Mathematical analysis predicts imbalanced IDH1/2 expression associates with 2-HG-inactivating β-oxygenation pathway in colorectal cancer

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Abstract. Bioinformatics and computational modeling offer innovative approaches to investigate cancer metabolism and predict the secondary and tertiary cellular responses. Dysregulation of metabolism has also been implicated in the pathophysiology of cancer. A significant proportion of patients with glioblastoma and hematological malignancies harbor the mutated forms of the oxidative phosphorylation (OxPhos) enzymes, isocitrate dehydrogenase (IDH) 1 or 2. The mutated forms of IDH1 and IDH2 produce an oncogenic metabolite, D-2-hydroxyglutarate (D2HG). A recent study of breast cancer patients showed that D2HG can also be produced in the absence of mutated IDH, through an alternative route involving over-activated MYC signaling. We developed a novel methodology to computationally analyze gene expression in colorectal cancer (CRC), and identified novel sets of genes that are associated with patient survival. The study of OxPhos-related genes revealed that an imbalance between the expression of IDH1 and IDH2, defined as overexpression of one isoform in relation to the other, was associated with worse prognosis in CRC patients. This effect was further accentuated by reduced expression of the β-oxygenation enzyme, 3-D-hydroxyacyl-CoA dehydratase (HCDH) 4, which has

been reported to contribute to metabolism of intracellular D2HG. The present computational analysis revealed a novel and potential mechanism of CRC development, through over-production of D2HG when there is an imbalance between IDH1 and IDH2 expression, resulting in decreased clearance of D2HG when the β-oxidization pathway is diminished. Additional validation analysis with another gene expression dataset resulted in IDH1/2 imbalanced expression with a shorter DFS compared with balanced expression. Altogether, these findings provide a strong rationale for studying this mechanism further in order to discover novel therapeutic targets for the treatment of CRC.

Introduction

In the last few decades, there has been an exponential rise of the available biological data including human genomic sequences, gene expression profiles, protein-protein interaction networks, and metabolomic data of physiologically active compounds. This vast amount of information provides us with a challenging opportunity to develop computational approaches for systematic analysis of various disorders, including cancers, in order to utilize the biomedical data for prediction of patient prognosis, disease modeling and biological systems analysis (1). The genetic diversity in cancer is vast, given cancers can arise from most of the cells in the body ($\sim 6 \times 10^{13}$ cells), and that tumors themselves also harbor heterogeneous cellular components ($\sim 10^{10}$ cells), including a subpopulation of cancer initiating cells ($\sim 10^{8}$ cells) (2,3). The gene expression between cancer cells is also highly variable, this variability being generated through numerous mechanisms, including chromosomal translocations, genetic mutations, epigenetic chromatin remodeling of the 3×10^{9} base paired DNA and alteration of histone structures (4-6). For example, sequencing analysis of liver cancer revealed that there are 11×10^{3} genetic mutations, of which 10-100 are driver mutations, as well as 21 chromosomal structural abnormalities (7). Genetic heterogeneity occurs in many tumors (http://cancergenome.nih.gov), indicating

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that mathematical and statistical analysis is indispensable to distinguish between driver and passenger mutations in order to identify druggable targets.

In 2008 and 2009, two independent cancer sequencing projects reported high frequencies of mutations of the gene encoding the cytosolic enzyme IDH1, a critical component of the tricarboxylic acid (TCA) cycle, in glioblastoma multiforme (GBM) (8) and acute myeloid leukemia (AML) (9). Further study reported that IDH2, the gene encoding the homologous enzyme in the mitochondria was also frequently mutated in GBM patients (10). Studies indicated that >75% of grade 2 and 3 GBM and 20% of AML harbor mutations of IDH1 at R132, or IDH2 at the homologous R172 residue (11). These oncogenic mutations in IDH1 and IDH2 enzymes reduce their native activity, and generate neomorphic activity that converts α-ketoglutarate (α-KG; also known as 2-oxyoglutarate) to D-2-hydroxygutarate (D2HG) (12,13). D2HG is an oncometabolite that can cause changes to the epigenetic landscape by inhibiting the activities of iron (II), α-KG-dependent dioxygenases, including: prolyl-hydroxylase-domains (PHDs) resulting in pseudo-hypoxic environment and activation of hypoxia-inducible-factor (HIF) pathway leading to aberrant cellular proliferation; ten-eleven translocation (TET)-family demethylases, AlkB-family dioxygenases, which protect nucleotides against methylating reactions by directly dealkylating bases (1mA, 3mC, 1mG and 3mT); and histone lysine demethylases (KDMs) (14).

Other oncometabolites that have been identified are succinate and fumarate, which are generated by mutations of succinate dehydrogenase (SDH) and fumarate hydratase (FH), respectively. These also competitively inhibit α-KG-dependent dioxygenases (14), and moreover provide a strong rationale that TCA enzymes as a family may be important in other malignancies.

Here we applied computational analysis to study the association between gene expression profiles of colorectal cancer (CRC) patients and their prognosis. Using this approach, we identified that the imbalances between the expressions of IDH1 and IHD2 was associated with poorer prognosis of patients with CRC. Subsequent unsupervised analysis of targeted genes identified low expression of hydroxyl-CoA dehydratase (HCDH) 4 to amplify the effect of imbalanced IDH1 and IDH2 expressions on the survival of CRC patients. HCDH4 is an enzyme that metabolizes D2HG in the β-oxygenation pathway, and is encoded by a gene that is located at chromosome 9p21.3, which is frequently deleted in cancers alongside tumor suppressor INK4A and microRNA-31 (miR-31). The present computational analysis linked the association combining expression of IDH1 and IDH2, rather than mutations, with β-oxidization pathway in the prognosis of CRC patients.

Materials and methods

Gene Expression database of colorectal cancer patients. We used the published GSE17536 database (http://www.ncbi.nlm. nih.gov/geo/query/acc.cgi?acc=GSE17536) (15) from the Gene Expression Omnibus in NCBI to analyze the effects of gene expression on disease-free survival (DFS) and overall survival (OS) of CRC patients. This database contains the microarray data of 177 patients, generated using the Affymetrix Human

Genome U133 Plus 2.0 Array. The microarray data were generated with multiple probes for one gene in some cases; if this was the case, the probe that demonstrated the widest variance of genetic expression amongst the cohort was selected for statistical analysis. Each selected gene was divided into low and high expression groups at the median point of expression. Additionally, to verify the relevance of our results with GSE17536 database, we used another reference dataset for CRC patients (16). The 91 patients with both CRC stage and DFS time information were selected from this reference dataset and used in validation analysis.

Analysis of the expression of genes of the oxidative phosphorylation pathway and their effect on the survival of colorectal cancer patients. Genes of the oxidative phosphorylation (OxPhos) pathway that are already known to be involved in neoplastic diseases were selected for initial analysis: TET 1-3 proteins, IDH1, IDH2, SDH, FH, MYC, glutaminase (GLS), and the mitochondrial alcohol dehydrogenase iron-containing 1 (ADHFE) (14). Genes in the above group were analyzed in all possible combinations as pairs in order to ascertain whether there were any associations between combined gene expressions and DFS or OS (Fig. 1). These analyses were conducted through the generation of Kaplan-Meier curves. Thirty-two patients were excluded from the analysis concerning the DFS given these were stated as being zero.

Screening for genes affecting the survival of colorectal cancer patients in a manner dependent on the expression of oxidative phosphorylation genes. Other genes that are associated with DFS and OS of CRC patients in a manner that is dependent on the selected genes of the OxPhos pathway were screened. The rationale for adopting this approach was to identify a biological pathway important in CRC, instead of emphasizing isolated gene expression. Only CRC patients with stages II or III were selected for this stage of analysis. Unbiased screening was performed for 'the third genes', searching for genes that do not have independent association with DFS or OS, but were nevertheless able to accentuate the effect of the selected OxPhos pathway genes on the DFS and OS (Fig. 1). Statistical analysis was performed; with statistical significance being defined when the p-value was <0.05. Kaplan-Meier curves were generated using the survival package (16) on R version 3.0.2 (17).

Results

Imbalance between the IDH1 and IDH2 expression is associated with decreased survival of colorectal cancer patients. The TCA cycle in mitochondria plays a critical role in OxPhos, which produces ATP via the electron transport chain (14). Given mutations of several of the OxPhos-related genes, such as families of TET proteins, IDH, SDH, and FH, are observed in human malignancies, we investigated whether the expression of these genes in combination as pairs would be associated with patient survival in CRC. Alterations of families of TETs, IDHs, SDHs, and FHs are expected to result in attenuation of α-KG-dependent dioxygenases, such as demethylases, which characterize cancers (14). There were no differences in the Kaplan-Meier curves for DFS or OS, when IDH1 or IDH2
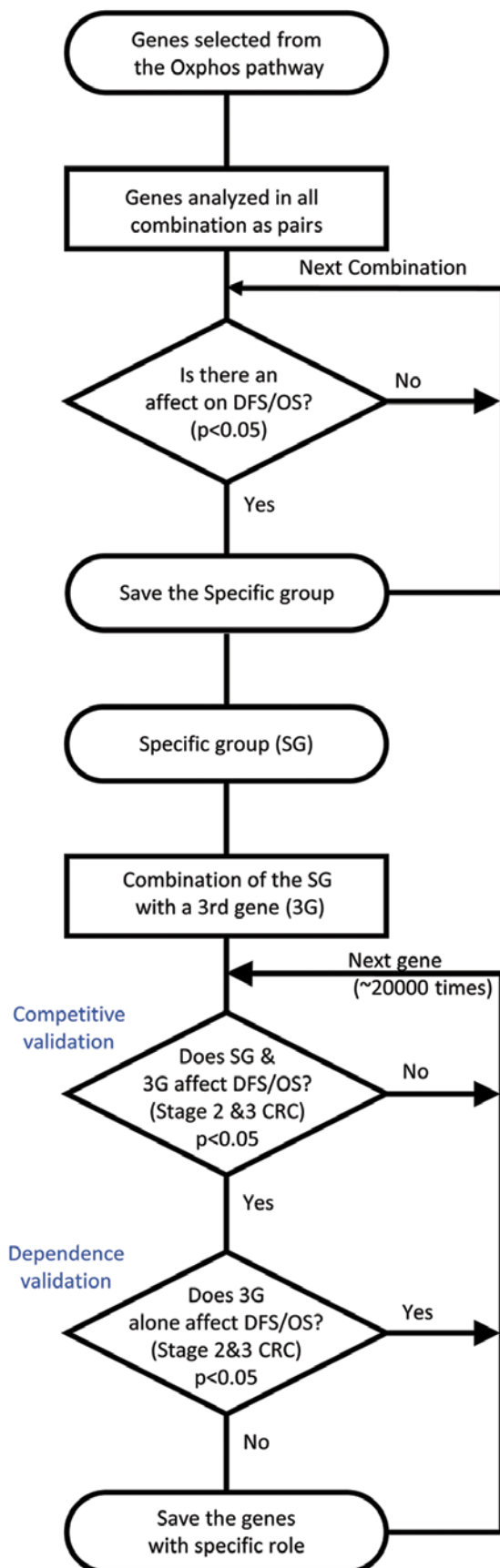
Figure 1. Flow chart of the computational analysis. First flow, pre-selected genes involved with the TCA cycle were analyzed in different combinations as pairs to identify gene combinations that were associated with reduced DFS/OS in CRC. Second flow, the search for 'third genes' that are associated with DFS and OS in a manner that is dependent on the above TCA cycle-related gene combinations.

expressions were analyzed in isolation. However, when the gene expression of IDH1 and IDH2 was analyzed together, the patients with an imbalance of IDH1 and IDH2 gene expression (i.e., IDH1^{high};IDH2^{low}, or IDH1^{low};IDH2^{high}), had a shorter DFS and OS compared to patients with a 'balanced' IDH gene expression (i.e., IDH1^{high};IDH2^{high}, or IDH1^{low};IDH2^{low}) (Fig. 2A and B).

We performed the same prognosis analysis for cancer recurrence with another reference dataset (16) to confirm that this biological behavior is not unique to the GSE17536 database. Also the validation results showed the same tendency that the patients with imbalanced expression of IDH1/2 had a shorter DFS compared to those with balanced expression (Fig. 2C), although this reference dataset is a small one, which has only 10 patients with relapsed cancer.

Reduced HCDH4 expression exacerbates the poor prognosis in colorectal cancer patients with imbalanced IDH expression. In order to identify novel partners of imbalanced IDH1 and IDH2 expression, we performed non-biased comparison against the total transcriptome data. Analysis was conducted in two steps, the first being to identify candidate genes that could separate the IDH1^{high}:IDH2^{low} and IDH1^{low}:IDH2^{high} patients from IDH1^{high}:IDH2^{high} and IDH1^{low}:IDH2^{low} patients in terms of the DFS and OS. Genes identified using this analysis were also checked for their ability to independently predict DFS or OS depending on their expression, and was excluded if this was the case. By using this approach, we identified 43 and 44 genes that had a significant effect on the DFS and OS, respectively (Table I). Of these ZNF91 (18,19) and HCDH4 [protein tyrosine phosphatase-like a domain containing 2 (PTPLAD2)] (20), were found to accentuate the reduction of both DFS and OS of CRC patients when IDH1 and IDH2 expression was imbalanced.

When the above analysis was conducted further for individual stages of CRC, reduced expression of HCDH4 was shown to exacerbate the poorer DFS and OS in CRC patients with stages II or III disease (Fig. 3A). However, ZNF91 expression showed discordant effect on DFS and OS between different stages of CRC, with high expression associated with worse prognosis in stage II disease, but low expression is associated with worse prognosis in stage III disease (Fig. 3B). This result indicates that HCDH4 was more likely to be a suitable candidate for further analysis. HCDH4 is an enzyme that metabolizes D2HG in the β-oxygenation pathway. The genome data-base (http://www.ncbi.nlm.nih.gov/guide/genomes-maps/) shows that the HCDH4 gene is located at chromosome 9p21.3, which is a genomic region frequently deleted in cancer, also containing the tumor suppressor gene p16/CDKN2A/INK4A. p16/CDKN2A/INK4A is inactivated by methylation of the gene promoter in early stages of gastrointestinal cancer (21). The gene locus containing p16/CDKN2A/INK4A and HCDH4 also harbors interferon genes and cancer-associated miR-31 (22), suggesting the alterations of this locus increases the susceptibility to cancer (Fig. 4A).

Prediction of a common pathway of IDH1, IDH2 and HCDH4 genes. Although the IDH expression changes analyzed in this study are likely to be associated with wild-type genes, we propose that an imbalance between the expressions of IDH1
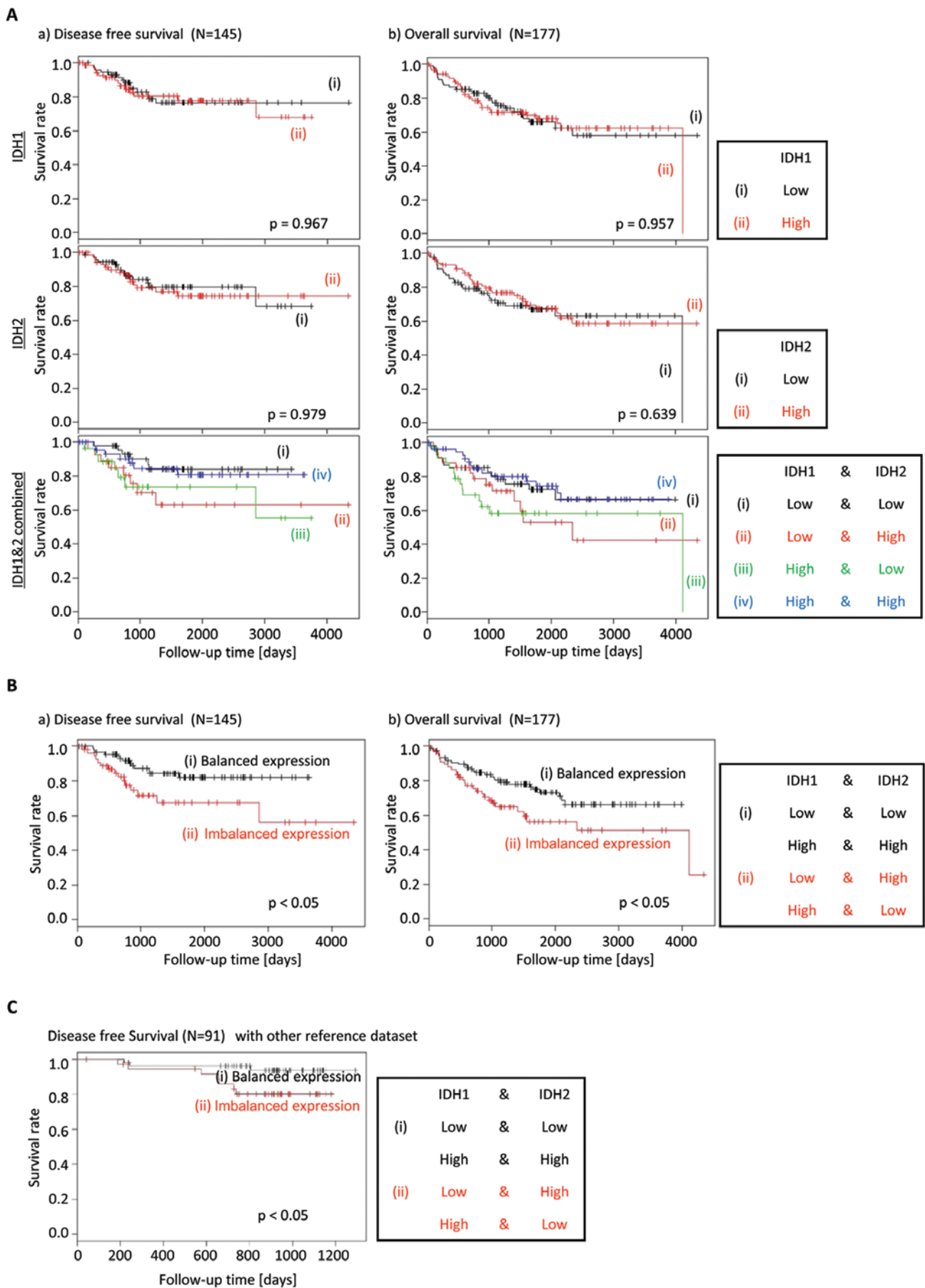
Figure 2. Expressions of IDH1 and IDH2 plotted against survival time. (A) Kaplan-Meier curves for IDH1, IDH2, and their combination with DFS and OS. (B) Kaplan-Meier curves for expression of IDH1 and IDH2 divided into balanced and imbalanced groups with DFS and OS. (C) Kaplan-Meier curve for combination of IDH1 and IDH2 with DFS time in the reference dataset (16).

Table I. Genes that have a significant effect on the DFS and OS, in a manner dependent on the imbalance between the expressions of IDH1 and IDH2.[a]

	DFS time			OS time	
Gene name	Stage 2	Stage 3	Gene name	Stage 2	Stage 3
AIM2	0.049	0.032	ADH5	0.029	0.000
ANKRD49	0.042	0.011	AK7	0.023	0.002
ARHGEF19	0.006	0.044	ANKRD49	0.029	0.007
C16orf73	0.042	0.000	ANKS1B	0.042	0.008
C20orf72	0.021	0.028	ANXA9	0.041	0.005
C2orf73	0.021	0.012	ATP5F1	0.029	0.014
C7	0.006	0.024	BMP2K	0.028	0.008
CA11	0.031	0.018	CHCHD2	0.034	0.010
CDH8	0.031	0.006	EMX2	0.023	0.023
CHEK2	0.021	0.039	FAM118B	0.016	0.011
DIO2	0.042	0.049	FAM124B	0.008	0.033
FAM54B	0.012	0.025	FAM54B	0.008	0.007
KIAA0355	0.042	0.028	FDX1	0.010	0.003
LELP1	0.031	0.049	GNN	0.023	0.037
LOC100129335	0.042	0.045	GOLT1A	0.029	0.042
LOC284379	0.031	0.005	GPR87	0.013	0.004
LOC388789	0.031	0.011	LOC100132354	0.042	0.023
LOC728804	0.014	0.009	LOC100132356	0.023	0.000
NCKAP5L	0.021	0.048	LOC100289086	0.034	0.046
OSBPL9	0.031	0.012	LOC389043	0.008	0.014
PHLDB3	0.014	0.039	LOC729420	0.034	0.037
POP4	0.042	0.002	LRRK1	0.029	0.025
PTP4A3	0.012	0.007	NINJ1	0.029	0.032
HCDH4	0.031	0.000	NR1H4	0.013	0.043
RPF2	0.021	0.035	OXA1L	0.023	0.036
SCN4B	0.021	0.045	PCMT1	0.029	0.015
SEC13	0.031	0.046	PEX16	0.029	0.033
SLC16A2	0.049	0.025	PRR13	0.029	0.003
SLC9A1	0.031	0.041	PTP4A3	0.023	0.041
SNTA1	0.021	0.025	**HCDH4**	0.029	0.002
STK25	0.012	0.022	RPS3	0.023	0.044
TBC1D19	0.014	0.037	SIX5	0.010	0.016
TGFBRAP1	0.031	0.004	SLC24A2	0.029	0.041
TGM6	0.042	0.029	SLITRK4	0.028	0.003
TLR1	0.042	0.035	SMARCAL1	0.034	0.029
UCHL3	0.042	0.044	TGFBRAP1	0.034	0.001
ULBP2	0.021	0.044	TMSB15B	0.029	0.020
WWP1	0.021	0.032	TRIM48	0.029	0.022
ZDHHC16	0.049	0.030	TUFM	0.029	0.029
ZNF142	0.042	0.044	UCHL3	0.034	0.016
ZNF573	0.021	0.004	WIPI1	0.029	0.033
ZNF814	0.042	0.015	YLPM1	0.018	0.004
ZNF91	0.042	0.015	ZNF362	0.010	0.040
-	-	-	**ZNF91**	0.029	0.011

[a]P-values between the combined imbalanced IDH1/2 group with high and low groups of the third gene expression in competitive analyses. Some genes observed in common with the DFS and OS results are written in bold.
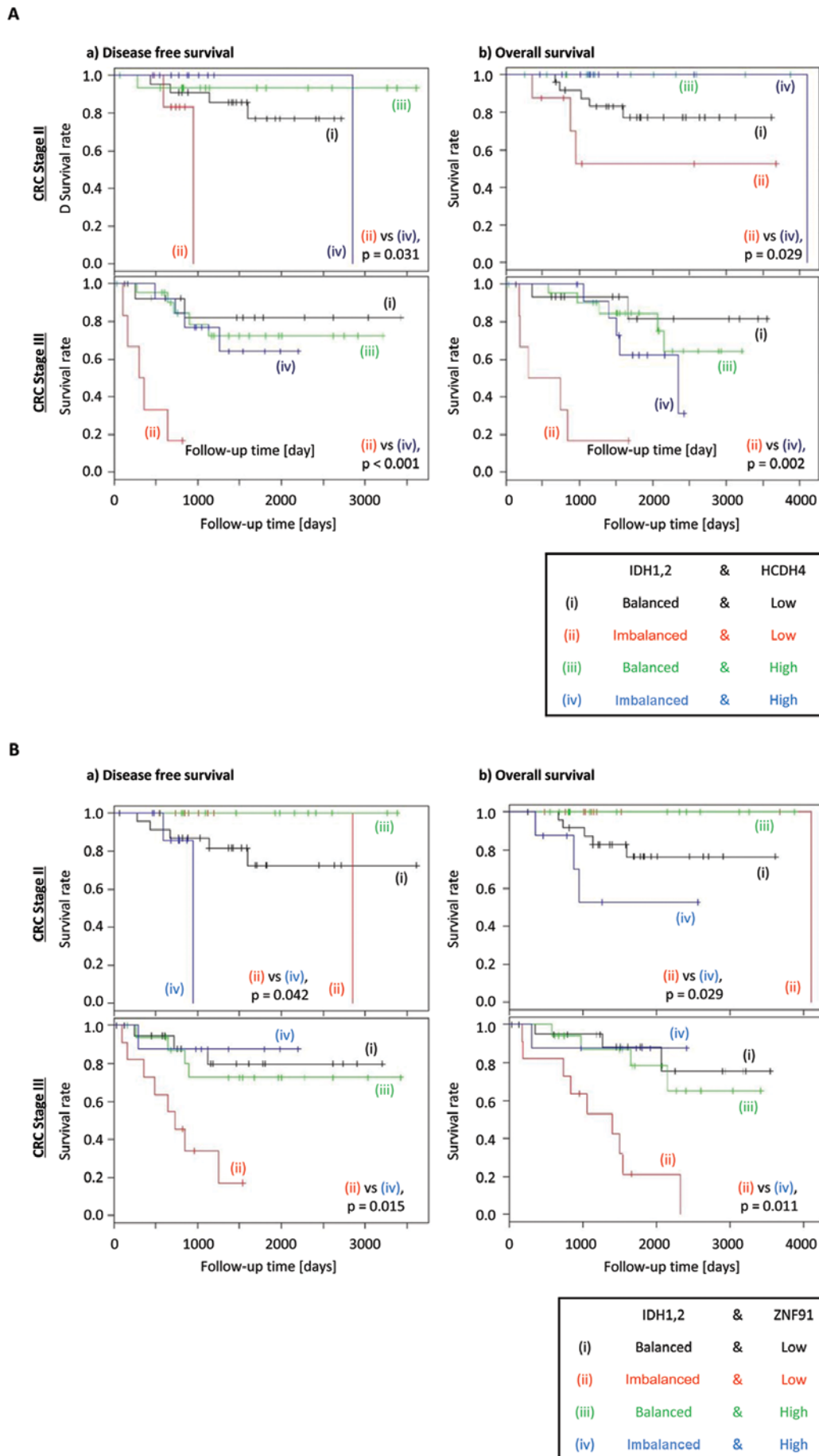
Figure 3. Kaplan-Meier curves for the combination of (A) HCDH4 and (B) ZNF91 with imbalanced expression of IDH1 and IDH2 using DFS and OS for CRC patients with stages II or III disease.
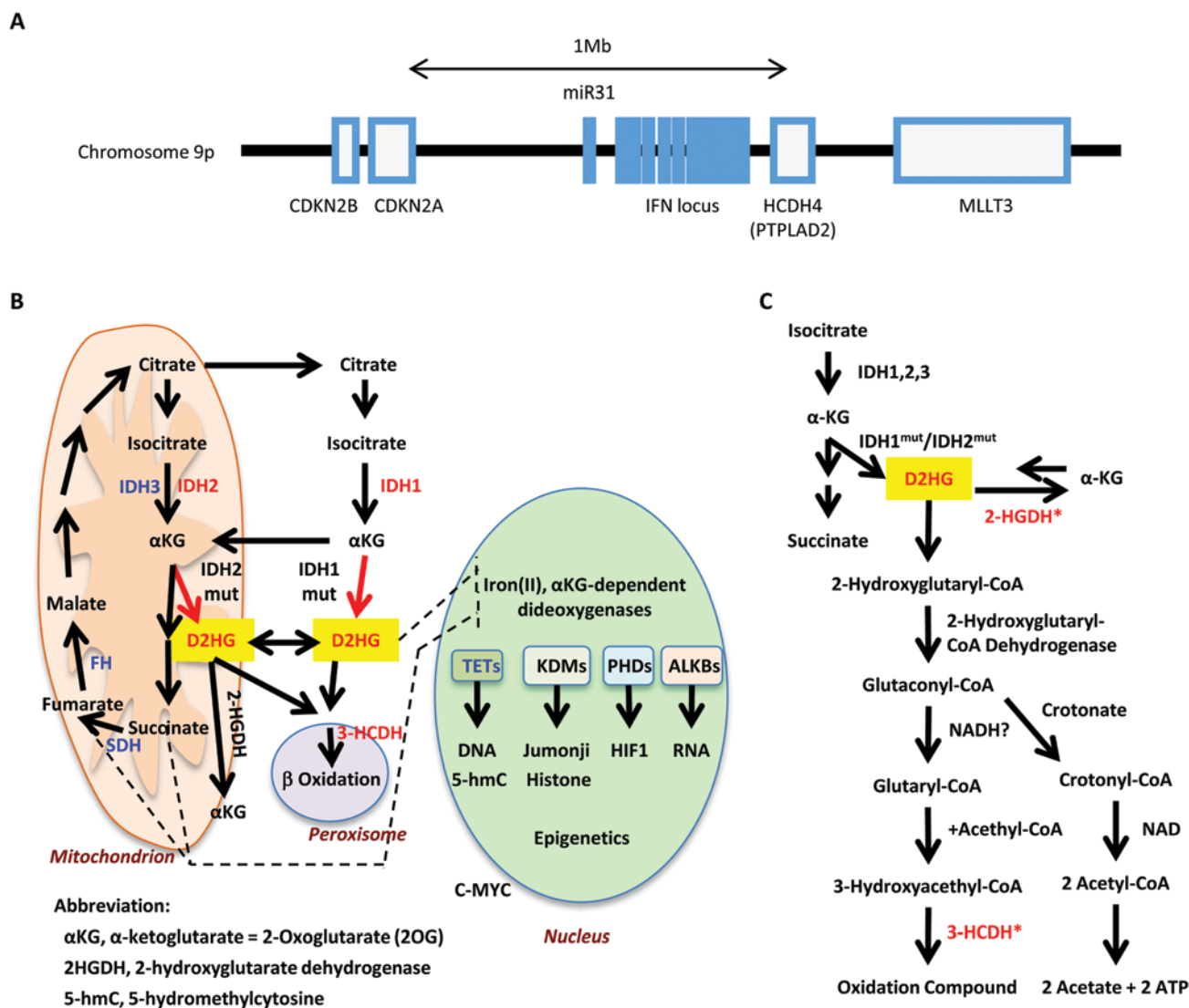
Figure 4. (A) The region of the genome encoding HCDH4, also contains CDKN2 and miR 31. (B) Schematic diagram of the TCA cycle and the origin of D2HG production, as well as the downstream effects of D2HG on the iron (II), α-KG-dependent di-deoxygenases. (C) The metabolic pathway of D2HG involving 2-HGDC or HCDH.

and IDH2 could lead to the overproduction of D2HG, of which more will be explained in Discussion. Increasing the levels of the oncometabolite D2HG is potentially a common denominator of IDH1, IDH2 and HCDH4, with the decreased expression of the latter enzyme leading to reduced metabolism of D2HG. Therefore, D2HG provides an explanation for poorer survival of patients with imbalanced IDH1 and IDH2 as well as low HCDH4 expression. Previous studies have shown that HCDH is involved with the metabolization of D2HG in the glutaconyl pathway (http://www.genome.jp/kegg/) (Fig. 4). The direction of chemical reaction generally depends on the ΔG value instead of the activation energy. Given that the conversion from D2HG to α-KG is a relatively small exothermic reaction of 0.791 (kcal/mol) (calculated with MP2/6-31G level in Gaussian 09 package) (23), this suggests that D2HG is produced under physiological conditions, with IDH1 and IDH2 expression determining rate of production and D2HGDH and HCDH governing metabolization (Figs. 4C and 6A).

MYC overexpression is not associated with decreased survival of colorectal cancer patients. Recently, overexpression of the MYC oncogene occurring in a sub-group of breast cancer patients was shown to be associated with higher levels of D2HG and decreased survival. D2HG production was demonstrated to be driven by MYC overexpression *in vitro*, demonstrating a novel mechanism of D2HG production given IDH1 or IDH2 mutations in breast cancers are rare (24). In this cohort of CRC patients, MYC expression was not associated with decreased DFS or OS. Interestingly, the Kaplan-Meier analysis showed that patients with high MYC expression have better DFS or OS, regardless of the expression status of FH, IDH2 and SDHB (Fig. 5), suggesting that the role of MYC in CRC is distinct from that in breast cancers.

Discussion

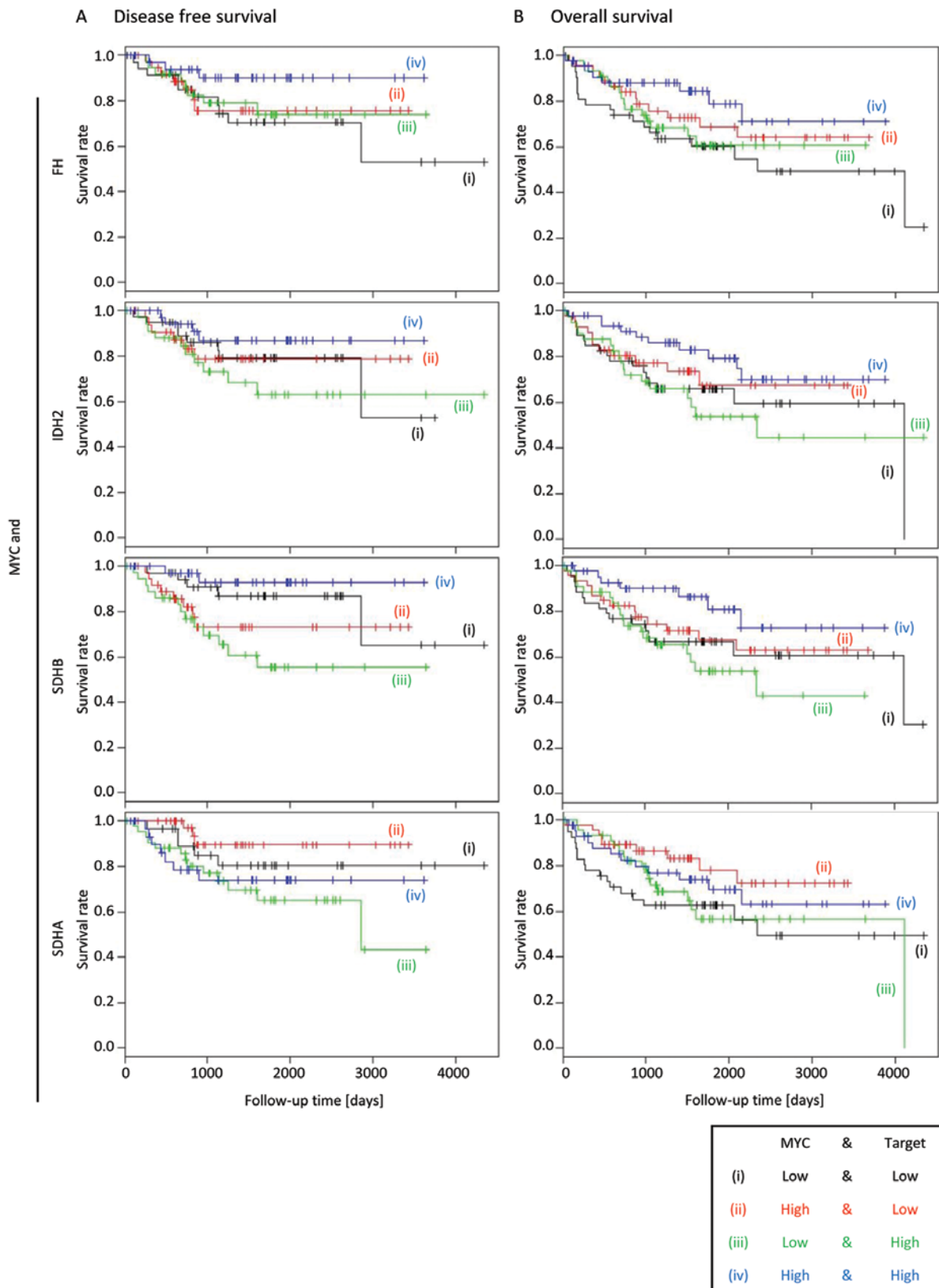In this study, an imbalance between the expression of IDH1 and IDH2 was associated with decreased DFS and OS in

Figure 5. Kaplan-Meier curve for the combination of FH, IDH2 SDHB or SDHA with MYC expression using DFS and OS.

CRC patients; these reductions in DFS and OS were further accentuated by reduced expression of HCDH4. While previous investigations support the role of mutated forms of IDH and its importance in the pathophysiology of cancer, the link with the relative levels of expression between wild-type IDH1 and IDH2 in cancer survival is unprecedented. Mutated IDH1 and IDH2 have been implicated in causing cancer through the production of the onco-metabolite D2HG (11). A recent study of hematopoietic malignancies show that the IDH2R^{140Q} mutation was necessary alongside overexpression of Hox9A and
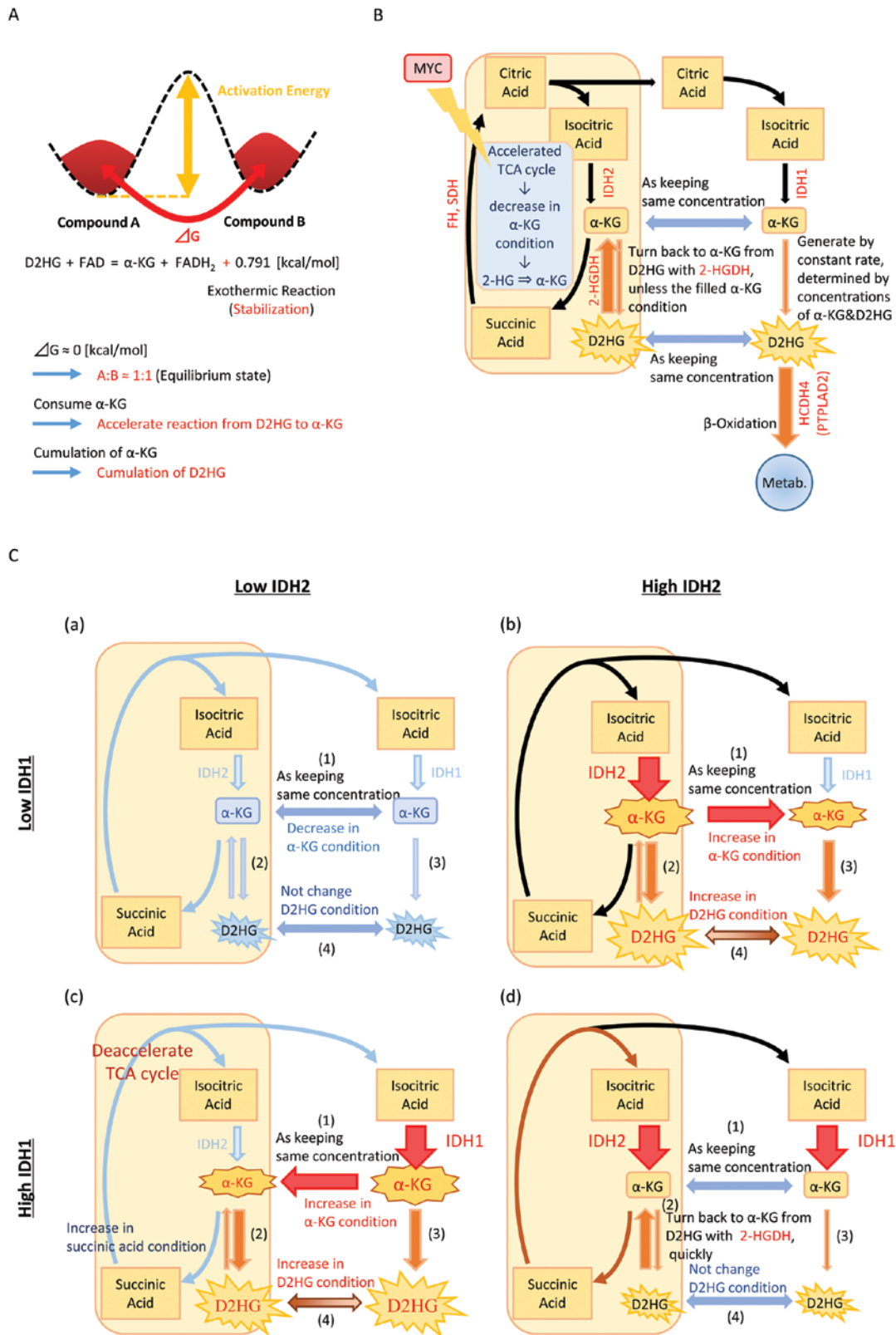
Figure 6. (A) At the state of equilibrium, the abundance of substance A and B depends not on the activation energy but the ΔG. (B) Schematic diagram of the relationship of IDH1, IDH2 and the other genes involved in the TCA cycle. Although, on a routine basis, α-KG and D2HG transfer to and from between mitochondria and cell cytoplasm keeping the same conditions, their conditions are changed by imbalanced IDH1/2 expression. High MYC expression makes improvements because of acceleration of TCA cycle. (C) Schematic diagram explaining how an imbalance between the expressions of IDH1 and IDH2 could lead to increased production of D2HG. (a) IDH1^{low};IDH2^{low}: (1) transfer to and from caused by the same condition, (2) insensitive to transfer to and from caused by each low concentration, (3) insensitive to D2HG in low concentration of α-KG, (4) transfer to and from caused by the same condition; (b) IDH1^{high};IDH2^{low}: (1) move to cell cytoplasm caused by high concentration in mitochondria, (2 and 3) transfer from α-KG to D2HG caused by high concentration of α-KG, (4) go and come keeping the same concentration; (c) IDH1^{low};IDH2^{high}: (1) move to mitochondria caused by high concentration in the cell cytoplasm, (2) a disproportionate emphasis on D2HG caused by high concentration of α-KG and deacceleration of TCA cycle, (3) transfer to D2HG caused by high concentration of α-KG, (4) go and come transfer keeping the same concentration; (d) IDH1^{high};IDH2^{high}: (1) transfer to and from caused by the same condition, (2) a disproportionate emphasis on α-KG caused by the consume α-KG of along with acceleration of TCA cycle, (3) insensitive to D2HG in low concentration of α-KG, (4) transfer to and from caused by the same condition.

Meis1a, or mutation of FMS-like tyrosine kinase 3 (FLT3), to initiate and maintain acute leukemia, most likely through increased levels of D2HG (25).

We postulate a role for D2HG in promoting CRC progression based on our current results, along with other recent published evidence. First, D2HG levels are raised in the tumors and plasma of mice with azoxymethane-induced intestinal cancer (26). Second, 5-hydroxymethylcytosine levels are decreased in human CRC, indicating decreased TET2 activity, which could reflect inhibition through high levels of D2HG (27).

We propose a model that explains how imbalanced IDH1 and IDH2 expression could lead to increased production of D2HG (Fig. 6B and C). In this scheme, we make several assumptions, including that both α-KG and D2HG are able to permeate between mitochondria and cell cytoplasm and equilibrate by osmosis. We also assume that the direction of chemical reaction depends on the difference of free energy between substances rather than the activation energy. When both IDH1 and IDH2 expressions are low, the production of α-KG is also low, leading to low levels of D2HG. When both IDH1 and IDH2 expressions are high, α-KG levels will rise due to increased production, but will either be metabolized effectively by the TCA cycle or be converted back to isocitric acid by the IDH enzymes, and the D2HG levels will remain low. When IDH1 expression is high and IDH2 expression is low, there will be increased production of α-KG in the cytoplasm, which will diffuse into the mitochondria by osmosis. In the absence of a high TCA cycle activity, the low level of IDH2 is not able to convert all of the α-KG into isocitric acid, resulting in the production of D2HG through a relatively small endothermic reaction. Similarly, when IDH2 expression is high and IDH1 expression is low, mitochondrial α-KG will diffuse into the cytoplasm, where the low levels of IDH1 is unable to convert all the α-KG into isocitric acid, instead leading to the production of D2HG.

We next focused on the relationship between MYC and components of the TCA cycle, given MYC has previously been reported to accelerate the TCA cycle. We proposed that the TCA cycle in an accelerated state is able to more efficiently metabolize α-KG, thereby preventing the production of D2HG and increasing the survival of CRC patients. As expected, high expression of MYC alongside high expression of the TCA cycle enzymes IDH2, FH and SDHB were associated with increased DFS and OS. However, the decreased DFS and OS when MYC and SDHA expression is high were inconsistent with the above notion. We are therefore preparing additional analysis concerning SDHA. The increased MYC expression associated with improved survival in this study is consistent with previous studies of CRC (28). In breast cancer however, increased MYC signaling was responsible for D2HG production and was associated with decreased survival (24), suggesting that MYC plays a different role to that in CRC.

HCDH4 is an enzyme that metabolizes D2HG in the β-oxygenation pathway and its mutation has been reported to cause hereditary peroxisomal disorders (20). Therefore the IDH enzymes together with HCDH4 affecting the prognosis of patients with CRC adds further support that D2HG is likely to play a role. Given that HCDH4 gene is located in the frequently deleted genomic region with tumor suppressor INK4A and miR-31 at chromosome 9p21.3, the inactivation of HCDH4 may contribute to CRC progression together with other oncogenic mutations.

The importance of D2HG in pathology is further demonstrated in a rare autosomal recessive neurometabolic disorder called D2HG aciduria (30). This condition occurs when D-2-hydroxyglutarate dehydrogenase (D2-HGHD), a mitochondrial enzyme belonging to the FAD-binding oxidoreductase/transferase type 4 family, is mutated. This enzyme, which is most active in liver and kidney but also active in heart and brain, converts D2HG to α-KG. The condition is characterized by developmental delay, epilepsy, hypotonia, and dysmorphic features (29). Given multiple mechanisms are involved in metabolizing D2HG, this suggests the importance of maintaining D2HG levels low. Biochemical studies indicated that HCDH functions to inactivate D2HG via the glutaconyl pathway, whereas D-2-hydroxyglutarate dehydrogenase (D2-HGHD) is involved in the inactivation of physiological level of D2HG (http://www.genome.jp/kegg/) (Fig. 4B and C).

In order to be certain of the significance of IDH1 and IDH2 expression on the prognosis of colorectal cancer patients, the data requires validation using other cohorts of CRC patients, ideally with matched levels of D2HG. We are presently planning to evaluate D2HG levels in CRC patients.

This study provides direction for future studies and has demonstrated the following: i) the expressions of both IDH1 and IDH2, critical enzymes involved in oxidative phosphorylation in mitochondria, are associated with patient survival in CRC; ii) the usefulness of computational analysis of high volume data to suggest novel biological mechanisms and predict patient survival; iii) the benefit of the gene expression microarray for identifying potential novel therapeutic targets; iv) a link between IDH1, IDH2 and D2HG-inactivating β-oxidization pathway in CRC.

Acknowledgements

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