



## Clinical trial results:

### A TACL Phase 1/2 Study of PO Ixazomib in Combination with Chemotherapy for Childhood Relapsed or Refractory Acute Lymphoblastic Leukemia and Lymphoblastic Lymphoma

#### Summary

EudraCT number	2019-001947-28
Trial protocol	Outside EU/EEA
Global end of trial date	

#### Results information

Result version number	v1
This version publication date	03 December 2021
First version publication date	03 December 2021

#### Trial information

##### Trial identification

Sponsor protocol code	T2017-002
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##### Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT03817320
WHO universal trial number (UTN)	-

Notes:

#### Sponsors

Sponsor organisation name	Takeda
Sponsor organisation address	95 Hayden Avenue, Lexington, MA, United States, 60015
Public contact	Study Director, Takeda, trialdisclosures@takeda.com
Scientific contact	Study Director, Takeda, trialdisclosures@takeda.com

Notes:

#### Paediatric regulatory details

Is trial part of an agreed paediatric investigation plan (PIP)	Yes
EMA paediatric investigation plan number(s)	EMA-001410-PIP02-17
Does article 45 of REGULATION (EC) No 1901/2006 apply to this trial?	No
Does article 46 of REGULATION (EC) No 1901/2006 apply to this trial?	No

Notes:

## Results analysis stage

Analysis stage	Interim
Date of interim/final analysis	18 May 2021
Is this the analysis of the primary completion data?	No

Global end of trial reached?	No
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Notes:

## General information about the trial

Main objective of the trial:

The main objective of this study was to determine the maximum tolerated dose (MTD) and/or recommended phase 2 dose (RP2D), define and describe toxicities, and characterized the pharmacokinetics of orally (PO) administered ixazomib in conjunction with block 1 re-induction chemotherapy in children with relapsed/refractory acute lymphoblastic leukaemia (ALL) or lymphoblastic lymphoma (LLy).

Protection of trial subjects:

All study participants were required to read and sign an Informed Consent Form.

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	12 February 2019
Long term follow-up planned	Yes
Long term follow-up rationale	Safety
Long term follow-up duration	2 Years
Independent data monitoring committee (IDMC) involvement?	Yes

Notes:

## Population of trial subjects

### Subjects enrolled per country

Country: Number of subjects enrolled	United States: 10
Worldwide total number of subjects	10
EEA total number of subjects	0

Notes:

### Subjects enrolled per age group

In utero	0
Preterm newborn - gestational age < 37 wk	0
Newborns (0-27 days)	0
Infants and toddlers (28 days-23 months)	0
Children (2-11 years)	6
Adolescents (12-17 years)	3
Adults (18-64 years)	1
From 65 to 84 years	0
85 years and over	0

## Subject disposition

### Recruitment

Recruitment details:

Participants took part in the study at 6 investigative sites in the United States from 12 February 2019 to 18 May 2021. The study is ongoing.

### Pre-assignment

Screening details:

Participants with childhood relapsed or refractory acute lymphoblastic leukemia and lymphoblastic lymphoma were enrolled to receive ixazomib 1.6 mg/m<sup>2</sup>/day or 2 mg/m<sup>2</sup>/day.

### Period 1

Period 1 title	Treatment Period: Day 1 to Week 4
Is this the baseline period?	Yes
Allocation method	Non-randomised - controlled
Blinding used	Not blinded

### Arms

Are arms mutually exclusive?	Yes
<b>Arm title</b>	Ixazomib 1.6 mg/m <sup>2</sup>

Arm description:

Ixazomib at dose level 1, 1.6 mg/m<sup>2</sup>/day, for participants 1 year of age or older (Strata A), and 0.05 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXL backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.

Arm type	Experimental
Investigational medicinal product name	Doxorubicin
Investigational medicinal product code	
Other name	Adriamycin
Pharmaceutical forms	Capsule
Routes of administration	Oral use

Dosage and administration details:

Doxorubicin solution for infusion.

Investigational medicinal product name	Ixazomib
Investigational medicinal product code	
Other name	MLN9708
Pharmaceutical forms	Capsule
Routes of administration	Oral use

Dosage and administration details:

Ixazomib capsules.

Investigational medicinal product name	Vincristine
Investigational medicinal product code	
Other name	Oncovin, VCR, LCR
Pharmaceutical forms	Solution for injection
Routes of administration	Intravenous use

Dosage and administration details:

Vincristine IV push

Investigational medicinal product name	Dexamethasone
Investigational medicinal product code	
Other name	Decadron, Hexadrol, Dexone, Dexameth
Pharmaceutical forms	Solution for injection

Routes of administration	Intravenous use, Oral use
Dosage and administration details: Dexamethasone PO or IV.	
Investigational medicinal product name	PEG-asparaginase
Investigational medicinal product code	
Other name	Oncaspar, Pegaspargase, Polyethylene Glycol Conjugated L-asparaginase-H
Pharmaceutical forms	Solution for injection/infusion
Routes of administration	Intramuscular and intravenous use
Dosage and administration details: PEG-asparaginase solution for injection/infusion.	
<b>Arm title</b>	Ixazomib 2 mg/m <sup>2</sup>
Arm description: Ixazomib at dose level 2, 2 mg/m <sup>2</sup> /day, for participants 1 year of age or older (Strata A), and 0.07 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXL backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.	
Arm type	Experimental
Investigational medicinal product name	Vincristine
Investigational medicinal product code	
Other name	Oncovin, VCR, LCR
Pharmaceutical forms	Solution for injection
Routes of administration	Intravenous use
Dosage and administration details: Vincristine IV push	
Investigational medicinal product name	Ixazomib
Investigational medicinal product code	
Other name	MLN9708
Pharmaceutical forms	Capsule
Routes of administration	Oral use
Dosage and administration details: Ixazomib capsules.	
Investigational medicinal product name	PEG-asparaginase
Investigational medicinal product code	
Other name	Oncaspar, Pegaspargase, Polyethylene Glycol Conjugated L-asparaginase-H
Pharmaceutical forms	Solution for injection/infusion
Routes of administration	Intramuscular and intravenous use
Dosage and administration details: PEG-asparaginase solution for injection/infusion.	
Investigational medicinal product name	Doxorubicin
Investigational medicinal product code	
Other name	Adriamycin
Pharmaceutical forms	Capsule
Routes of administration	Oral use
Dosage and administration details: Doxorubicin solution for infusion.	
Investigational medicinal product name	Dexamethasone
Investigational medicinal product code	
Other name	Decadron, Hexadrol, Dexone, Dexameth
Pharmaceutical forms	Solution for injection
Routes of administration	Intravenous use, Oral use

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**Dosage and administration details:**

Dexamethasone PO or IV.

<b>Number of subjects in period 1</b>	<b>Ixazomib 1.6 mg/m<sup>2</sup></b>	<b>Ixazomib 2 mg/m<sup>2</sup></b>
Started	4	6
Completed	4	6

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**Period 2**

Period 2 title	Follow-up Period: Week 5 to Week 104
Is this the baseline period?	No
Allocation method	Non-randomised - controlled
Blinding used	Not blinded

**Arms**

Are arms mutually exclusive?	Yes
<b>Arm title</b>	Ixazomib 1.6 mg/m <sup>2</sup>

## Arm description:

Ixazomib at dose level 1, 1.6 mg/m<sup>2</sup>/day, for participants 1 year of age or older (Strata A), and 0.05 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXL backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.

Arm type	Experimental
Investigational medicinal product name	Vincristine
Investigational medicinal product code	
Other name	Oncovin, VCR, LCR
Pharmaceutical forms	Solution for injection
Routes of administration	Intravenous use

## Dosage and administration details:

Vincristine solution for infusion.

Investigational medicinal product name	Ixazomib
Investigational medicinal product code	
Other name	MLN9708
Pharmaceutical forms	Capsule
Routes of administration	Oral use

## Dosage and administration details:

Ixazomib capsules.

Investigational medicinal product name	Dexamethasone
Investigational medicinal product code	
Other name	Decadron, Hexadrol, Dexone, Dexameth
Pharmaceutical forms	Solution for injection
Routes of administration	Intravenous use, Oral use
Dosage and administration details: Dexamethasone PO or IV.	
Investigational medicinal product name	Doxorubicin
Investigational medicinal product code	
Other name	Adriamycin
Pharmaceutical forms	Capsule
Routes of administration	Oral use
Dosage and administration details: Doxorubicin solution for infusion.	
Investigational medicinal product name	PEG-asparaginase
Investigational medicinal product code	
Other name	Oncaspar, Pegaspargase, Polyethylene Glycol Conjugated L-asparaginase-H
Pharmaceutical forms	Solution for injection/infusion
Routes of administration	Intramuscular and intravenous use
Dosage and administration details: PEG-asparaginase solution for injection/infusion.	
<b>Arm title</b>	Ixazomib 2 mg/m <sup>2</sup>
Arm description: Ixazomib at dose level 2, 2 mg/m <sup>2</sup> /day, for participants 1 year of age or older (Strata A), and 0.07 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXLD backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.	
Arm type	Experimental
Investigational medicinal product name	Ixazomib
Investigational medicinal product code	
Other name	MLN9708
Pharmaceutical forms	Capsule
Routes of administration	Oral use
Dosage and administration details: Ixazomib capsules.	
Investigational medicinal product name	Doxorubicin
Investigational medicinal product code	
Other name	Adriamycin
Pharmaceutical forms	Capsule
Routes of administration	Oral use
Dosage and administration details: Doxorubicin solution for infusion.	
Investigational medicinal product name	PEG-asparaginase
Investigational medicinal product code	
Other name	Oncaspar, Pegaspargase, Polyethylene Glycol Conjugated L-asparaginase-H
Pharmaceutical forms	Solution for injection/infusion
Routes of administration	Intramuscular and intravenous use
Dosage and administration details: PEG-asparaginase solution for injection/infusion.	

Investigational medicinal product name	Vincristine
Investigational medicinal product code	
Other name	Oncovin, VCR, LCR
Pharmaceutical forms	Solution for injection
Routes of administration	Intravenous use

Dosage and administration details:

Vincristine solution for infusion.

Investigational medicinal product name	Dexamethasone
Investigational medicinal product code	
Other name	Decadron, Hexadrol, Dexone, Dexameth
Pharmaceutical forms	Solution for injection
Routes of administration	Intravenous use, Oral use

Dosage and administration details:

Dexamethasone PO or IV.

<b>Number of subjects in period 2</b>	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>
Started	4	6
Completed	4	6

## Baseline characteristics

### Reporting groups

Reporting group title	Ixazomib 1.6 mg/m <sup>2</sup>
Reporting group description:	
Ixazomib at dose level 1, 1.6 mg/m <sup>2</sup> /day, for participants 1 year of age or older (Strata A), and 0.05 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXLD backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.	
Reporting group title	Ixazomib 2 mg/m <sup>2</sup>
Reporting group description:	
Ixazomib at dose level 2, 2 mg/m <sup>2</sup> /day, for participants 1 year of age or older (Strata A), and 0.07 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXLD backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.	

Reporting group values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>	Total
Number of subjects	4	6	10
Age categorical			
Units: Subjects			
Age Continuous			
Units: years			
median	8.7	12.1	
full range (min-max)	4.6 to 15.7	6.1 to 20.5	-
Sex: Female, Male			
Units: Subjects			
Female	3	3	6
Male	1	3	4
Race/Ethnicity, Customized			
Units: Subjects			
American Indian or Alaska Native	0	0	0
Asian	0	0	0
Native Hawaiian or Other Pacific Islander	0	0	0
Black or African American	0	1	1
White	2	5	7
More than one race	0	0	0
Unknown or Not Reported	2	0	2
Ethnicity (NIH/OMB)			
Units: Subjects			
Hispanic or Latino	3	5	8
Not Hispanic or Latino	0	1	1
Unknown or Not Reported	1	0	1
Region of Enrollment			
Units: Subjects			
United States	4	6	10



## End points

### End points reporting groups

Reporting group title	Ixazomib 1.6 mg/m <sup>2</sup>
Reporting group description: Ixazomib at dose level 1, 1.6 mg/m <sup>2</sup> /day, for participants 1 year of age or older (Strata A), and 0.05 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXLD backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.	
Reporting group title	Ixazomib 2 mg/m <sup>2</sup>
Reporting group description: Ixazomib at dose level 2, 2 mg/m <sup>2</sup> /day, for participants 1 year of age or older (Strata A), and 0.07 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXLD backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.	
Reporting group title	Ixazomib 1.6 mg/m <sup>2</sup>
Reporting group description: Ixazomib at dose level 1, 1.6 mg/m <sup>2</sup> /day, for participants 1 year of age or older (Strata A), and 0.05 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXLD backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.	
Reporting group title	Ixazomib 2 mg/m <sup>2</sup>
Reporting group description: Ixazomib at dose level 2, 2 mg/m <sup>2</sup> /day, for participants 1 year of age or older (Strata A), and 0.07 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXLD backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.	

### Primary: Number of Participants With Dose Limiting Toxicity (DLT) During Block (Cycle) 1 of Chemotherapy

End point title	Number of Participants With Dose Limiting Toxicity (DLT) During Block (Cycle) 1 of Chemotherapy <sup>[1]</sup>
End point description: DLT: defined as per National Cancer Institute Common Toxicity Criteria (NCI CTC) version 5.0 as follows: 1) Any Grade 4/3 non-hematologic toxicity that occurs after first dose of ixazomib and results in omission of subsequent block of chemotherapy or delay of beginning of subsequent block of chemotherapy for >7 days, with exception of fever or infection. 2) Hematologic toxicities: Failure to recover a peripheral absolute neutrophil count (ANC) $\geq 500/\mu\text{L}$ and platelet (PLT) $> 20,000/\mu\text{L}$ , PLT infusion independent, due to documented bone marrow hypoplasia (cellularity < 10-20%) within 49 days of beginning of systemic chemotherapy without evidence of active disease or infection by bone marrow aspiration. Participants from evaluable response set who were evaluable for safety. Evaluable response set: all participants enrolled and received all or part of protocol therapy, are under follow-up for a sufficient period to evaluate disease at end of one treatment cycle or meet definition of progressive	
End point type	Primary
End point timeframe: Up to Cycle 1 (28 days)	
Notes: [1] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point. Justification: Only descriptive statistics were reported for this endpoint.	

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	3	6		
Units: participants	0	0		

## Statistical analyses

No statistical analyses for this end point

### Primary: Number of Participants with Grade 3 or More Treatment Emergent Adverse Events (TEAEs) Graded Using Common Toxicity Criteria for Adverse Event (CTCAE) Criteria Version 5.0 During Block (Cycle) 1 of Chemotherapy

End point title	Number of Participants with Grade 3 or More Treatment Emergent Adverse Events (TEAEs) Graded Using Common Toxicity Criteria for Adverse Event (CTCAE) Criteria Version 5.0 During Block (Cycle) 1 of Chemotherapy <sup>[2]</sup>
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End point description:

TEAEs are defined as any AEs that occurred or worsened during the on-treatment period. TEAEs were graded using National Cancer Institute (NCI) CTCAE Version 5.0. SAEs are generally defined in this Phase 1 study as all Grade 3 and 4 events both unexpected and expected that are possibly, probably, or definitely related to Ixazomib or the chemotherapy backbone, excluding hematologic toxicities unless the event meets the criteria for a DLT. Participants from evaluable response set who were evaluable for safety.

End point type	Primary
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End point timeframe:

Up to 104 weeks

Notes:

[2] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	3	6		
Units: participants	3	6		

## Statistical analyses

No statistical analyses for this end point

### Primary: Tmax: Time to Reach the Maximum Plasma Concentration (Cmax) for Ixazomib at Day 1

End point title	Tmax: Time to Reach the Maximum Plasma Concentration (Cmax) for Ixazomib at Day 1 <sup>[3]</sup>
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End point description:

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

End point type	Primary
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End point timeframe:

Day 1 at multiple time points (up to 72 hours) post-dose for participants weighing <20 kg and ≥20 kg

Notes:

[3] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

<b>End point values</b>	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	2	4		
Units: hours (h)				
median (full range (min-max))	4.50 (1.00 to 8.00)	1.50 (1.00 to 4.00)		

## Statistical analyses

No statistical analyses for this end point

## Primary: Cmax: Maximum Observed Plasma Concentration for Ixazomib at Day 1

End point title	Cmax: Maximum Observed Plasma Concentration for Ixazomib at Day 1 <sup>[4]</sup>
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End point description:

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

End point type	Primary
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End point timeframe:

Day 1 at multiple time points (up to 72 hours) post-dose for participants weighing <20 kg and ≥20 kg

Notes:

[4] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

<b>End point values</b>	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	2	4		
Units: ng/mL				
geometric mean (geometric coefficient of variation)	10.0 (± 809.5)	28.1 (± 30.5)		

## Statistical analyses

No statistical analyses for this end point

**Primary: T1/2: Terminal Elimination Half-Life of Ixazomib at Day 1**

End point title	T1/2: Terminal Elimination Half-Life of Ixazomib at Day 1 <sup>[5]</sup>
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End point description:

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease. 99999 indicates geometric coefficient of variation was not estimable for one participant.

End point type	Primary
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End point timeframe:

Day 1 at multiple time points (up to 72 hours) post-dose for participants weighing <20 kg and ≥20 kg

Notes:

[5] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	1	4		
Units: hours (h)				
geometric mean (geometric coefficient of variation)	67.7 (± 99999)	36.4 (± 41.9)		

**Statistical analyses**

No statistical analyses for this end point

**Primary: Adjusted R<sup>2</sup>: Coefficient of Determination for the Terminal Disposition Phase Slope Adjusted for the Number of Data Points Used in the Analysis at Day 1**

End point title	Adjusted R <sup>2</sup> : Coefficient of Determination for the Terminal Disposition Phase Slope Adjusted for the Number of Data Points Used in the Analysis at Day 1 <sup>[6]</sup>
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End point description:

Adjusted Rsq describes the goodness of fit for the terminal phase slope (log concentration vs time, used to then calculate the half-life). Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease. 99999 indicates geometric coefficient of variation was not estimable for one participant.

End point type	Primary
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End point timeframe:

Day 1 at multiple time points (up to 72 hours) post-dose for participants weighing <20 kg and ≥20 kg

Notes:

[6] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	1	4		
Units: Unitless				
geometric mean (geometric coefficient of variation)	0.587 (± 99999)	0.495 (± 58.5)		

## Statistical analyses

No statistical analyses for this end point

### Primary: AUC%extrap,obs: Percent of Area Under the Plasma Concentration-time Curve from Dosing Time Infinity due to Extrapolation From the Last Observed Concentration for Ixazomib at Day 1

End point title	AUC%extrap,obs: Percent of Area Under the Plasma Concentration-time Curve from Dosing Time Infinity due to Extrapolation From the Last Observed Concentration for Ixazomib at Day 1 <sup>[7]</sup>
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End point description:

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease. 99999 indicates geometric coefficient of variation was not estimable for one participant.

End point type	Primary
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End point timeframe:

Day 1 at multiple time points (up to 72 hours) post-dose for participants weighing <20 kg and ≥20 kg

Notes:

[7] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	1	4		
Units: percentage of AUC				
geometric mean (geometric coefficient of variation)	39.7 (± 99999)	26.8 (± 45.7)		

## Statistical analyses

No statistical analyses for this end point

### Primary: AUC(0-72): Area Under the Plasma Concentration-Time Curve From Time 0 to 72 Hours Postdose for Ixazomib at Day 1

End point title	AUC(0-72): Area Under the Plasma Concentration-Time Curve From Time 0 to 72 Hours Postdose for Ixazomib at Day 1 <sup>[8]</sup>
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End point description:

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

End point type	Primary
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End point timeframe:

Day 1 at multiple time points (up to 72 hours) post-dose for participants weighing <20 kg and ≥20 kg

Notes:

[8] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	2	4		
Units: h*ng/mL				
geometric mean (geometric coefficient of variation)	178 (± 125.7)	331 (± 61.7)		

## Statistical analyses

No statistical analyses for this end point

### Primary: Tmax: Time to Reach Maximum Observed Plasma Concentration for Ixazomib at Day 11

End point title	Tmax: Time to Reach Maximum Observed Plasma Concentration for Ixazomib at Day 11 <sup>[9]</sup>
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End point description:

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

End point type	Primary
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End point timeframe:

Day 11 pre-dose and at multiple time points post-dose (up to 72 hours for participants weighing <20 kg and up to 264 hours for participants weighing ≥20 kg)

Notes:

[9] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	3	6		
Units: ng/mL				
median (full range (min-max))	4.00 (0.50 to 4.00)	3.00 (0.50 to 72.00)		

## Statistical analyses

No statistical analyses for this end point

### Primary: Cmax: Maximum Plasma Concentration (Cmax) for Ixazomib at Day 11

End point title	Cmax: Maximum Plasma Concentration (Cmax) for Ixazomib at Day 11 <sup>[10]</sup>
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End point description:

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

End point type	Primary
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End point timeframe:

Day 11 pre-dose and at multiple time points post-dose (up to 72 hours for participants weighing <20 kg and up to 264 hours for participants weighing ≥20 kg)

Notes:

[10] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	3	6		
Units: hours (hr)				
geometric mean (geometric coefficient of variation)	22.8 (± 169.1)	73.6 (± 75.1)		

## Statistical analyses

No statistical analyses for this end point

### Primary: T1/2: Terminal Elimination Half-Life of Ixazomib at Day 11

End point title	T1/2: Terminal Elimination Half-Life of Ixazomib at Day 11 <sup>[11]</sup>
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End point description:

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

End point type	Primary
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End point timeframe:

Day 11 pre-dose and at multiple time points post-dose (up to 72 hours for participants weighing <20 kg and up to 264 hours for participants weighing ≥20 kg)

Notes:

[11] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	2	5		
Units: hours (h)				
geometric mean (geometric coefficient of variation)	111 (± 59.8)	99.4 (± 37.4)		

## Statistical analyses

No statistical analyses for this end point

### Primary: Adjusted R<sup>2</sup>: Coefficient of Determination for the Terminal Disposition Phase Slope Adjusted for the Number of Data Points Used in the Analysis at Day 11

End point title	Adjusted R <sup>2</sup> : Coefficient of Determination for the Terminal Disposition Phase Slope Adjusted for the Number of Data Points Used in the Analysis at Day 11 <sup>[12]</sup>
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End point description:

Adjusted Rsq describes the goodness of fit for the terminal phase slope (log concentration vs time, used to then calculate the half-life). Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

End point type	Primary
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End point timeframe:

Day 11 pre-dose and at multiple time points post-dose (up to 72 hours for participants weighing <20 kg and up to 264 hours for participants weighing ≥20 kg)

Notes:

[12] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	2	5		
Units: Unitless				
geometric mean (geometric coefficient of variation)	0.969 (± 0.5)	0.940 (± 3.8)		

## Statistical analyses

No statistical analyses for this end point

### Primary: AUC%extrap,obs: Percent of Area Under the Plasma Concentration-time Curve from Dosing Time Infinity due to Extrapolation From the Last Observed Concentration for Ixazomib at Day 11

End point title	AUC%extrap,obs: Percent of Area Under the Plasma Concentration-time Curve from Dosing Time Infinity due to Extrapolation From the Last Observed Concentration for Ixazomib at Day 11 <sup>[13]</sup>
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**End point description:**

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

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End point type	Primary
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**End point timeframe:**

Day 11 pre-dose and at multiple time points post-dose (up to 72 hours for participants weighing <20 kg and up to 264 hours for participants weighing ≥20 kg)

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**Notes:**

[13] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	2	5		
Units: percentage of AUC				
geometric mean (geometric coefficient of variation)	14.6 (± 130.5)	12.8 (± 105.6)		

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**Statistical analyses**

No statistical analyses for this end point

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**Primary: AUC(0-72): Area Under the Plasma Concentration-Time Curve From Time 0 to 72 Hours Postdose for Ixazomib at Day 11**

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End point title	AUC(0-72): Area Under the Plasma Concentration-Time Curve From Time 0 to 72 Hours Postdose for Ixazomib at Day 11 <sup>[14]</sup>
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**End point description:**

Evaluable response set included all participants enrolled and who received all or part of protocol therapy and are under follow-up for a sufficient period to evaluate the disease at the end of the one treatment cycle or meet the definition of progressive disease.

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End point type	Primary
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**End point timeframe:**

Day 11 pre-dose and at multiple time points (up to 72 hours) post-dose for participants weighing <20 kg and ≥20 kg

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**Notes:**

[14] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	3	6		
Units: h*ng/mL				
geometric mean (geometric coefficient of variation)	586 (± 58.2)	1650 (± 35.9)		

## Statistical analyses

No statistical analyses for this end point

### Primary: Percentage of Participants With Treatment Emergent Adverse Events (TEAEs), and Serious Adverse Events (SAEs)

End point title	Percentage of Participants With Treatment Emergent Adverse Events (TEAEs), and Serious Adverse Events (SAEs) <sup>[15]</sup>
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End point description:

AE: any untoward medical occurrence in a clinical investigation participant administered a investigational drug; it does not necessarily have to have a causal relationship with trial drug administration. SAE: any untoward medical occurrence that: 1) results in death, 2) is life-threatening, 3) requires inpatient hospitalization or prolongation of existing hospitalization, 4) results in persistent or significant disability/incapacity, 5) leads to a congenital anomaly/birth defect in offspring of participant or 6) is a medically important event that satisfies any of the following: a) May require intervention to prevent items 1 to 5 above. b) May expose participant to danger, even though event is not immediately life threatening or fatal or does not result in hospitalization. Participants from evaluable response set who were evaluable for safety.

End point type	Primary
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End point timeframe:

Up to 104 weeks

Notes:

[15] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	3	6		
Units: percentage of participants				
number (not applicable)				
TEAEs	100	100		
SAEs	100	100		

## Statistical analyses

No statistical analyses for this end point

### Primary: Phase 2: Response (CR and CR+CRp) After Block 1 Chemotherapy

End point title	Phase 2: Response (CR and CR+CRp) After Block 1 Chemotherapy <sup>[16]</sup>
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End point description:

Best response: measured by bone marrow status as percentage of participants with CR: a bone marrow with <5% blasts by morphology; no evidence of circulating blasts or extramedullary disease; recovery of

peripheral counts(ANC $\geq$ 500/ $\mu$ L, PLT $\geq$ 20,000/ $\mu$ L,platelet infusion independent) or CR MRD-:a bone marrow with<5% blasts by morphology;MRD<0.1%by flow or molecular testing(e.g.PCR);no evidence of circulating blasts or extramedullary disease;recovery of peripheral counts(ANC $\geq$ 500/ $\mu$ L,PLT $\geq$ 20,000/ $\mu$ L,platelet infusion independent) or CRi:all CR criteria except for insufficient recovery of ANC (<500/ $\mu$ L), and/or PLT counts (<20,000/ $\mu$ L) are reported. Evaluable response set: all participants enrolled and received all/part of protocol therapy and are under follow-up for a sufficient period to evaluate disease at end of 1 treatment cycle/meet definition of progressive disease.

End point type	Primary
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End point timeframe:

Up to Cycle 1 (28 days)

Notes:

[16] - No statistical analyses have been specified for this primary end point. It is expected there is at least one statistical analysis for each primary end point.

Justification: Only descriptive statistics were reported for this endpoint.

End point values	Ixazomib 1.6 mg/m <sup>2</sup>	Ixazomib 2 mg/m <sup>2</sup>		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	0 <sup>[17]</sup>	0 <sup>[18]</sup>		
Units: percentage of participants				
number (not applicable)				

Notes:

[17] - Data is not available for this endpoint as the Phase 2 of the study is ongoing.

[18] - Data is not available for this endpoint as the Phase 2 of the study is ongoing.

## Statistical analyses

No statistical analyses for this end point

## Adverse events

### Adverse events information

Timeframe for reporting adverse events:

Up to 104 weeks

Adverse event reporting additional description:

At each visit investigator had to document any occurrence of AE and abnormal laboratory findings. Any event reported by participant or by investigator was recorded, irrespective of the relation to study treatment. All-cause mortality: Evaluable response set(n=4,6). SAE and non-SAEs: Participants from evaluable response set with data for safety.

Assessment type	Systematic
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### Dictionary used

Dictionary name	MedDRA
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Dictionary version	Unknown
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### Reporting groups

Reporting group title	Ixazomib 2 mg/m <sup>2</sup>
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Reporting group description:

Ixazomib at dose level 2, 2 mg/m<sup>2</sup>/day, for participants 1 year of age or older (Strata A), and 0.07 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXL backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.

Reporting group title	Ixazomib 1.6 mg/m <sup>2</sup>
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Reporting group description:

Ixazomib at dose level 1, 1.6 mg/m<sup>2</sup>/day, for participants 1 year of age or older (Strata A), and 0.05 mg/kg/day, for infants less than 1 year of age (Strata B), orally, once on Days 1, 4, 8 and 11 of each 28-day cycle, given in combination with VXL backbone chemotherapy (vincristine, dexamethasone, asparaginase, and doxorubicin), up to 4 weeks. Participants were followed for safety up to 100 weeks after last dose.

Serious adverse events	Ixazomib 2 mg/m <sup>2</sup>	Ixazomib 1.6 mg/m <sup>2</sup>	
Total subjects affected by serious adverse events			
subjects affected / exposed	6 / 6 (100.00%)	3 / 3 (100.00%)	
number of deaths (all causes)	2	1	
number of deaths resulting from adverse events	0	0	
Nervous system disorders			
Syncope			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Blood and lymphatic system disorders			
Febrile neutropenia			
subjects affected / exposed	4 / 6 (66.67%)	3 / 3 (100.00%)	
occurrences causally related to treatment / all	5 / 6	3 / 3	
deaths causally related to treatment / all	0 / 0	0 / 0	

Generalized edema			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Immune system disorders			
Allergic reaction			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Gastrointestinal disorders			
Abdominal pain			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Nausea			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Respiratory, thoracic and mediastinal disorders			
Hypoxia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Mucositis oral			
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	2 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Infections and infestations			
Catheter related infection			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Sepsis			

subjects affected / exposed	1 / 6 (16.67%)	1 / 3 (33.33%)	
occurrences causally related to treatment / all	2 / 2	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Metabolism and nutrition disorders			
Hyperglycemia			
subjects affected / exposed	1 / 6 (16.67%)	1 / 3 (33.33%)	
occurrences causally related to treatment / all	1 / 1	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
ALT increased			
subjects affected / exposed	4 / 6 (66.67%)	1 / 3 (33.33%)	
occurrences causally related to treatment / all	5 / 5	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
AST increased			
subjects affected / exposed	3 / 6 (50.00%)	1 / 3 (33.33%)	
occurrences causally related to treatment / all	3 / 3	1 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypertriglyceridemia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypoalbuminemia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypokalemia			
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	2 / 2	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Hypophosphatemia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences causally related to treatment / all	1 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	

Frequency threshold for reporting non-serious adverse events: 5 %

Non-serious adverse events	Ixazomib 2 mg/m <sup>2</sup>	Ixazomib 1.6 mg/m <sup>2</sup>	
Total subjects affected by non-serious adverse events			
subjects affected / exposed	6 / 6 (100.00%)	3 / 3 (100.00%)	
General disorders and administration site conditions			
Alopecia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Dehydration			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Chills			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Epistaxis			
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences (all)	2	0	
Fatigue			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Pain			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	2	0	
Oral pain			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Fever			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Respiratory, thoracic and mediastinal disorders			
Cough			
subjects affected / exposed	1 / 6 (16.67%)	1 / 3 (33.33%)	
occurrences (all)	1	1	
Mucositis oral			

subjects affected / exposed occurrences (all)	2 / 6 (33.33%) 2	0 / 3 (0.00%) 0	
Psychiatric disorders			
Anxiety			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Insomnia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Anorexia			
subjects affected / exposed	2 / 6 (33.33%)	1 / 3 (33.33%)	
occurrences (all)	2	1	
Irritability			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Injury, poisoning and procedural complications			
Infusion related reaction			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Cardiac disorders			
Hypertension			
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences (all)	2	0	
Electrocardiogram QT corrected interval prolonged			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Sinus tachycardia			
subjects affected / exposed	4 / 6 (66.67%)	0 / 3 (0.00%)	
occurrences (all)	5	0	
Nervous system disorders			
Dizziness			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Peripheral sensory neuropathy			
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences (all)	2	0	



Blood and lymphatic system disorders			
Activated partial thromboplastin time prolonged			
subjects affected / exposed	1 / 6 (16.67%)	1 / 3 (33.33%)	
occurrences (all)	1	1	
Anemia			
subjects affected / exposed	5 / 6 (83.33%)	3 / 3 (100.00%)	
occurrences (all)	7	3	
Blood bicarbonate decreased			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	2	0	
Febrile neutropenia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Blood bilirubin increased			
subjects affected / exposed	2 / 6 (33.33%)	1 / 3 (33.33%)	
occurrences (all)	6	3	
Platelet count decreased			
subjects affected / exposed	4 / 6 (66.67%)	3 / 3 (100.00%)	
occurrences (all)	6	3	
Neutrophil count decreased			
subjects affected / exposed	5 / 6 (83.33%)	3 / 3 (100.00%)	
occurrences (all)	6	3	
Methemoglobinemia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Lymphocyte count decreased			
subjects affected / exposed	5 / 6 (83.33%)	0 / 3 (0.00%)	
occurrences (all)	5	0	
White blood cell decreased			
subjects affected / exposed	5 / 6 (83.33%)	3 / 3 (100.00%)	
occurrences (all)	6	3	
Thrombus			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Edema face			

subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences (all)	2	0	
Edema limbs			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Eye disorders			
Eye pain			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Gastrointestinal disorders			
Dyspepsia			
subjects affected / exposed	3 / 6 (50.00%)	0 / 3 (0.00%)	
occurrences (all)	3	0	
Gastroesophageal reflux disease			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Abdominal distension			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Abdominal pain			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Diarrhea			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Constipation			
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences (all)	3	0	
Bloating			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Pain in groin			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Vomiting			

subjects affected / exposed	2 / 6 (33.33%)	2 / 3 (66.67%)	
occurrences (all)	4	2	
Nausea			
subjects affected / exposed	3 / 6 (50.00%)	1 / 3 (33.33%)	
occurrences (all)	4	1	
Increased hunger			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Skin and subcutaneous tissue disorders			
Dry skin			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Erythema			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Palmar-plantar erythrodysesthesia syndrome			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Rash acneiform			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Rash maculo-papular			
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences (all)	2	0	
Renal and urinary disorders			
Bladder spasm			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Cystitis noninfective			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Dysuria			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Hematuria			

subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Proteinuria			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Hyperuricemia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Rectal pain			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Renal colic			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Urinary tract pain			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Endocrine disorders			
Syndrome of inappropriate antidiuretic hormone secretion			
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)	
occurrences (all)	0	1	
Musculoskeletal and connective tissue disorders			
Muscle cramp			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Pain in extremity			
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)	
occurrences (all)	2	0	
Back pain			
subjects affected / exposed	2 / 6 (33.33%)	1 / 3 (33.33%)	
occurrences (all)	2	1	
Arthralgia			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
Infections and infestations			

Adenovirus infection subjects affected / exposed occurrences (all)	1 / 6 (16.67%) 1	0 / 3 (0.00%) 0	
Parainfluenza 1 subjects affected / exposed occurrences (all)	0 / 6 (0.00%) 0	1 / 3 (33.33%) 1	
Enterocolitis infectious subjects affected / exposed occurrences (all)	1 / 6 (16.67%) 1	0 / 3 (0.00%) 0	
Herpes simplex reactivation subjects affected / exposed occurrences (all)	0 / 6 (0.00%) 0	1 / 3 (33.33%) 1	
Upper respiratory infection subjects affected / exposed occurrences (all)	0 / 6 (0.00%) 0	2 / 3 (66.67%) 2	
Thrush subjects affected / exposed occurrences (all)	1 / 6 (16.67%) 1	0 / 3 (0.00%) 0	
RHINOVIRUS subjects affected / exposed occurrences (all)	1 / 6 (16.67%) 1	0 / 3 (0.00%) 0	
Sepsis subjects affected / exposed occurrences (all)	0 / 6 (0.00%) 0	1 / 3 (33.33%) 1	
UTI R/T BK/ADENOVIRUS subjects affected / exposed occurrences (all)	1 / 6 (16.67%) 1	0 / 3 (0.00%) 0	
Metabolism and nutrition disorders ALT increased subjects affected / exposed occurrences (all)	2 / 6 (33.33%) 3	3 / 3 (100.00%) 4	
Alkaline phosphatase increased subjects affected / exposed occurrences (all)	1 / 6 (16.67%) 1	3 / 3 (100.00%) 4	
AST increased			

subjects affected / exposed	3 / 6 (50.00%)	2 / 3 (66.67%)
occurrences (all)	7	4
GGT increased		
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)
occurrences (all)	1	0
Hypercalcemia		
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)
occurrences (all)	1	0
Hyperglycemia		
subjects affected / exposed	3 / 6 (50.00%)	3 / 3 (100.00%)
occurrences (all)	9	8
Hyperkalemia		
subjects affected / exposed	2 / 6 (33.33%)	0 / 3 (0.00%)
occurrences (all)	2	0
Hypoalbuminemia		
subjects affected / exposed	4 / 6 (66.67%)	3 / 3 (100.00%)
occurrences (all)	4	4
Hypertriglyceridemia		
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)
occurrences (all)	0	1
Hyperphosphatemia		
subjects affected / exposed	0 / 6 (0.00%)	1 / 3 (33.33%)
occurrences (all)	0	1
Hypermagnesemia		
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)
occurrences (all)	2	0
Hypernatremia		
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)
occurrences (all)	1	0
Hypocalcemia		
subjects affected / exposed	3 / 6 (50.00%)	1 / 3 (33.33%)
occurrences (all)	6	3
Hypomagnesemia		
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)
occurrences (all)	2	0
Hypokalemia		

subjects affected / exposed	1 / 6 (16.67%)	1 / 3 (33.33%)	
occurrences (all)	1	4	
Hypoglycemia			
subjects affected / exposed	2 / 6 (33.33%)	1 / 3 (33.33%)	
occurrences (all)	2	2	
Hyponatremia			
subjects affected / exposed	4 / 6 (66.67%)	2 / 3 (66.67%)	
occurrences (all)	4	5	
Hypophosphatemia			
subjects affected / exposed	3 / 6 (50.00%)	1 / 3 (33.33%)	
occurrences (all)	5	1	
Weight loss			
subjects affected / exposed	1 / 6 (16.67%)	1 / 3 (33.33%)	
occurrences (all)	1	1	
Weight gain			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	
INR increased			
subjects affected / exposed	1 / 6 (16.67%)	0 / 3 (0.00%)	
occurrences (all)	1	0	

## More information

### Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
19 December 2019	The primary purpose of the amendment 1 was to make following changes: <ul style="list-style-type: none"><li>• Dose Modification for Intrathecal Methotrexate/Triple Intrathecal Therapy toxicities: Mercaptopurine dose adjustments made to allow for genotype variations in either TPMT or NUDT15</li><li>• Dose Modification for Intermediate-Dose Methotrexate toxicities: Added subsection 4.7.7.5 to provide guidance in the event of excess extravasation of fluids into tissues (third spacing)</li><li>• Clinical and Laboratory Studies: Added neurological exam to be performed prior to each Ixazomib dosing</li><li>• Bone Marrow Response Criteria for participants with leukemia: amended response definitions to replace CRp category with CRi and divide the CR category to include CR MRD-.</li></ul>
16 September 2020	The primary purpose of the amendment 2 was to make following changes: <ul style="list-style-type: none"><li>• Exclusion criteria: updated the list of excluded CYP3A4 agents</li><li>• Treatment program: under Block 1, 2, and Maintenance Block, revised timing of Leucovorin for Down syndrome Participants to be received at hours 24 and 30 after intrathecal methotrexate (IT MTX) or intrathecal triple therapy (ITT)</li><li>• Dose Limiting Toxicity: changed platelet criteria for hematological toxicity definition to platelet <math>\geq 20,000/\mu\text{L}</math>, platelet infusion independent and length of evaluation time to 49 days.</li></ul>
11 May 2021	The primary purpose of the amendment 3 was to make following changes: <ul style="list-style-type: none"><li>• Inclusion criteria: clarify the definition of what chemotherapy regimens and drugs and dosages (i.e. maintenance therapy drugs) are allowed prior to enrollment and what length of washout is required for the drugs and biologics</li><li>• Treatment program: added clarification to allow for flexibility in the timing of chemotherapy administration, including the timing of Day 29 IT therapy, allowing for up to 72 hours flexibility for scheduling or other issues.</li></ul>

Notes:

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### Interruptions (globally)

Were there any global interruptions to the trial? No

### Limitations and caveats

None reported