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Sample Information

Patient Name: 張家恩 Gender: Male ID No.: P120343983 History No.: 41643612

Age: 53

Ordering Doctor: DOC1326K 簡聖軒

Ordering REQ.: H489M84 Signing in Date: 2023/11/10

Path No.: M112-00289 MP No.: MY23071

Assay: Oncomine Myeloid Assay **Sample Type:** Bone Marrow

Bone Marrow Aspirating Date: 2023/11/02

Reporting Doctor: DOC5444B 楊靜芬 (Phone: 8#5444)

Note:

Sample Cancer Type: Acute Myeloid Leukemia

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Report Highlights

0 Clinical Trials

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Relevant Biomarkers

Tier	Genomic Alteration	Relevant Therapies (In this cancer type)	Relevant Therapies (In other cancer type)	Clinical Trials
IA	IDH2 p.(R172K) c.515G>A isocitrate dehydrogenase (NADP(+)) 2 Allele Frequency: 21.70%	enasidenib ¹ azacitidine decitabine enasidenib + chemotherapy venetoclax + chemotherapy	None	0

Public data sources included in relevant therapies: FDA1, NCCN, EMA2, ESMO

Variants (Exclude variant in Taiwan BioBank with >1% allele frequency)

DNA Sequence variants								
Allele								
Gene	Amino Acid Change	Coding	Variant ID	Locus	Frequency	Transcript	Variant Effect	Coverage
IDH2	p.(R172K)	c.515G>A	COSM33733	chr15:90631838	21.70%	NM_002168.4	missense	1995

Biomarker Descriptions

IDH2 (isocitrate dehydrogenase (NADP(+)) 2)

Background: The IDH1 and IDH2 genes encode homologous isocitrate dehydrogenase enzymes that catalyze the conversion of isocitrate to α-ketoglutarate (α-KG) 1 . The IDH1 gene encodes the NADP+ dependent cytoplasmic isocitrate dehydrogenase enzyme; IDH2 encodes the mitochondrial isoform.

Alterations and prevalence: Recurrent somatic mutations in IDH1 and IDH2 are mutually exclusive and observed in several malignancies including glioma, chondrosarcoma, intrahepatic cholangiocarcinoma, acute myeloid leukemia (AML), and myelodysplastic syndrome (MDS)². Recurrent IDH2 variants include predominately R140Q and R172K plus other substitutions at lower frequencies. These gain of function variants confer neomorphic enzyme activity³. Although wild-type enzymatic activity is ablated, recurrent IDH2 variants catalyze the conversion of α -KG to D-2-hydroxyglutarate, an oncometabolite with diverse effects on cellular metabolism, epigenetic regulation, redox states, and DNA repair¹.4. Recurrent IDH2 mutations are present in 10-20% of patients with AML and 5% of patients with MDS⁵.6.7.

Potential relevance: Enasidenib⁸ is FDA approved (2017) for the treatment of AML patients with IDH2 R140G/L/Q/W and R172G/K/M/S/W mutations. In AML, acquired resistance to enasidenib has been associated with the emergence of Q316E or I319M mutations⁹. IDH2 R172 and R140Q variants are associated with poor prognosis in MDS but have been shown to confer improved prognosis in lower grade gliomas^{10,11,12}. Additionally, IDH2 mutations are associated with inferior overall survival in polycythemia vera (PV) and essential thrombocythemia (ET) as well as inferior leukemia-free survival in primary myelofibrosis (PMF)¹³. Mutations in IDH2 are diagnostic of astrocytoma IDH-mutant and oligodendroglioma IDH-mutant and 1p/19q-codeleted subtypes of central nervous system (CNS) tumors¹⁴

Relevant Therapy Summary

In this cancer type In oth	er cancer type	In this cancer type and other cancer types	×	No evidence
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IDH2 p.(R172K) c.515G>A					
Relevant Therapy	FDA	NCCN	ЕМА	ESMO	Clinical Trials*
enasidenib	•		×		×
azacitidine	×	•	×	×	×
decitabine	×	•	×	×	×
enasidenib + azacitidine	×	•	×	×	×
venetoclax + azacitidine	×	•	×	×	×
venetoclax + cytarabine	×	•	×	×	×
venetoclax + decitabine	×	•	×	×	×

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Relevant Therapy Details

Current FDA Information

In this cancer type

O In other cancer type

In this cancer type and other cancer types

FDA information is current as of 2023-09-13. For the most up-to-date information, search www.fda.gov.

IDH2 p.(R172K) c.515G>A

enasidenib

Cancer type: Acute Myeloid Leukemia Label as of: 2020-11-24 Variant class: IDH2 R172K mutation

Indications and usage:

IDHIFA® is an isocitrate dehydrogenase-2 inhibitor indicated for the treatment of adult patients with relapsed or refractory acute myeloid leukemia (AML) with an isocitrate dehydrogenase-2 (IDH2) mutation as detected by an FDA-approved test.

Reference:

https://www.accessdata.fda.gov/drugsatfda_docs/label/2020/209606s004lbl.pdf

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Current NCCN Information

In this cancer type

O In other cancer type

In this cancer type and other cancer types

NCCN information is current as of 2023-09-01. For the most up-to-date information, search www.nccn.org. For NCCN International Adaptations & Translations, search www.nccn.org/global/what-we-do/international-adaptations.

Some variant specific evidence in this report may be associated with a broader set of alterations from the NCCN Guidelines. Specific variants listed in this report were sourced from approved therapies or scientific literature. These therapeutic options are appropriate for certain population segments with cancer. Refer to the NCCN Guidelines® for full recommendation.

IDH2 p.(R172K) c.515G>A

venetoclax + azacitidine

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 R172 mutation

NCCN Recommendation category: 1

Population segment (Line of therapy):

(Induction therapy); Preferred intervention

Reference: NCCN Guidelines® - NCCN-Acute Myeloid Leukemia [Version 4.2023]

enasidenib

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 R172 mutation

NCCN Recommendation category: 2A

Population segment (Line of therapy):

Relapsed, Refractory (Line of therapy not specified)

Reference: NCCN Guidelines® - NCCN-Acute Myeloid Leukemia [Version 4.2023]

venetoclax + decitabine

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 R172 mutation

NCCN Recommendation category: 2A

Population segment (Line of therapy):

(Induction therapy); Preferred intervention

Reference: NCCN Guidelines® - NCCN-Acute Myeloid Leukemia [Version 4.2023]

azacitidine

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 mutation

NCCN Recommendation category: 2A

Population segment (Line of therapy):

(Induction therapy); Other recommended intervention

Reference: NCCN Guidelines® - NCCN-Acute Myeloid Leukemia [Version 4.2023]

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IDH2 p.(R172K) c.515G>A (continued)

decitabine

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 mutation

NCCN Recommendation category: 2A

Population segment (Line of therapy):

(Induction therapy); Other recommended intervention

Reference: NCCN Guidelines® - NCCN-Acute Myeloid Leukemia [Version 4.2023]

enasidenib

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 mutation

NCCN Recommendation category: 2A

Population segment (Line of therapy):

(Induction therapy); Preferred intervention

Reference: NCCN Guidelines® - NCCN-Acute Myeloid Leukemia [Version 4.2023]

venetoclax + cytarabine

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 mutation

NCCN Recommendation category: 2A

Population segment (Line of therapy):

■ (Induction therapy); Other recommended intervention

Reference: NCCN Guidelines® - NCCN-Acute Myeloid Leukemia [Version 4.2023]

enasidenib + azacitidine

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 mutation

NCCN Recommendation category: 2B

Population segment (Line of therapy):

(Induction therapy); Useful in certain circumstances

Reference: NCCN Guidelines® - NCCN-Acute Myeloid Leukemia [Version 4.2023]

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Current ESMO Information

■ In this cancer type
O In other cancer type
In this cancer type and other cancer types

ESMO information is current as of 2023-09-01. For the most up-to-date information, search www.esmo.org.

IDH2 p.(R172K) c.515G>A

enasidenib

Cancer type: Acute Myeloid Leukemia Variant class: IDH2 mutation

ESMO Level of Evidence/Grade of Recommendation: IV / B

Population segment (Line of therapy):

■ Relapsed, Refractory (Second-line therapy)

Reference: ESMO Clinical Practice Guidelines - ESMO-Acute Myeloblastic Leukaemia in Adult Patients [Ann Oncol (2020); 31(6): 697-712.]

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