PATIENT
EXAMPLE-LUNG-01
REPORT DATE
17-Sep-2025

Gender: Female | Birth year: 1975 | WHO: 1

Tumor: Lung adenocarcinoma | Lesions: Liver, Lung | Stage: IV

Summary

Clinical summary

Relevant systemic treatment history 6/2023-1/2025 Osimertinib

Relevant other oncological history

Previous primary tumor

None

Relevant non-oncological history 2023 Rheumatoid arthritis

Recent molecular results

Hartwig WGS (22-Feb-2025)

Biopsy location Lung (purity 50%)

Molecular tissue of origin prediction Lung: Non-small cell: LUAD (98%)

Tumor mutational load / burden TML 160 / TMB 14 mut/Mb

Microsatellite (in)stability Stable

HR status Proficient (0)

Driver mutations EGFR C797S, EGFR L858R, KRAS G12C, KRAS G12D

Amplified genes None

Deleted genes TP53

Homozygously disrupted genes None

Gene fusions MET(exon13)::MET(exon15) fusion

Virus None

Trial-relevant IHC results

PD-L1 Score > 50%

Trials in NL that are open and potentially eligible (5 cohorts from 5 trials)

Trial	Cohort	Molecular	Sites	Warnings
METC 04 TEDR1	Lung cancer C797S cohort	EGFR C797S	NKI-AvL	None
METC 02 KAYRAS	Dose expansion - monotherapy - NSCLC	KRAS G12D, PD-L1 >= 50.0	Erasmus MC	Variant(s) G12D in KRAS but subclonal likelihood of > 50%
METC 01 IEMOEN	Dose escalation - monotherapy (no slots)	None		Has not exhausted SOC (at least platinum doublet remaining)
EGFR-C797S- TRIAL	EGFR C797S	EGFR C797S	Elisabeth- TweeSteden Ziekenhuis	
EGFR-L858R- TRIAL	EGFR L858R	EGFR L858R	Elisabeth- TweeSteden Ziekenhuis	

Trials matched solely on molecular event and tumor type (no clinical data used) are shown in italicized, smaller font.

International trials that are open and potentially eligible (2 cohorts from 2 trials)

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Trial	Cohort	Molecular	Sites
EGFR-BE	EGFR L858R	EGFR L858R	Belgium: Brussels
KRAS-G12C-TRIAL-DE	KRAS G12C	KRAS G12C	Germany: Stuttgart

International trials are matched solely on molecular event and tumor type (clinical data excluded).

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Molecular Details

Hartwig WGS (EXAMPLE-LUNG-01-T, 22-Feb-2025)

General

Purity	Ploidy	TML Status	TMB Status	MS Stability	HR Status	DPYD	UGT1A1	
50%	2.3	High (160)	High (14)	Stable	Proficient (0)	*1_HOM (Normal function)	*1_HOM (Normal function)	
Predicted tum	nor origin							
				1. Lung: Non-sm	all cell: LUAD			
Combined prediction score			98%					
This score is c	This score is calculated by combining information on:							

60%

70%

80%

Other cohorts have a combined prediction of 2% or lower

(3) Driver genes and passenger characteristics

(2) SNV genomic localisation distribution

Key drivers

(1) SNV types

Туре	Driver	Trials (Locations)	Trials in Hartwig	Best evidence in External	Resistance in External
Mutation (gain of function)	EGFR L858R (2/4 copies)		NCT00000006, NCT00000007	Approved	
Mutation (gain of function)	EGFR C797S (1/4 copies)	TEDR1 (NKI-AvL)	NCT00000008	Pre-clinical	
Mutation (gain of function)	KRAS G12D (0.3/2 copies)*	KAYRAS (Erasmus MC)			
Mutation (gain of function)	KRAS G12C (0.3/2 copies)*		NCT00000009		
Deletion	TP53 del, 0 copies				

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External

External

Continued from the previous page

Type Driver Trials (Locations) Trials in Hartwig Best evidence in Resistance in External External Known fusion MET(exon13)::MET(exon15) fusion * Variant has > 50% likelihood of being sub-clonal Other drivers or relevant events Type Driver Trials (Locations) Trials in Hartwig Best evidence in Resistance in

IHC results

None

PD-L1 Score > 50%

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Relevant other oncological history None

Previous primary tumor None

Relevant non-oncological history 2023 Rheumatoid arthritis

Patient current details (20-Feb-2025)

Unresolved toxicities grade => 2

LVEF

50%

Cancer-related complications

Known allergies

None

Recent surgeries 01-Aug-2024 Cholecystectomy

Tumor details (20-Feb-2025)

Measurable disease Yes

Known lesions Liver, Lung
Unknown lesions None

No lesions present CNS, Brain, Bone, Lymph node

Active medication details

Medication	Administration route	Start date	Stop date	Dosage	Frequency
St. John's Wort	Oral	01-Feb-2023		300 MILLIGRAMS	1 / 2 DAYS

Blood transfusions

Product	Date
ERTHROCYTES_FILTERED	20-Sep-2024

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Other Trial Matching Results

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

Trial	Cohort	Molecular	Sites	Warnings
METC 01	Dose expansion - monotherapy	None		Has not exhausted SOC (at least platinum doublet remaining)

Trials and cohorts that are considered ineligible (4)

Trial	Cohort	Molecular	Ineligibility reasons
METC 03 NO-SEE797ES	Dose escalation - monotherapy	EGFR C797S	C797S in EGFR in canonical transcript
METC 02 KAYRAS	Dose expansion - monotherapy - Colorectum	KRAS G12D, PD-L1 >= 50.0	No colorectal cancer
METC 05 PICKME3CA	Applies to all cohorts below	None	No PIK3CA activating mutation(s)
	Dose expansion - monotherapy - NSCLC (closed)		
	Dose expansion - monotherapy - Other cancer types (closed)		Tumor belongs to DOID term(s) lung non-small cell carcinoma

Trials and cohorts that are not evaluable or ignored (0)

Trial	Cohort	Molecular	Sites	Configuration
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
Reference	Evaluation
I-03	WARN
	Has not exhausted SOC (at least platinum doublet remaining)
E-02	UNDETERMINED
	No measurement found for hemoglobin
E-03	UNDETERMINED
	No measurement found for absolute neutrophil count
E-01	PASS
	Has no other condition belonging to category autoimmune disease
I-01	PASS
	Patient is at least 18 years old
I-02	PASS
	Has solid primary tumor
	Stage IV is considered metastatic

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 02

Potentially eligible	Yes
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
Reference	Evaluation
I-04	WARN
	Variant(s) G12D in KRAS but subclonal likelihood of > 50%
I-03	UNDETERMINED
	ASAT and ALAT are not present or cannot be evaluated
I-01	PASS
	Patient is at least 18 years old
I-02	PASS
	Stage IV is considered metastatic
I-05	PASS
	PD-L1 expression above minimum of 50.0

METC 02 - Dose expansion - monotherapy - NSCLC

Cohort ID	A
Potentially eligible?	Yes
Open for inclusion?	Yes
Has slots available?	Yes
Reference	Evaluation
I-02	PASS
	Tumor belongs to DOID term(s) lung non-small cell carcinoma

METC 02 - Dose expansion - monotherapy - Colorectum

No colorectal cancer

Cohort ID	В
Potentially eligible?	No
Open for inclusion?	Yes
Has slots available?	Yes
Reference	Evaluation
I-02	FAIL

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METC 04

Potentially eligible	Yes
Acronym	TEDR1
Title	TEDR1 Trial: A phase II trial to evaluate efficacy of specific EGFR inhibitors in lung cancer
Reference	Evaluation
I-1	PASS
	Patient is at least 18 years old
I-2	PASS
	Stage IV is considered metastatic
	Tumor belongs to DOID term(s) lung cancer
I-3	PASS
	C797S in EGFR in canonical transcript

METC 04 - Lung cancer C797S cohort

Cohort ID	Α
Potentially eligible?	Yes
Open for inclusion?	Yes
Has slots available?	Yes

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Other trials & cohorts

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

Reference Evaluation

I-03 FAIL

C797S in EGFR in canonical transcript

METC 03 - Dose escalation - monotherapy

Cohort ID A

Potentially eligible? No

Open for inclusion? Yes

Has slots available? Yes

METC 05

Potentially eligible No

Acronym PICKME3CA

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

Reference Evaluation

I-04 FAIL

No PIK3CA activating mutation(s)

METC 05 - Dose expansion - monotherapy - NSCLC

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

METC 05 - Dose expansion - monotherapy - Other cancer types

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

Reference Evaluation

I-03 FAIL

Tumor belongs to DOID term(s) lung non-small cell carcinoma