PATIENT ACTN-01-02-9999

REPORT DATE

10-Mar-2022

Sample ID: ACTN01029999T | Gender: Male | Birth year: 1950 | WHO: 1

Tumor: Skin - Melanoma | Lesions: Abdominal, CNS, Liver, Pulmonal | Stage: IV

Summary

Patient clinical history (28-Feb-2022)

Relevant systemic treatment history Vemurafenib (3/2021), Ipilimumab (2022)

Other oncological history Resection (2020)

Previous primary tumor: Bone/Soft tissue Schwannoma (3/2019, considered active)

Relevant non-oncological history Pancreatitis, Myocardial infarction

Molecular results

Previous relevant molecular results ERBB2 3 + (IHC)

WGS molecular results (05-Mar-2022)

Biopsy location Liver

Events with approved treatment evidence in evidence

BRAF V600E, High TML

Events with trial eligibility in local database

BRAF V600E, High TML

Additional events with experimental evidence (evidence)

None

Additional events with other responsive evidence in evidence

None

Events with resistance evidence in evidence BRAF V600E: Erlotinib

Approved treatments considered eligible

Treatment

Not yet determined

local trials and cohorts considered potentially eligible (1)

Trial ID Acronym Cohort Cohort

Test Trial TEST-TRIAL Cohort B Yes

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Clinical Details

Patient current details (28-Feb-2022)

Unresolved toxicities grade => 2 Fatigue (2), Dizziness
Significant aberration on latest ECG Atrial arrhythmia

Cancer-related complications Ascites

Known allergies Wasps (Environment), Pembrolizumab (Medication)

Recent surgeries 08-Feb-2022

Tumor details (28-Feb-2022)

Measurable disease Yes

CNS lesion status CNS (active)

Brain lesion status No known brain lesions

Laboratory results (28-Feb-2022)

Liver function Total bilirubin

ASAT (< 33 U/L) 36 U/L (no trend information available)

ALAT ALP

Albumin

Kidney function Creatinine

CKD-EPI eGFR (> 100 mL/min) > 100 mL/min (08-Feb-2022)

Other Hemoglobin (6.5 - 9.5 mmol/L) 5.5 mmol/L (no trend information available)

Thrombocytes (155 - 350 10^9/L) 150 10^9/L (out of range for 2 cons. measurements, trend down)

LDH (< 245 U/L) **240 U/L (18-Feb-2022)**

Medication details

MedicationCategoriesStart dateStop dateActive?DosageFrequencyIbuprofenNSAIDs08-Feb-202225-Mar-2022Yes750 - 1000 mg1 / day

Blood transfusions

Product Date

Thrombocyte concentrate 23-Feb-2022

ACTIN Report

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Trial Matching Summary

Trial counts

Trials evaluated 1
Cohorts evaluated 3

Evaluation results per trial & cohort

Trial / Cohort	# Criteria	# Pass	# Warn	# Fail	# Undet.	# No eval
Test Trial (TEST-TRIAL)	3	2	0	0	0	1
Test Trial - Cohort A	1	0	0	1	0	0
Test Trial - Cohort B	0	0	0	0	0	0
Test Trial - Cohort C	0	0	0	0	0	0

ACTIN Report

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Trial Matching Details

Potentially eligible trials & cohorts

Test Trial

Evaluation PASS

Acronym TEST-TRIAL

Title This is an ACTIN test trial

RuleReferenceEvaluationI-01Is adultPASS

pass

E-01 This rule has 2 conditions: NOT_EVALUATED

Patient has no active brain metastases.

pass

pass

2. Patient has exhausted SOC.

Test Trial - Cohort A

Cohort ID A

Evaluation FAIL

Open for inclusion? Yes

Rule Reference Evaluation

I-02 Has no active CNS metastases FAIL

fail

Test Trial - Cohort B

Cohort ID B
Evaluation PASS
Open for inclusion? Yes

Test Trial - Cohort C

Cohort ID C
Evaluation PASS
Open for inclusion? Yes
Blacklisted for eligibility? Yes