

PARC

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PARC: a phase I/II study evaluating the safety and activity of pegylated recombinant human arginase BCT-100 in relapsed/refractory cancers of children and young adults

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Background: The survival for many children with relapsed/refractory cancers remains poor despite advances in therapies. Arginine metabolism plays a key role in the pathophysiology of a number of pediatric cancers. We report the first in child study of a recombinant human arginase, BCT-100, in children with relapsed/refractory hematological, solid or CNS cancers.

Procedure: PARC was a single arm, Phase I/II, international, open label study. BCT-100 was given intravenously over one hour at weekly intervals. The Phase I section utilized a modified 3 + 3 design where escalation/de-escalation was based on both the safety profile and the complete depletion of arginine (defined as adequate arginine depletion; AAD <8μM arginine in the blood after 4 doses of BCT-100). The Phase II section was designed to further evaluate the clinical activity of BCT-100 at the pediatric RP2D determined in the Phase I section, by recruitment of patients with pediatric cancers into 4 individual groups. A primary evaluation of response was conducted at eight weeks with patients continuing to receive treatment until disease progression or unacceptable toxicity.

Results: 49 children were recruited globally. The Phase I cohort of the trial established the Recommended Phase II Dose of 1600U/kg iv weekly in children, matching that of adults. BCT-100 was very well tolerated. No responses defined as a CR, CRi or PR were seen in any cohort within the defined 8 week primary evaluation period. However a number of these relapsed/refractory patients experienced prolonged radiological SD.

Conclusion: Arginine depletion is a clinically safe and achievable strategy in children with cancer. The RP2D of BCT-100 in children with relapsed/refractory cancers is established at 1600U/kg intravenously weekly and can lead to sustained disease stability in this hard to treat population.

Clinical trial registration: EudraCT, 2017-002762-44; ISRCTN, 21727048; and [ClinicalTrials.gov](https://www.clinicaltrials.gov/ct2/show/study/NCT03455140), NCT03455140.

KEYWORDS

arginase, arginine, pediatric, cancer, relapse

Introduction

Despite significant improvements in the survival of children with cancer, certain tumor types, notably sarcomas, neuroblastoma, high grade gliomas and relapsed/high risk leukemias, still have poor outcomes with current treatment approaches (1). Therapeutic strategies for relapsed and refractory malignancies are limited, frequently using chemotherapy drugs with similar mechanisms of cytotoxicity to those in frontline. The benefits of dose-intensification of these agents has similarly been maximized leading to significant acute and chronic toxicities for children (2). Where possible targeted, immune and cellular therapies are also being utilized but their application may be limited to biomarker selected subgroups. Therapeutic strategies, which target malignancies through new mechanisms, and that do not add to the burden of toxicity are urgently needed.

Arginine is a semi-essential amino acid required for protein synthesis, cell division and a number of intracellular pathways that maintain cell survival (3). Both normal and malignant cells import arginine from the blood and microenvironment via the Cationic

Amino Acid (CAT/SLC7A) family of transporters and catabolize arginine through Arginase I, Arginase II, and nitric oxide synthetase (NOS2) enzyme activity. Although whole body arginine levels are maintained through dietary intake and tissue-specific re-synthesis, under conditions of high demand such as inflammation, pregnancy and cancer, arginine availability is limited. In the majority of non-malignant cells, precursors are recycled back into arginine through the expression of enzymes ornithine transcarbamylase (OTC – converting ornithine into citrulline), argininosuccinate synthase (ASS – converting citrulline into argininosuccinate), and argininosuccinate lyase (ASL – converting argininosuccinate back to arginine) (4). However in many cases both solid and haematological cancer cells are dependent on extracellular arginine for survival (arginine auxotrophism) due to the loss of ASS or OTC recycling enzyme expression; making them vulnerable to therapeutic arginine depletion (5). To date, absent OTC expression and low ASS expression has been reported in a number of pediatric cancers through *in vitro* and *in vivo* modelling suggesting they are auxotrophic for arginine (6–9).

The most clinically relevant approach to targeting tumor arginine metabolism is through therapeutic arginine depletion with a recombinant enzyme. BCT-100 is a pegylated recombinant human arginase that leads to a rapid depletion of arginine in pre-clinical models and in clinical trials of adult solid and haematological patients (10–13). In patients with adult malignancies, trials to date demonstrate a very encouraging safety profile as a single agent or in combination with chemotherapy. The frequency of Grade 3 or above toxicity has been low in trials to date. The most frequent reported adverse events include Grade 1-2 diarrhea, abdominal discomfort, or nausea. The optimal biological dose has been determined as 1600U/kg administered intravenously weekly. In Phase I trials the time taken for BCT-100 to achieve

Abbreviations: AAD, Adequate arginine depletion; AE, Adverse Event; ALL, Acute Lymphoblastic Leukemia; AML, Acute Myeloid Leukemia; ASL, Argininosuccinate lyase; ASS, Argininosuccinate synthase; CAT, Cationic Amino Acid; CR, Complete response; CRi, Complete response with incomplete count recovery; CT, Computerize Tomography Scan; DLT, Dose Limiting Toxicity; HGG, High Grade Glioma; MITT, Modified Intention to Treat; MRI, Magnetic Resonance Imaging scan; NOS2, Nitric Oxide Synthetase; OS, Overall Survival; OTC, Ornithine transcarbamylase; PFS, Progression Free Survival; PR, Partial Response; RP2D, Recommended Phase 2 Dose; SAE, Serious Adverse Event; SD, Stable Disease.

undetectable plasma arginine was 2 hours for all patients following a single dose. At 1600U/kg, adequate arginine depletion was achieved after the second weekly dose and was maintained throughout the treatment period. Here we report the findings from the PARC trial - the first to explore the safety of BCT-100 in a pediatric population (first in child) and activity in pediatric cancer populations.

Methods

Design and eligibility

PARC was a single arm, Phase I/II, international, open label study in the UK (7 sites), Netherlands (1 site) and Australia (5 sites). Key eligibility criteria included being less than 25 years old, histologically confirmed ALL/AML (Group 1), neuroblastoma (Group 2), sarcoma (Group 3), or high grade glioma (group 4) with radiological or laboratory evidence of disease progression during or after prior therapy, adequate organ function and no prior treatment with an arginine depleting drug.

BCT-100 was given intravenously over one hour at weekly intervals. The Phase I section utilized a modified 3 + 3 design where escalation/de-escalation was based on both the safety profile (occurrence of dose limiting toxicity; DLT) and the complete depletion of arginine (defined as adequate arginine depletion; AAD <8μM arginine in the blood after 4 doses of BCT-100). Patients were recruited in groups of 3 and followed for the occurrence of DLT. The standard 3 + 3 design was modified such that after each group of 3 patients had been assessed for safety a further evaluation of arginine depletion would take place. Dose de-escalation was to be advised as per the standard 3 + 3 algorithm, if 2 DLTs are observed at any dose level. Dose escalation would only take place if the dose was deemed safe but arginine depletion did not occur in all patients. The first cohort of patients were treated at 1600U/kg intravenously days 1,8,15, 22 of every 28 day course, the equivalent of the adult Recommended Phase II Dose (RP2D). Ultimately no patients required recruitment to the additional dose levels at 2000 U/kg (Level +1), 2500U/kg (Level +2), or 1200U/kg (Level -1).

The Phase II section was designed to further evaluate the clinical activity of BCT-100 at the pediatric RP2D determined in the Phase I section, by recruitment of patients with pediatric cancers into 4 individual groups: Group 1- Leukemias (Acute lymphoblastic leukemia (ALL) and acute myeloid leukemia (AML), Group 2 - Neuroblastoma, Group 3 - Sarcomas, and Group 4 - High Grade Gliomas (as defined by WHO central nervous system classification). 13 patients were planned to be recruited per group, patients who were treated at the selected Phase II dose in the Phase I component contributed to this 13 patient requirement. A primary evaluation of response was conducted at eight weeks with patients continuing to receive treatment until disease progression or

unacceptable toxicity. Tumor response was measured every 8 weeks thereafter whilst on therapy.

All patients were treated following written informed consent. The trial was sponsored by the University of Birmingham and registered with EudraCT, 2017-002762-44, ISRCTN, 21727048 and ClinicalTrials.gov NCT03455140.

Outcome measures and toxicity

For the Phase I section, the primary outcome measure was the safe and optimal (in terms of arginine depletion) RP2D of BCT-100 as determined by both the safety profile as measured by the occurrence/non-occurrence of DLT within 28 days of treatment with BCT-100 and the optimal dose as measured by the complete depletion of arginine (defined as <8μM arginine in the blood after 4 doses of BCT-100). For the Phase II section, the primary outcome measure was disease response (Complete Response - CR, or Partial Response - PR) after 8 weeks of treatment with BCT-100. The protocol defined responses were as follows: Group 1 (Leukaemia) - CR, Complete response with incomplete count recovery (CRi), Complete response without platelet recovery (CRp; ALL only), or PR determined by bone marrow, peripheral blood count/blasts and extramedullary disease (AML CR/PR criteria based on Cheson et al., 2003) (14); Group 2 (Neuroblastoma) - CR/PR determined by cross-sectional imaging by CT or MRI, MIBG scan and bone marrow evaluation using the International Neuroblastoma Response Criteria (INRC) (15); Group 3 (Sarcoma) - CR/PR determined by cross-sectional imaging by CT or MRI using RECIST version 1.1 (16); and Group 4 (High Grade Glioma) - CR/PR determined by cross-sectional imaging by MRI using RANO criteria (17). Secondary outcome measures were the incidence and severity of Adverse Events (AEs) defined by National Cancer Institute Common Terminology Criteria for Adverse Events (NCI CTCAE version 4), disease response (CR/PR) at any time during treatment with BCT-100, Progression Free Survival (PFS), Overall Survival (OS), and arginine concentrations in the blood, bone marrow, and cerebrospinal fluid. Exploratory outcome measures were disease response or stability (CR, PR, or Stable Disease - SD) after treatment with BCT-100, and the duration of disease response or stability after treatment with BCT-100.

Statistical analysis

All analyses were on a Modified Intention-To-treat Population (MITT), where those patients who withdrew or died prior to starting treatment are considered non-evaluable. Patients who had no recorded response at week 8 or confirmed progression/death prior to, are included as non-responders. A true response rate greater than 20% was of interest in any of the four disease groups. Response definitions were different for

each disease group and are based on specific criteria related to that disease e.g. Leukemia, Solid tumors as specified in the Methods section. Evaluable patients with no response data are included as non-responders. Each group had the response rate individually assessed using Bayesian posterior probability plots and highest posterior density intervals (HPD intervals). Posterior Probabilities were calculated for the true response rate in each arm using a non-informative prior Beta(0.5, 0.5).[19] The same design was implemented for each arm individually, that is, each arm had its assessment of response calculated separately.

Arginine ELISA

Blood samples were taken immediately prior to the administration of doses of BCT-100. The concentration of arginine was quantified using a competitive enzyme linked immunoassay (K7733, Immunodiagnostik) according to the manufacturers' instructions. In brief, the assay uses a competitive enzyme immunoassay in which L-arginine is derivatized from samples and competes with L-arginine tracer for binding to antibodies bound in the microtiter wells.

Immunohistochemistry/flow cytometry

To determine any effect of arginine depletion on circulating T and myeloid cell frequency, whole blood underwent red cell lysis (Qiagen) and then staining with anti-human CD3, or anti-human CD14 or anti-human CD15 antibodies (Biolegend) on ice for 30 min. Cells were resuspended in fluorescence-activated cell sorting buffer. Propidium (Biolegend) was used to assess viability. Cells were analyzed using a Beckman Coulter CytOflex flow cytometer and analyzed using FlowJo and CytExpert software (Tree Star Inc).

Results

In total, 49 relapsed/refractory patients were recruited globally to the trial across Phase I and Phase II from August 2018 to July 2022. (Supplementary Table 1; Figure 1) Recruitment to the Leukemia cohort (Group 1) was stopped early based on the recommendation of the Data Monitoring Committee, due to poor recruitment (7 out of 13 patients recruited). Of the 49 patients recruited, 45 commenced trial treatment and were evaluable (13 Sarcoma, 13 High Grade Gliomas, 12 Neuroblastoma and 7 Leukaemia) All patients who started treatment were dosed at 1600U/kg weekly. A median of 2 weeks treatment was administered to patients in the Leukaemia cohort (range 1-10), median of 7 weeks treatment in the neuroblastoma cohort (1-40), median of 7 weeks in the sarcoma cohort (2-41), and a median of 6

weeks in the high grade glioma cohort (2-68). (Supplementary Table 2) All patients were followed up until their death.

Phase I outcome measures

In the Phase I section of the trial 5 patients were recruited: 1 leukemia, 1 sarcoma, and 3 HGG, of which 3 were evaluable. Two patients did not complete 28 days of treatment due to disease progression. 1600U/kg BCT-100 led to a depletion of plasma arginine <8uM in the 3 evaluable patients and no DLTs were reported. Thus the Data Monitoring Committee confirmed the Phase I cohort of the trial met the primary outcome measures and the RP2D of 1600U/kg, matching that of adults.

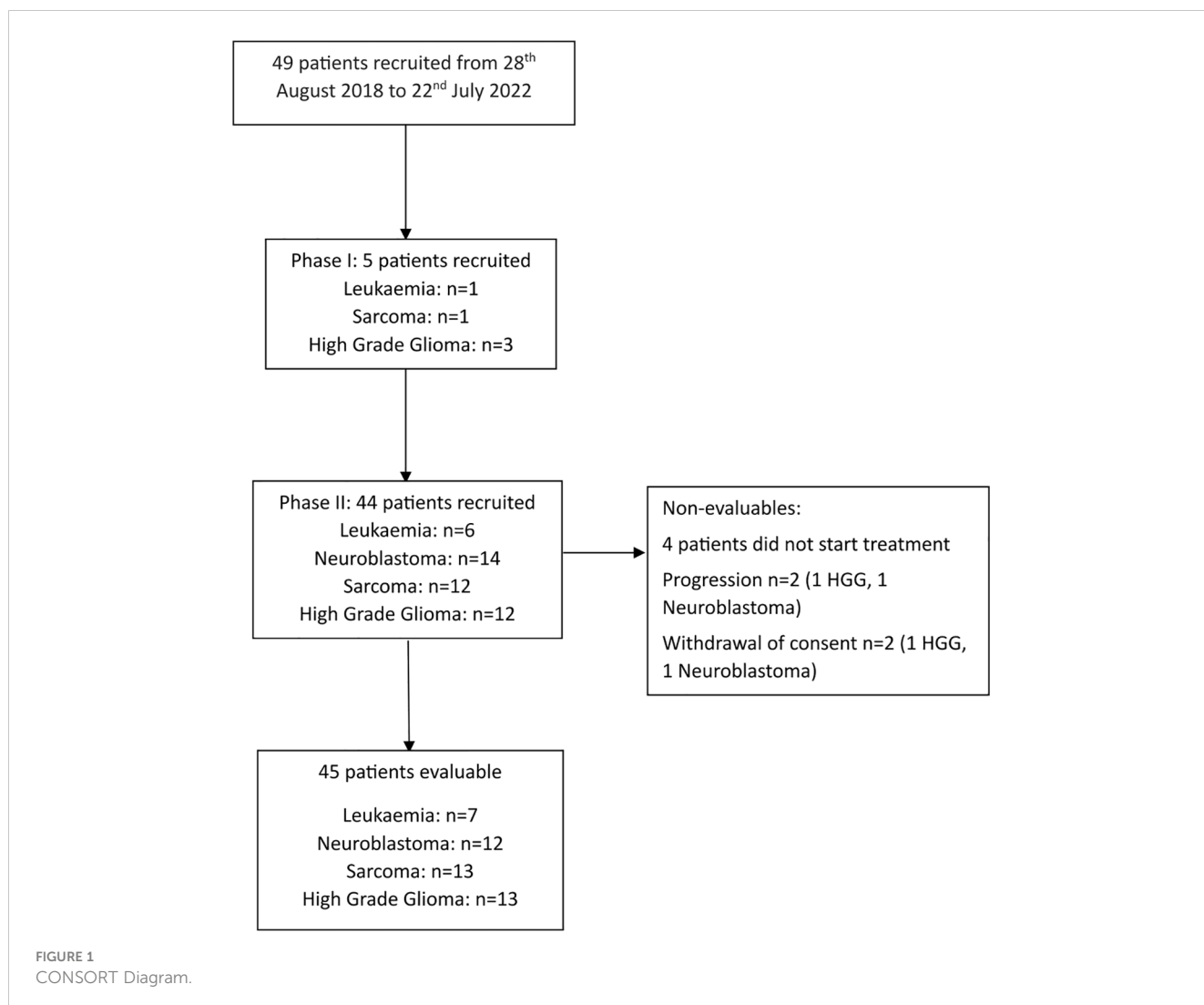
Phase II outcome measures

No responses defined as a CR, CRi or PR were seen in any cohort within the defined 8 week primary evaluation period. However a number of these relapsed/refractory patients experienced prolonged radiological SD (Figure 2A). In the HGG cohort one patient remained on treatment for 14 weeks and another patient for 68 weeks (PR at Week 16 reassessments). In the Neuroblastoma cohort 2 patients remained on treatment for 16 and 40 weeks, with a further patient receiving treatment for 8 weeks and remaining stable until week 24. In the Sarcoma cohort 3 patients remained on treatment for 24, 40, and 41 weeks. In the Leukaemia cohort one patient remained on treatment for 10 weeks (SD at week 8 bone marrow assessment). The OS per group is shown in Figure 2B and Supplementary Table 3. Progression Free survival per group is shown in Figure 2C and Supplementary Table 4.

In the majority of Phase II patients arginine was depleted to less than 8uM and sustained for the duration of their time on therapy (Figure 3A). In 7 patients (1 leukemia, 5 neuroblastoma, and 1 HGG) plasma arginine concentrations greater than 8uM were recorded whilst on therapy. Arginine concentrations greater than 8uM did not correlate with the length of time patients experienced disease response. Unfortunately samples were not provided by sites on all patients at all time points, limiting further interpretation of the plasma arginine profile over time.

Toxicity

Rates of Grade 3 or higher toxicity were very low in all groups (Supplementary Tables 5, 6). Only two serious adverse events (SAEs) were deemed associated with trial treatment: one patient with HGG was hospitalized with a seizure that resolved with no sequelae; one patient with neuroblastoma was hospitalized with an allergic reaction that resolved with no sequelae. A low number of Adverse Events were recorded as summarized in Supplementary Tables 7, 8. No significant



changes in the frequencies of CD3+ T cells, or CD14+ or CD15+ myeloid cells were seen in the blood over time (Figures 3B–D).

Discussion

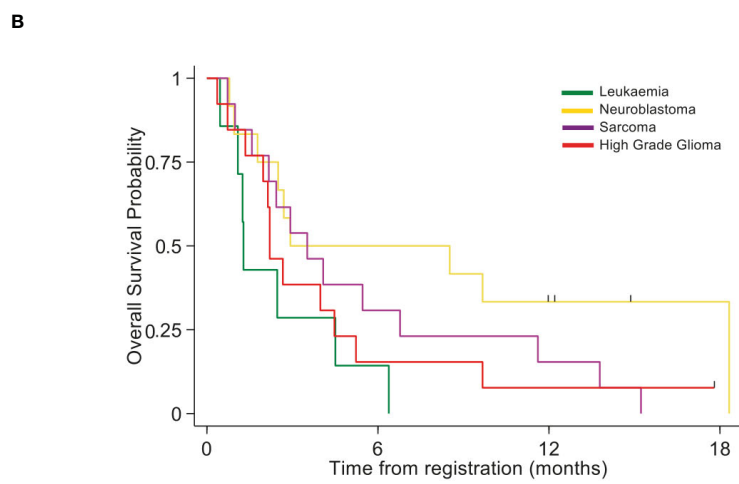
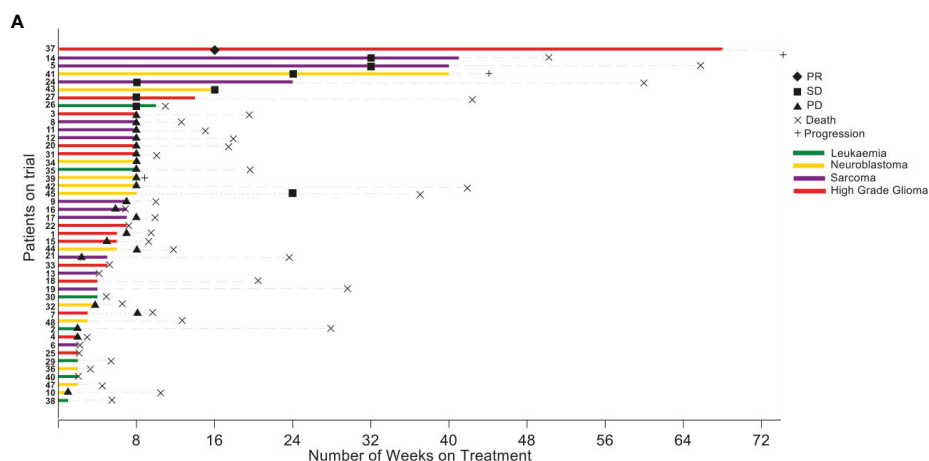
We demonstrated that BCT-100 recombinant arginase can be administered to children with relapsed/refractory solid, haematological or Central Nervous System malignancies, with an acceptable toxicity profile. The RP2D was confirmed as 1600U/kg weekly via intravenous infusion, consistent with the adult RP2D previously identified (12, 13). The low toxicity in this patient population, despite heavy prior treatment, is encouraging as a basis for future combination therapeutic approaches.

Although no patients exhibited a CR, 1 patient with a HGG experienced a PR. Furthermore 8 patients maintained SD (1 HGG, 1 Leukaemia, 3 Neuroblastoma, and 3 Sarcomas) – notable as all patients had confirmed disease progression at the time of trial enrolment. 8 patients remained on treatment

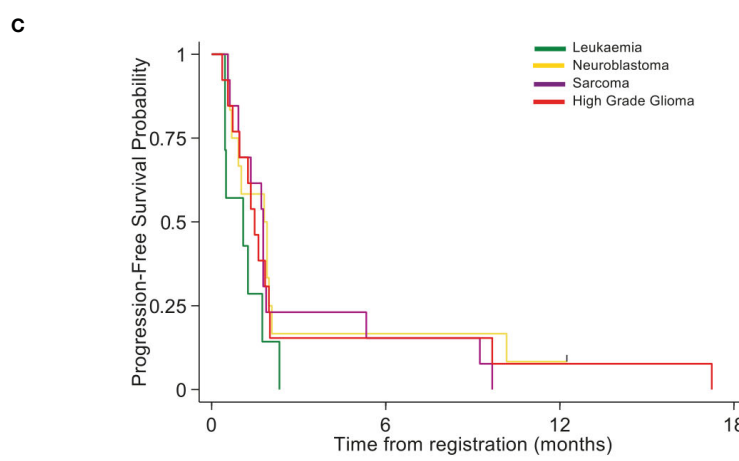
beyond the initial 8 weeks evaluation time point (range: 10–68 weeks), suggesting BCT-100 can provide sustained tumor growth inhibition.

In some adult trials, the depth or duration of arginine depletion in the serum correlated with patient outcome (12, 18), however tumor cell expression of ASS or OTC are more often used as predictive biomarkers of response to therapeutic arginine depletion. We and others have previously shown that the majority of pediatric cancers have low to absent ASS or OTC expression, thus enrichment for ASS/OTC negative patients could be one strategy to enhance response. Similar biomarker-enriched approaches were taken for BCT-100 and other arginine depleting enzymes still under clinical investigation (18, 19).

Overall, the findings in the PARC trial are similar to those of single-agent BCT-100 in adult cancers. We identified that the adult RP2D of 1600U/kg iv weekly is the RP2D in children, and that some patients with different tumor types can experience prolonged disease stability with BCT-100 as a single agent. We acknowledge that this study is limited by the lack of samples to conduct a



	0	6	12	18			
Number at risk	7	6	1	(1)	0	(0)	0
Leukaemia	7	(6)	1	(1)	0	(0)	0
Neuroblastoma	12	(6)	6	(2)	3	(0)	1
Sarcoma	13	(9)	4	(2)	2	(2)	0
High Grade Glioma	13	(11)	2	(1)	1	(0)	0



	0	6	12	18			
Number at risk	7	(7)	0	(0)	0	(0)	0
Leukaemia	7	(7)	0	(0)	0	(0)	0
Neuroblastoma	12	(10)	2	(1)	1	(0)	0
Sarcoma	13	(11)	2	(2)	0	(0)	0
High Grade Glioma	13	(11)	2	(1)	1	(1)	0

FIGURE 2 (A) Swimmer plot indicating the number of weeks patients received treatment with BCT-100. Disease cohort and response rates indicated by colors and symbols in the key. (B) Overall survival (OS) (C) Progression-free survival (PFS).

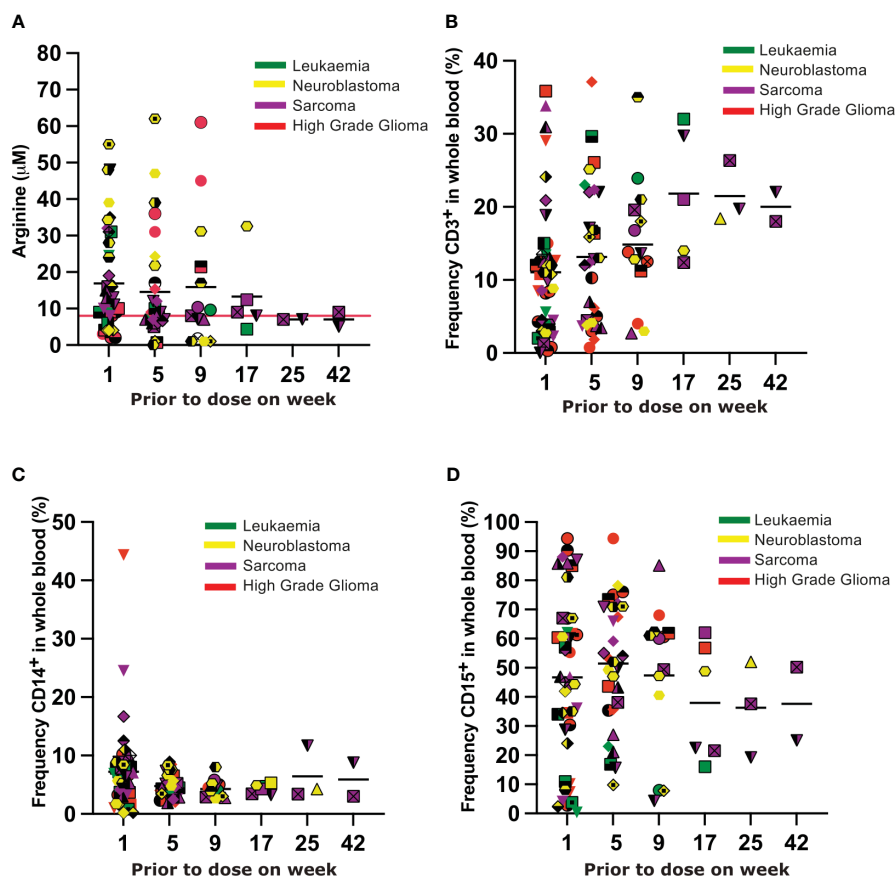


FIGURE 3

(A) Plasma arginine concentrations prior to each dose of BCT-100. Each unique symbol represents an individual patient. Each color represents the tumour subtype, as indicated in the legend. (B–D) Frequency of CD3+ T cells, CD14+ myeloid cells and CD15+ myeloid cells respectively in the blood of patients prior to each dose of BCT-100. Each unique symbol represents an individual patient. Each color represents the tumour subtype, as indicated in the legend.

comprehensive analysis into biomarkers of response which would have shed important biological insights into similarities and differences between adult and pediatric responding tumours. However, with the low toxicity profile, BCT-100 remains an attractive molecule for further clinical development in combination with chemotherapy regimens. Pre-clinical testing suggests that combinations with Dintuximab/Ironotecan/Temozolomide for relapsed neuroblastoma or irinotecan/temozolomide for relapsed sarcomas could be synergistic, and form the basis for a subsequent clinical trial.

Data availability statement

The dataset supporting the conclusions of this article is available by application to the Cancer Research UK Clinical Trials Unit at the University of Birmingham.

Ethics statement

All patients were treated following written informed consent, following IRB review within each country participating in the trial.

Author contributions

NF: Conceptualization, Data curation, Funding acquisition, Methodology, Project administration, Resources, Supervision, Writing – original draft, Writing – review & editing. RW: Formal analysis, Methodology, Supervision, Writing – original draft, Writing – review & editing. KW: Data curation, Formal analysis, Funding acquisition, Investigation, Methodology, Resources, Supervision, Writing – review & editing. JH: Data curation, Resources, Writing – review & editing. LM: Investigation, Resources, Writing – review & editing. ME: Investigation, Resources, Writing – review & editing. GM: Investigation, Resources, Writing – review & editing. AN: Investigation, Resources, Visualization, Writing – review & editing. BB: Investigation, Resources, Writing – review & editing. SL: Investigation, Resources, Writing – review & editing. JA: Investigation, Resources, Writing – review & editing. JK: Investigation, Resources, Writing – review & editing. RC: Investigation, Resources, Writing – review & editing. MG: Investigation, Resources, Writing – review & editing. AM: Investigation, Resources, Writing – review & editing. TT: Investigation, Resources, Writing – review & editing. MR: Investigation, Resources, Writing – review & editing. MC: Investigation, Resources, Writing – review & editing.

HC: Investigation, Resources, Writing – review & editing. MW: Investigation, Resources, Writing – review & editing. MK: Investigation, Resources, Writing – review & editing. NV: Investigation, Resources, Writing – review & editing. JK: Investigation, Resources, Writing – review & editing. US: Data curation, Formal analysis, Investigation, Methodology, Writing – review & editing. VS: Data curation, Formal analysis, Investigation, Methodology, Writing – review & editing. LF: Data curation, Formal analysis, Investigation, Methodology, Writing – review & editing. SB: Data curation, Formal analysis, Investigation, Methodology, Writing – review & editing. PC: Funding acquisition, Investigation, Methodology, Project administration, Resources, Supervision, Writing – review & editing. CD: Conceptualization, Formal analysis, Funding acquisition, Investigation, Methodology, Project administration, Resources, Supervision, Writing – original draft, Writing – review & editing. FM: Conceptualization, Data curation, Funding acquisition, Investigation, Methodology, Project administration, Resources, Supervision, Writing – original draft, Writing – review & editing.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fonc.2024.1296576/full#supplementary-material>

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