Integrative Analysis Defines Distinct Prognostic Subgroups of Intrahepatic Cholangiocarcinoma

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Intrahepatic cholangiocarcinoma (iCCA) is the second most common primary liver cancer. It is defined by cholinergic differentiation and has poor prognosis. Recently, epigenetic processes have been shown to play an important role in cholangiocarcinogenesis. We performed an integrative analysis on 52 iCCAs using both genetic and epigenetic data with a specific focus on DNA methylation components. We found recurrent isocitrate dehydrogenase 1 (IDH1) and IDH2 (28%) gene mutations, recurrent arm-length copy number alterations (CNAs), and focal alterations such as deletion of 3p21 or amplification of 12q15, which affect BRCA1 Associated Protein 1, polybromo 1, and mouse double minute 2 homolog. DNA methylome analysis revealed excessive hypermethylation of iCCA, affecting primarily the bivalent genomic regions marked with both active and repressive histone modifications. Integrative clustering of genetic and epigenetic data identified four iCCA subgroups with prognostic relevance further designated as IDH, high (H), medium (M), and low (L) alteration groups. The IDH group consisted of all samples with IDH1 or IDH2 mutations and showed, together with the H group, a highly disrupted genome, characterized by frequent deletions of chromosome arms 3p and 6q. Both groups showed excessive hypermethylation with distinct patterns. The M group showed intermediate characteristics regarding both genetic and epigenetic marks, whereas the L group exhibited few methylation changes and mutations and a lack of CNAs. Methylation-based latent component analysis of cell-type composition identified differences among these four groups. Prognosis of the H and M groups was significantly worse than that of the L group. Conclusion: Using an integrative genomic and epigenomic analysis approach, we identified four major iCCA subgroups with widespread genomic and epigenomic differences and prognostic implications. Furthermore, our data suggest differences in the cell-of-origin of the iCCA subtypes. (Hepatology 2019;69:2091-2106).

Cholangiocarcinoma (CCA) is a rare malignancy of the intrahepatic or extrahepatic bile ducts with very limited treatment options and poor prognosis.1 CCA is classified based on the anatomical location as intrahepatic (iCCA), perihilar (pCCA), and distal CCA (dCCA). The incidence and

Abbreviations: CCA, cholangiocarcinoma; CNA, copy number alteration; CpG, cytosine-guanine dinucleotide; dCCA, distal cholangiocarcinoma; FDR, false discovery rate; FFPE, formalin-fixed paraffin-embedded; HCC, hepatocellular carcinoma; iCCA, intrahepatic cholangiocarcinoma; IDH, isocitrate dehydrogenase; LMC, latent methylation component; LUMP, leukocytes unmethylation for purity; Myc, myelocytomatosis; OS, overall survival; pCCA, perihilar cholangiocarcinoma; PDAC, pancreatic adenocarcinoma; TCGA-CHOL, The Cancer Genome Atlas Cholangiocarcinoma Consortium; TSS, transcription start site.

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etiologic factors of CCA vary in different geographic locations. In Southeast Asia, CCA is frequently caused by liver fluke infections, whereas the etiology is less clear in Western countries. Chronic inflammation and injury of bile duct cells are known CCA promoting conditions. Based on histology, iCCAs are subdivided into two groups: a bile duct type that resembles extrahepatic CCA with columnar cells with mucin production, and a cholangiolar type that recapitulates a genuine small-duct iCCA morphological pattern with cell-rich tubuli formed by cuboidal cells without extracellular mucin. (2) The bile duct type has a higher frequency of KRAS mutations, whereas the cholangiolar type shows a higher frequency of IDH mutations. (2) In addition, it was shown that the mutational landscape is partly subtype-specific, particularly displaying discriminating differences between iCCA versus pCCA and dCCA with, for example, isocitrate dehydrogenase (IDH) mutations almost exclusively detected in iCCA. (3,4) Frequent genetic alterations of epigenetic key players indicate a high impact of epigenetic processes in cholangiocarcinogenesis. (5) Deletions and mutations of genes encoding the chromatin remodeling enzymes BAP1, ARID1A, and PBRM1 (6,7) and gain-of-function mutations of IDH1 and IDH2 (8) are the most common alterations perturbing the epigenetic landscape of iCCA.

Most epigenetic and genetic analyses were performed on mixed iCCA, pCCA, and dCCA cohorts, which may result in failure to detect variation within the iCCA subtype. In addition, the phenotypic and molecular heterogeneity of CCA in general, and iCCA in particular, is suspected to be a result of diverse cellular origins. (9,10) Potential cells-of-origin for iCCA are cholangiocytes, peribiliary glands, and hepatic stem/progenitor cells. (9,11-13) The methylation pattern of a tumor not only reflects tumorigenesis but also the methylation profile of the tumor-initiating cell types. (14) Therefore, we hypothesized that an integrative approach with special attention to cell type–composition differences may result in the identification of tumor subgroups with distinct clinical behavior. Here, we present a comprehensive integrated analysis on the genetic and epigenetic data of 52 iCCA patients of European descent with a non-liver fluke–associated etiology. We identified four distinct iCCA subgroups with prognostic relevance: an IDH group, a low (L group), a medium (M group), and a high genetic and epigenetic alteration group (H group). These four iCCA groups differ in the degree and pattern of DNA methylation, in their genomic alterations and gene mutations. Thus, these four iCCA subgroups might be clinically relevant for patient prognosis and treatment.
Materials and Methods

STUDY POPULATION AND HISTOMORPHOLOGICAL SUBCLASSIFICATION

The study consisted of 52 iCCA patients (Table 1 and Supporting Table S1) and 12 nonneoplastic samples of cholangiocytes originating from the cystic duct of nonneoplastic cholecystectomies. All tissue samples were provided by the Tissue Bank of the National Center for Tumor Diseases (NCT, Heidelberg, Germany) in accordance with the regulations of the NCT Tissue Bank. Informed consent in writing was obtained from each patient and the study protocol conformed to the ethical guidelines of the 1975 Declaration of Helsinki as reflected by a priori approval of the ethics committee of the University of Heidelberg (S-206/2005, S-207/2015, and S-539/2012). Each iCCA tumor sample was histologically confirmed by at least two experienced pathologists (B.G., S.S., and P.S.). In addition, a histomorphological subtyping into bile duct type or cholangiolar type according to Liau et al.\(^2\) was performed (Supporting Fig. S1).

GENOMIC DNA ISOLATION

Genomic DNA was isolated from fresh frozen tissue using the QIAamp DNA micro kit (Qiagen, Hilden, Germany) for whole exome sequencing according to the manufacturer’s instructions (Supporting Table S1). From formalin-fixed paraffin-embedded (FFPE) samples, genomic DNA was extracted using the AllPrep DNA/RNA FFPE Kit (Qiagen), as recommended by the manufacturer with the following modifications: After addition of xylene, samples were incubated at 56°C for 2 minutes followed by two ethanol washes. The first proteinase K digestion was performed with 20 µL at 56°C for 30 minutes. The DNA was eluted twice with 30 µL of H\(_2\)O.

EXOME SEQUENCING

Whole exome sequencing libraries were prepared from DNA isolated from fresh frozen tissue and from microdissected surrounding normal tissue to distinguish somatic from germline mutations. Sequencing of the libraries was done at the German Cancer Research Center (DKFZ) Genomics and Proteomics Core Facility using the Agilent SureSelectXT Human all Exon V4 kit and a HiSeq2000 instrument (Illumina, San Diego, CA) using the 100-bp paired-end mode.

Panel Sequencing

To analyze the samples for genetic variations, a custom gene panel for massive parallel sequencing was used. This panel consisted of 285 amplicons covering 165 exons within 40 genes frequently mutated in biliary tract cancers (Supporting Table S2). For detailed information see the Supporting Information.

DNA METHYLATION ANALYSIS

DNA methylation profiles were determined by the Genomics and Proteomics Core Facility (DKFZ Heidelberg) using the Infinium HumanMethylation450 BeadChip from FFPE tissue–derived genomic DNA, as described previously.\(^15\)

DATA ANALYSIS

Available data and the process of data analysis are shown in Fig. 1A.

Methylation data were processed using the R platform for statistical computing (see Supporting Information). Copy number alterations were assessed based on the signal intensities measured in the methylation array. Differential methylation analyses (between tumor and normal or between the identified subgroups) were conducted using linear models. Adjustments were made for patient age, gender, and the source of tissue in tumor-normal comparison and for age in the group-wise comparison among subgroups. A difference was considered to be significant if the false discovery rate (FDR) corrected \(P\) value (q value) was less than 0.05. Tumor purity was estimated using the LUMP (leukocytes unmethylation for purity) method.\(^16\) To assess cell type heterogeneity and to trace the cell-of-origin in tumor samples, we used MeDeCom\(^17\) to decompose methylation data into latent methylation components (LMCs) and to assess their proportions in each sample. The integrative clustering was confined to tumor samples and performed using the copy number, DNA methylation, and LMC data with the iClusterPlus R package.\(^18\)

To identify possible candidate genes located in the respective chromosomal arms or focal
regions with significant copy number alterations (CNAs), we used the gene expression and copy number data of The Cancer Genome Atlas Cholangiocarcinoma Consortium (TCGA-CHOL) (https://portal.gdc.cancer.gov/projects/TCGA-CHOL). Pearson correlation was performed in chromosomal arm level analysis and Spearman correlation in the focal alterations. FDR values less than 0.05 were considered

### TABLE 1. Clinical Characteristics of the iCCA Study Population (n = 52)

| Characteristics | Value | P Value* |
|-----------------|-------|----------|
| Age             | 59.91 | 0.51     |
| Tumor size      | 7.19  | 0.034    |
| Number of patients | **n** | **Percentage** |
| Gender          |       |          |
| Male            | 25    | 48.1 Ref. |
| Female          | 27    | 51.9     |
| Histological type |     |          |
| Cholangiolar type | 40   | 76.9 Ref. |
| Bile duct type  | 12    | 23.1     |
| T stage         |       |          |
| pT1             | 16    | 30.8 Ref. |
| pT2             | 28    | 53.8     |
| pT3 (n = 7)/pT4 (n = 1) | 8    | 15.4 0.661 |
| Histologic grade |     |          |
| G1              | 4     | 7.7 Ref. |
| G2              | 36    | 69.2     |
| G3              | 12    | 23.1     |
| N               |       |          |
| N0              | 17    | 32.7 Ref. |
| N1              | 11    | 21.2     |
| N.A.            | 24    | 46.2 N.A. |
| M               |       |          |
| M0              | 52    | 100.0 N.A. |
| L               |       |          |
| L0              | 31    | 59.6 Ref. |
| L1              | 16    | 30.8     |
| N.A.            | 5     | 9.6 N.A. |
| V               |       |          |
| V0              | 33    | 63.5 Ref. |
| V1              | 14    | 26.9     |
| N.A.            | 5     | 9.6 N.A. |
| R               |       |          |
| R0              | 28    | 53.8 Ref. |
| R1              | 16    | 30.8     |
| R2              | 3     | 5.8      |
| N.A.            | 5     | 9.6 N.A. |
| UICC            |       |          |
| UICC 1          | 4     | 7.7 Ref. |
| UICC 2          | 10    | 19.2     |
| UICC 3          | 15    | 28.8     |
| N.A.            | 23    | 44.2 N.A. |
| Hepatobiliary disease |     |          |
| Hepatitis B virus | 3    | 5.8     |
| Hepatitis C virus | 3    | 5.8     |
| Hepatic steatosis | 4    | 7.7     |
| NAFLD           | 4     | 7.7     |
| Pre-existing inflammatory biliary tract disease† | 8    | 15.4 0.421 |
| Chronic pancreatitis | 1    | 1.9 N.A. |
| M. Wilson       | 1     | 1.9 N.A. |
| Thorotrast      | 1     | 1.9 N.A. |
| None detected   | 27    | 51.9 Ref. |

* Cox regression P value.
† Cholecystitis and/or choledocholithiasis.
Abbreviations: N.A., not available; NAFLD, nonalcoholic fatty liver disease; Ref., reference; and UICC, International Union Against Cancer.
FIG. 1. DNA methylome landscape of iCCA. (A) Overview of the study. (B) Clustering of the methylation values of tumor and normal samples (columns), ranging from 0 to 1, of the 10,000 most variable CpG sites (rows). Tumor purity is represented by LUMP values ranging from 0 (low purity) to 1 (high purity). (C) Enrichment analysis of the hypomethylated and hypermethylated CpG sites compared with normal controls, using the 18-state ChromHMM model of Roadmap Epigenomics. The log2 fold change is represented by the color of the dots, whereas the size reflects the \(-\log_{10}(P\text{ value})\). The border of the dots shows whether the result is significant. (D) Enrichment analysis using the GO Molecular functions database for the hypermethylated CpG sites (|beta difference| > 0.2). FDR q values < \(10^{-300}\) were set to \(10^{-300}\). (E) Clustering results of the MeDeCom analysis for tumor and normal samples (columns) with respect to the five LMCs (LMC1 to LMC5) (rows). Color represents the contribution of the given LMC to the respective sample. Abbreviations: 1 TssA, active TSS; 2 TssFlnk, flanking TSS; 3 TssFlnkU, flanking TSS upstream; 4 TssFlnkD, flanking TSS downstream; 5 Tx, strong transcription; 6 TxWk, weak transcription; 7 EnhG1, genic enhancer1; 8 EnhG2, genic enhancer 2; 9 EnhA1, active enhancer 1; 10 EnhA2, active enhancer 2; 11 EnhWk, weak enhancer; 12 ZNF/Rpts, ZNF genes and repeats; 13 Het, heterochromatin; 14 TssBiv, bivalent/poised TSS; 15 EnhBiv, bivalent enhancer; 16 ReprPC, repressed PolyComb; 17 ReprPCWk, weak repressed PolyComb; and 18 Quies, quiescent/low.
to be significant. Survival analysis was performed using Cox proportional hazards regression model. Detailed information on the methods used is described in the Supporting Information.

**Results**

**iCCA IS CHARACTERIZED BY FREQUENT TP53 AND IDH MUTATIONS AND LARGE CHROMOSOMAL ABERRATIONS**

To genetically characterize our 52 iCCA samples, we performed whole exome or targeted panel sequencing of 40 genes commonly mutated in CCA in a subset of 36 iCCA tumor samples (Supporting Table S1), which revealed 24 mutations (Table 2 and Supporting Table S2). Recurrent missense mutations in *IDH1* were observed in 6 of 36 (17%) patients (Table 2). All *IDH1* mutations were located in a mutation hotspot and altered the same codon: p.R132C, p.R132G, or p.R132L. IDH2 encoding the mitochondrial isozyme of *IDH1* displayed three different missense mutations in the same mutation hotspot, leading to p.R172W, p.R172M, or p.R172S. *IDH1* and *IDH2* mutations lead to the production of the oncometabolite 2-hydroxylutarate, which was shown to inhibit histone and DNA demethylation. TP53 was the second most affected gene with two nonsense and two missense mutations in 4 patients. These mutations all impaired the DNA binding domain of the TP53 protein, as has been observed frequently in iCCA. The mutations in *IDH1*, *IDH2*, and *TP53* were mutually exclusive with each other. In addition, two missense mutations were found in *TGFB2* and one mutation was found in *ARID1A, SF3B1, ROBO1, ROBO2, FBXW7, BRAF, CDKN2A, FGFR2, KDM5A, KRAS, SMARC4*, and *GNAS* each (Table 2).

We used HumanMethylation450 BeadChip data to identify recurrent DNA methylation and genomic

| Gene  | Amino Acid Change | Nucleotide Change | Type       | Location |
|-------|-------------------|-------------------|------------|----------|
| ARID1A| p.M890V           | NM_006015:c.2668A>G | Missense   | 1p36.11  |
| BRAF  | p.V600E           | NM_00433.4:c.1799T>A | Missense   | 7q34     |
| CDKN2A| p.A97V            | NM_058195:c.290C>T | Missense   | 9p21.3   |
| FBXW7 | p.R505H           | NM_033632:c.1514G>A | Missense   | 4q31.3   |
| FGFR2 | p.C382R           | NM_000141:c.1144T>C | Missense   | 10q26.13 |
| GNAS  | p.R201C           | NM_000516:c.601C>T | Missense   | 20q13.32 |
| IDH1  | p.R132C           | NM_001282386:c.394C>T | Missense   | 2q34     |
|       | p.R132G           | NM_001282386:c.394G>C | Missense   |          |
|       | p.R132L           | NM_001282386:c.395G>T | Missense   |          |
| IDH2  | p.R172W           | NM_001282386:c.514A>T | Missense   | 15q26.1  |
|       | p.R172S           | NM_001290114:c.1266G>T | Missense   |          |
|       | p.R172M           | NM_001290114:c.1256G>T | Missense   |          |
| KDM5A | p.R604C           | NM_000495:c.1810C>T | Missense   | 12p13.33 |
| KRAS  | p.G12delinsGAG    | NM_00433.4:39_40delGAG | Insertion  | 12p12.1  |
| ROBO1 | p.V1454L          | NM_002941:c.4366G>T | Missense   | 3p12.3   |
| ROBO2 | p.G866D           | NM_002942:c.2597G>A | Missense   | 3p12.3   |
| SF3B1 | p.A702_S705delRTI | NM_012433:c.2104_2112del | Deletion   | 2p33.1   |
| SMARC4 | p.E882D         | NM_00128845:c.2646A>C | Missense   | 19p13.2  |
| TGFB2 | p.R528C           | NM_004333:c.1799T>A | Missense   | 3p24.1   |
|       | p.V412M           | NM_001024847:c.1234G>A | Missense   | 17p13.1  |
| TP53  | p.R248Q           | NM_000546:c.743G>A | Missense   |          |
|       | p.E171*           | NM_000546:c.511G>T | Nonsense   |          |
|       | p.Y163*           | NM_000546:c.489C>G | Nonsense   |          |
|       | p.R290C           | NM_000546:c.868C>T | Missense   |          |

Table 2. Mutations in the Study Population (n = 36)

Note: All changes occurred once (2.8%) except for IDH1 p.R132C, which occurred 4 times (11.1%).
We found recurrent deletions and amplifications of entire chromosome arms (Table 3) and of focal genomic regions (Table 4 and Supporting Fig. S2). Chromosome arms 1p, 3p, 6q, 8p, 9p, 9q, 12q, 13q, 14q, 16q, and 17p were affected by large deletions consisting of between 422 genes on 9p and up to 2,121 genes on 1p (Table 3). Some genes were found in both lists of deleted and mutated genes (namely, ARID1A on 1p; ROBO1, ROBO2, and TGFBR2 on 3p; CDKN2A on 9p; and TP53 on 17p). Chromosome arm 1q showed long-range amplifications covering up to 1,955 genes (Table 3).

We identified genes whose expression might be affected by focal deletions or amplifications in our study population by correlation analyses of publicly available CNA and gene expression data from TCGA-CHOL (n = 51) and considered information on known driver genes (22) (Table 4). For example, the focally deleted genes BAP1 and PBRM1 on 3p21.1 were also commonly affected by an arm-length deletion in our study population (38%).

### Table 3. Recurrent Arm-Length CNAs and Affected Genes (n = 52)

| Arm | Genes (n) | Frequency | FDR q Value | Deleted Genes With Mutation in Study* |
|-----|-----------|-----------|-------------|-------------------------------------|
| 1p  | 2,121     | 0.18      | 0.0002      | ARID1A                              |
| 3p  | 1,062     | 0.38      | 0           | ROBO1, ROBO2, TGFBR2                |
| 6q  | 839       | 0.48      | 0           |                                     |
| 8p  | 580       | 0.16      | 0.0002      |                                     |
| 9p  | 422       | 0.25      | 9.64*10⁻¹³ | CDKN2A                              |
| 9q  | 1,113     | 0.25      | 2.56*10⁻¹¹ |                                     |
| 12q | 1,447     | 0.12      | 0.0493      |                                     |
| 13q | 654       | 0.21      | 1.44*10⁻⁰⁸ |                                     |
| 14q | 1,341     | 0.23      | 4.00*10⁻⁰⁹ |                                     |
| 16q | 702       | 0.12      | 0.0222      |                                     |
| 17p | 683       | 0.12      | 0.0222      |                                     |

*Mutated genes are listed in Table 2.

### Table 4. Recurrent Focal CNAs and Possible Candidate Genes in the Region

| Chromosome | Cytoband | Coordinates       | q Value | Candidate Genes |
|------------|----------|-------------------|---------|-----------------|
| Deleted Regions |
| 1          | 1p13.2   | 100598444-116907532 | 0.0873  | SLC30A7, CTNBP2NL, PHF1, KCNC4, RHOC, SARS, PSMA5, NRAS, MOV10, MDM1678, DDX20, GNAI3, CSDE1, CLCC1, BCAS2 |
| 1          | 1p36.23  | 8014651-8874983    | 0.0045  | BAP1, SPNS1, PBRM1*† |
| 3          | 3p21.1   | 52318567-52812032  | 0.0089  | CASP3, DCTD, FAT1, FRG1, ING2, FBXO8, UFSP2, LRIP28P, SNX25 |
| 4          | 4q35.2   | 171011578-191154276 | 0.0289  | RASA1, TBCA, XRC4C |
| 5          | 5q14.3   | 76116089-95991890  | 0.0029  | CASP1†           |
| 7          | 7q21.11  | 79881211-85911765  | 0.0002  |                |
| 11         | 11q22.3  | 104769088-105947697 | 0.0002  |                |
| 17         | 17q21.2  | 38808305-39465712  | 0.0289  |                |

Amplified Regions

| Chromosome | Cytoband | Coordinates       | q Value | Candidate Genes |
|------------|----------|-------------------|---------|-----------------|
| 1          | 1q21.3   | 150554876-152815842 | 0.0106  | ARNT, ENSA, PI4KB, VPS72, PIK5K1A, SETDB1, POGZ, GOLPH3L, CDC42SE1, ZNF687, PRUNE, GABPB2, LYSMD1 |
| 2          | 2q31.1   | 176936398-177001961 | 0.0023  |                |
| 4          | 4p15.32  | 16791107-17785836  | 1.57*10⁻³¹ |                |
| 12         | 12q5     | 66220166-71083782  | 0.0023  | MDM2, NUP107I, CNOT2 |

Note: Bold indicates a significant correlation between gene expression and CNA in TCGA-CHOL.
*Identified as possible driver gene in Ref. (50).
†Identified as driver gene in Ref. (22).
Table 3) and by known inactivating mutations in CCA and renal cell carcinoma.\(^6,\)\(^23\) As an example of a candidate gene residing in the amplified 12q15 region, the \(MDM2\) oncogene showed a significant positive correlation \((P = 0.016, r = 0.40)\) with its expression in the TCGA-CHOL cohort.

**iCCA SHOWS EXTENSIVE HYPERMETHYLATION**

Clustering of the 10,000 most variably methylated CpGs (cytosine-guanine dinucleotides) revealed distinct clusters for the normal samples and the iCCAs (Fig. 1B). One cluster of iCCA samples displayed DNA methylation levels similar to the nonneoplastic cholangiocyte samples, whereas the remaining iCCA cases displayed different patterns (Fig. 1B). The similarity in DNA methylation profiles between the group of nonneoplastic cholangiocyte samples and the normal-like iCCA group might in part be ascribed to a higher immune cell infiltration in both groups, as suggested by LUMP analysis (Fig. 1B). Primarily CpG hypermethylation was seen in the iCCA cohort with approximately 37,600 hypermethylated CpGs exhibiting increased methylation by more than 20%. In contrast, only 2,217 CpG sites were hypomethylated by more than 20%. We overlapped the hypermethylated and hypomethylated CpG sites with publicly available 18-state chromatin segmentation data of H1 ES cells and found that hypermethylated regions were enriched at transcription start sites (TSSs) and enhancers but depleted in heterochromatic regions and gene bodies. Hypomethylated regions were enriched in gene bodies, heterochromatic and quiescent regions, but depleted at TSSs (Fig. 1C).

To further characterize the hypermethylated regions, we used the GREAT tool and found genes involved in transcription factor and HMG (3-hydroxy-3-methylglutaryl) box domain binding to be enriched (Fig. 1D). We also performed MeDeCom analysis to dissect methylation patterns into LMCs to incorporate information on possible cell-type composition differences (Fig. 1E). Five major LMCs were identifiable, of which LMC1 appeared to be characteristic for normal bile duct samples (Supporting Fig. S3). LMC1 and LUMP values were strongly negatively correlated \((r = -0.799, P < 0.01;\) Supporting Fig. S4).

**FOUR iCCA SUBGROUPS ARE IDENTIFIED WITH INTEGRATIVE CLUSTERING**

Using iCluster\(^{18}\) on the tumor samples, we performed an integrative clustering combining CNA, methylation, and MeDeCom analysis data and identified four iCCA groups, named the IDH group, L (low alteration) group, M (medium alteration) group, and H (high alteration) group. These four groups are characterized by IDH mutation, the degree of acquired genetic and epigenetic alterations, and by LMC profiles (Fig. 2A, Supporting Fig. S3). The enrichment analyses of the LMC-specific hypermethylation show the involvement of developmental and differentiation processes for LMC3 and LMC4, whereas in LMC2, primarily metabolic processes are affected (Supporting Fig. S5). The L group partially overlaps with the normal-like samples (Fig. 1A) and is characterized by the highest proportion of LMC1 (48% versus 23%, 17% and 17%, respectively) and LMC5 (28% versus 22%, 2%, and 8%), low mutation number (4 mutations in 4 of 11 samples sequenced), low frequency of CNA (approximately 40% for 6q deletion and less for the other alterations), and generally low methylation levels. The high ratio of LMC1 may be partially explained by high infiltration of immune cells, as shown by the correlation of LMC1 with LUMP (Supporting Fig. S4).

Most of the samples in the IDH group are characterized by mutations in \(IDH1\) or \(IDH2\) (9 out of 10 sequenced), a unique pattern of relatively high methylation values, a high level of CNAs (up to 80% for 3p deletions), and high values of LMC2 (Fig. 2A and Supporting Fig. S6). The high level of methylation in the IDH group is consistent with the neo-enzymatic function of affected IDH proteins, resulting in the inhibition of TET-enzymes and ultimately in inhibition of active DNA demethylation.\(^{20}\) The iCluster H group has high methylation levels, frequent CNAs (up to 70% for some chromosome arms), and high LMC4 values. Finally, the M group showed a mixed LMC composition, low frequency of deletions and amplifications, and a group-specific gain of the chromosome arm 8q harboring the \(Myc\) (myelocytomatosis) oncogene. Thus, iCluster analysis revealed four distinct iCCA subgroups with specific methylation patterns, LMCs, CNAs, and mutations.
The four iCCA subgroups did not show any significant associations with basic clinical characteristics, including age, gender, tumor staging, and underlying hepatobiliary disease (Supporting Table S3). In the IDH group, the only histological type was the cholangiolar type, whereas the other three groups included 18% to 36% samples with bile duct type (Fisher’s exact test $P = 0.047$; comparison of IDH versus L, M, and H groups combined). In addition, we found that the molecular subgroups showed significantly different outcomes (Fig. 2B). Overall survival (OS) of the M group was the worst ($P = 0.005$, OS M group versus L group), followed by the H group ($P = 0.034$, OS H group versus L group). The L and the IDH groups showed the most favorable prognosis and did not significantly differ from each other (Fig. 2B).

Further analyses of known cancer driver genes showed that $RPL22$, $ROBO1$, $ROBO2$, $TGFBR1$, and $TGFBR2$ were most frequently altered either by deletion or promoter hypermethylation in all groups (Fig. 3A). The IDH and H groups frequently showed deletions on chromosome arms 3p and 6q, the former harboring the tumor-suppressor genes $BAP1$, $PBRM1$, $TGFBR2$, $ROBO1$ and $ROBO2$. 

FIG. 2. Integrative cluster analysis identifying four subgroups in iCCA. (A) Integrative clustering using CNA, DNA methylation, and MeDeCom data splits the iCCA samples into four groups. In addition, the histological types of the iCCA samples, the mutation status of the three most commonly mutated genes, tumor size, and tumor purity according to LUMP are indicated. (B) Kaplan-Meier survival plots of the four groups. The tables indicate the number of patients at risk at given time points and the results of the Cox proportional hazard regression model using the L group as reference.
the latter ZNF292 and EEF1A1. Both the M and H groups shared amplification of chromosome 1q, whereas amplification of 8q, which harbors the MYC oncogene, predominated in the M group (Fig. 3B and Supporting Table S4). We narrowed down the amplification to a focal region in 1q21.3 containing a number of candidate genes (Table 4).

To better define candidate driver genes affected by genomic alteration in these regions, we correlated copy number gain with expression data obtained from TCGA-CHOL. Among the genes amplified on 1q, a strong correlation was observed for PI4KB ($r = 0.71$, FDR $q = 3.66 \times 10^{-5}$) and PIP5K1A ($r = 0.69$, FDR $q = 5.91 \times 10^{-5}$), which, together with AKT3 ($r = 0.28$, FDR $q = 0.19$) (Supporting Fig. S7A), belong to the cancer-relevant phosphoinositide signaling pathway. (24) Another candidate was YY1AP1 ($r = 0.65$, FDR $q = 2.36 \times 10^{-4}$), known as an oncopgenic driver in EpCAM(+) AFP(+) hepatocellular carcinoma, (25) which shows features of hepatic stem/progenitor cells. (26) By high correlation ($r = 0.82$, FDR $q = 1.09 \times 10^{-6}$), SETDB1, encoding a histone lysine methyltransferase and known to be involved in breast cancer, (27) proved a candidate driver gene on the focally amplified 1q21.3 (Supporting Fig. S7B). MYC presented as a candidate on 8q, yet this role was not underscored by correlation with expression. However, potential candidates on 8q are CHRAC1 ($r = 0.61$, FDR $q = 0.010$), RAD21 ($r = 0.51$, FDR $q = 0.023$), and TRAPPCC9 ($r = 0.50$, FDR $q = 0.030$) (Supporting Fig. S7C), all known to be involved in breast cancer. (28,29) These results suggest the importance of CNA-driven gene-expression changes.

As the epigenomic landscape contributes to the properties and specification of tumor subgroups, (30,31) we searched for characteristics of the four iCCA groups regarding their methylation patterns. Globally, all four subgroups show a significant increase in their methylation levels, with significant differences between the L and the M versus the H and IDH mutant groups (Supporting Fig. S6). To decipher the group-specific changes, we performed differential methylation analysis comparing each group to all other tumor samples (Supporting Fig. S8). The IDH and H groups exhibited broad hypermethylation with highly group-specific patterns. This finding is supported by the number of unique and overlapping differentially methylated CpG sites using the L group as reference (Fig. 4A). In contrast, the M group showed only few CpGs with group-specific hypermethylation (Fig. 4A and Supporting Fig. S8).

Enrichment analysis of the specific hypermethylated sites in the IDH group showed increased appearance in bivalent TSSs and enhancers, but the enrichment is even more pronounced in the flanking regions of the TSSs (Fig. 4B). The H group exhibited an even stronger enrichment at the bivalent enhancers and TSSs, whereas the TSS flanking regions were depleted. Pathway analysis revealed that most frequently affected genes encode mostly homeobox proteins and transcription factors (Fig. 4C). However, the IDH group-specific alterations did not show enrichment of cancer-related pathways (Supporting Fig. S9) but were aggregated in CTCF and RAD21 binding sites and depleted in the binding sites of other transcription factors (Fig. 4D).

To compare the resemblance and the potential cell-of-origin of the four iCCA groups with hepatocellular carcinoma (HCC) and pancreatic adenocarcinoma (PDAC), we applied our MeDeCom model to TCGA DNA methylation data (see Supporting Information). We included 100 randomly selected HCCs and added all IDH-mutant HCCs that were not randomly selected (TCGA-LIHC, $n = 106$, including 9 IDH-mutant HCCs), 100 randomly selected PDACs and added all IDH-mutant PDACs (TCGA-PAAD, $n = 101$, including 1 IDH mutant), and all available CCAs (TCGA-CHOL; $n = 45$; Supporting Table S5). Clustering analysis of the tumor-specific LMC2–5 profiles showed that LMC2 co-occurred with IDH1 and IDH2 gain-of-function mutation of HCC and CCA but did not include any PDAC cases (Fig. 4E and Supporting Fig. S10). Furthermore, the TCGA-CHOL samples exhibited similar subgroups as our iCCA study population. A subgroup of PDAC cases formed a distinct cluster with primarily M group iCCA samples with high LMC3 ratio, whereas clusters with LMC4 and LMC5 included only HCC and CCA cases. H group iCCA shared similarities with LMC4-high HCC and CCA, but this cluster did not include any PDAC cases. Consistently, the bile duct–type iCCA clustered together with M group iCCA and PDAC, whereas the cholangiolar-type iCCA showed similar patterns to a subgroup of HCC high LMC4 or LMC5. In addition, iCCA cases with amplification of YY1AP1, a potential marker of HCC with hepatic stem/progenitor features, did not cluster together.
with the IDH-mutant cases that are LMC2-high (Fig. 4E). YY1AP1-amplified iCCA showed high levels of LMC4 or LMC5 and clustered primarily with a subgroup of IDH wild-type HCC and CCA but not with PDAC. The remaining HCC and PDAC samples showed very low ratios of LMC2 to LMC5. Taken together, using three TCGA data sets consisting of HCC, CCA and PDAC cases, we found that IDH gain-of-function mutations resulted in distinct LMC2-high profiles, M group iCCA were enriched for bile duct–type iCCA and exhibited similarities with LMC3-high PDAC, whereas H group iCCA had higher LMC4 and the LMC4-high cluster was enriched for cholangiolar-type iCCA.
Recent studies in cholangiocarcinoma (CCA). These genome-wide studies focused primarily on understanding the mechanisms underlying iCCA, ear-lier understood; some studies showed that hypermethyl-
ation of bivalent regions continues the repression of
transcription factors and developmental processes. Hypermethylation of the regions show-

ting bivalent characteristics in embryonic stem cells is

ing bivalent regulatory regions in this iCCA cohort

Our sequencing analysis confirmed the recurrence of 

Discussion
As a reflection of the growing urgency to better understand the mechanisms underlying iCCA, earlier omic studies analyzed the molecular landscape of CCA. These genome-wide studies focused primarily on the mutational and genomic CNA landscape of CCA.\(^{(6,7,21,32)}\) Recent studies in cholangiocarcinoma used integrative analysis of genomic, transcriptomic, and epigenomic data in mixed cohorts of iCCA, pCCA, and dCCA.\(^{(33,34)}\) Our present study included a clinically well-characterized European cohort of iCCA patients only (n = 52) with different underlying hepatobiliary disease but without liver fluke association (Supporting Table S3). Consistent with previous studies, pre-existing cholecystitis and/or choledocholithiasis was associated with bile duct–type iCCA (Supporting Table S6). Here, we integrated mutational data from 40 candidate genes, CNAs, whole genome DNA methylation changes, and publicly available TCGA-CHOL transcriptomes. We also harnessed the methylome data for tumor purity assessment and the identification of latent methylation components as proxies for different cell types, and thereby identified four distinct iCCA groups with prognostic significance. Importantly, the four iCCA groups resulting from the iCluster analysis are largely different in the proportions of LMCs, which recover cell type–specific hidden methylation patterns.\(^{(17)}\)

Our sequencing analysis confirmed the recurrence of \textit{IDH1} and \textit{IDH2} gain-of-function mutations (28%) and \textit{TP53} loss-of-function mutations (9%) in carcinogenesis of iCCA. Moreover, additional mutations in epigenetic genes other than \textit{IDH1} and \textit{IDH2} included \textit{ARID1A} and \textit{SMARCA}, both involved in chromatin remodeling, and \textit{KDM5A}, encoding a histone demethylase, highlighting a strong epigenetic component in iCCA carcinogenesis. Recurrent long-range genomic alterations additionally underscored the importance of known iCCA candidate genes such as \textit{BAP1} and \textit{PBRM1} on deleted 3p21.1\(^{(6)}\) or \textit{MDM2} on amplified 12q15. The latter gene may initiate tumor onset and progression by negatively regulating tumor suppressor p53\(^{(35)}\) or by a p53-independent mechanism.\(^{(36)}\) In addition, \textit{ARID1A}, \textit{ROBO1}, \textit{ROBO2}, \textit{TGFBR2}, \textit{CDKN2A}, and \textit{TP53} were found to be affected by recurrent deletions or amplifications.

Methylome analysis showed excessive hypermethylation of iCCA. Hypermethylation disproportionally affected bivalent regulatory regions in this iCCA cohort and was enriched for differentiation and developmental processes. Hypermethylation of the regions showing bivalent characteristics in embryonic stem cells is a general phenomenon in many different cancers.\(^{(37,38)}\) It usually affects transcription factors and developmental genes, especially those from the homeobox gene family. Our data show high concordance with these previous findings. Despite its widespread occurrence, the role of this specific pattern is still not completely understood; some studies showed that hypermethylation of bivalent regions continues the repression of
Integrative clustering is a powerful tool to identify patient subgroups. Our iCluster analysis revealed four different molecular iCCA groups, designated as the IDH, L, M, and H groups. The most conspicuous subgroup is the IDH group characterized by gain-of-function IDH1 and IDH2 hotspot mutations in 9 of 10 tumor samples sequenced, many CNAs, DNA hypermethylation, and high LMC2 values. The H group has a similarly disrupted genome as the IDH group with frequent deletions of chromosome arms 3p and 6q. The 3p deletions include tumor-suppressor genes like BAP1 and PBRM1, found to be frequently inactivated in iCCA. Amplification of 1q, which is seen in the M and H groups, is common in many different cancers and often associated with bad prognosis. In HCC, this amplification occurs in more than 70% of cases and has been described in iCCA as well. It harbors the stem cell–related oncogene YY1AP1 and genes of the phosphoinositide signaling pathway known to be involved in breast cancer. A possible candidate gene of the 1q region, PIPS14, is a potential therapeutic target of kinase inhibitors, of which some are already in clinical trials. SETDB1, which is amplified in almost 50% of the patients in the H and M groups, is discussed as a therapeutic target, showing sensitivity against mithramycin.

Gain–of–function mutations of IDH1 and IDH2 are known to lead to DNA hypermethylation by the enrichment of the oncometabolite 2-hydroxyglutarate, which inhibits TET enzymes involved in active demethylation. IDH mutations alone were shown to be sufficient to induce a hypermethylator phenotype, and they tend to appear more frequently in recurrent tumors with gene–expression traits of epithelial–mesenchymal transition. Targeted therapies for IDH–mutant tumors are already in clinical trials.

Our pathway analyses, consistent with the general hypermethylation of IDH–mutant tumors, showed no strong enrichment in any pathways (Supporting Fig. S9), but the transcription factor binding site–enrichment analysis revealed enrichment of hypermethylation at CTCF and RAD21 binding sites. This is largely in line with the recent finding of Flavahan et al., showing that hypermethylation led to disruption of the insulator function of the CTCF protein in IDH–mutant gliomas and subsequent overexpression of the PDGFRα candidate oncogene. Our results suggest a similar mechanism in the IDH–mutant iCCAs. Both the H and M groups have a specific hypermethylation profile that is less pronounced in the M group and different from the pattern associated with the IDH group. This profile is enriched in embryonic stem cell–related bivalent regulatory regions that are indicators of pluripotency in stem cells and whose methylation manifests silencing during differentiation. A strong difference between the M and H group is the enrichment of LMC4 in the H group. This enrichment, combined with the similar methylation profile, suggests related mechanisms of tumorigenesis, but probably a different cell–of–origin. Whether the L group and the IDH group with enrichment in high LMC1 and LMC2 values, respectively, may be traced to distinct cells–of–origin, or whether these profiles reflect properties independent thereof, remains to be clarified. Multiple cells–of–origin have been proposed for CCA, and the distinct cells–of–origin is supported by a recent pan–cancer integrative clustering that showed diverse cluster memberships for cholangiocarcinoma, where the clusters dominantly reflected on cells–of–origin–associated signals.

The hypothesis that the cells–of–origin of cholangiocarcinoma, where the clusters dominantly reflected on cells–of–origin–associated signals, may be traced to distinct cells–of–origin, or whether these profiles reflect properties independent thereof, remains to be clarified. Multiple cells–of–origin have been proposed for CCA, and the distinct cells–of–origin is supported by a recent pan–cancer integrative clustering that showed diverse cluster memberships for cholangiocarcinoma, where the clusters dominantly reflected on cells–of–origin–associated signals.

In this study, we used biliary epithelia microdissected from nonneoplastic cystic ducts as normal control tissues. Comparative methylation data analyses showed similar DNA methylation patterns and LMC profiles of L group iCCA and the normal control samples, indicating a large bile duct cholangiocyte as the cell–of–origin of the L group iCCA. However, due to the lack of direct matched–pair samples, direct comparisons cannot be conducted here. In addition, it would be very interesting to analyze matched–pair DNA methylation profiles of biliary precursor lesions.
and invasive CCAs in future studies. Consistent with the low burden of mutations and epigenetic alterations compared with the normal cholangiocytes, the L group exhibited the most favorable outcome with a 3-year survival rate of 91%, compared with 65%, 50%, and 36% for the IDH, H, and M groups, respectively.

Further analyses are needed to reveal alterations in the background of the poor survival of the M group. A limitation of our study is that whole genome sequencing and gene-expression data that could reveal a hidden candidate gene affecting the prognosis in this group are missing. Our observation of group-specific clinical outcomes requires particular consideration: Both the L and IDH groups appear to have a better outcome than the other two groups. The favorable outcome of the IDH group is in agreement with two other studies. (19,34)

In summary, we identified four different iCCA groups that probably differ by their cell-of-origin, underlying tumorigenic mechanism, and clinical outcome. Thus, the four iCCA subgroups presented in this study demonstrate options for the stratification of patients according to our molecular profiling and may lead to subgroup-specific treatment modalities in the future.

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