A Novel Predictive Equation for Potential Diagnosis of Cholangiocarcinoma

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Abstract
Cholangiocarcinoma (CCA) is the second most common-primary liver cancer. The difficulties in diagnosis limit successful treatment of CCA. At present, histological investigation is the standard diagnosis for CCA. However, there are some poorly-defined tumor masses which cannot be definitively diagnosed by general histopathology. As molecular signatures can define molecular phenotypes related to diagnosis, prognosis, or treatment outcome, and CCA is the second most common cancer found after hepatocellular carcinoma (HCC), the aim of this study was to develop a predictive model which differentiates CCA from HCC and normal liver tissues. An in-house PCR array containing 176 putative CCA marker genes was tested with the training set tissues of 20 CCA and 10 HCC cases. The molecular signature of CCA revealed the prominent expression of genes involved in cell adhesion and cell movement, whereas HCC showed elevated expression of genes related to cell proliferation/differentiation and metabolisms. A total of 69 genes differentially expressed in CCA and HCC were optimized statistically to formulate a diagnostic equation which distinguished CCA cases from HCC cases. Finally, a four-gene diagnostic equation (CLDN4, HOXB7, TMSB4 and TTR) was formulated and then successfully validated using real-time PCR in an independent testing set of 68 CCA samples and 77 non-CCA controls. Discrimination analysis showed that a combination of these genes could be used as a diagnostic marker for CCA with better diagnostic parameters with high sensitivity and specificity than using a single gene marker or the usual serum markers (CA19-9 and CEA). This new combination marker may help physicians to identify CCA in liver tissues when the histopathology is uncertain.

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Introduction
Cholangiocarcinomas (CCA) are bile duct tumors which can be classified into intrahepatic CCA (ICC) and extrahepatic CCA. CCA is the second most common primary hepatobiliary malignancy, and its incidence is increasing globally [1]. The incidence of CCA is geographically unequal. It is high in Southeast Asia and low in Western countries [2]. The highest incidence of CCA was found in the Northeast Thailand where the liver fluke, Opisthorchis viverrini (Ov), is a major risk factor for CCA [3,4]. In Western and East Asian countries, the reported risk factors are chronic inflammation and cholestatic conditions, such as primary sclerosing cholangitis, cholelithiasis, and hepatitis C infection [5]. Complete resection is the current therapy of choice. However, most cases of CCA are diagnosed at advanced stages when surgery is no longer a feasible option. The accurate interpretation of a definite diagnosis is necessary so that a medical specialist can assess the severity of the disease and select the most suitable therapy for patients. At present, histological investigation is the standard diagnosis. However, there are some biopsy specimens and poorly-defined tumor tissues which cannot be definitively diagnosed by general histopathology. Hence, searching for a new diagnostic tool for these specimens is necessary.

In the past decade, many investigators have focused on the molecular and cellular perturbations which characterize the malignant phenotype. The power of a molecular signature in defining molecular phenotypes related to diagnosis, prognosis or treatment outcome was clearly seen in many studies. Several gene expression signatures have been reported for the tracking of true molecular phenotypes correlated with diseases, for example, in the classification of multiple sarcoma [6], in the outcome and chemotherapy response of ovarian cancer [7], and in the prediction of patient survival of gastric cancer [6,8].

At present, the availability of a rapid and formal proof of malignancy is still a constant goal in the diagnosis of CCA. In the current study, we sought to develop and validate a predictive model which can differentiate tumor mass commonly found in liver, ICC and hilar CCA with liver mass from HCC and normal liver tissues. An in-house PCR array containing 176 putative CCA marker genes was tested with the training set tissues of 20 CCA and 10 HCC cases, and 69 differentially expressed genes were optimized statistically to formulate a four-gene diagnostic equation...
which could distinguish CCA cases from HCC cases. Finally, we validated this equation in an independent testing set of 68 CCA samples and 77 non-CCA controls. This equation was successfully validated with a high sensitivity and specificity.

**Materials and Methods**

**Tissue Samples**

Frozen and paraffin embedded liver tissue-microarrays from patients with histologically confirmed CCA, HCC and chronic liver diseases were obtained from a specimen bank of the Liver Fluke and Cholangiocarcinoma Research Center, Faculty of Medicine, Khon Kaen University, Thailand. Written informed consent was obtained from each subject, and the study protocol was approved by the Ethics Committee for Human Research, Khon Kaen University. The diagnosis of benign hepatobiliary disease was based on clinical and histological records.

Frozen tumor tissues from CCA (n = 20) and HCC (n = 10) cases were used as the training set and the expression profiles were examined using the in-house PCR array. The characteristics of the CCA and HCC patients are summarized in Table S1. The testing set comprised 68 cases of CCA, 47 cases of HCC (Table S2), 21 cases of non-cancerous liver tissues, and nine cases with chronic biliary-liver diseases which were biliary hyperplasia (n = 2), haemangioma (n = 2), cystadenoma (n = 2), chronic inflammation (n = 2) and hepatolithiasis (n = 1).

**In-house PCR array and Primer Design**

An in-house PCR array with two duplicate sets of 191 genes was used as a single training dataset in a 348-well microplate. Each set of 191 genes contained 176 CCA associated genes, five internal controls (18S rRNA, ACTB, B2M, GAPDH and FDFT1) and two for HCC markers (AFP and GPC3). To ensure that the majority of cells in the tissue tested were CCA cells, several cell-type markers were included in the array; these were genes for biliary cells (KRT7 and KRT19), fibroblasts (ACTA2 and MME), hepatocytes (ALB and FGF), and white blood cells (ITGAL and PTPRC). These markers were selectively expressed for each cell type based on the SAGE database (http://cgap.nci.nih.gov/SAGE), B2M of pooled cDNA from normal liver tissues, CCA, and CCA cell lines, was used as an inter-refrence gene.

All specific primers were designed using the following guidelines: 1) for a gene which has more than one transcript variant, the design of the primer was based on the conserved region, 2) the length of the primer was 18–25 bp, 3) the length of the designed PCR product was 75–200 bp, and 4) the optimal melting temperature was 55°C. The specificity of the primers was tested using Primer-BLAST [9] and the conventional PCR for a single PCR product verification.

Approximately 2 μg of total RNA was reverse transcribed to cDNA by the High Capacity cDNA Reverse Transcription Kit (Applied Biosystems, Foster City, CA) according to the manufacturer's protocol. An in-house PCR array was prepared using a Biomek® NXP® Laboratory Automatic Workstation (Beckman Coulter, Fullerton, CA). Briefly, each primer pair was transferred into a 384-well white PCR plate. cDNA from each sample was mixed with 2× LightCycler® 480 SYBR Green I Master (Roche Applied Science, Mannheim, Germany) and then added to the plate. Real-time RT-PCR was performed using the LightCycler 480 II (Roche Diagnostics, Rotkreuz, Switzerland). All data were analyzed using the LightCycler® 480 SW 1.5 software. A combination of B2M and GAPDH were selected as the reference genes by NormFinder [10] and the geometric mean was used for normalising the quantities of mRNA species in each sample.

Hierarchical Cluster Analysis

Unsupervised hierarchical cluster analysis was used to explore the differential gene expression between the CCA and HCC samples in the training set. The expression level of each gene after normalization was transformed into a 2^-dCp value. The unsupervised hierarchical analysis was performed using dChip software [11]. Independent t-tests were performed to identify genes whose expressions in the CCA samples were significantly different from those in the HCC samples. Only genes whose expressions were found to be different at the P value <0.05 level were selected to formulate an equation for differentiating CCA from HCCs.

**Immunohistochemistry**

Two tissue-microarrays, one of triplicate samples from CCA patients (n = 28) and one of quadruplicate samples from HCC patients (n = 24) were subjected to a standard immunohistochemical staining according to manufacturers’ recommendations with the Envision Plus Detection Kit (Dako, Carpinteria, CA) for HOXB7, TMSB4, and TTR, and with the Histofine® Immunohistochemical staining reagent (Nichirei Biosciences Inc., Tokyo, Japan) for CLDN4. Tissue microarrays were treated with 1:50 anti-CLDN4 (Santa cruz biotechnology, CA), 1:100 anti-HOXB7 (Abnova, Taipei, Taiwan), 1:20000 anti-TMSB4 (abcam, Cambridge, MA), and 1:100 anti-TTR (Abnova, Taipei, Taiwan). Since almost all tissues had a similar positive frequency for immunoreactivity, the immunostaining was semi-quantitatively scored on the basis of intensity as: 0 = negative; 1+ = weak; 2+ = moderate; and 3+ = strong.

**Statistical analysis**

To select candidate genes for verification and the formulation of a diagnostic equation, the normalized gene expression data which were differentially expressed between CCA and HCC were subjected to a multiple linear regression analysis using STATA version 8.0 (Stata Corporation, College Station, TX).

**Results**

To prepare a gene set for the in-house PCR array, a search for CCA associated genes was conducted in 32 studies published between 2000 and 2009; three studies reported publicly available microarrays [12–14], one study reported SAGE data [15], and another an expression sequence tag [16]. Sixteen of twenty upregulated genes reported in microarray database obtained from Thai patients [13] were included in this study and finally, a total of 176 genes retrieved from 1,154 reported CCA cases were selected for an in-house PCR array. As advanced hilar CCA always invade the liver parenchyma and form large focal liver mass similar to HCC, therefore the hilar extrahepatic cholangiocarcinoma with liver mass were included in this study. In CCA cases, all tested samples exhibited a high expression of biliary cell markers and a low expression of hepatocyte markers; the reverse occurred for the HCC specimens.

Unsupervised Hierarchical Clustering of CCA and HCC

Using the in-house PCR array, we first examined the differential gene expression of tumor tissues from the CCA and HCC cases in the training set. An unsupervised hierarchical cluster analysis using the normalized gene expression data classified the 30 samples into two distinct groups (Fig. 1): one group contained nine HCCs and the other contained 20 CCA and one HCC. There were 69 differentially expressed genes: 26 genes in the CCA cases and 43 genes in the HCC cases (Table 1). The overexpressed genes in the CCA were associated with cell adhesion (e.g., SPP1, MMP7 and...
CLDN4) and cell movement (e.g., S100P, TMSB4 and S100A11). In contrast, the overexpressed genes in the HCCs were associated with xenobiotic metabolisms (ADH1B, ADH1C and ALDH1A1), biomolecule metabolisms (APOF, DPYD and GC), cell proliferation (IGF1 and ARID3A), differentiation (EGR1 and GPC3) and the transport of small molecules (ALB, AKR1C4 and TTR). The primer sequences of these genes are summarized in Table S3.

Discrimination Analysis

The next goal was to identify the individual genes or combinations of genes which were related to diagnosis in the training set. A multiple linear regression analysis was used to find the best models composed of the fewest number of genes for use as a diagnostic equation for discriminating between CCA and HCC tissues. From the 69 differentially expressed genes obtained in the hierarchical cluster analysis, an equation involving a combination of four genes, \[ Z = 1.232 - 0.761(\text{CLDN4}) - 7.09(\text{HOXB7}) + 0.221(\text{TMSB4}) + 0.055(\text{TTR}) \], gave the best discriminating power. In a receiver operating characteristic curve (ROC) analysis, this diagnostic equation yielded an area under the curve (AUC) of 0.98, and, when a Z-score of 1.23 was used as the optimal cut-off point to discriminate between CCA and HCC, the sensitivity and specificity were 90% and 100%, respectively (Figs. 2A–B). A Z-score of less than 1.23 indicated CCA rather than HCC. Hence the four-gene diagnostic equation was designated as “CCA diagnostic equation”.

Figure 1. Unsupervised hierarchical clustering of CCAs and HCCs.
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Validation of the “CCA diagnostic equation” in the testing set

To validate the accuracy of the “CCA diagnostic equation” in the discrimination of liver masses of CCA from HCC and noncancerous liver tissues, the expressions of CLDN4, HOXB7, TMSB4 and TTR were verified in the larger testing set by real-time PCR. Using the Z-score of 1.23 as the cut-off point, the “CCA diagnostic equation” could distinguish CCA patients from non-CCA cases with a sensitivity of 95.6% and specificity of 88.3%, and the AUC was 0.94 (Fig. 2C). Finally, we sought to determine whether the “CCA diagnostic equation” developed in the current study was a more powerful discriminator than an individual gene marker or the serum markers routinely used for the diagnosis of CCA, namely carbohydrate antigen19-9 (CA19-9) and carcinoembryonic antigen (CEA). We first inspected the ROC analysis of the individual gene marker to obtain the best cut-off value from the training set, and then verified the diagnostic value in the testing set (n = 145). For serum markers, the diagnostic parameters were compared in the CCA, HCC and chronic biliary-liver cases whose serum CA19-9 and CEA were recorded (n = 66). As shown in Table 2, the “CCA diagnostic equation” gave better diagnostic results than a single gene marker. In addition, the equation also yielded a much better sensitivity, negative predictive value and false negative rate than the use of serum CA19-9 and CEA (Table 3).

Validation by immunohistochemical staining

To establish that the four genes in the diagnostic equation reflected CCA and HCC tissue, we verified the expression levels of CLDN4, HOXB7, TMSB4 and TTR in the tumor tissues of CCA (n = 28) and HCC (n = 24) using immunohistochemistry. Compared with HCC, CCA expressed significantly higher levels of

| Table 1. List of top 20 overexpressed and 10 underexpressed genes in 20 CCA tissues. |
|-----------------------------------|------------------|------------------|
| No | UniGene ID | Symbol | Title | Mean* | CCA As | HCCs | P value |
|----|------------|--------|-------|-------|--------|-------|---------|
| **Overexpressed genes** | | | | | | |
| 1 | BG571732 | S100P | S100 calcium binding protein P | 2.2859 | 0.5445 | 0.0060 |
| 2 | NM_001251830 | SPP1 | secreted phosphoprotein 1 | 1.7711 | 0.1281 | 0.0002 |
| 3 | BQ688566 | MMP7 | matrix metallopeptidase 7 (matrilysin, uterine) | 1.3315 | 0.0151 | 0.0003 |
| 4 | BM923753 | TFF1 | trefoil factor 1 | 1.3116 | 0.0001 | 0.0001 |
| 5 | BC023552 | SFN | Stratifin | 1.1101 | 0.1443 | 0.0017 |
| 6 | BM926728 | GSTP1 | glutathione S-transferase pi1 | 0.9891 | 0.0630 | <0.0001 |
| 7 | BF680512 | TMSB4 | thymosin beta 4, X-linked | 0.3832 | 0.2333 | 0.0453 |
| 8 | BQ683841 | S100A11 | S100 calcium binding protein A11 | 0.2753 | 0.0469 | 0.0004 |
| 9 | AK074480 | ANXA1 | annexin A1 | 0.1859 | 0.0361 | 0.0019 |
| 10 | NM_001305 | CLDN4 | claudin 4 | 0.1632 | 0.0260 | 0.0009 |
| 11 | NM_002483 | CEACAM6 | carcinoembryonic antigen-related cell adhesion molecule 6 (non-specific cross reacting antigen) | 0.1330 | 0.0005 | <0.0001 |
| 12 | AL832780 | TM4SF1 | transmembrane 4 L six family member 1 | 0.0723 | 0.0072 | 0.0002 |
| 13 | NM_003816 | ADAM9 | ADAM metallopeptidase domain 9 | 0.0721 | 0.0120 | 0.0045 |
| 14 | NM_021102 | SPINT2 | serine peptidase inhibitor, Kunitz type, 2 | 0.0370 | 0.0004 | <0.0001 |
| 15 | XS2228 | MUC1 | mucin 1, cell surface associated | 0.0365 | 0.0006 | <0.0001 |
| 16 | NM_003870 | IQGAP1 | IQ motif containing GTPase activating protein 1 | 0.0308 | 0.0066 | 0.0002 |
| 17 | AK128505 | KRT7 | keratin 7 | 0.0150 | 0.0018 | 0.0012 |
| 18 | AK223249 | HOXB7 | homeobox B7 | 0.0057 | 0.0033 | 0.0453 |
| 19 | BM904612 | S100A6 | S100 calcium binding protein A6 | 0.0013 | 0.0008 | 0.0165 |
| 20 | NM_182848 | CLDN10 | claudin 10 | 0.0012 | 0.0000 | 0.0366 |
| **Underexpressed genes** | | | | | | |
| 1 | NM_000477 | ALB | albumin | 3.7085 | 71.6658 | <0.0001 |
| 2 | NM_021870 | FGG | fibrinogen gamma chain | 0.9397 | 20.2895 | <0.0001 |
| 3 | BM942013 | TTR | transthyretin | 0.2005 | 7.9316 | <0.0001 |
| 4 | NM_001904 | CTNNB1 | catenin (cadherin-associated protein), beta 1, 88kDa | 1.7146 | 3.4423 | 0.0453 |
| 5 | NM_001633 | AMBP | alpha-1-microglobulin/bikunin precursor | 0.0746 | 2.1106 | <0.0001 |
| 6 | AF130100 | SERPINC1 | serpin peptidase inhibitor, clade C (antithrombin), member 1 | 0.0414 | 1.8204 | <0.0001 |
| 7 | MS58569 | FGA | fibrinogen alpha chain | 0.1090 | 1.8176 | 0.0001 |
| 8 | NM_001689 | ALDH1A1 | aldehyde dehydrogenase 1 family, member A1 | 0.1674 | 1.7379 | 0.0001 |
| 9 | NM_001164617 | GPC3 | glypican 3 | 0.0049 | 0.3247 | 0.0008 |
| 10 | BC027881 | AFP | alpha-fetoprotein | 0.0002 | 0.2638 | 0.0002 |

*mean of expression level of individual gene after normalization with the geometric mean of B2M and GAPDH and transformed into a 2^-dCp.

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CLDN4 ($P<0.001$) and HOXB7 ($P<0.01$), a similar expression of TMSB4, but a lower level of TTR ($P<0.02$) (Fig. 3).

**Table 2.** Comparison of diagnostic indices between each single marker and “CCA Diagnostic equation”.

| Diagnostic indices (%) | Tissue mRNA marker (n = 145) | “CCA Diagnostic eq.” |
|------------------------|-------------------------------|----------------------|
|                        | CLDN4 | HOXB7 | TMSB4 | TTR | Sensitivity | Specificity | Positive predictive value | Negative predictive value | Fault positive | Fault negative | AUC   |
| Sensitivity            | 86.8  | 85.3  | 39.7  | 95.6 | 95.6        | 72.7       | 53.2               | 70.1               | 79.2          | 88.3         | 0.87  |
| Specificity            | 73.8  | 61.7  | 54.0  | 80.2 | 87.8        | 72.7       | 53.2               | 70.1               | 79.2          | 88.3         | 0.77  |
| Positive predictive value | 86.2  | 80.4  | 56.8  | 95.3 | 95.8        | 73.8       | 61.7               | 54.0               | 80.2          | 87.8         | 0.56  |
| Negative predictive value | 14.5  | 24.8  | 15.9  | 11.0 | 6.2         | 6.2        | 6.9                | 28.3               | 2.1           | 2.1          | 0.91  |
| Fault positive         | 6.2   | 6.9   | 28.3  | 2.1  | 2.1         | 6.2        | 6.9                | 28.3               | 2.1           | 2.1          | 0.91  |
| Fault negative         | 0.87  | 0.77  | 0.56  | 0.91 | 0.94        | 0.87       | 0.77               | 0.56               | 0.91          | 0.94         | 0.94  |

AUC = area under curve of the receiver operating characteristic curve; CLDN4, claudin 4 (cut-off value was 0.01); HOXB7, homeobox B7 (cut-off value was 0.001); TMSB4, thymosin beta 4, X-linked (cut-off value was 0.08); TTR, transthyretin (cut-off value was 0.30); eq., equation.

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**Table 3.** Comparison of diagnostic indices between serum markers and “CCA Diagnostic equation”.

| Diagnostic indices (%) | Serum marker (n = 66) | “CCA Diagnostic eq.” |
|------------------------|-----------------------|----------------------|
|                        | CA19-9 | CEA | Sensitivity | 44.4 | 38.9 | 97.2 |
| Specificity            | 100.0  | 100.0 | 86.7 |
| Positive predictive value | 100.0 | 100.0 | 89.7 |
| Negative predictive value | 65.2  | 57.7 | 96.3 |
| Fault positive         | 0.0    | 0.0   | 6.1 |
| Fault negative         | 54.5   | 54.5  | 1.5 |
| AUC                    | 0.67   | 0.88  | 0.94 |

AUC = area under curve of the receiver operating characteristic curve; eq., equation; CA19-9, carbohydrate antigen 19-9 (cut-off value was 100 U/ml); CEA, carcinoembryonic antigen (cut-off value was 22 μg/ml).

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**Discussion**

The accurate and definitive diagnosis of a tumor is necessary for a medical specialist to assess the severity of the disease and select appropriate treatment options.
the most suitable therapy. Histopathological diagnosis is the routine, standard method of diagnosing a solid tumor. However, there are some biopsy specimens and ill-defined tumor tissues which cannot be definitively diagnosed by general histopathology. Currently, the molecular signature characterized the malignant phenotype is usually reported. In this study, we developed and validated a model of gene expression which can distinguish liver tissues of CCA from HCC and benign biliary-liver diseases. A “CCA diagnostic equation” involving four genes (CLDN4, HOXB7, TMSB4 and TTR) was formulated for the diagnosis of CCA with high sensitivity and specificity.

The unsupervised hierarchical cluster analysis retrieved from the in-house PCR array of the well-defined samples in the training set could initially separate patients into two groups: CCA and HCC cases. The results indicated that CCA cases share a characteristic gene expression profile which is distinguishable from HCC by a small subset of genes. The molecular functions and biological processes of the overexpressed genes in CCA were involved in the regulation of cell adhesion and migration. Conversely, the overexpressed genes in HCC were associated with xenobiotic and biomolecule metabolisms. Our PCR array profiling data are consistent with those reported using SAGE and oligonucleotide microarray analyses which specified the upregulation of genes associated with cell adhesion molecules in CCA [14,15].

The gene expression profiling results have also yielded lists of genes which are potential biomarkers for diagnosis. We first identified the best models for discriminating CCA from HCC in the training set using multiple linear regression analysis. After extensive cross-validation, a combination of CLDN4, HOXB7, TMSB4 and TTR designated as “CCA diagnostic equation”, yielded the best equation for differentiating CCA from HCC (Fig. 2, Table 2). In our experience, majority of the advanced hilar CCAs always invade the liver parenchyma and form large focal liver mass indistinguishable from ICC and HCC. In this study, the liver masses from hilar CCAs were included in both training set and testing set. Regardless to CCA origin, the “CCA diagnostic equation” can differentiate the tumor masses of ICC and hilar CCA from HCC.

Figure 3. Immunohistochemistry of CLDN4, HOXB7, TMSB4, and TTR in tumor tissues from CCA and HCC patients. (A) CLDN4, HOXB7 and TMSB4 were obviously expressed in CCA tissues while TTR was strongly expressed in HCC tissues. (B) The expressions of CLDN4, HOXB7, TMSB4, and TTR were quantified based on the intensity. *$P$ value<0.05. HCC = hepatocellular carcinoma; CCA = cholangiocarcinoma.

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The immunohistochemistry of the four diagnostic genes expressed in CCA and HCC tissues revealed a higher expression of CLDN4, HOXB7 in CCA tissues and a higher expression of TTR in HCC tissues. The formulated equation was further validated in the testing set with a larger sample size and a variety of non-CCA controls. Sensitivity (the true positive rate) and specificity (the true negative rate) are statistical measures of a binary classification test performance. A perfect predictor would be described as 100% sensitivity and 100% specificity which can be represented graphically as a ROC with AUC = 1. In the present study, the AUC of each test demonstrated that the “CCA diagnostic equation” could effectively diagnose CCA cases from the controls with a higher sensitivity and specificity compared with those of individual gene analysis. The use of the “CCA diagnostic equation” was also superior to the use of known serum markers; CA19-9 and CEA as the “CCA diagnostic equation” gave the highest AUC compared to serum those of CA19-9 and CEA. Since no correlation was found between these serum markers and the “CCA diagnostic equation” (data not shown), the inclusion of serum markers and molecular markers “CCA diagnostic equation” may increase its diagnostic power, and this needs to be investigated.

The better diagnostic parameters obtained from the “CCA diagnostic equation” formulated in the present study are consistent with the finding of a previous study [15] in which a different diagnostic equation was reported to improve both the sensitivity and specificity of the diagnosis of ICC when compared with the use of CA19-9 and CEA. A comparison of the gene expression profiles in parasite-associated (Thai patients) and non-parasite-associated (Japanese patients) human ICCs demonstrated different molecular signatures between the two sample groups [13]. An elevated expression of genes involved in xenobiotic metabolism was found in the parasite-associated ICCs whereas genes related to growth factor signaling were shown in the non-parasite-associated ICCs. These findings may explain the difference in the set of genes formulated for the diagnostic equation in our study and that reported by Nishino [15]. As a consequence, one should aware of possible limitation of our study, namely that the vast majority of reported by Nishino [15]. As a consequence, one should aware of possible limitation of our study, namely that the vast majority of CCA from other countries. An additional challenge is to explore the possibility of using the “CCA diagnostic equation” to diagnose the non-liver fluke ICC from other countries.

Supporting Information
Table S1 Clinicopathological features of samples used for training set.
(DOC)
Table S2 Clinicopathological features of samples used for testing set.
(DOC)
Table S3 Primer sequences data.
(DOC)

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Author Contributions
Conceived and designed the experiments: RK SW CW. Performed the experiments: RK KL. Analyzed the data: RK SW CW. Contributed reagents/materials/analysis tools: NK CW SW. Wrote the paper: RK KL NK CP SW CW.

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