Supplementary Material

# Supplementary Tables

**Table S1| Antibody clones, catalogue numbers and manufacturer’s details**

| **Antibody** | **Clone** | **Catalogue #** | **Species** | **Company** |
| --- | --- | --- | --- | --- |
| **ALDH1** | 44 | 611195 | Mouse | BD Biosciences™ |
| **AR** | AR441 | M3562 | Mouse | Dako | Agilent |
| **Bcl2** | 124 | MO887 | Mouse | Dako | Agilent |
| **CD44** | DF1485 | M7082 | Mouse | Dako | Agilent |
| **CD24** | SN3b | MS1279 | Mouse | ThermoFisher Scientific |
| **CK5** | XM26 | NCL-L-CK5 | Mouse | Leica Biosystems |
| **CK14** | LL002 | NCL-L-LL002 | Mouse | Leica Biosystems |
| **COX2** | CX-294 | M3617 | Mouse | Dako | Agilent |
| **Ki-67** | MIB-1 | M7240 | Mouse | Dako | Agilent |
| **Nestin** | 10c2 | sc-23927 | Mouse | Santa Cruz Biotechnology, Inc |

**Table S2| Details of the positive control tissues, retrieval buffer, dilution and incubation time of the antibodies**

| **Antibody** | **Positive control tissues** | **Target Retrieval Solution\***  | **Dilution** | **Incubation time (min)** |
| --- | --- | --- | --- | --- |
| **ALDH1** | IDC\*\* /liver | Low pH | 1:750 | 60 |
| **AR** | Luminal cells of BPH\*\*\* | Low pH | 1:100 | 60 |
| **Bcl2** | Tonsil | High pH | 1:250 | 60 |
| **CD44** | IDC | Low pH | 1:100 | 60 |
| **CD24** | Normal epithelium of small and large bowel /colon Ca | High pH | 1:100 | 60 |
| **CK5** | Myoepithelial cells of normal breast ducts | Low pH | 1:100 | 60 |
| **CD14** | Myoepithelial cells of normal breast ducts | Low pH | 1:100 | 60 |
| **COX2** | Basal cells in ductal hyperplasia | High pH | 1:100 | 60 |
| **Ki-67** | IDC nuclei | High pH | 1:100 | 60 |
| **Nestin** | Myoepithelial cells of normal breast ducts/IDC | Low pH | 1:100 | 60 |

\*Agilent |Dako; \*\*IDC= Infiltrating ductal carcinoma; \*\*\* BPH=benign prostatic hyperplasia

**Table S3| Scoring criteria for IHC expression**

| **Marker**  | **Expression pattern**  | **Scoring criteria**  | **Reference**  |
| --- | --- | --- | --- |
| **ALDH1**  | Cytoplasmic  | Percentage (***P)*** and intensity (***I***) of cytoplasmic expression was recorded:***Intensity (I) of Expression***: 0=negative; 1=weak; 2=moderate; 3=strong ***ALDH1 Score (S) = P x I***For statistical analysis: Score =0 (Negative) Score >0 (Positive) | [1] |
| **AR** | Nuclear | **Allred Score:*****Proportion Score (PS):***0=0; 1=1/100; 2=1/10; 3=1/3; 4=2/3; 5= 1***Intensity Score (IS):***0= Negative; 1= Weak; 2 = Intermediate; 4= Strong ***Allred Score= PS + IS*** Negative: ≤ 2Weak: 3-4Intermediate: 5-6Strong: 7-8 | [2] |
| **Bcl-2** | Cytoplasmic | Positive: ≥10% cytoplasmic expression Negative: <10% cytoplasmic expression | [3] |
| **CD44** | Membranous | 0=no expression; 1=1-10% positive tumor cells; 2=11-50%; 3=51-75%; 4=76-100% | [4] |
| **CD24** | Cytoplasmic | 0=no expression; 1=1-10% positive tumor cells; 2=11-50%; 3=51-75%; 4=76-100% | [4] |
| **CK5** | Cytoplasmic | Positive expression: Any proportion of cytoplasmic expression in tumor cells  | [5] |
| **CK14** | Cytoplasmic | Positive expression: Any proportion of cytoplasmic expression in tumor cells  | [5] |
| **COX2** | Cytoplasmic  | Weighted score was computed based upon the percentage of tumor positivity and intensity as follows: | [6] |
| ***Score Percent Staining (%)***0 <51 5-252 26-503 51-754 >75***Score Intensity***0 Absent1+ Weak2+ Medium3+ Strong ***Weighted Score = % of tumor cell positivity x intensity:***Negative: 0-4Intermediate: 5-8High: 9-12 |
| **Ki-67** | Nuclear | At least 500 tumor cells were counted, and a threshold of >25% nuclear expression was considered to be positive, irrespective of the staining intensity.  | [7] |
| **Nestin** | Cytoplasmic /membranous  | Expression was defined as either positive (at least 3 clearly positive tumor cells) or negative. | [8] |

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4. Honeth, G., et al., *The CD44+/CD24- phenotype is enriched in basal-like breast tumors.* Breast Cancer Res, 2008. **10**(3): p. R53.

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8. Kruger, K., et al., *Expression of Nestin associates with BRCA1 mutations, a basal-like phenotype and aggressive breast cancer.* Sci Rep, 2017. **7**(1): p. 1089.