

SUPPLEMENTARY MATERIAL

Table S1. *In vitro* selection for high-affinity RNA binding sequences of mouse Dazl

7 round sample
UCGUUCUAAGUUUAUCAUAGA
UUGUGUAGUCCAAGUUAUGUA
UUUCCCCAGUAAAGUUUGAUU
UUGGCUCUCUGUCGUUUUGA
UUUGGCUCCAUUGGUUGUGU
UCGCCAUUGUUUUUGUUAGA
UGCUGUGCUGUUGUAGUUUA
UAGCCCACAUGUUUUUGUUUA
UUGUUUGCCUGUUAGUAGUUA
GUGCUUGUCAAGUUUGUUUU
UGUCAUUAGUUUCGUAGUUUA
UUCUCUUCGCCUUGUUUUUA
UUUCGUUUUGUGUUUAGUUUA
UUUUUCGUUCUUGUUUGUUU
UUUGUUUCGGUUUUUGUUUA
9 round sample
UUGCCUGUUGCAAGUCGGUUA
UCCUGUUGUCCCAGUUGUAA
UUGCCUGUUGCAGUGUUGUAA
UUGCAGGCUCUUGUUGGUUU
UGCCUGUUGCUUGUUUGUUUA
UCCUGGUCCAUUCGUUUAAUU
UUCGGUCUGUUAUAAGUUUAA
UUAGUUUGUACGUUCAGUUUA
UUAGUGUCAUUAGUUUAGUUGA
CUUUUGUGUGGUGUCCGGGAA
UGGUCCUGUUCGUUAGUUUA
UUUGUUUUUCGUUUAGUU
UUUGUUUUUGCUGUUGUUUGA
UUUGUUUGUCCGUUUUGUUUU
UUUGUUUUAGUUUUAGUUUUA
UGUUAGGUUUUGUUUUUGUUUU

Samples of Dazl-binding sequences were taken after seven and nine rounds of selection. Residues flanked by three Us on either side are shown in bold.

Table S2. Genetic screening for high-affinity RNA binding sequences of mouse Dazl by a yeast three-hybrid screen

1.	UGAGAUUUUU	UUUUUUUUUU	GUUUUUUGGU	UUUUUUUUUU	UGUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
2.	GUUUUGAUCC	AAUUCCUGGG	CCUAGGUUUU	GUUUUGUUUG	UUUUUGUGUG	UUUUUGUGUG	UUUUUGUGUG
3.	UUUUUUUUUU	UUUUUUAAAU	CUGUUUGUUU	UCUGUCCUC	UUUUUUAAUC	UUUUUUAAUC	UUUUUUAAUC
4.	AUUUGGAGUU	CUGCAUGCUU	CUUGUAUGUU	CAUGGACAUC	UCUUUCAUUA	GGUUAGGAAA	GGUUUUUUUU
5.	UCAUUUUCCA	CGUUUUUCAG	UGAUUUUUUC	AUUUUUCAAG	UCGUCAAGUG	GAUGUUUCUC	AUUUUCCAUG
6.	AUUUGUUUGU	GUUUUUUUUG	UUUGUUUUUG	UUUUUUUUUU	UUUGGCAAAG	GAGUCCCAAU	AUUAAGUAAA
7.	UUGAUUUUUC	ACUUUUUAJU	UUUGUUUGUU	UUUUUUUUUC	ACUUAUUUCC	AUUAGUCUUU	UUUUUUUUUG
8.	UUUGUUUUUU	GAGUUUUUCG	GUUUUUUUUG	CUGUUUUUUU	UUUUUUUUUC	GGGUUUUUUG	CCUGUUUUUC
9.	UUUUUUUUUC	AUGUUUGAAU	UCACCAUUCU	UUUUUUUUUA	UGGCUUUUUG	AUCUCUUCCA	UGGUUUUUUC
10.	UGCUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
11.	AUUUGAGUUA	AUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
12.	GUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
13.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
14.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
15.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
16.	AGUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
17.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
18.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
19.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
20.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
21.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
22.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
23.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
24.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
25.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
26.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
27.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
28.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
29.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
30.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
31.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
32.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
33.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
34.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
35.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
36.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
37.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
38.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
39.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
40.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
41.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
42.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
43.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
44.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU
45.	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU	UUUUUUUUUU

Seven contiguous blocks of 10 nt containing the highest U content were selected for presentation. The first 45 strongly binding sequences sequenced are shown, sequence no. 45 corresponds to PRL-1 tyrosine phosphatase. Residues flanked by three Us on either side are in bold.