

# MLiB: Interpretable model-based data integration for cancer type classification

## Supplementary Material

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**Table 1:** Permutation importances of features which appeared in the 500 most important genes of both GE and CN. ACSM3 was the only feature for which has a positive feature importance in both datasets. Each feature was permuted ten times. The cancer associations column shows a short description of some papers which found evidence linking each feature to cancer. The GE and CN columns show the permutation importance of each feature as the decrease in  $F_1$ -score (%). \* = Non-cancer research. Unless marked otherwise, all research was done *in vitro*.

Gene	GE	CN	Cancer associations
ACSM3	$0.269 \pm 0.135$	$0.059 \pm 0.048$	Deregulated in hepatocellular carcinoma [1].
ALG12	$0.000 \pm 0.000$	$-0.126 \pm 0.127$	Elevated expression in ovarian cancer [2].
ATP2B2	$0.000 \pm 0.000$	$-0.058 \pm 0.047$	Can be upregulated in breast cancer [3]–[5].
CCL23	$0.135 \pm 0.165$	$0.000 \pm 0.000$	Has pro- and anti-cancer properties w.r.t. to cancer cell interaction [6].
CDH15	$-0.233 \pm 0.233$	$0.071 \pm 0.058$	Expression level serves as a breast cancer prognosis marker [7], [8].
C20orf201	$0.000 \pm 0.000$	$0.000 \pm 0.000$	Cancer testis antigen that could be a useful cancer biomarker [9].
LDLRAD3	$0.338 \pm 0.391$	$-0.003 \pm 0.010$	Biomarker for pancreatic cancer diagnosis [10].
NCLN	$0.303 \pm 0.101$	$-0.045 \pm 0.054$	*Knock-out of <i>C. elegans</i> homolog enhances induced neuronal death [11].
NLGN1	$0.000 \pm 0.000$	$-0.504 \pm 0.331$	Upregulation in colorectal cancer [12].
PTPN1	$0.000 \pm 0.000$	$0.037 \pm 0.037$	( <i>in silico</i> ) expression levels correlated with incidence of esophageal cancer [13].
SF3A2	$0.255 \pm 0.354$	$0.000 \pm 0.000$	Abnormal expression in bladder cancer [14].
SPPL2B	$0.034 \pm 0.101$	$-0.019 \pm 0.056$	Expression levels associated with breast cancer [15].
UBE2E2	$0.135 \pm 0.165$	$-0.003 \pm 0.010$	Upregulation in ovarian cancer [16].

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