

Table S1. Sorting of the 26 HCC studied according to HBsAg status.

| HBsAg status | NT/T | SEX | AGE | PATHOLOGY | | TP53 status | p53 immuno-staining | CTNNB1 status | HCV RNA | HBV | | | |
|--------------|------|-----|-----|-------------|----------------|-------------|---------------------|---------------|---------|-----|--------------------|-----------------|---------|
| | | | | TUMOR GRADE | META VIR SCORE | | | | | DNA | Genotype, serotype | Mutation (NT/T) | |
| | | | | | | | | | | | | BCP | PreCore |
| - | 1 | M | 46 | G4 | A2 F4 | G331H | - | WT | + | + | C, adr | WT+DV/DV | WT/WT |
| | 2 | M | 40 | G1 | A0 F2 | WT | - | WT | - | + | NA | WT+DV/WT+DV | WT/WT |
| | 3 | F | 37 | G2 | A0 F1 | WT | - | WT | - | + | B, adw2 | NA | NA |
| | 4 | M | 61 | G2-3 | A2 F4 | WT | - | T41A | - | + | C, adr | NA/WT | WT/WT |
| | 5 | M | 73 | G3 | NA | WT | - | S37F | - | + | NA | NA | NA |
| | 6 | M | 70 | G2 | NA | WT | - | WT | - | - | - | - | - |
| | 7 | M | 39 | G3-4 | A0 F3 | WT | - | WT | - | - | - | - | - |
| | 8 | F | 50 | G2-3 | A0 F1 | WT | 20-50% | WT | - | - | - | - | - |
| | 9 | F | 35 | G2-3 | NA | WT | <10% | NA | - | - | - | - | - |
| | 10 | M | 53 | G3 | A0 F2 | WT | 10-20% | WT | + | - | - | - | - |
| + | 11 | M | 34 | G4 | A1 F4 | R249S | <10% | WT | + | + | C, adr | DV/DV | WT/WT |
| | 12 | M | 43 | G1-2 | A2 F4 | R249S | <10% | S41F | - | + | C, adr | NA | NA |
| | 13 | M | 50 | G3-4 | A0 F1 | R249S | <10% | WT | - | + | B, adw2 | NA | NA |
| | 14 | M | 44 | G3 | NA | R249S | <10% | WT | - | + | C, adr | NA | NA |
| | 15 | M | 60 | G3-4 | A2 F4 | P278R | 10-20% | S33Y | - | + | C, NA | DV/WT+DV | WT/WT |
| | 16* | M | 42 | G2 | A0 F2 | WT | - | NA | - | + | NA | WT+DV/WT+DV | WT/WT |
| | 17 | M | 39 | G2 | A1 F4 | WT | - | WT | + | + | NA | NA/DV | WT/WT |
| | 18 | M | 62 | G2-3 | A1 F3 | WT | - | WT | - | + | C, adr | DV/WT+DV | MUT/WT |
| | 19* | M | 36 | G4 | A1 F2 | WT | - | NA | - | + | C, NA | NA | NA |
| | 20 | M | 37 | G3-4 | A0 F1 | WT | - | WT | - | + | C, NA | NA | NA |
| | 21 | M | 57 | G2-3 | A1 F3 | WT | - | WT | - | + | NA | NA | NA |
| | 22 | M | 17 | G3 | NA | R249S | <10% | S33Y | - | - | NA | NA | NA |
| | 23* | M | 40 | G2-3 | NA | WT | - | NA | - | - | NA | NA | NA |
| NA | 24* | NA | NA | G3-4 | A0 F2 | R249S | <10% | NA | - | + | NA | NA | NA |
| | 25 | NA | NA | G1 | A0 F4 | R249S | <10% | S37F | - | + | NA | NA | NA |
| | 26 | F | 48 | NA | A0 F1 | WT | - | WT | - | + | NA | WT/WT | WT/WT |

-=negative, +=positive, NA=not available, NT=non-tumoral tissue, T=tumoral tissue, WT=wild-type, DV=dual variant (A1764G/T1762A), /=no sample, *=four HBV-positive cases with non-available CTNNB1 status were excluded in Fig. 3. p53 immunostaining is given for T samples as it was negative for all the NT tissues.#

Table S2. DHPLC conditions for exon 3 of *CTNNB1* gene.

| Exon | Temperature (°C) | Acetonitrile gradient (%B) | Positive controls |
|------|------------------|----------------------------|--|
| 3 | 60 | 52-60 | H358S (codon 75: <u>A</u> CT- <u>G</u> CT) |
| | 60 | 52-60 | HCT116 (codon 45: deleted) |
| | 63 | 49-57 | SW48 (codon 33: T <u>C</u> T- T <u>A</u> T) |

Temperature and gradient analyzes were determined by Transgenomic software.

Forward primer (β_f) 5'-ccaatctactaatgctaatactg-3', reverse primer (β_r) 5'-
ctgcattctgactttcagtaagg-3'

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| | 120 | 130 | 140 | 150 |
|---------|----------|-------------------|------------|------------------|
| THAI6 | GPCKTCTI | PAQGTSMF | PSCCCTKPSD | GNCTCIPI |
| THAI10 | GPCKTCT | N PAQNTSMF | PSCCCTKPSD | R NCTCTPI |
| THAI26* | GPCKTCT | A TAQGTSMF | PSCCCTKPTD | GNCTCIPI |
| THAI27 | GPCKTCTI | PAQGTSMF | PSCCCTKPSD | R NCTCIPI |
| THAI30 | GPCKTCTI | PAQGTSMF | PSCCCTKPSD | GNCTCIPI |
| THAI34 | GPCKTCTI | PAQGTSMF | PSCCCTKPSD | GNCTCIPI |
| THAI36 | GPCKTCTI | PAQGTSMF | PSCCCTKPSD | GNCTCIPI |
| THAI55 | GPCKTCTI | PAQGTSMF | PSCCCTKPSD | GNCTCIPI |
| THAI66 | GPCKTCTI | PAQGTSMF | PSCCCTKPSD | GNCTCIPI |
| THAI89* | GPCKTCTT | PAQGTSMF | PSCCCTKPTD | GNCTCIPI |
| THAI96 | GPCKTCTT | PAQGTSMF | PSCCCTKPSD | GNCTCIPI |

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Figure S3. Alignments of HBsAg in determinant “a” region from 11 HBV patients (classified as genotype C except samples marked with stars which are genotype B). Samples 10, 26 and 27 are occult HBV infections. *R145G* is a common mutation causing false negative results for HBV serological testing using first generation antibodies. Double mutant *A126I / T127P* was detected in case 26.

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