ACTN01029999

REPORT DATE
06-Oct-2023

Gender: Male | Birth year: 1950 | WHO: 1

Tumor: Skin - Melanoma | Lesions: CNS, Liver, Lymph Node | Stage: IV

## Summary

## **Clinical summary**

Relevant systemic treatment history	8/2020-3/2021	Therapy2	
	2020	Therapy1	
	2022	Clinical trial: Trial1	
	2022	Clinical trial: Trial1 (3 cycles, stop reason: toxicity)	
	2022	Clinical trial (tr2): Trial2 (3 cycles, stop reason: toxicity)	
	2022	Clinical trial: Trial4 (adjuvant, 3 cycles, stop reason: toxicity)	
	2022	Clinical trial: Trial5 (adjuvant and consolidation, 3 cycles, stop reason:	
		toxicity)	
Relevant other oncological history	2022	Clinical trial (details unknown)	
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Previous primary tumor	Lung adenocarcinoma (diagnosed 10/2020, considered non-active)		
Relevant non-oncological history	Pancreatitis		

#### Recent molecular results

## WHOLE\_GENOME of ACTN01029999 (01-Oct-2023)

Biopsy location Liver (purity 98%) Molecular tissue of origin prediction Melanoma (100%) Tumor mutational load / burden TML high (185) / TMB high (13.7) Microsatellite (in)stability Stable HR status **Proficient** Genes with high driver mutation BRAF Amplified genes MYC Deleted genes **PTEN** Homozygously disrupted genes **PTEN** EML4 - ALK fusion Gene fusions HPV (3 integrations detected) Virus detection

Potentially actionable events with medium/low driver: PTEN disruption

IHC results V600e positive (Panel NGS): BRAF

## Approved treatments considered eligible

Treatment

Not yet determined

## EMC trials that are open and considered eligible and currently have slots available (1)

Trial	Cohort	Molecular	Warnings
Test Trial 1 TEST-1	Cohort B	None	Undetermined SOC exhaustion

## EMC trials that are open and considered eligible but currently have no slots available (2)

Trial	Cohort	Molecular	Warnings
Test Trial 1	Cohort A	BRAF V600E	Undetermined SOC exhaustion
TEST-1			

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## Continued from the previous page

Trial	Cohort	Molecular	Warnings
Test Trial 2 TEST-2	Cohort A	BRAF V600E	None

## trial kb trials potentially eligible based on molecular results (8)

Event	Trials
EML4 - ALK fusion	external trial
HPV positive	external trial
MYC amp	external trial
PTEN del	Trial 1
PTEN disruption	external trial
PTEN hom disruption	external trial
TMB High	external trial
TML High	external trial

PATIENT ACTN01029999

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## **Molecular Details**

IHC results V600e positive (Panel NGS): BRAF

## WHOLE\_GENOME (ACTN01029999T, 01-Oct-2023)

#### General

Purity	Sufficient Quality	TML Status	TMB Status	MS Stability	HR Status	DPYD	UGT1A1
98%	Yes	High (185)	High (13.7)	Stable	Proficient	*1 HOM (Normal function)	*1 HET (Normal function), *28 HET (Reduced
							function)

#### Predicted tumor origin

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Combined avadiation seems	1000/	
Combined prediction score	100%	
This score is calculated by combining information on:		
(1) SNV types	98%	
(2) SNV genomic localisation distribution	99%	
(3) Driver genes and passenger characteristics	97%	

Other cohorts have a combined prediction of 0% or lower

#### **Drivers**

Туре	Driver	Driver likelihood	Trials in EMC	Trials in trial kb	Best evidence in kb	Resistance in kb
Mutation (Hotspot)	BRAF V600E (4/6 copies)	High	TEST-1, TEST-2		Approved	
Amplification	MYC amp, 38 copies	High		external trial	Approved	Known resistance
Loss	PTEN del, 0 copies	High		Trial 1		
Known fusion	EML4 - ALK fusion, exon 6 - exon 20	High		external trial	Approved	Known resistance
Disruption (homozygous)	PTEN	High		external trial	Approved	Known resistance
Virus	HPV positive, 3 integrations detected	High		external trial	Approved	Known resistance
Disruption	PTEN, DEL (1.1 disr. / 1.8 undisr. copies)	Low		external trial	Approved	Known resistance

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## **Clinical Details**

#### **Clinical summary**

Relevant systemic treatment history 8/2020-3/2021 Therapy2 2020 Therapy1

2022 Clinical trial: Trial1

2022 Clinical trial: Trial1 (3 cycles, stop reason: toxicity)
2022 Clinical trial (tr2): Trial2 (3 cycles, stop reason: toxicity)
2022 Clinical trial: Trial4 (adjuvant, 3 cycles, stop reason: toxicity)

2022 Clinical trial: Trial5 (adjuvant and consolidation, 3 cycles, stop reason:

toxicity)

Relevant other oncological history 2022 Clinical trial (details unknown)

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Previous primary tumor Lung adenocarcinoma (diagnosed 10/2020, considered non-active)

Relevant non-oncological history Pancreatitis

## Patient current details (26-Sep-2023)

Unresolved toxicities grade => 2 Fatigue (2)

Cancer-related complications Ascites

Known allergies Wasps (Environment)

Recent surgeries 06-Sep-2023

## Tumor details (26-Sep-2023)

Measurable disease Yes

CNS lesion status

Present CNS lesions (active)

Brain lesion status

No known brain lesions

## **Active medication details**

Medication	Administration route	Start date	Stop date	Dosage	Frequency
Ibuprofen		06-Sep-2023	21-Oct-2023	750 - 1000 mg	1 / day
Prednison		06-Sep-2023	21-Oct-2023	750 - 1000 mg	1 / 3 months

## **Blood transfusions**

Product	Date
Thrombocyte concentrate	21-Sep-2023

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# **Trial Matching Summary**

EMC trials and cohorts that meet molecular requirements and may be eligible, but are closed (0)

None

EMC trials and cohorts that are open but considered ineligible (1)

Trial	Cohort	Molecular	Ineligibility reasons
Test Trial 2 TEST-2	Cohort B		Vemurafenib treatment