# REPORT

Analysis of binding pattern of commercially available antibodies on hEXselect Arrays



### Your Question. Our Service. Your Result.





**DETERMINATION OF ANTIBODY SPECIFICITY** by analysis of >21,000 antigens in one shot. In an internal project we have examined three commercially available antibodies with the engine hEXselect arrays (product No. 1003).

We analyse following antibodies:

- (polyclonal goat) anti-mouse IgG (Fc specific)-AP, Sigma A2429
- $\rho$  (monoclonal mouse) anti-human PP2A-A $\alpha$ / $\beta$  (clone A5), Santa Cruz sc74580
- (monoclonal mouse) anti-human Karyopherin β1, Santa Cruz sc365299

We identified specific and several unspecific bindings.

### Properties of our hEXselect Array

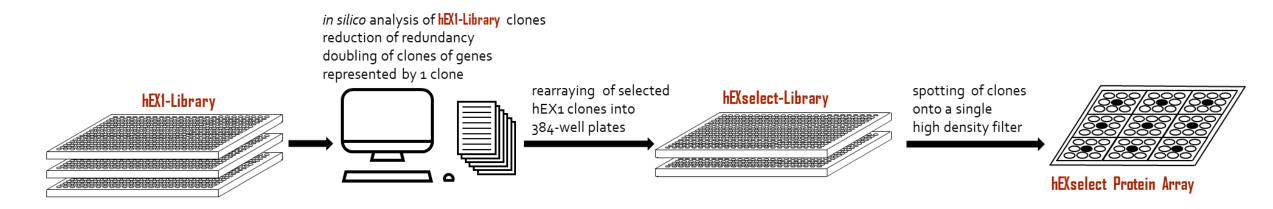




- Proteins are expressed by clones derived from a human fetal brain cDNA library
- Mixture of partially de- and renatured, full length proteins and peptides
- ORF: Mix of in-frame and out-of-frame proteins, 5' and 3' UTRs
- Proteins and peptides are expressed with an n-terminal RGS-His6-tag
- Spotting in 5x5 pattern
- 23,806 clones representing 6,909 different human proteins
- Product Number: 1003

### Creation of hEXselect Array

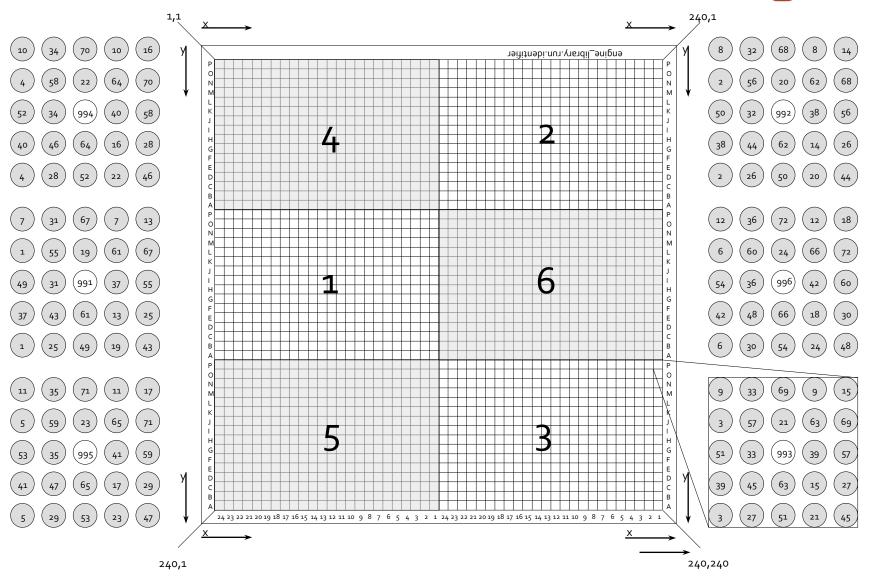




- hEX1-Library clones were analysed in silico
- Redundancy was reduced
- For unique Genes (represented by only one clone), the clone was doubled
- In order to fit onto one Array, the total number of clones was reduced.
- Clones were rearrayed into the final hEXselect-Library

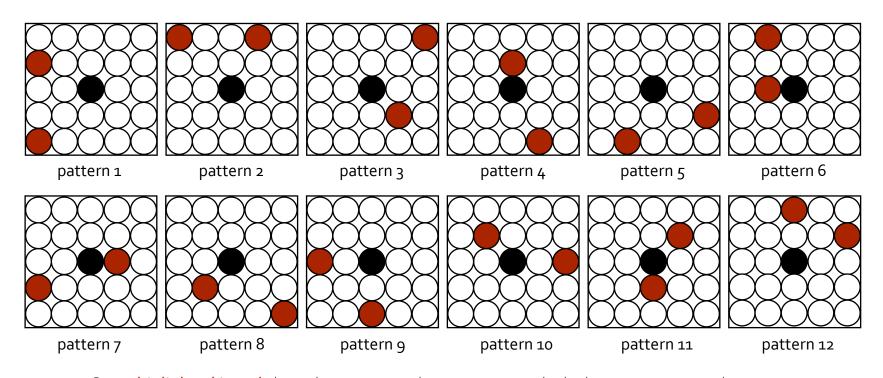
### ArrayDesign\_240x240





### Scoring Pattern on hEXselect Protein Arrays





- Spots higlighted in red show the corresponding pattern, in which clones were spotted
- A clone is only regarded as positive, if both spots can be detected
- Black spots are guiding dots

### engine Protein Arrays – as simple as a Western Blot

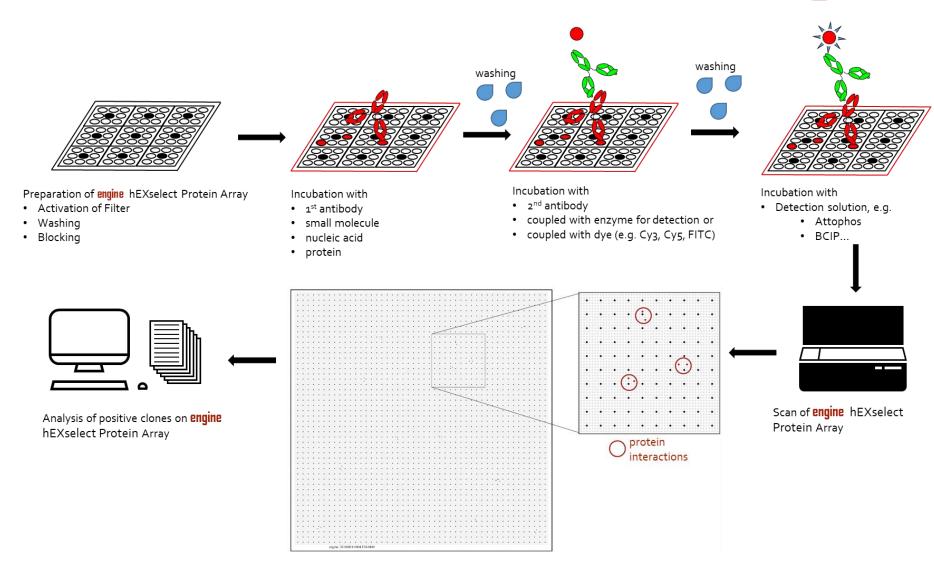




- Activation of array (PDVF membrane)
- Removal of E. coli debris with Kimwipe tissue
- Washing
- Blocking
- Incubation with sample (serum, antibody etc.)
- Washing
- Incubation with secondary antibody (AP-labeled)
- Detection with AttoPhos® substrate
- Software-based evaluation

# Detection of positives hits at a glance





### Detection of positives hits at a glance



#### Scoring (circle colour):

Red - 3 = strong

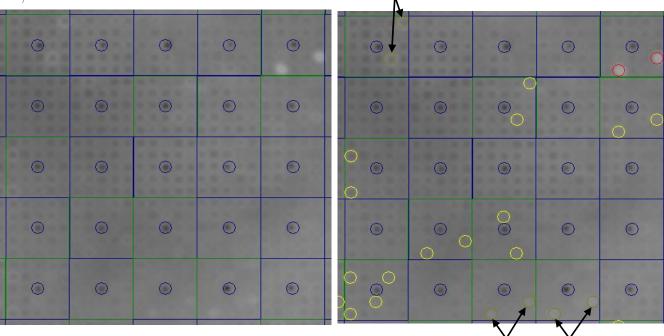
Brown -2 = moderate

Yellow - 1 = low

As brown circles of score 2 are poorly visible, these are marked by arrows in addition

#### Example for Scoring:

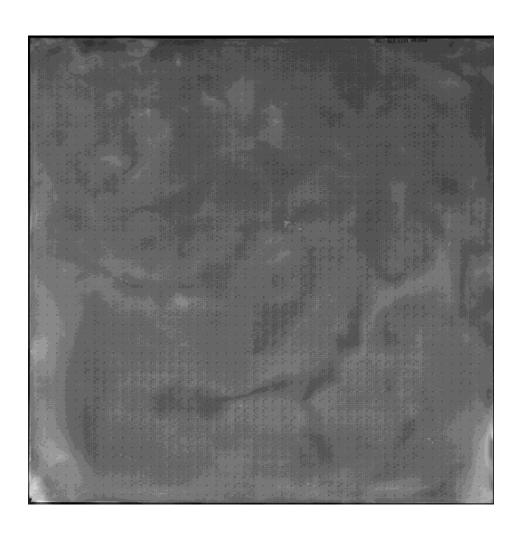
The same section of an Array was cropped and is shown without (left) and with (right) scoring of positive hits (bright spots), as displayed by the software.



- Hits, which were clearly detectable by the analysis software at a zoom of 50%, were marked with a score of "3"
- Hits, which were clearly detectable by the analysis software at a zoom of 200%, were marked with a score of "2"
- Hits, which were detectable by the analysis software but not standing out prominently at a zoom of 200%, were marked with a score of "1"

### Results – Secondary antibody control





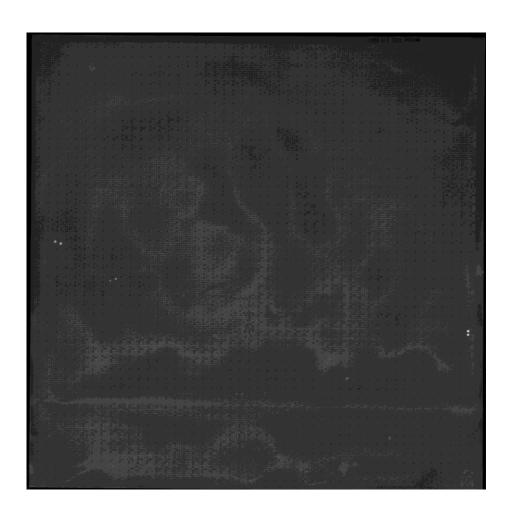
- Sample: none secondary antibody control
- Sec antibody: anti mouse IgG (Fc spec)-AP 1:10.000
- Array: engine\_1003\_179\_0700

#### Positives total: 33

visual score of spot intensity		#
Intensity 1	low	24
Intensity 2	moderate	9
Intensity 3	strong	0

### Results – anti human KPNB1





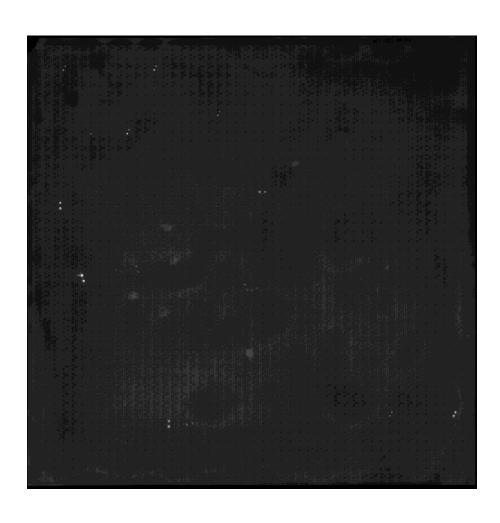
- Sample: anti human KPNB1 antibody,
   3 μg total (15 μl/50 ml)
- Sec antibody: anti mouse IgG (Fc spec)-AP 1:10.000
- Array: engine\_1003\_179\_0600

#### Positives total: 17

visual score of spot intensity		#
Intensity 1	low	12
Intensity 2	moderate	2
Intensity 3	strong	3

### Results – anti human PPP2R1A





- Sample: anti human PPP2R1A (clone A5) antibody, 3 μg total (15 μl/50 ml)
- Sec antibody: anti mouse IgG (Fc spec)-AP 1:10.000
- Array: engine\_1003\_179\_0300

#### Positives total: 39

visual score of spot intensity		#
Intensity 1	low	21
Intensity 2	moderate	10
Intensity 3	strong	8

### Summary





For all samples, antibody binding to antigens at the protein array could be detected.

#### Distribution of hits and intensity of hits:

		visual score of spot intensity		
	total	1 low	2 moderate	3 strong
anti mouse IgG (Fc-spec)	33	24	9	0
anti Karyopherin β1	17	12	2	3
anti PP2A-Aα/β	39	21	10	8

# Summary anti mouse IgG



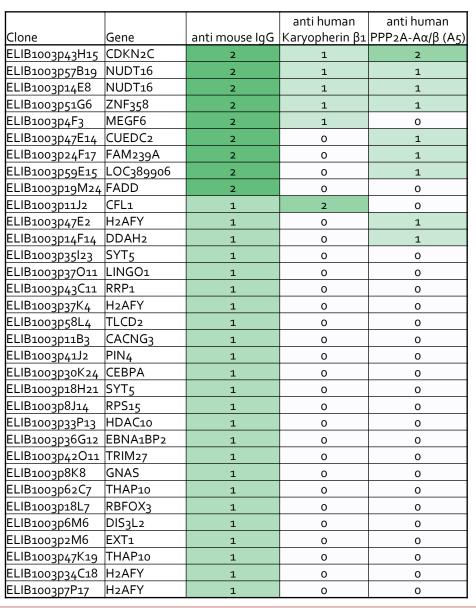




Table shows all antigens which reacts with secondary anti mouse IgG **and** overlapping reactions with anti KPNB1 and anti PPP2R1A

Determination of 33 off-target activities for commercial Fc specific anti mouse IgG!

# Summary anti KPNB1





		anti human	anti human
Clone	Gene	Karyopherin b1	PPP2A-Aa/b (A5)
ELIB1003p36A	1 KPNB1	3	0
ELIB1003p25G	16 <b>KPNB1</b>	3	0
ELIB1003p16C	2 KPNB1	1	0
ELIB1003p43K	22 DNAJC2	3	0
ELIB1003p9H2	2 ZNF511	2	0
ELIB1003p44H	l2 ANKRD16	1	1
ELIB1003p35l1	4 CCDC130	1	0
ELIB1003p20H	l24 RTN1	1	0
ELIB1003p21l2	1 WASHC1	1	0
ELIB1003p33l2	o HDAC10	1	0
ELIB1003p20E	3 HIST1H1C	1	0
ELIB1003p64L	11 KPNB1	0	2
ELIB1003p41G	7 KPNB1	0	2

Table shows all antigens which reacts with anti KPNB1 antibody and overlapping reactions with anti PPP2R1A, excluding secondary antibody binding.

#### Determination of

- 3 specific reactions (three different KPNB1 clones at the array) but there is no reaction with 2 other KPNB1 clones
- 8 off-target activities (strong spot intensity with DNAJC2 and moderate spot intensity with ZNF511)
- 1 overlapping reaction with anti PPP2R1A (ANKRD16)

# Summary anti PPP2R1A



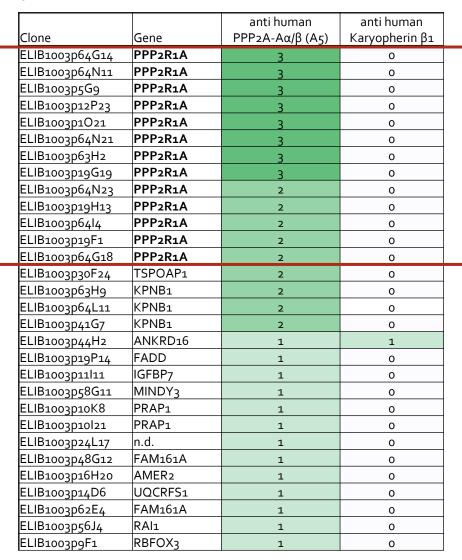




Table shows all antigens which reacts with anti PPP2R1A antibody and overlapping reactions with anti KPNB1, excluding secondary antibody binding.

#### Determination of

- 13 specific reactions

   (13 different PPP2R1A clones at the array)
- 17 off-target activities (4 with moderate spot intensity)
- 1 overlapping reaction with anti KPNB1 (ANKRD16)

### Conclusion





We could detect **specific reactions** for anti KPNB1 and anti PPP2R1A.

All antibodies have **off-target activities**, which can lead to non-specific reactions, e.g., in an ELISA, because the antibodies do not only bind their specific antigen. Therefore, the investigation of antibodies via the protein arrays is a good tool for the selection of antibodies to **avoid false-positive** results e.g., in diagnostics.

The hEXselect Array (product no. 1003) has >45,000 protein-coding E. coli clones which represents > 19,000 human antigens and > 4,000 different human genes. Because we have the proteins **not only in full-length, but also as protein isoforms & peptides** (different clones, same gene) we can specify the binding domain in more detail, like **epitope mapping**.





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