate the pharmacokinetic (PK) interactions between GSK3640254 and dolutegravir (DTG). Treatment of human immunodeficiency virus (HIV) infection frequently involves combination therapy. It is important to understand any interactions and resulting changes in exposure (if any) when HIV medications are given in combination. Data from this study will contribute to dosing recommendations when GSK3640254 and DTG are given in combination.

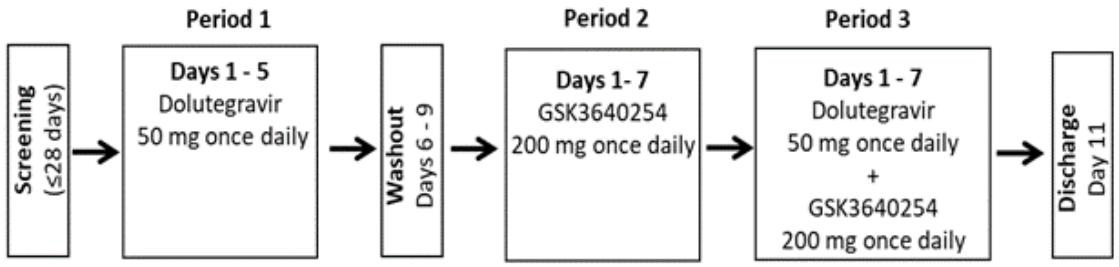
2.1. Changes to the Protocol Defined Statistical Analysis Plan

There were no changes or deviations to the originally planned statistical analysis specified in the protocol (Dated: 12/DEC/2018).

2.2. Study Objective(s) and Endpoint(s)

Objectives	Endpoints
Primary Objectives	**Primary Endpoints**
• To assess the effect of GSK3640254 on the steady state PK of DTG under fed conditions in healthy participants	• Area under the plasma concentration-time curve (AUC) from time 0 to the end of the dosing interval at steady state [AUC(0-τ)], maximum observed concentration (Cmax), and plasma concentration at the end of the dosing interval (Cτ) for DTG
• To assess the effect of DTG on the steady state PK of GSK3640254 under fed conditions in healthy participants	• AUC(0-τ), Cmax, and Cτ for GSK3640254
Secondary Objectives	**Secondary Endpoints**
• To assess the safety and tolerability of GSK3640254 and DTG administered alone and when given in combination in healthy participants	• Safety and tolerability parameters for adverse events (AEs)/serious AEs (SAEs), observed and change from baseline clinical laboratory assessments, electrocardiograms (ECGs), and vital sign measurements
• To characterize the steady state PK of GSK3640254 alone and when given in combination with DTG under fed conditions in healthy participants	• Time of maximum observed concentration (Tmax) and apparent terminal phase half-life (t1/2) for GSK3640254
• To characterize the steady state PK of DTG alone and when given in combination with GSK3640254 under fed conditions in healthy participants	• Tmax and t1/2 for DTG

2.3. Study Design

Overview of Study Design and Key Features	
Figure 1 Study Design Schematic  	
Design Features	• A phase 1, open-label, fixed-sequence, 3-period, two-way drug interaction study. • The study will consist of a screening period, 3 sequential treatment periods with a washout period between treatment periods 1 and 2, and a follow-up period. ○ Screening period: up to 28 days before the first dose ○ Treatment period 1: 5 days ○ Washout period: 4 days between the last dose of study intervention in Period 1 and the initiation of study intervention in Period 2 ○ Treatment period 2: 7 days ○ Treatment period 3: 7 days ○ Follow-up period: 4 days • Approximately 16 participants will be treated to ensure that 12 evaluable participants complete the study.
Dosing	• Treatment A: DTG 50 mg once daily (QD) (conventional 50 mg tablet) administered orally on Days 1 through 5 in Period 1 • Treatment B: GSK3640254 200 mg QD (2 × 100 mg capsules) administered orally on Days 1 through 7 in Period 2 • Treatment C: DTG 50 mg QD (conventional 50 mg tablet) coadministered with GSK3640254 200 mg QD (2 × 100 mg capsules) administered orally on Days 1 through 7 in Period 3
Time & Events	• Refer to Appendix 1: Schedule of Activities
Treatment Assignment	• This is an open-label study. All eligible participants will receive the same treatment.
Interim Analysis	• No interim analysis is planned for this study

2.4. Statistical Hypotheses

No formal hypotheses will be statistically tested for this study.

Analyses will be performed to assess the effect of GSK3640254 on the PK of DTG, and the effect of DTG on the PK of GSK3640254, as appropriate.

Analyses will be performed on the natural logarithms of the primary plasma PK parameters (AUC(0-τ), Cτ, and Cmax) using linear mixed-effect models with period as a

fixed effect and measurements within participant as repeated measures. Effects will be estimated, and confidence intervals (CIs) will be constructed for the following treatment comparisons:

- Period 3 versus Period 1 (DTG)
- Period 3 versus Period 2 (GSK3640254)

3. PLANNED ANALYSES

3.1. Final Analyses

The final planned primary analyses will be performed after the completion of the following sequential steps:

1. All participants have completed the study as defined in the protocol.
2. All required database cleaning activities have been completed and final database release (DBR) and database freeze (DBF) has been declared by Data Management.

4. ANALYSIS POPULATIONS

Population	Definition / Criteria	Analyses Evaluated
Screened	• All participants who signed the informed consent form	• Study Population
Safety	• All participants who received at least 1 dose of study medication. • This population will be used for all demographic and safety summaries	• Study Population • Safety
Pharmacokinetic Concentration	• All participants who underwent plasma PK sampling and had evaluable PK assay results. • This population will be used for the PK concentration listings, summary tables, and plotting of concentration-time data.	• PK Concentration
Pharmacokinetic Parameter	• All participants who underwent plasma PK sampling and had evaluable PK parameters estimated. • This population will be used for PK parameter listings, summary tables, and statistical analysis tables.	• PK Parameter • PK statistical analysis

Refer to Appendix 9: List of Data Displays which details the population used for each display.

4.1. Protocol Deviations

Significant protocol deviations (including deviations related to study inclusion/exclusion criteria, conduct of the trial, participant management or participant assessment) will be summarised and listed.

Protocol deviations will be tracked by the study team throughout the conduct of the study in accordance with the Protocol Deviation Management Plan dated 12FEB2019.

- Data will be reviewed prior to freezing the database to ensure all significant deviations and deviations which may lead to exclusion from the analysis are captured and categorised on the protocol deviations dataset.
- This dataset will be the basis for the summaries and listings of protocol deviations.

A separate listing of all inclusion/exclusion criteria deviations will also be provided. This listing will be based on data as recorded on the inclusion/exclusion page of the electronic case record form (eCRF).

5. CONSIDERATIONS FOR DATA ANALYSES AND DATA HANDLING CONVENTIONS

5.1. Study Treatment & Sub-group Display Descriptors

Treatment Group Descriptions		
Data Displays for Reporting		
Description	**Code**	**Order in TLF**
DTG 50 mg QD on Days 1 through 5 in Period 1	DTG 50 mg	1
GSK3640254 200 mg QD on Days 1 through 7 in Period 2	GSK3640254 200 mg	2
DTG 50 mg QD + GSK3640254 200 mg QD on Days 1 through 7 in Period 3	DTG 50 mg + GSK3640254 200 mg	3

5.2. Baseline Definitions

For all endpoints (except as noted in baseline definitions), the baseline value will be the latest pre-dose assessment with a non-missing value, including those from unscheduled visits. If time is not collected, Day 1 assessments are assumed to be taken prior to first dose and used as baseline. Baseline definitions in the tables below are applicable to each period.

Parameter	Study Assessments Considered as Baseline			Baseline Used in Data Display
	Screening	Day -1	Day 1 (Pre-Dose)	
Safety				
Vital Sign	X	X	X	Day 1 (Pre-Dose)
12-Lead ECG	X	X	X	Day 1 (Pre-Dose)[1]

[1] Baseline for 12-lead ECGs are applied to each period. The average (for quantitative assessments) or the worst case (for interpretation) of the predose triplicate assessments on Day 1 of Period 1 will be used as the baseline for Period 1. The average (for quantitative assessments) or the worst case (for interpretation) of the predose triplicate assessments on Day 1 of Period 2 will be used as the baseline for Period 2.

Parameter	Study Assessments Considered as Baseline			Baseline Used in Data Display
	Day -1	Period 1 Day 9	Period 2 Day 7	
Safety				
Hematology, Clinical Chemistry, Urinalysis	X	X	X	Day -1 for Period 1; Period 1 Day 9 for Period 2; Period 2 Day 7 for Period 3

Unless otherwise stated, if baseline data is missing no derivation will be performed and baseline will be set to missing.

5.3. Other Considerations for Data Analyses and Data Handling Conventions

Other considerations for data analyses and data handling conventions are outlined in the appendices:

Section	Component
10.2	Appendix 2: Study Phases and Treatment Emergent Adverse Events
10.3	Appendix 3: Data Display Standards & Handling Conventions
10.4	Appendix 4: Derived and Transformed Data
10.5	Appendix 5: Reporting Standards for Missing Data
10.6	Appendix 6: Division of AIDS (DAIDS) Table for Grading the Severity of Adult and Pediatric Adverse Events
10.7	Appendix 7: Values of Potential Importance

6. STUDY POPULATION ANALYSES

6.1. Overview of Planned Study Population Analyses

The study population analyses will be based on the "Safety" population, unless otherwise specified.

Study population analyses including analyses of participant's disposition, protocol deviations (including inclusion/exclusion criteria deviations), demographic and baseline characteristics, prior and concomitant medications, and exposure will be based on GSK Core Data Standards. Details of the planned displays are presented in Appendix 9: List of Data Displays.

7. PHARMACOKINETIC ANALYSES

7.1. Primary Pharmacokinetic Analyses

7.1.1. Endpoint / Variables

7.1.1.1. Drug Concentration Measures

Refer to Appendix 3: Data Display Standards & Handling Conventions (Section 10.3.3 Reporting Standards for Pharmacokinetics). Plasma concentrations of DTG and GSK3640254 will be measured and reported.

7.1.1.2. Derived Pharmacokinetic Parameters

Pharmacokinetic parameters will be calculated by standard non-compartmental analysis according to current working practices and using the currently supported version of WinNonlin (6.4 or higher). All calculations of non-compartmental parameters will be based on actual sampling times. Pharmacokinetic parameters listed will be determined from the plasma concentration-time data, as data permits.

Parameter	Parameter Description
Cmax	Maximum observed concentration, determined directly from the concentration-time data.
C_τ	Plasma concentration at the end of the dosing interval
AUC(0-τ)	Area under the plasma concentration-time curve from time 0 to the end of the dosing interval at steady state, to be calculated using the linear trapezoidal rule for each incremental trapezoid and the log trapezoidal rule for each decremental trapezoid.

NOTES:

- Additional parameters may be included as required.

7.1.2. Summary Measure

Area under concentration-time curve [AUC(0- τ)], $C\tau$, and Cmax at steady state following doses of DTG 50 mg QD Days 1 through 5 in Period 1, GSK3640254 200 mg QD Days 1 through 7 in Period 2, and DTG 50 mg QD coadministered with GSK3640254 200 mg QD on Days 1 through 7 in Period 3 in healthy participants.

7.1.3. Population of Interest

The primary PK analyses will be based on the PK concentration population for plasma PK concentrations and the PK parameter population for plasma PK parameters and statistical analysis.

7.1.4. Statistical Analyses / Methods

Details of the planned displays are provided in Appendix 9: List of Data Displays and will be based on GSK Data Standards and statistical principles.

Unless otherwise specified, endpoints / variables defined in Section 7.1.1 will be summarised using descriptive statistics, graphically presented (where appropriate), and listed.

7.1.4.1. Statistical Methodology Specification

The following PK statistical analyses will only be performed if sufficient data is available (i.e. if participants have well defined plasma profiles).

Endpoint / Variables
• Plasma primary PK endpoints include AUC(0-τ), Cτ, and Cmax, as data permit
Model Specification
• Analyses will be performed on the natural logarithms of AUC(0-τ), Cτ, and Cmax using linear mixed-effect models with period as a fixed effect, participants as random effect, and measurements within participant as repeated measures. • Effects will be estimated, and CIs will be constructed for the following treatment comparisons: Period 3 versus Period 1 (DTG) Period 3 versus Period 2 (GSK3640254) • Point estimates and 90% CIs for treatment differences on the log scale derived from the model will be exponentiated to obtain estimates for geometric mean ratios and CIs on the original scale.
Model Checking & Diagnostics
• Based on the data.
Model Results Presentation
• Statistical analysis by analysis of variance (ANOVA) will be presented in tabular format with geometric mean ratios for: Period 3 versus Period 1 (DTG) Period 3 versus Period 2 (GSK3640254)

7.2. Secondary Pharmacokinetic Analyses

7.2.1. Endpoint / Variables

7.2.1.1. Drug Concentration Measures

Refer to Appendix 3: Data Display Standards & Handling Conventions (Section 10.3.3 Reporting Standards for Pharmacokinetics)

7.2.1.2. Derived Pharmacokinetic Parameters

Pharmacokinetic parameters will be calculated by standard non-compartmental analysis according to current working practices and using the currently supported version of WinNonlin (6.4 or higher). All calculations of non-compartmental parameters will be based on actual sampling times.

Plasma PK parameters listed below will be determined from the total plasma concentration-time data, as data permits.

Parameter	Parameter Description
Tmax	Time to first occurrence of Cmax
t½	Terminal half-life will be calculated as: t½ = ln2 / λz
λz	Terminal-phase rate constant

NOTES:

- Additional parameters may be included as required.

7.2.2. Summary Measure

Tmax and t1/2 at steady state following doses of DTG 50 mg QD Days 1 through 5 in Period 1, GSK3640254 200 mg QD Days 1 through 7 in Period 2, and DTG 50 mg QD coadministered with GSK3640254 200 mg QD on Days 1 through 7 in Period 3 in healthy participants.

7.2.3. Population of Interest

The secondary PK analyses will be based on the PK concentration population for plasma PK concentrations, and the PK parameter population for plasma and statistical analysis, unless otherwise specified.

7.2.4. Statistical Analyses / Methods

Details of the planned displays are provided in Appendix 9: List of Data Displays and will be based on GSK Data Standards and statistical principles.

Unless otherwise specified, endpoints / variables defined in Section 7.2.1 will be summarised using descriptive statistics, graphically presented (where appropriate) and listed.

Secondary plasma PK parameters (Tmax and t1/2) will be estimated for DTG (Periods 1 and 3) and GSK3640254 (Periods 2 and 3). Summary statistics (arithmetic mean, geometric mean, median, standard deviation (SD), minimum, maximum, and coefficient of variation) for plasma GSK3640254 and DTG PK parameter values will be summarized by treatment.

Predose (trough) PK plasma concentrations (DTG: Days 2 through 5 of Period 1 and Days 2 through 7 of Period 3; GSK3640254: Days 4 through 7 of both Period 2 and Period 3) will be summarized using the PK Parameter Population and used to assess achievement of steady state.

8. SAFETY ANALYSES

The safety analyses will be based on the "Safety" population, unless otherwise specified.

8.1. Adverse Events Analyses

Adverse events analyses including the analysis of AEs, SAEs and other significant AEs will be based on GSK Core Data Standards. The details of the planned displays are provided in Appendix 9: List of Data Displays.

8.2. Clinical Laboratory Analyses

Laboratory evaluations including the analyses of chemistry laboratory tests, hematology laboratory tests, urinalysis, and liver function tests will be based on GSK Core Data Standards and will be graded using the Division of AIDS (DAIDS) Table for Grading the Severity of Adult and Pediatric Adverse Events (Version 2.1, July 2017). The details of the planned displays are in Appendix 9: List of Data Displays.

8.3. Other Safety Analyses

The analyses of non-laboratory safety test results including ECGs, vital signs, and Columbia Suicide Severity Rating Scale (C-SSRS) will be based on GSK Core Data Standards, unless otherwise specified. A figure of mean change from baseline in QTcF interval along with the 2-sided 95% CI using Student's t distribution will be presented by treatment and visit. The details of the planned displays are presented in Appendix 9: List of Data Displays.

9. REFERENCES

ViiV Healthcare group of companies Document Number 2018N383395_01 (12-DEC-2018): An Open-Label Two-way Interaction Clinical Trial to Evaluate the Pharmacokinetic Interactions Between GSK3640254 and Dolutegravir in Healthy Subjects

10. APPENDICES

10.1. Appendix 1: Schedule of Activities

10.1.1. Protocol Defined Schedule of Events

Screening Visit

Procedure	Screening (up to 28 days before Day 1)
Outpatient visit	X
Informed consent	X
Inclusion and exclusion criteria	X
Demography	X
Full physical examination including height and weight [1]	X
Laboratory assessments (hematology, chemistry, urinalysis)	X
12-lead electrocardiogram	X
Vital sign measurements	X
Medication/drug/alcohol history	X
Past and current medical conditions	X
Columbia Suicide Severity Rating Scale	X
Serum pregnancy test	X
Follicle-stimulating hormone (as needed, to confirm postmenopausal status)	X
Drug, alcohol, and cotinine screen	X
HIV, Hepatitis B and C screening	X

HIV = human immunodeficiency virus.

1 A full physical examination will include at a minimum, assessments of the skin, cardiovascular, respiratory, gastrointestinal, and neurological systems.

Time and Events Table

Procedure	Period 1		W	Period 2			Period 3							Follow-up				Notes
	Day –1	Day 1-5	Day 6-9	Day 1-5	Day 6	Day 7	Day 1	Day 2	Day 3	Day 4	Day 5	Day 6	Day 7	Day 8	Day 9	Day 10[1]	Day 11	
Admit to clinic	X																	
Discharge from clinic																	X	
Brief physical examination	X															X		A brief physical examination will include, at a minimum, assessments of the skin, lungs, cardiovascular system, and abdomen (liver and spleen).
Vital sign measurements	X	X		X	X	X	X	X	X	X	X	X	X	X	X	X	X	
12-Lead ECG	X	D1 and D5		D1 and D5			X			X	X		X			X		All ECGs in Period 1-3 will be pre-dose, post-dose at 2 hours, and post-dose at 4 hours. The predose ECGs on Day 1 of both Period 1 and Period 2 will be taken in triplicate.
Drug, alcohol, and cotinine screen	X																	See Protocol Appendix 2 for specific tests to be performed.
Laboratory assessments (hematology, chemistry, urinalysis)	X		D9			X				X			X			X		See Protocol Appendix 2 for specific tests to be performed.
Pregnancy test	X															X		
Columbia Suicide Severity Rating Scale				D1									X					
Genetic sample	X																	

Procedure	Period 1		W	Period 2			Period 3							Follow-up				Notes
	Day –1	Day 1-5	Day 6-9	Day 1-5	Day 6	Day 7	Day 1	Day 2	Day 3	Day 4	Day 5	Day 6	Day 7	Day 8	Day 9	Day 10[1]	Day 11	
Study intervention: Dolutegravir 50 mg		X					X	X	X	X	X	X	X					
Study intervention: GSK3640254 200 mg				X	X	X	X	X	X	X	X	X	X					
DTG trough PK sampling		D2-4						X	X	X	X	X						Blood collection for DTG trough PK samples will be collected on Days 2 through 4 in Period 1, Days 2 through 6 in Period 3.
DTG serial PK sampling		D5	D6-8										X	X	X	X		Blood collection for PK analysis of DTG will be collected predose, and after dosing at 1, 1.5, 2, 3, 3.5, 4, 4.5, 5, 6, 8, 12, 24, 48, and 72 hours relative to Day 5 dosing in Period 1, and relative to Day 7 dosing in Period 3.
GSK3640254 trough PK sampling				D 4-5	X					X	X	X						Blood collection for GSK3640254 trough PK samples will be collected on, Days 4 through 6 in Period 2; and Days 4 through 6 in Period 3.

Procedure	Period 1		W	Period 2			Period 3							Follow-up				Notes
	Day –1	Day 1-5	Day 6-9	Day 1-5	Day 6	Day 7	Day 1	Day 2	Day 3	Day 4	Day 5	Day 6	Day 7	Day 8	Day 9	Day 10[1]	Day 11	
GSK3640254 serial PK sampling						X	X						X	X	X	X	X	Relative to Day 7 dosing in Period 2, blood for PK analysis of GSK3640254 will be collected predose, and after dosing at 1, 2, 2.5, 3, 3.5, 4, 4.5, 5, 6, 8, 12, and 24 hours. Relative to Day 7 dosing in Period 3, the PK samples will be collected at predose, and after dosing at 1, 2, 2.5, 3, 3.5, 4, 4.5, 5, 6, 8, 12, 24, 48, 72, and 96 hours.
AE review		←==========→																
SAE review	←==========→																	
Concomitant medication review	←==========→																	

AE = adverse event; D = Day; DTG = dolutegravir; ECG = electrocardiogram; PK = pharmacokinetic; SAE = serious adverse event; W = washout.

1 Evaluations scheduled for Day 10 in Period 3 will also be performed for participants who discontinue early.

10.2. Appendix 2: Study Phases and Treatment Emergent Adverse Events

10.2.1. Study Phases

Assessments and events will be classified according to the time of occurrence relative to study treatment start date(/time) and stop date(/time).

Study Phase	Definition
Pre-Treatment	Date and Time ≤ Study Treatment Start Date and Time
On-Treatment	Study Treatment Start Date and Time < Date and Time ≤ Study Treatment Stop Date and Time + 5 days
Post-Treatment	Date and Time > Study Treatment Stop Date and Time + 5 days

10.2.1.1. Study Phases for Concomitant Medication

Study Phase	Definition
Prior	If medication end date is not missing and is before Day -1 of Period 1
Concomitant	Any medication that is not a prior

NOTES:

- Please refer to Appendix 5: Reporting Standards for Missing Data for handling of missing and partial dates for concomitant medication. Use the rules in this table if concomitant medication date is completely missing.

10.2.2. Treatment Emergent Flag for Adverse Events

Flag	Definition
Treatment Emergent	• If AE onset date and time is on or after treatment start date and time & on or before treatment stop date and time + 5 days. • Study Treatment Start Date and Time ≤ AE Start Date and Time ≤ Study Treatment Stop Date and Time + 5 days. • If the AE onset date is completely missing, the AE is considered as treatment emergent.

NOTES:

- If the study treatment stop date is missing, then the AE will be considered to be On-Treatment.
- Please refer to Appendix 5: Reporting Standards for Missing Data for handling of missing and partial dates for adverse events. Use the rules in this table if the adverse event onset date is completely missing.

10.3. Appendix 3: Data Display Standards & Handling Conventions

10.3.1. Reporting Process

Software	
• The currently supported versions of SAS software (9.4 or higher) will be used.	
Reporting Area	
HARP Server	\\us1salx00259.corpnet2.com
HARP Compound	\gsk3640254\mid209712\final_01
Analysis Datasets	
• Analysis datasets will be created according to CDISC standards (SDTM IG Version 3.2 & ADaM IG Version 1.0). • For creation of ADaM datasets (ADC1/ADCM/ADAE), the same version of dictionary datasets will be implemented for conversion from SI to SDTM.	
Generation of RTF Files	
• RTF files will be generated for all reporting efforts described in the RAP.	

10.3.2. Reporting Standards

General

• The current GSK Integrated Data Standards Library (IDSL) will be applied for reporting, unless otherwise stated (IDSL Standards Location: https://spope.gsk.com/sites/IDSLLibrary/SitePages/Home.aspx):
  • 4.03 to 4.23: General Principles
  • 5.01 to 5.08: Principles Related to Data Listings
  • 6.01 to 6.11: Principles Related to Summary Tables
  • 7.01 to 7.13: Principles Related to Graphics
• Do not include participant level listings in the main body of the GSK Clinical Study Report. All participant level listings should be located in the modular appendices as ICH or non-ICH listings.

Formats
• All data will be reported according to the actual treatment the participant received unless otherwise stated. • GSK IDSL Statistical Principles (5.03 & 6.06.3) for decimal places (DP's) will be adopted for reporting of data based on the raw data collected, unless otherwise stated. • Numeric data will be reported at the precision collected on the eCRF. • The reported precision from non eCRF sources will follow the IDSL statistical principles but may be adjusted to a clinically interpretable number of DP's.
Planned and Actual Time

• Reporting for tables, figures, and formal statistical analyses:
  • Planned time relative to dosing will be used in figures, summaries, statistical analyses and calculation of any derived parameters, unless otherwise stated.
  • The impact of any major deviation from the planned assessment times and/or scheduled visit days on the analyses and interpretation of the results will be assessed as appropriate.
• Reporting for Data Listings:
  • Planned and actual time relative to study drug dosing will be shown in listings (Refer to IDSL Statistical Principle 5.05.1).
  • Unscheduled or unplanned readings will be presented within the participant's listings.

• Visits outside the protocol defined time-windows (i.e. recorded as protocol deviations) will be included in listings but omitted from figures (mean figures only for PK concentrations), summaries and statistical analyses (excluding statistical analyses of PK parameters).	
Unscheduled Visits	
• Unscheduled visits will not be included in summary tables except for determining the worst-case values. • Unscheduled visits will not be included in figures. • All unscheduled visits will be included in listings.	
Descriptive Summary Statistics	
Continuous Data	Refer to IDSL Statistical Principle 6.06.1
Categorical Data	N, n, frequency, %
Graphical Displays	
• Refer to IDSL Statistical Principals 7.01 to 7.13.	

10.3.3. Reporting Standards for Pharmacokinetics

Pharmacokinetic Concentration Data	

Descriptive Summary Statistics, Graphical Displays and Listings	Refer to IDSL PK Display Standards.
	Refer to IDSL Statistical Principle 6.06.1.
	For continuous data:
	• NQs at the beginning of a participant profile (i.e. before the first incidence of a measurable concentration) are deemed to be zero as it is assumed that in this circumstance no drug is yet measurable in the blood.
	• For NQs at the end of the participant profile (i.e. after the last incidence of a measurable concentration);
	  • for individual plots and pharmacokinetic analyses these are dropped (set to missing) as they do not provide any useful information (and can erroneously indicate that absolutely no drug is present)
	  • for summary statistics, these are set to 0 (to avoid skewing of the summary statistics)
	• Individual NQs which fall between two measurable concentrations are set to missing (individual values of this nature are assumed to be an anomaly)
	If two or more NQ values occur in succession between measurable concentrations, the profile will be deemed to have terminated at the last measurable concentration prior to these NQs. For the purpose of individual participant plots, these NQs will be set to 0, and the subsequent measurable concentrations will be retained. For the derivation of pharmacokinetic parameters, these NQs and any subsequent measurable concentrations will be omitted (set to missing).
	Note: Concentration values will be imputed as per GUI_51487 for descriptive summary statistics/analysis and summarized graphical displays only.

Pharmacokinetic Parameter Data	
Descriptive Summary Statistics, Graphical Displays and Listings	N, n, geometric mean, 95% CI of geometric mean, SD of logged data and between-subject geometric coefficient of variation (CVb (%)) will be reported. CV_b (%) = $\sqrt{}$ (exp(SD^2) - 1) * 100 (SD = SD of Ln-Transformed data)
Parameters Not Being Ln-Transformed	Tmax, λz, λz lower, λz upper, and λz no. of points.
Parameters Not Being Summarized	λz, λz lower, λz upper, and λz no. of points.
Listings	Include the first point, last point and number of points used in the determination of λz and Rsq_adjusted for listings.

10.4. Appendix 4: Derived and Transformed Data

10.4.1. General

Multiple Measurements at One Analysis Time Point
• Mean of the measurements will be calculated and used in any derivation of summary statistics but if listed, all data will be presented. • The worst finding/interpretation associated with multiple measurements as the finding/interpretation for that time point. • Participants having both High and Low values for Normal Ranges at any post-baseline visit for safety parameters will be counted in both the High and Low categories of "Any visit post-baseline" row of related summary tables. This will also be applicable to relevant Potential Clinical Importance summary tables.
Study Day

• Calculated as the number of days from First Dose Date in Period 1:
  • Ref Date = Missing → Study Day = Missing
  • Ref Date < First Dose Date in Period 1 → Study Day = Ref Date – First Dose Date
  • Ref Date >= First Dose Date in Period 1 → Study Day = Ref Date – (First Dose Date in Period 1) + 1

Period Day

• Calculated as the number of days from First Dose Date for the respective period:
  • Ref Date = Missing → Study Day = Missing
  • Ref Date < First Dose Date in Period 1 → Study Day = Ref Date – First Dose Date
  • First Dose Date in Period 1 <= Ref Date < First Dose Date in Period 2 → Study Day = Ref Date – (First Dose Date in Period 1) + 1
  • First Dose Date in Period 2 <= Ref Date < First Dose Date in Period 3 → Study Day = Ref Date – (First Dose Date in Period 2) + 1
  • Ref Date >= First Dose Date in Period 3 → Study Day = Ref Date – (First Dose Date in Period 3) + 1

10.4.2. Study Population

Age

• GSK standard IDSL algorithms will be used for calculating age where birth date will be imputed as follows:
  o Any participant with a missing day will have this imputed as day '15'.
  o Any participant with a missing day and month will have this imputed as '30th June'.
• Birth date will be presented in listings as 'YYYY'.

Body Mass Index (BMI)
• Calculated as Weight (kg) / [Height $(m)^2$]

10.4.3. Safety

Adverse Events
AEs of Special Interest
• No analysis for AEs of Special Interest will be performed

10.5. Appendix 5: Reporting Standards for Missing Data

10.5.1. Premature Withdrawals

Element	Reporting Detail
General	• Participant study completion (i.e. as specified in the protocol) was defined as the participant has completed all phases of the study including the final date on which data were or are expected to be collected. • Withdrawn participants were not replaced in the study. • All available data from participants who were withdrawn from the study will be listed and all available planned data will be included in summary tables and figures, unless otherwise specified.

10.5.2. Handling of Missing Data

Element	Reporting Detail
General	• Missing data occurs when any requested data is not provided, leading to blank fields on the collection instrument: o These data will be indicated by the use of a “blank” in participant listing displays. Unless all data for a specific visit are missing in which case the data is excluded from the table. o Answers such as “Not applicable” and “Not evaluable” are not considered to be missing data and should be displayed as such.
Outliers	• Any participants excluded from the summaries and/or statistical analyses will be documented along with the reason for exclusion in the clinical study report.

10.5.2.1. Handling of Missing and Partial Dates

Element	Reporting Detail
General	• Partial dates will be displayed as captured in participant listing displays.
Adverse Events	• The eCRF allows for the possibility of partial dates (i.e., only month and year) to be recorded for AE start and end dates; that is, the day of the month may be missing. In such a case, the following conventions will be applied for calculating the time to onset and the duration of the event: ○ Missing Start Day: First of the month will be used unless this is before the start date of study treatment; in this case the study treatment start date will be used and hence the event is considered On-treatment as per Appendix 4: Study Phases and Treatment Emergent Adverse Events. ○ Missing Stop Day: Last day of the month will be used, unless this is after the stop date of study treatment; in this case the study treatment stop date will be used. • Completely missing start or end dates will remain missing, with no imputation applied. Consequently, time to onset and duration of such events will be missing.
Concomitant Medications	• Partial dates for any concomitant medications recorded in the eCRF will be imputed using the following convention: ○ If the partial date is a start date, a '01' will be used for the day and 'Jan' will be used for the month ○ If the partial date is a stop date, a '28/29/30/31' will be used for the day (dependent on the month and year) and 'Dec' will be used for the month. • The recorded partial date will be displayed in listings.

10.6. Appendix 6: Division of AIDS (DAIDS) Table for Grading the Severity of Adult and Pediatric Adverse Events

10.6.1. Laboratory Values

Laboratory abnormalities will be graded according to the DAIDS grading table Version 2.1, July 2017. Laboratory results are converted to use SI units; only the numeric part of the criteria will be used. If for a laboratory parameter there are multiple grades sharing the same criteria, the maximum grade will be used.

HEMATOLOGY				
	Grade 1	**Grade 2**	**Grade 3**	**Grade 4**
Absolute Lymphocyte Count, Low (cell/mm^3; cells/L) > 5 years of age (not HIV infected)	600 to < 650 0.600 × 10^9 to < 0.650 × 10^9	500 to < 600 0.500 × 10^9 to < 0.600 × 10^9	350 to < 500 0.350 × 10^9 to < 0.500 × 10^9	< 350 < 0.350 × 10^9
Absolute Neutrophil Count, Low (cells/mm^3; cells/L) > 7 days of age	800 to 1,000 0.800 × 10^9 to 1.000 × 10^9	600 to 799 0.600 × 10^9 to 0.799 × 10^9	400 to 599 0.400 × 10^9 to 0.599 × 10^9	< 400 < 0.400 × 10^9
Hemoglobin, Low (g/dL; mmol/L) ≥ 13 years of age (male only)	10.0 to 10.9 6.19 to 6.76	9.0 to < 10.0 5.57 to < 6.19	7.0 to < 9.0 4.34 to < 5.57	< 7.0 <4.34
Hemoglobin, Low (g/dL; mmol/L) ≥ 13 years of age (female only)	9.5 to 10.4 5.88 to 6.48	8.5 to < 9.5 5.25 to < 5.88	6.5 to < 8.5 4.03 to < 5.25	< 6.5 < 4.03
Platelets, Decreased (cells/mm^3; cells/L)	100,000 to < 125,000 100.000 × 10^9 to < 125.000 × 10^9	50,000 to < 100,000 50.000 × 10^9 to < 100.000 × 10^9	25,000 to < 50,000 25.000 × 10^9 to < 50.000 × 10^9	< 25,000 < 25.000 × 10^9
White Blood Cell, Decreased (cells/mm^3; cells/L) > 7 days of age	2,000 to 2,499 2.000 × 10^9 to 2.499 × 10^9	1,500 to 1,999 1.500 × 10^9 to 1.999 × 10^9	1,000 to 1,499 1.000 × 10^9 to 1.499 × 10^9	< 1,000 < 1.000 × 10^9

Clinical Chemistry				
	Grade 1	**Grade 2**	**Grade 3**	**Grade 4**
Albumin, Low (g/dL; g/L)	3.0 to < LLN 30 to < LLN	≥ 2.0 to < 3.0 ≥ 20 to < 30	< 2.0 < 20	NA
Alkaline Phosphatase, High	1.25 to < 2.5 × ULN	2.5 to < 5.0 × ULN	5.0 to < 10.0 × ULN	≥ 10.0 × ULN
Alanine Aminotransferase, High	1.25 to < 2.5 × ULN	2.5 to < 5.0 × ULN	5.0 to < 10.0 × ULN	≥ 10.0 ULN
Amylase (Total), High	1.1 to < 1.5 × ULN	1.5 to < 3.0 × ULN	3.0 to < 5.0 × ULN	≥ 5.0 × ULN
Aspartate Aminotransferase, High	1.25 to < 2.5 × ULN	2.5 to < 5.0 × ULN	5.0 to < 10.0 × ULN	≥ 10.0 × ULN
Bicarbonate, Low (mEq/L; mmol/L)	16.0 to < LLN 16.0 to < LLN	11.0 to < 16.0 11.0 to < 16.0	8.0 to < 11.0 8.0 to < 110.	< 8.0 < 8.0
Direct Bilirubin, High > 28 days of age	NA	NA	> ULN with other signs and symptoms of hepatotoxicity	> ULN with life-threatening consequences (e.g., signs and symptoms of liver failure)
Total Bilirubin, High > 28 days of age	1.1 to < 1.6 × ULN	1.6 to < 2.6 × ULN	2.6 to < 5.0 × ULN	≥ 5.0 × ULN
Calcium, High (mg/dL; mmol/L) ≥ 7 days of age	10.6 to < 11.5 2.65 to < 2.88	11.5 to < 12.5 2.88 to < 3.13	12.5 to < 13.5 3.13 to < 3.38	≥ 13.5 ≥ 3.38
Calcium, Low (mg/dL; mmol/L) ≥ 7 days of age	7.8 to < 8.4 1.95 to < 2.10	7.0 to < 7.8 1.75 to < 1.95	6.1 to < 7.0 1.53 to < 1.75	< 6.1 < 1.53
Creatine Kinase, High	3 to < 6 × ULN	6 to < 10 × ULN	10 to < 20 × ULN	≥ 20 × ULN
Creatinine, High *Choose the method that selects for the higher grade*	1.1 to 1.3 × ULN	> 1.3 to 1.8 × ULN OR Increase to 1.3 to < 1.5 × participant's baseline	> 1.8 to < 3.5 ULN OR Increase to 1.5 to < 2.0 × participant's baseline	≥ 3.5 × ULN OR Increase of ≥ 2.0 × participant's baseline
Glucose Fasting, High (mg/dL; mmol/L)	110 to 125 6.11 to < 6.95	> 125 to 250 6.95 to < 13.89	> 250 to 500 13.89 to < 27.75	≥ 500 ≥ 27.75
Glucose, Low (mg/dL; mmol/L) ≥ 1 month of age	55 to 64 3.05 to < 3.55	40 to < 55 2.22 to < 3.05	30 to < 40 1.67 to < 2.22	< 30 < 1.67
Lipase, High	1.1 to < 1.5 × ULN	1.5 to < 3.0 × ULN	3.0 to < 5.0 × ULN	≥ 5.0 × ULN
Cholesterol, Fasting, High (mg/dL; mmol/L) ≥ 18 years of age	200 to < 240 5.18 to < 6.19	240 to < 300 6.19 to < 7.77	≥ 300 ≥ 7.77	NA
Triglycerides, Fasting, High (mg/dL; mmol/L)	150 to 300 1.71 to 3.42	> 300 to 500 > 3.42 to 5.7	> 500 to < 1.000 > 5.7 to 11.4	> 1,000 > 11.4
Phosphate, Low (mg/dL; mmol/L) > 14 years of age	2.0 to < LLN 0.65 to < LLN	1.4 to < 2.0 0.45 to < 0.65	1.0 to < 1.4 0.32 to < 0.45	< 1.0 < 0.32

Clinical Chemistry				
	Grade 1	**Grade 2**	**Grade 3**	**Grade 4**
Potassium, High (mEq/L; mmol/L)	5.6 to < 6.0 5.6 to < 6.0	6.0 to < 6.5 6.0 to < 6.5	6.5 to < 7.0 6.5 to < 7.0	≥ 7.0 ≥ 7.0
Potassium, Low (mEq/L; mmol/L)	3.0 to < 3.4 3.0 to < 3.4	2.5 to < 3.0 2.5 to < 3.0	2.0 to < 2.5 2.0 to < 2.5	< 2.0 < 2.0
Sodium, High (mEq/L; mmol/L)	146 to < 150 146 to < 150	150 to < 154 150 to < 154	154 to < 160 154 to < 160	≥ 160 ≥ 160
Sodium, Low (mEq/L; mmol/L)	130 to < 135 130 to < 135	125 to < 130 125 to < 130	121 to < 125 121 to < 125	≤ 120 ≤ 120
Uric Acid, High (mEq/L; mmol/L)	7.5 to < 10.0 0.45 to < 0.59	10.0 to < 12.0 0.59 to < 0.71	12.0 to < 15.0 0.71 to < 0.89	≥ 15.0 ≥ 0.89

NA=not applicable; LLN = lower limit of normal; ULN=upper limit of normal.

Urinalysis				
	Grade 1	**Grade 2**	**Grade 3**	**Grade 4**
Glucose/Glycosuria (random collection tested by dipstick)	Trace to 1+ or ≤ 250 mg	2+ or > 250 to ≤ 500 mg	> 2+ or > 500 mg	NA
Protein/Proteinuria (random collection tested by dipstick)	1+	2+	3+ or higher	NA
Red Blood Cells (RBCs)/Hematuria (not to be reported based on dipstick findings or on blood believed to be of menstrual origin)	6 to < 10 RBCs per high power field	≥ 10 RBCs per high power field	Gross, with or without clots OR with RBC casts OR intervention indicated	Life-threatening consequences

NA=not applicable

10.7. Appendix 7: Values of Potential Clinical Importance

10.7.1. ECG

ECG Parameter	Units	Potential Clinically Important Range	
		Lower	Upper
Absolute			
Absolute QTc Interval	msec	< 320[1]	> 450[1]
		> 450[2]	≤ 480[2]
		> 480[2]	≤ 500[2]
		> 500[2]	
Absolute PR Interval	msec	< 120[1]	> 200[1]
Absolute QRS Interval	msec	< 60[1]	> 120[1]
Change from Baseline			
Increase from Baseline QTc	msec	≤ 30[2]	
	msec	> 30[1]	≤ 60[2]
	msec	> 60[2]	

NOTES:

1. Represent standard ECG values of potential clinical importance for HV studies.
2. Represent further subdivisions of ECG values for analysis.

10.7.2. Vital Signs

Vital Sign Parameter (Absolute)	Units	Potential Clinically Important Range	
		Lower	Upper
Systolic Blood Pressure	mmHg	< 85	> 140
Diastolic Blood Pressure	mmHg	< 45	> 90
Heart Rate	bpm	< 40	> 100

10.8. Appendix 8: Abbreviations & Trade Marks

10.8.1. Abbreviations

Abbreviation	Description
ADaM	Analysis Data Model
AE	Adverse Event
AUC	Area under the Plasma Concentration-Time Curve
AUC(0-τ)	AUC from Time 0 to the End of the Dosing Interval at Steady State
CDISC	Clinical Data Interchange Standards Consortium
CI	Confidence Interval
Cmax	Maximum Observed Concentration
C-SSRS	Columbia Suicide Severity Rating Scale
Cτ	Plasma Concentration at the End of the Dosing Interval
CV_b	Coefficient of Variation (Between)
DBF	Database Freeze
DBR	Database Release
DP	Decimal Places
DTG	Dolutegravir
ECG	Electrocardiogram
eCRF	Electronic Case Record Form
GSK	GlaxoSmithKline
HIV	human immunodeficiency virus
ICH	International Conference on Harmonization
IDSL	Integrated Data Standards Library
LLN	Lower Limit of Normal
PK	Pharmacokinetic
QD	Once Daily
RAP	Reporting & Analysis Plan
SAC	Statistical Analysis Complete
SAE	Serious Adverse Event
SD	Standard Deviation
SDTM	Study Data Tabulation Model
Tmax	Time of Maximum Observed Concentration
t1/2	Apparent Terminal Phase Half-Life
ULN	Upper Limit of Normal

10.8.2. Trademarks

Trademarks of the GlaxoSmithKline Group of Companies
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10.9. Appendix 9: List of Data Displays

10.9.1. Data Display Numbering

The following numbering will be applied for RAP generated displays:

Section	Tables	Figures
Study Population	1.1 to 1.7	
Safety	2.1 to 2.19	2.1
Pharmacokinetic	3.1 to 3.10	3.1 to 3.10
Section	**Listings**	
ICH Listings	1 to 30	
Other Listings	31 to 34	

10.9.2. Mock Example Shell Referencing

Non-IDSL specifications will be referenced as indicated and if required example mock-up displays provided in the Table/Listing/Figure Shells.

Section	Figure	Table	Listing
Study Population	POP_Fn	POP_Tn	POP_Ln
Safety	SAFE_Fn	SAFE_Tn	SAFE_Ln
Pharmacokinetic	PK_Fn	PK_Tn	PK_Ln

NOTES:

- Non-Standard displays are indicated in the 'IDSL / Example Shell' or 'Programming Notes' column as '[Non-Standard] + Reference.'

10.9.3. Deliverables

Delivery	Description
SAC	Final Statistical Analysis Complete

10.9.4. Study Population Tables

Study Population Tables					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
Subject Disposition					
1.1.	Safety	NS1	Summary of Number of Subjects Enrolled by Country and Site ID		SAC
1.2.	Safety	ES1A	Summary of Subject Disposition for the Subject Conclusion Record		SAC
1.3.	Screened	SD1	Summary of Screening Status and Reasons for Screen Failures		SAC
Protocol Deviation					
1.4.	Safety	DV1	Summary of Important Protocol Deviations		SAC
Demographic and Baseline Characteristics					
1.5.	Safety	DM3	Summary of Demographic Characteristics		SAC
1.6.	Safety	DM5	Summary of Race and Racial Combinations		SAC
1.7.	Safety	DM11	Summary of Age Ranges		SAC

10.9.5. Safety Tables

Safety: Tables					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
Adverse Events (AEs)					
2.1.	Safety	AE1CP	Summary of Adverse Events by System Organ Class and Preferred Term		SAC
2.2.	Safety	AE1CP	Summary of Drug-Related Adverse Events by System Organ Class and Preferred Term		SAC
2.3.	Safety	AE3	Summary of Common (>=5%) Adverse Events by Overall Frequency		SAC
2.4.	Safety	AE15	Summary of Common (>=5%) Non-serious Adverse Events by System Organ Class and Preferred Term (Number of Subjects and Occurrences)		SAC
2.5.	Safety	AE16	Summary of Serious Adverse Events by System Organ Class and Preferred Term (Number of Subjects and Occurrences)		SAC
2.6.	Safety	AE5A	Summary of Adverse Events by System Organ Class and Preferred Term and Maximum Intensity		SAC
Laboratory: Chemistry					
2.7.	Safety	LB1	Summary of Clinical Chemistry Changes from Baseline		SAC
2.8.	Safety	LB16	Summary of Clinical Chemistry Results by Maximum Grade Increase Post-Baseline Relative to Baseline		SAC
Laboratory: Hematology					
2.9.	Safety	LB1	Summary of Hematology Changes from Baseline		SAC
2.10.	Safety	LB16	Summary of Hematology Results by Maximum Grade Increase Post-Baseline Relative to Baseline		SAC

Safety: Tables					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
Laboratory: Urinalysis					
2.11.	Safety	UR3	Summary of Urinalysis Dipstick Results		SAC
2.12.	Safety	LB1	Summary of Urine Concentration Changes from Baseline		SAC
2.13.	Safety	LB16	Summary of Urinalysis by Maximum Grade Increase Post-Baseline Relative to Baseline		SAC
ECG					
2.14.	Safety	EG1	Summary of ECG Findings		SAC
2.15.	Safety	EG2	Summary of ECG Changes from Baseline		SAC
2.16.	Safety	EG10	Summary of Maximum QTc Values Post-Baseline Relative to Baseline by Category		SAC
2.17.	Safety	EG11	Summary of Maximum Increase in QTc Values Post-Baseline Relative to Baseline by Category		SAC
Vital Signs					
2.18.	Safety	VS1	Summary of Vital Sign Changes from Baseline		SAC
C-SSRS					
2.19.	Safety	CSSRS4	Listing of C-SSRS Suicidal Ideation and Behavior Data		SAC

10.9.6. Safety Figures

Safety: Tables					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
ECG					
2.1.	Safety	EG9	Mean (95% CI) Change from Baseline in QTcF Interval by Timepoint and Treatment		SAC

10.9.7. Pharmacokinetic Tables

Pharmacokinetic: Tables					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
PK Concentration Data					
3.1.	PK Concentration	PKCT1	Summary of DTG Plasma Pharmacokinetic Concentration-Time Data by Treatment		SAC
3.2.	PK Concentration	PKCT1	Summary of GSK3640254 Plasma Pharmacokinetic Concentration-Time Data by Treatment		SAC
3.3.	PK Concentration	PKCT1	Summary of Predose (trough) DTG Plasma Concentrations by Treatment		SAC
3.4.	PK Concentration	PKCT1	Summary of Predose (trough) GSK3640254 Plasma Concentrations by Treatment		SAC
PK Derived Parameters					
3.5.	PK Parameter	PKPT4	Summary of Derived DTG Plasma Pharmacokinetic Parameters (Non-Transformed) Based on Actual Time by Treatment	Parameters with units	SAC

Pharmacokinetic: Tables					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
3.6.	PK Parameter	PKPT4	Summary of Derived DTG Plasma Pharmacokinetic Parameters (Ln-Transformed) Based on Actual Time by Treatment	Parameters with units	SAC
3.7.	PK Parameter	PKPT4	Summary of Derived GSK3640254 Plasma Pharmacokinetic Parameters (Non-Transformed) Based on Actual Time by Treatment	Parameters with units - use ug instead of ng where applicable for GSK3640254	SAC
3.8.	PK Parameter	PKPT4	Summary of Derived GSK3640254 Plasma Pharmacokinetic Parameters (Ln-Transformed) Based on Actual Time by Treatment	Parameters with units - use ug instead of ng where applicable for GSK3640254	SAC
PK Analysis Tables					
3.9.	PK Parameter	PKPT3	Statistical Analysis of DTG Plasma Pharmacokinetic Parameters	AUC(0-τ), Cτ, and Cmax	SAC
3.10.	PK Parameter	PKPT3	Statistical Analysis of GSK3640254 Plasma Pharmacokinetic Parameters	AUC(0-τ), Cτ, and Cmax	SAC

10.9.8. Pharmacokinetic Figures

Pharmacokinetic: Figures					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable [Priority]**
Individual Concentration Plots					
3.1.	PK Concentration	PKCF1	Individual DTG Plasma Concentration-Time Plots by Participant (Linear and Semi-Logarithmic)	Paginate by Participant Dashed line represents the LLQ Treatments Overlaid	SAC
3.2.	PK Concentration	PKCF1	Individual GSK3640254 Plasma Concentration-Time Plots by Participant (Linear and Semi-Logarithmic)	Paginate by Participant Dashed line represents the LLQ Treatments Overlaid	SAC
3.3.	PK Concentration	PKCF1	Individual DTG Plasma Concentration-Time Plots by Treatment (Linear and Semi-Logarithmic)	Paginate by Treatment Dashed line represents the LLQ Participants Overlaid	SAC
3.4.	PK Concentration	PKCF1	Individual GSK3640254 Plasma Concentration-Time Plots by Treatment (Linear and Semi-Logarithmic)	Paginate by Treatment Dashed line represents the LLQ Participants Overlaid	SAC
Mean / Median Concentration Plots					
3.5.	PK Concentration	PKCF2	Mean (± Standard Deviation) DTG Plasma Concentration-Time Plots by Treatment (Linear and Semi-Logarithmic)	Treatments Overlaid	SAC
3.6.	PK Concentration	PKCF2	Mean (± Standard Deviation) GSK3640254 Plasma Concentration-Time Plots by Treatment (Linear and Semi-Logarithmic)	Treatments Overlaid	SAC
3.7.	PK Concentration	PKCF3	Median (Range) DTG Plasma Concentration-Time Plots by Treatment (Linear and Semi-Logarithmic)	Treatments Overlaid	SAC

Pharmacokinetic: Figures					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable [Priority]**
3.8.	PK Concentration	PKCF3	Median (Range) GSK3640254 Plasma Concentration-Time Plots by Treatment (Linear and Semi-Logarithmic)	Treatments Overlaid	SAC
3.9.	PK Concentration	PKCF2	Mean (± Standard Deviation) Predose (Trough) DTG Plasma Concentration Plots by Treatment (Linear and Semi-Logarithmic)		SAC
3.10.	PK Concentration	PKCF2	Mean (± Standard Deviation) Predose (Trough) GSK3640254 Plasma Concentration-Time Plots by Treatment (Linear and Semi-Logarithmic)		SAC

10.9.9. ICH Listings

ICH: Listings					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
Subject Disposition					
1.	Safety	ES3	Listing of Reasons for Study Withdrawal		SAC
2.	Screened	DS7	Listing of Reasons for Screen Failure		SAC
Protocol Deviations					
3.	Safety	DV2	Listing of Important Protocol Deviations		SAC
4.	Safety	IE3	Listing of Subjects with Inclusion/Exclusion Criteria Deviations		SAC
Populations Analysed					
5.	Safety	SP3A	Listing of Subjects Excluded from Any Population		SAC
Demographic and Baseline Characteristics					
6.	Safety	DM2	Listing of Demographic Characteristics		SAC
7.	Safety	DM9	Listing of Race		SAC
Prior and Concomitant Medications					
8.	Safety	CM5	Listing of Concomitant Medications	Based on GSK Drug Dictionary	SAC
Exposure and Treatment Compliance					
9.	Safety	EX4	Listing of Exposure Data		SAC
10.	Safety	POP_L1	Listing of Meal Data		SAC
Adverse Events					
11.	Safety	AE2	Listing of Relationship Between System Organ Class and Verbatim Text		SAC
12.	Safety	AE7	Listing of Subject Numbers for Individual Adverse Events		SAC

ICH: Listings					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
13.	Safety	AE9CP	Listing of All Adverse Events		SAC
Serious and Other Significant Adverse Events					
14.	Safety	AE9CP	Listing of Study Drug Related Adverse Events		SAC
15.	Safety	AE9CP	Listing of Serious Adverse Events (Fatal & Non-Fatal)		SAC
16.	Safety	AE14	Listing of Reasons for Considering as a Serious Adverse Event		SAC
17.	Safety	AE9CP	Listing of Adverse Events Leading to Withdrawal from Study		SAC
Hepatobiliary (Liver)					
18.	Safety	MH2	Listing of Medical Conditions for Subjects with Liver Stopping Events		SAC
19.	Safety	SU2	Listing of Substance Use for Subjects with Liver Stopping Events		SAC
All Laboratory					
20.	Safety	LB5A	Listing of Clinical Chemistry with any Toxicities		SAC
21.	Safety	LB5A	Listing of All Clinical Chemistry Data for Subjects with any Toxicities		SAC
22.	Safety	LB5A	Listing of Hematology with any Toxicities		SAC
23.	Safety	LB5A	Listing of All Hematology Data for Subjects with any Toxicities		SAC
24.	Safety	LB5A	Listing of Urinalysis with any Toxicities		SAC
25.	Safety	LB5A	Listing of All Urinalysis Data for Subjects with any Toxicities		SAC
ECG					
26.	Safety	EG6	Listing of All ECG Findings		SAC
27.	Safety	EG6	Listing of All Abnormal ECG Findings		

ICH: Listings					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
28.	Safety	EG4	Listing of All ECG Values for Subjects with Any Value of Potential Clinical Importance		SAC
Vital Signs					
29.	Safety	VS5	Listing of All Vital Signs of Potential Clinical Importance		SAC
30.	Safety	VS5	Listing of All Vital Signs for Subjects with any Value of Potential Clinical Importance		SAC

10.9.10. Non-ICH Listings

Non-ICH: Listings					
No.	**Population**	**IDSL / Example Shell**	**Title**	**Programming Notes**	**Deliverable**
Pharmacokinetics					
31.	PK Concentration	PKCL1P	Listing of Dolutegravir Plasma Concentrations (units) by Treatment		SAC
32.	PK Concentration	PKCL1P	Listing of GSK3640254 Plasma Concentrations (units) by Treatment		SAC
33.	PK Parameter	PKPL1P	Listing of Dolutegravir Plasma Pharmacokinetic Parameters Based on Actual Time by Treatment		SAC
34.	PK Parameter	PKPL1P	Listing of GSK3640254 Plasma Pharmacokinetic Parameters Based on Actual Time by Treatment		SAC