

Supplementary appendix

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Nivolumab + BV for pts aged 5 to 30 y with R/R cHL

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Nivolumab + BV for pts aged 5 to 30 y with R/R cHL

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Nivolumab + BV for pts aged 5 to 30 y with R/R cHL

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Supplementary tables

Supplementary table S1. Risk stratification algorithm for CheckMate 744

Stage at diagnosis	Time to relapse from end of therapy (months)	B symptoms ^a or extranodal disease at relapse, extensive disease where radiation therapy was contraindicated at relapse, or relapse in a prior radiation field
Standard risk: R2 cohort		
Any	<3	Yes/No
IA, IIA	3–12 >3 cycles and/or RT	Yes/No
IB, IIB, IIIA	<12	
IIIB, IV	Any	Yes
Any	Any	
Low risk: R1 cohort		
IA, IIA	≥12	No
	3–12 ≤3 cycles and no RT	No
IB, IIB, IIIA	>12	No

^aB symptoms include fever (≥101 °F or ≥38.3 °C), drenching night sweats, and unexplained weight loss (≥10% of body weight over 6 months).

RT, radiation therapy.

Supplementary table S2. Treatment summary for patients who proceeded to consolidation off protocol

Patient	Completed treatment	Response after 4 cycles of induction		Best response prior to consolidation		Off-protocol therapies, auto-HCT, subsequent therapies
		Per BICR	Per INV	Per BICR	Per INV	
1	4 cycles of induction	PMR	CMR	CMR	CMR	<ul style="list-style-type: none"> • BEAM (HDCT) • Auto-HCT
2	4 cycles of induction	PMR	PMR	CMR	CMR	<ul style="list-style-type: none"> • Gemcitabine, ifosfamide, prednisolone, vinorelbine (second salvage) • Lomustine, cytarabine, etoposide, melphalan (HDCT) • Auto-HCT
3 ^a	4 cycles of induction	NE	CMR	NE	CMR	<ul style="list-style-type: none"> • BEAM (HDCT) • Auto-HCT
4	4 cycles of induction	CMR	CMR	CMR	CMR	<ul style="list-style-type: none"> • Fotemustine, cytarabine, etoposide, melphalan (HDCT) • Auto-HCT
5	4 cycles of induction + 4 cycles of intensification	PMR	PMR	PMR	CMR	<ul style="list-style-type: none"> • BEAM (HDCT) • Auto-HCT • Surgery
6	4 cycles of induction	PMR	CMR	PMR	CMR	<ul style="list-style-type: none"> • BEAM (HDCT) • Auto-HCT
7	4 cycles of induction	CMR	CMR	CMR	CMR	<ul style="list-style-type: none"> • BEAM (HDCT) • Auto-HCT
8	4 cycles of induction	PMD	PMR	CMR	PMR	<ul style="list-style-type: none"> • Ifosfamide, carboplatin, etoposide (second salvage) • Auto-HCT

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9 ^b	2 cycles of induction	NR	NR	NR	NR	<ul style="list-style-type: none">• Bendamustine (second salvage)• BEAM (HDCT)• Auto-HCT• BV (post-transplant immunotherapy)
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^aThis patient was not evaluable for response because he/she did not achieve PMR or CMR per BICR at any time, or completion of 6 cycles of therapy (4 cycles of nivolumab plus BV and 2 cycles of BV plus bendamustine).

^bThis patient discontinued after 2 cycles of induction due to toxicity, but was evaluable for response.

BEAM, carmustine, cytarabine, etoposide, and melphalan; INV, investigator; NE, not evaluable; NR, no response; PMD, progressive metabolic disease.

Supplementary table S3. Subsequent cancer therapy

	R2 cohort (n=44)
Any subsequent therapy ^a	27 (61.4)
Radiation therapy ^b	9 (20.5)
Curative intent	2 (4.5)
Other ^c	7 (15.9)
Systemic therapy	21 (47.7)
Reason for subsequent systemic therapy	
Maintenance with BV after auto-HCT (without disease progression or clinical deterioration)	10 (22.7)
Disease progression	3 (6.8)
Other/not reported	9 (20.5)

Data are n (%).

^aPatients could receive a combination of therapies.

^bEight patients received radiation therapy as consolidation after auto-HCT; 1 patient received radiation therapy as a subsequent therapy after progression.

^cOther reasons included local consolidation, standard of care, and follow-up treatment.

Auto-HCT, autologous hematopoietic cell transplantation; BV, brentuximab vedotin.

Supplementary table S4. Stem cell mobilization summary

	R2 cohort (n=40)
Median (range) number of apheresis sessions per patient	1.0 (1–5)
Median (range) number of CD34+ cells collected per session, $\times 10^6$ cells/kg (n=59 sessions)	4.0 (0.3–268.0)
Mobilization agents (≥ 2 patients), n (%)	
G-CSF ^a	22 (55)
Plerixafor	4 (10.0)
Calcium	3 (7.5)
Carmustine	2 (5.0)
Cyclophosphamide	2 (5.0)
Cytarabine	2 (5.0)
Etoposide	2 (5.0)
Melphalan	2 (5.0)

^aIncludes filgrastim, G-CSF, and lenograstim.

G-CSF, granulocyte colony stimulating factor.

