PATIENT ACTN-01-02-9999

REPORT DATE

22-Mar-2022

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

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

Summary

Patient clinical history (12-Mar-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 (17-Mar-2022)

Biopsy location Liver

Events with approved treatment evidence in general BRAF V600E, High TML

Events with trial eligibility in local database BRAF V600E, High TML

Additional events with trial eligibility in NL (external)

Additional events with experimental evidence (general)

Additional events with other responsive evidence in general

None

Events with resistance evidence in general BRAF V600E: Erlotinib

Approved treatments considered eligible

Treatment

Not yet determined

local trials that are open and considered eligible (1)

Trial	Acronym Cohor	t Molecular	Warnings
Test Trial 1	TEST-TRIAL-1 Cohort	t B No	None

external trials potentially eligible based on molecular results

Event	Trials
BRAF V600E	Trial 1
High TML	Trial 1

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Clinical Details

Patient current details (12-Mar-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 20-Feb-2022

Tumor details (12-Mar-2022)

Measurable disease Yes

CNS lesion status CNS (active)

Brain lesion status No known brain lesions

Laboratory results (12-Mar-2022)

Liver function Total bilirubin

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

ALAT

Albumin

Kidney function Creatinine

CKD-EPI eGFR (> 100 mL/min) > 100 mL/min (20-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 (02-Mar-2022)**

Medication details

MedicationCategoriesStart dateStop dateActive?DosageFrequencyIbuprofenNSAIDs20-Feb-202206-Apr-2022Yes750 - 1000 mg1 / day

Blood transfusions

Product Date

Thrombocyte concentrate 07-Mar-2022

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Trial Matching Summary

local trials that are closed but considered eligible (0)

Trial	Acronym	Cohort	Molecular	Warnings
None				

local trials not considered eligible (3)

Trial	Acronym	Cohort	Open	Ineligibility reasons
Test Trial 1	TEST-TRIAL-1	Cohort A	Yes	fail general
Test Trial 1	TEST-TRIAL-1	Cohort C	Yes	Cohort blacklisted
Test Trial 2	TEST-TRIAL-2		Yes	fail general

ACTIN Report

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Trial Matching Details

Potentially eligible trials & cohorts

Test Trial 1

Potentially eligible Yes

Acronym TEST-TRIAL-1

Title This is the first ACTIN test trial

RuleReferenceEvaluationI-01Is adultPASS

pass specific

E-01 This rule has 2 conditions: NOT_EVALUATED

Patient has no active brain metastases.
 Patient has exhausted SOC.

not evaluated specific

Test Trial 1 - Cohort A

Cohort ID A

Potentially eligible No

Open for inclusion? Yes

Rule Reference Evaluation

I-02 Has no active CNS metastases FAIL

fail specific

Test Trial 1 - Cohort B

Cohort ID B
Potentially eligible Yes
Open for inclusion? Yes

Test Trial 1 - Cohort C

Cohort ID C
Potentially eligible No
Open for inclusion? Yes
Blacklisted for eligibility? Yes

ACTIN Report

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Other trials & cohorts

Test Trial 2

Potentially eligible No

Acronym TEST-TRIAL-2

Title This is the second ACTIN test trial

Rule Reference Evaluation

I-01 Should have active infection FAIL

fail specific