PATIENT
EXAMPLE-LUNG-02
REPORT DATE
01-Nov-2024

Gender: Male | Birth year: 1965 | WHO: 1

Tumor: Lung - Squamous cell carcinoma | Lesions: Liver, Lung | Stage: III

Summary

Clinical summary

Relevant systemic treatment history 6/2023-6/2024 Carboplatin+Pemetrexed continued with Pemetrexed

maintenance

7/2024-8/2024 Docetaxel

Relevant other oncological history None

Previous primary tumor None

Relevant non-oncological history 1/2020 Acute myocardial infarction

1/2000 Bipolar disorder

Recent molecular results

NGS Lung Panel 1.2 (15-Sep-2024)

Biopsy location Lung

Molecular tissue of origin prediction Unknown

Tumor mutational load / burden Unknown / TMB low (8)

Microsatellite (in)stability

Stable

HR status

Proficient

High driver mutations PIK3CA E545K

Amplified genes None

Deleted genes TP53, KEAP1

Homozygously disrupted genes

Gene fusions

None

Virus detection

None

Potentially actionable events with medium/low driver:

None

IHC results PD-L1: Score < 1%

Example trials that are open and potentially eligible (1 cohort from 1 trial)

Trial	Cohort	Molecular	Warnings
METC 05	Dose expansion - monotherapy -	PIK3CA E545K	History of Acute myocardial infarction
PICKME3CA	NSCLC		

Example trials that are open and potentially eligible but currently have no slots available (1 cohort from 1 trial)

Trial	Cohort	Molecular	Warnings
METC 01	Dose escalation - monotherapy		None
IEMOEN			

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Trial Matching Summary

Example trials and cohorts that are potentially eligible, but are closed (1)

Trial	Cohort	Molecular	Warnings
METC 01 IEMOEN	Dose expansion - monotherapy		None
Example trials and cohorts that are considered ineligible (5)			

Trial	Cohort	Molecular	Ineligibility reasons
METC 05 PICKME3CA	Dose expansion - monotherapy - Other cancer types	PIK3CA E545K	Has tumor belonging to DOID term(s) lung non-small cell carcinoma
METC 02	Applies to all cohorts below	None	G12D not detected in KRAS
KAYRAS	Dose expansion - monotherapy - Colorectum		No colorectal cancer
	Dose expansion - monotherapy - NSCLC		
METC 03 NO-SEE797ES	Dose escalation - monotherapy		No variants in codon(s) C797 in EGFR
METC 04 TEDR1	Lung cancer C797S cohort		C797S not detected in EGFR

Open cohorts with no slots available are shown in grey.

Example trials and cohorts that are not evaluable or ignored (0)

None

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Trial Matching Details

Potentially eligible open trials & cohorts

METC 01

Potentially eligible Yes

Acronym IEMOEN

Title Phase I first-in-human study to evaluate safety of IEMOEN, a new PD-L1 inhibitor in advanced solid tumors

Rule	Reference	Evaluation
E-02	Has hemoglobin below 6 mmol/l	WARN
		Hemoglobin 8.2 mmol/L exceeds minimum of 6.0 mmol/L, but measurement occurred before 03-Aug- 2024
E-03	Has absolute neutrophil count below 1.5 x 10^9/l	UNDETERMINED
		No measurement found for absolute neutrophil count
I-01	Patients must be ≥18 years old.	PASS
		Patient is at least 18 years old
I-02	IEMOEN monotherapy is indicated for the treatment of adults with advanced/metastatic solid tumors.	PASS
		Patient has solid primary tumor
		Tumor stage III is considered locally advanced
		Tumor stage III is considered metastatic
I-03	Patient should have exhausted applicable available standard-of-care treatments.	PASS
		SOC considered exhausted since platinum doublet in treatment history
E-01	Has an active autoimmune disease that requires systemic treatment and poses a risk according	PASS
	to the investigator.	Patient has no other condition belonging to category autoimmune disease

METC 01 - Dose escalation - monotherapy

Cohort ID A

Potentially eligible? Yes

Open for inclusion? Yes

Has slots available? No

METC 01 - Dose expansion - monotherapy

Cohort ID B
Potentially eligible? Yes
Open for inclusion? No
Has slots available? No

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METC 05

Potentially eligible Yes

Acronym PICKME3CA

Title A phase 1/2 trial of ABC123 +/- platinum doublet in PIK3CA-mutated solid cancer

Rule	Reference	Evaluation
I-05	Patient can not have a severe cardiac condition.	WARN
		Patient has history of condition(s) Acute myocardial infarction, which is indicative of coronary artery disease
I-03	ALAT and ASAT should be at most 3*ULN, or at most 5*ULN in case of liver metastases.	UNDETERMINED
		ASAT and ALAT are not present or cannot be evaluated
I-01	Patient is ≥18 years old.	PASS
		Patient is at least 18 years old
I-02	Has locally advanced or metastatic cancer.	PASS
		Tumor stage III is considered locally advanced
		Tumor stage III is considered metastatic
I-04	Patient has confirmed activating PIK3CA mutation	PASS
		Activating mutation(s) detected in gene + PIK3CA: PIK3CA E545K

METC 05 - Dose expansion - monotherapy - NSCLC

Cohort ID A

Potentially eligible? Yes

Open for inclusion? Yes

Has slots available? Yes

Rule	Reference	Evaluation
I-03	Has non small cell lung cancer.	PASS
		Patient has tumor belonging to DOID
		term(s) lung non-small cell carcinoma

METC 05 - Dose expansion - monotherapy - Other cancer types

Cohort ID B
Potentially eligible? No
Open for inclusion? Yes
Has slots available? Yes

Has	s slots available? Yes	
Rule	Reference	Evaluation
I-03	Has solid cancer, other than non small cell lung cancer.	FAIL
		Patient has tumor belonging to DOID term(s) lung non-small cell carcinoma

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Other trials & cohorts

METC 02

Potentially eligible No

Acronym KAYRAS

Title A phase 1/2 trial for first in-human usage of KAYRAS, a new specific KRAS G12D inhibitor in NSCLC and

colorectal cancer

Rule	Reference	Evaluation
I-04	Patient has confirmed KRAS G12D mutation	FAIL

None of G12D detected in gene KRAS

METC 02 - Dose expansion - monotherapy - NSCLC

Cohort ID A

Potentially eligible? No

Open for inclusion? Yes

Has slots available? Yes

METC 02 - Dose expansion - monotherapy - Colorectum

Cohort ID B

Potentially eligible? No

Open for inclusion? Yes

Has slots available? Yes

Rule	Reference	Evaluation
I-02	KAYRAS monotherapy is indicated for the treatment of NSCLC or colorectal patients with	FAIL
	metastatic cancer.	Patient has no colorectal cancer

METC 03

Potentially eligible No

Acronym NO-SEE797ES

Title Phase I trial for development of NO-SEE797ES, a specific inhibitor for EGFR with C797 mutations but not C797S

in solid tumors

Rule	Reference	Evaluation
I-03	Tumors should contain EGFR C797 but not EGFR C797S mutation.	FAIL
		No variants in codon(s) C797 detected in gene EGFR

METC 03 - Dose escalation - monotherapy

Cohort ID A

Potentially eligible? No

Open for inclusion? Yes

Has slots available? Yes

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METC 04

Potentially eligible No
Acronym TEDR1

Title TEDR1 Trial: A phase II trial to evaluate efficacy of specific EGFR inhibitors in lung cancer

 Rule
 Reference
 Evaluation

 I-3
 Patient has a confirmed EGFR C797S mutation
 FAIL

None of C797S detected in gene EGFR

METC 04 - Lung cancer C797S cohort

Cohort ID A

Potentially eligible? No

Open for inclusion? Yes

Has slots available? Yes