

Supplementary Data

UTexas Aptamer Database: The Collection and Long-Term Preservation of Aptamer Sequence Information

AUTHORS

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Disclaimer:

The information provided in this analysis is based on data available as of July 2023 and may not reflect the most current data. Therefore, any decisions or actions taken based on this information should be carefully evaluated in light of more recent and relevant data.

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Database Name	Institution	Aptamer Sequences	Current Status	First launched or date of publication	Reference
SELEX_DB	Institute of Cytology and Genetics	116	Last updated 2015	2000	Ponomarenko, et al., 2000 & 2001
Aptamer Database*	The University of Texas at Austin	239	Inactive	2004	Lee, et al., 2004
RiboaptDB*	University of Southern Mississippi	3842	Inactive	2006	Thodima, et al., 2006
HTPSELEX	Swiss Institute for Experimental Cancer Research and Swiss Institute of Bioinformatics	Not determined	Last updated September 2012	2006	Jagannathan, et al., 2006
Apta-Index™	Aptagen	783 (as of April 2023)	Active and updated	2008	No paper to date.
Aptamer Base*	Carleton University	2334	Inactive (CSV file with aptamer sequence information is available on GitHub)	2012	Cruz-Toledo, et al., 2012
Aptabase	Indian Institute of Technology Guwahati	605	Periodically updated	2021	No paper to date.

Table S1: A summary of aptamer databases in order of launch date.

*Indicates the database management included mechanisms to check the accuracy of the sequences (e.g., “Aptamer Base” relied on users' wiki-contributions to correct or augment the dataset).

Function	Formula	Field used	Conditional formatting or Cell
Finding duplicates	=countif(D:D,D1)>1	Link to PubMed Entry (green), Journal DOI (light orange), Citation (Light blur), Aptamer sequence (yellow)	Conditional formatting
Finding duplicate in across multiple sheets/column.	Info Collect-setup =match(\$E1:\$E, indirect("Database!D2:D"),0) Database Sheet =match(\$E2:\$E, indirect("Database!D2:D"),0)	DOI. This was used to see if the doi in Info collect-setup sheet is the same as main database.	Conditional formatting
Sequence Length	=(LEN(I2)- LEN(SUBSTITUTE(I2,"G","")))+ (LEN(I2)-LEN(SUBSTITUTE(I2,"C","")) +(LEN(I2)-LEN(SUBSTITUTE(I2,"A","")) +(LEN(I2)-LEN(SUBSTITUTE(I2,"T","")))+ (LEN(I2)-LEN(SUBSTITUTE(I2,"U","")) + (LEN(I2)- LEN(SUBSTITUTE(I2,"g","")))+ (LEN(I2)-LEN(SUBSTITUTE(I2,"c","")) +(LEN(I2)-LEN(SUBSTITUTE(I2,"a","")) +(LEN(I2)-LEN(SUBSTITUTE(I2,"t","")))+ (LEN(I2)-LEN(SUBSTITUTE(I2,"u","")))	Sequence length	Cell
GC content calculation	=((LEN(I35)- LEN(SUBSTITUTE(I35,"G","")))+ (LEN(I35)- LEN(SUBSTITUTE(I35,"C","")))+ (LEN(I35)- LEN(SUBSTITUTE(I35,"g","")))+ (LEN(I35)- LEN(SUBSTITUTE(I35,"c",""))))/J35	GC content	Cell

Table S2: Formulas used in the Google Sheets (e.g., calculate GC content, sequence length, etc.).

a.

Name of Information Collectors [Initials of reviewers]	Year of Paper	Link to PubMed Entry	Journals	Journal DOI	Citation	Type of Nucleic Acid	Name of Aptamer	Target	Aptamer Sequence	Sequence Length	GC Content	Affinity	Kd (nM)	Pool Type	Pool Random Region	Binding Buffer/Conditions
Ali Askari [ISC]	2008	https://pubmed.ncbi.nlm.nih.gov/18111111/	J Agric Food Chem	https://doi.org/10.1021/jf071634h	Cruz-Aguado	ssDNA	112	Ochratoxin A	5'TGGTGCGCTAGGTCA	61	59%	Kd: 0.36	0.36	5-TGGTGCGGTAC	30	Selection buffer (SB)
Isaac Weislow [AA] [KG]	2000	https://pubmed.ncbi.nlm.nih.gov/10811111/	Bioorg Med Chem	https://doi.org/10.1016/S0958-2019(00)00111-1	Fukusaki, E.	ssDNA	Chi No 52	Poly-beta-1,4	5'TAGGGAATTCGTCG	105	60%	Binding Efficiency	N/A	5'-TAGGGAATTCG	59	100 mM NaCl, 100 mM MgCl
Sumedha Kota [PA]	2000	https://pubmed.ncbi.nlm.nih.gov/10811111/	Bioorg Med Chem	https://doi.org/10.1016/S0958-2019(00)00111-1	Okazawa, A.	ssDNA	26	Hematoporphyrin	5'TAGGGAATTCGTCG	102	67%	Kd: 1.6x10 ⁻⁶	1600	5'-TAGGGAATTCG	59	100 mM Trisacetate
Sumedha Kota [KG]	1996	https://pubmed.ncbi.nlm.nih.gov/8811111/	J Clin Invest	https://doi.org/10.1172/JCI111111	Hicke, B. J., V	ssDNA	LD20111	L-selectin-IgG	5'TAGCCAAGGTAACCAG	49	47%	Not reported	N/A	5'-CTACCTACGATC	40	20 mM Hepes, pH 7.4

b.

Pool Random Region	Binding Buffer/Conditions	Divalent Salt	Type of the buffer	pH	Molecular weight of target	Application as quoted in the referenced paper	Post-selex modifications to the aptamer	Additional Relevant Information	Serial Number	Corresponding Author Name, email address	please fill out the form for any feedbacks/comments	Aptamer Cross Referencing (Check Aptamer Chemistry, Affinity, Length, GC content, sequence)
30	Selection buffer (SB MgCl/CaCl)		Tris Buffers	7.0	Not reported	Research and Detection: "This work describes the identification of an aptamer that binds to the library with a K _d of 0.36 nM." [Cruz-Aguado et al., 2008]			10,000,654	Penner, G, E-mail: gpenner@uconn.edu	https://forms.gle/5dTpApdGpdT	5'dTpApdGpdT
59	100 mM NaCl, 100 mM MgCl		Tris Buffers	8.0	Not reported	Diagnostic: "Oligosaccharide antigens play essential biological roles in many biological processes." [Fukusaki et al., 2000]	Not applicable	G-cluster motif	10,000,317	Fukusaki E, fukusaki@uconn.edu	https://forms.gle/5dTpApdGpdT	5'dTpApdGpdT
59	100 mM Trisacetate	None	Tris Buffers	8.0	Not reported	Detection: "In the present study, we selected single-stranded DNA aptamers that bind to hemaphysalis with a K _d of 1.6 x 10 ⁻⁶ M." [Okazawa et al., 2000]	Not applicable		10,000,301	Kobayashi, A, kobayashi@uconn.edu	https://forms.gle/5dTpApdGpdT	5'dTpApdGpdT
40	20 mM Hepes, pH 7.4	MgCl/CaCl	Other Buffers	7.5	Not reported	Therapeutic: "Aptamers that bind with nanomolar affinity to L-selectin's lectin domain, prevent L-selectin-mediated cell adhesion." [Hicke et al., 1996]	Not applicable		10,000,139	Parma, D, parma@uconn.edu	https://forms.gle/5dTpApdGpdT	5'dGpdTpdGpc
35	CE Buffer (30 mM NaCl, 10 mM Tris, 1 mM EDTA)	None	PBS/pho	7.5	Not reported	Diagnostic and Therapeutic: "Alpha-fetoprotein (AFP) is a liver cancer marker." [Guo et al., 2010]	Not applicable		10,001,063	Guo, W, guowei@uconn.edu	https://forms.gle/5dTpApdGpdT	5'dGpdTpdGpc

Figure S1: A screenshot of the UTexas Aptamer dataset in Google Sheets.

The name of information collectors and initials of reviewers were recorded in the first column to track systematic errors across the database. Bright blue indicate there is more than one entry from one publication. Figure 1 B is the continuation of Figure 1 A.

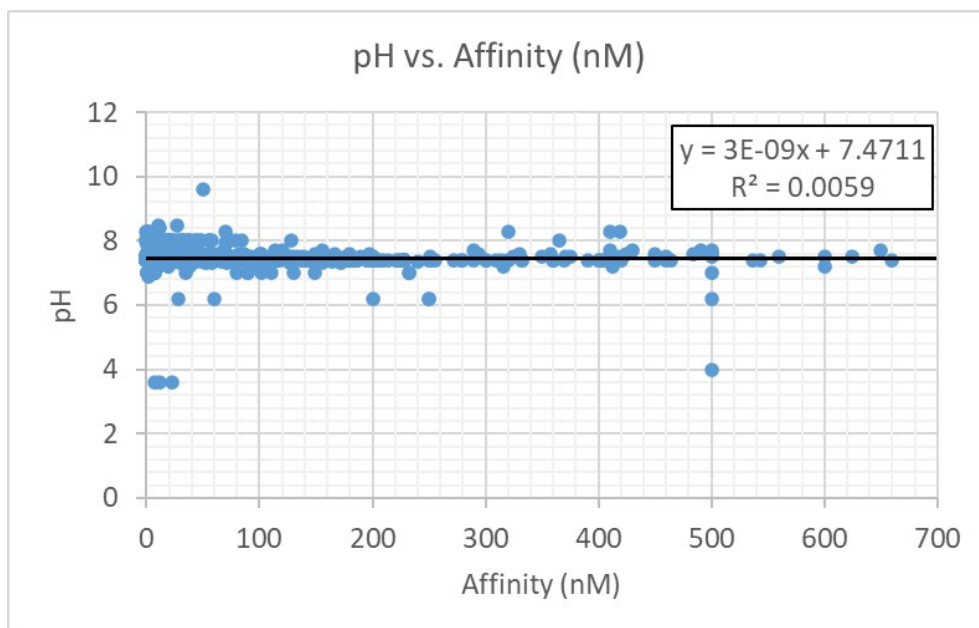


Figure S2: Aptamer binding affinity vs. pH.

The relationship between these two variables is assessed using the line of best fit. With an R^2 value of 0.0059, there is no correlation between the two variables. The y-axis was adjusted to a zoom level that selectively displays data up to 700 nM.

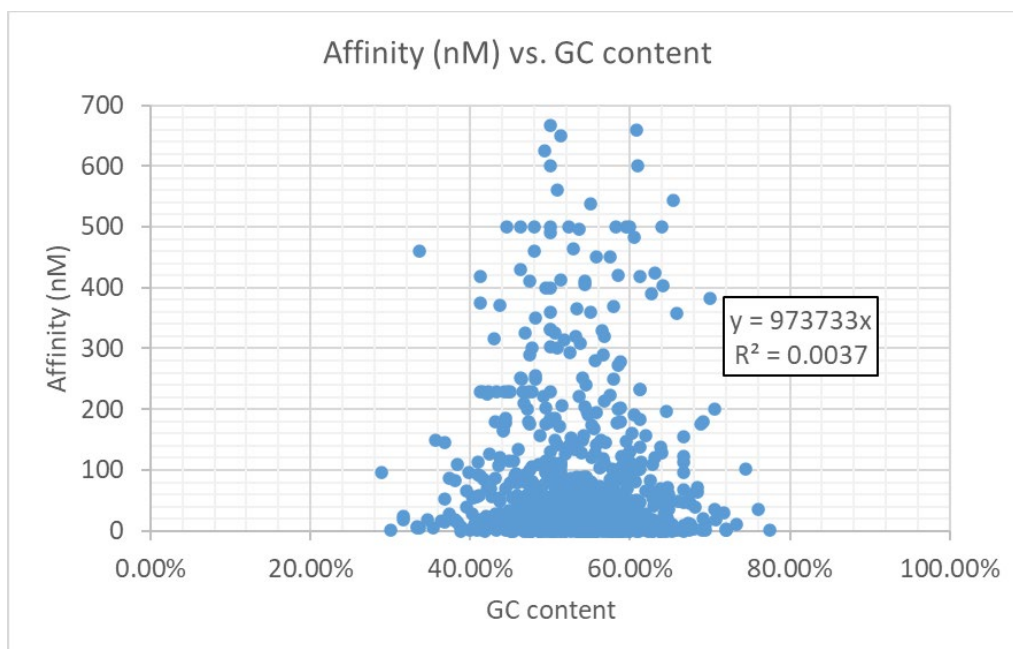


Figure S3: Aptamer binding affinity in nM vs. GC%.

The relationship between these two variables is assessed using the line of best fit. With an R^2 value of 0.0037, there is no correlation between the two variables. The y-axis was adjusted to a zoom level that selectively displays data up to 700 nM, and the intercept was set at zero.

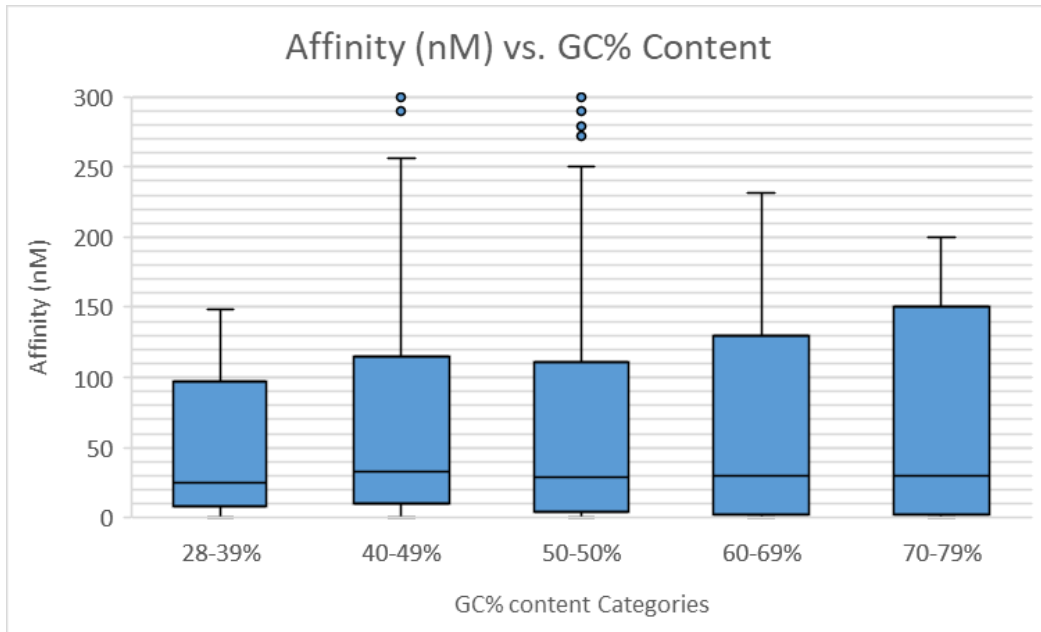


Figure S4: Aptamer binding affinity in nM vs. GC% (boxplot). There are five GC% categories to assess whether the specific range of GC% affect affinity. The y-axis was adjusted to a zoom level that selectively displays data up to 300 nM.

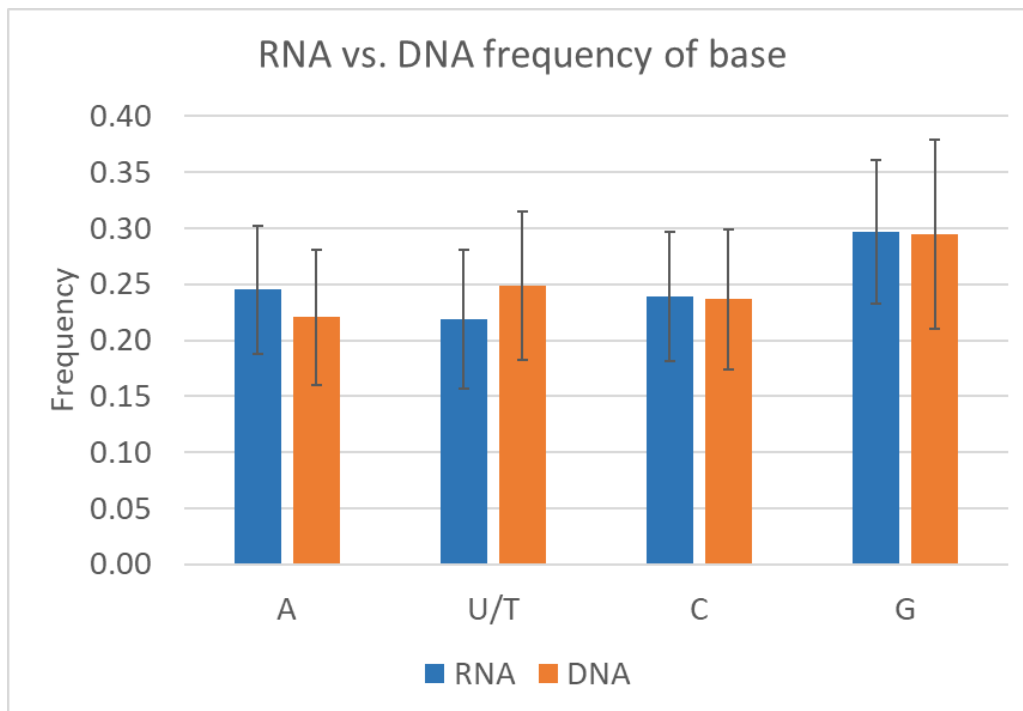


Figure S5: Frequency of each base per aptamer sequence.

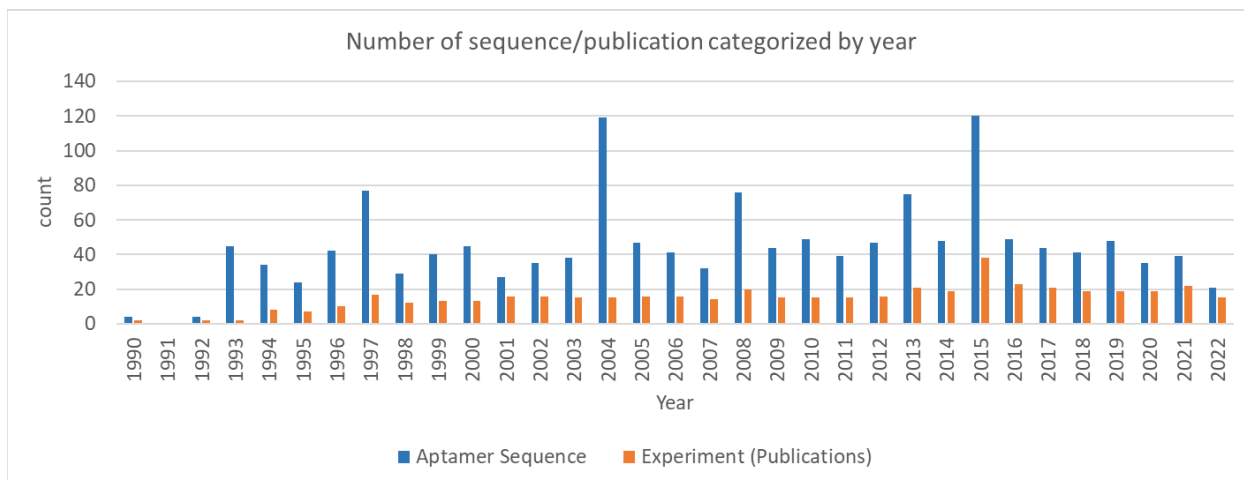


Figure S6: Number of aptamer sequences and aptamer publications per year.

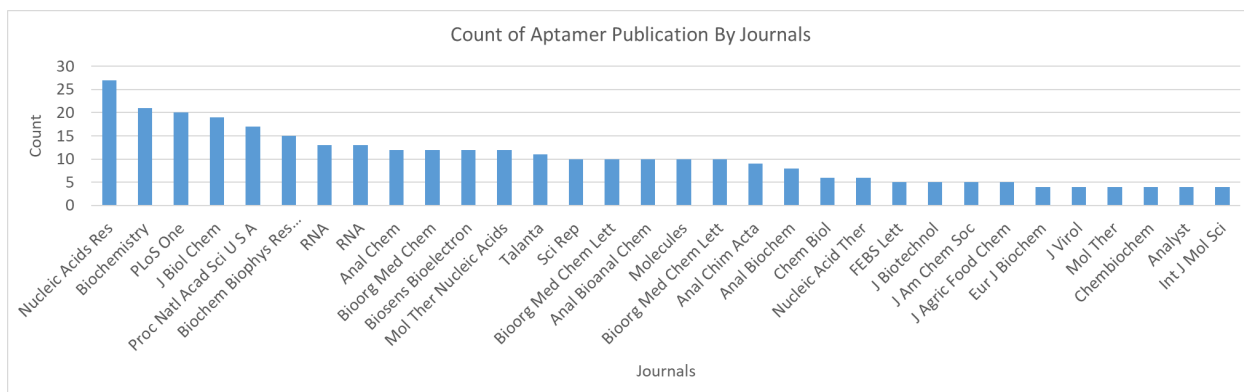


Figure S7: Count of aptamer publications by journals, according to the UTexas Aptamer dataset.

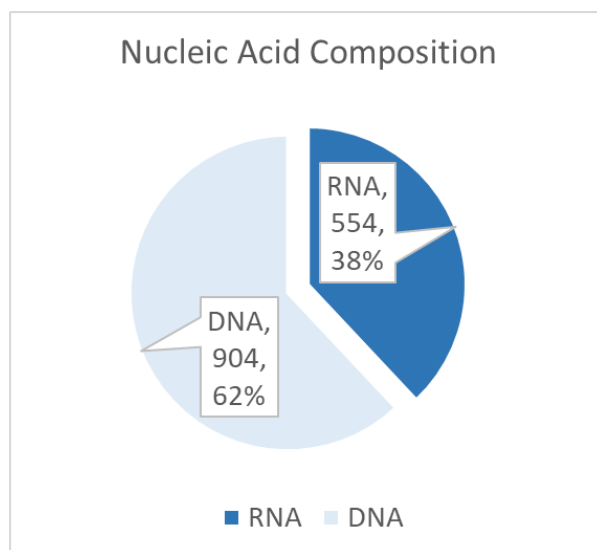


Figure S8: Summary of the nucleic acid type of aptamer sequence found in the UTexas Aptamer Database.

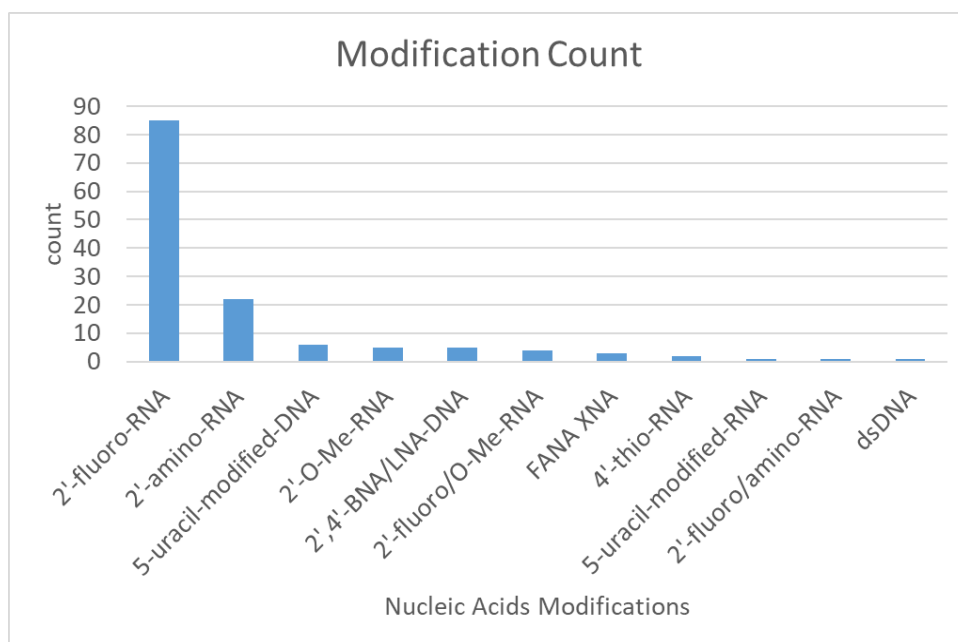


Figure S9: Count of each nucleic acid type in the UTexas Aptamer Database.

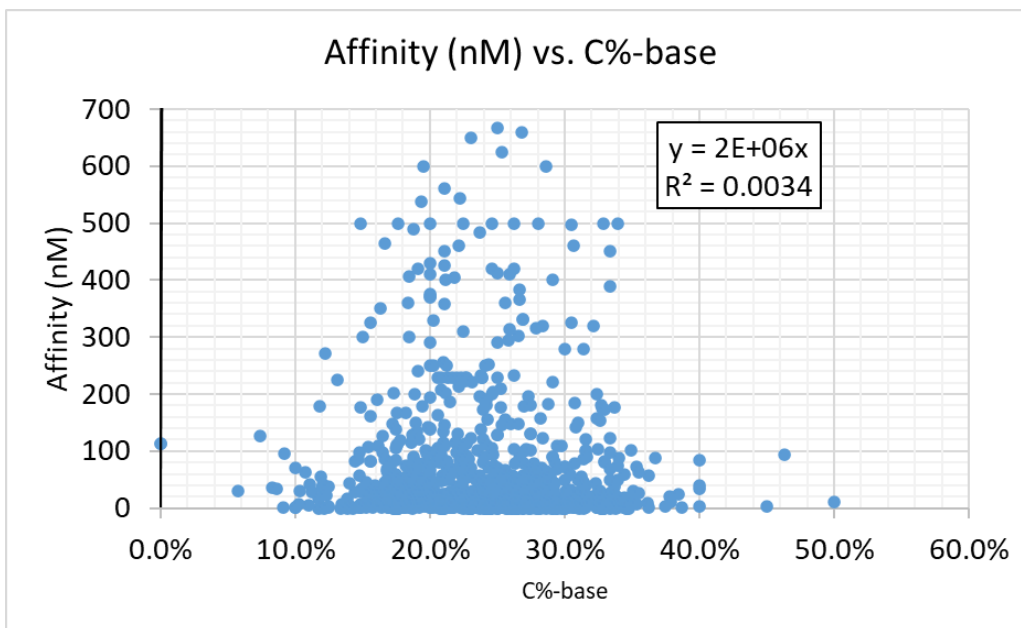


Figure S10: Aptamer binding affinity vs. aptamer C%-base count.

The relationship between these two variables is assessed using the line of best fit. With an R^2 value of 0.0034, there is no correlation between the two variables. The y-axis was adjusted to a zoom level that selectively displays data up to 700 nM, and the intercept was set at zero.

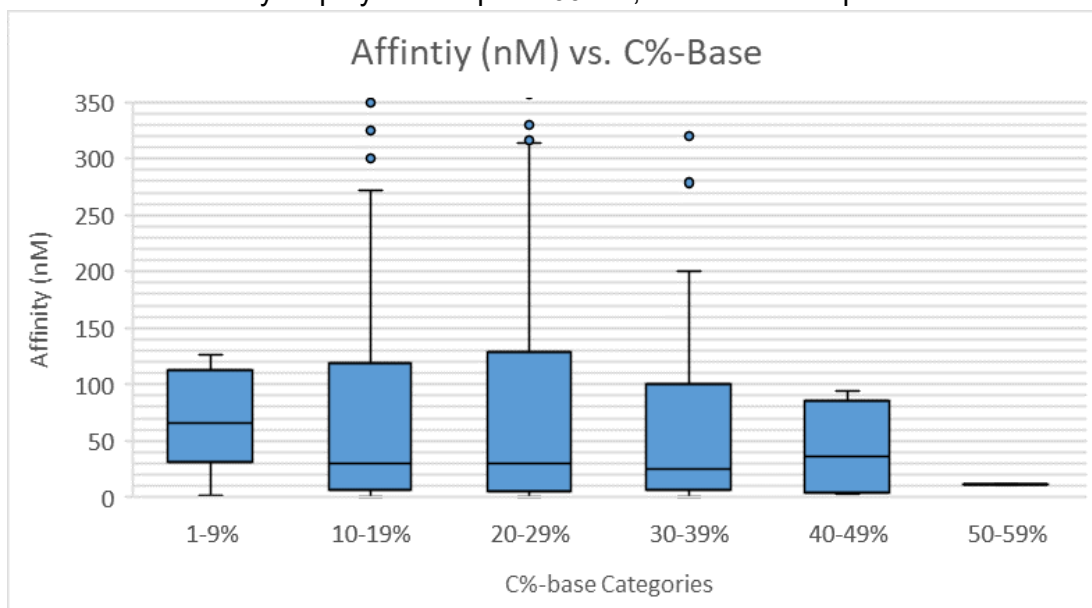


Figure S11: Aptamer binding affinity in nM vs. C% (boxplot).

There are six C% categories to asses whether the specific range of C% affect affinity. Categories 1-9%, 40-49%, and 50-59% have less than 20 sample sizes. The y-axis was adjusted to a zoom level that selectively displays data up to 350 nM.

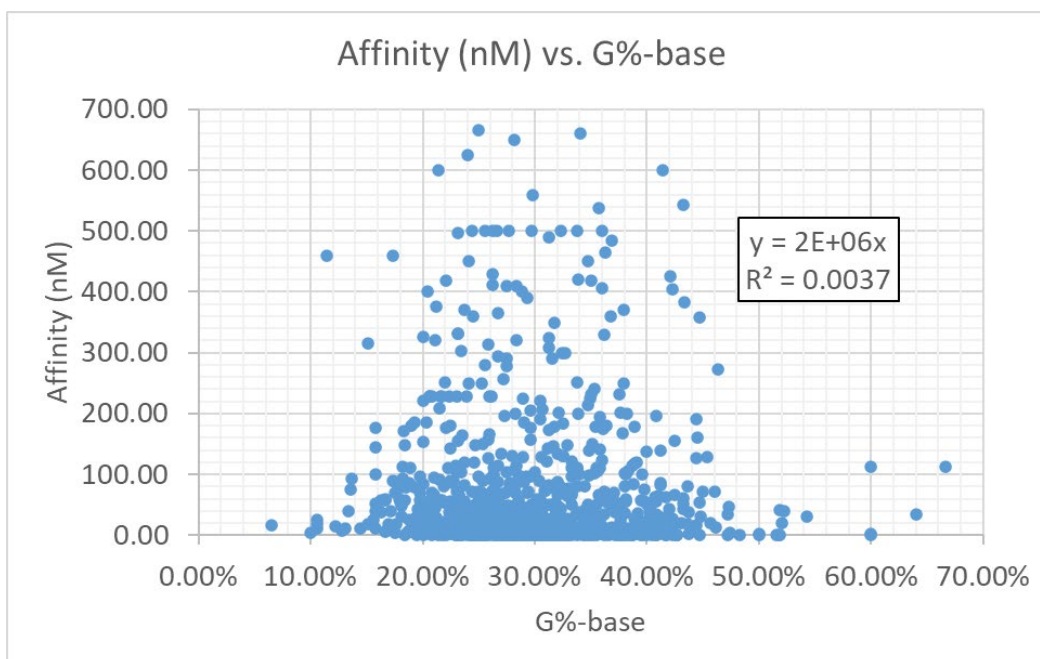


Figure S12: Aptamer binding affinity vs G%-Base count in the aptamer sequence. The relationship between these two variables is assessed using the line of best fit. With an R^2 value of 0.0037, there is no correlation between the two variables. The y-axis was adjusted to a zoom level that selectively displays data up to 700 nM, and the intercept was set at zero.

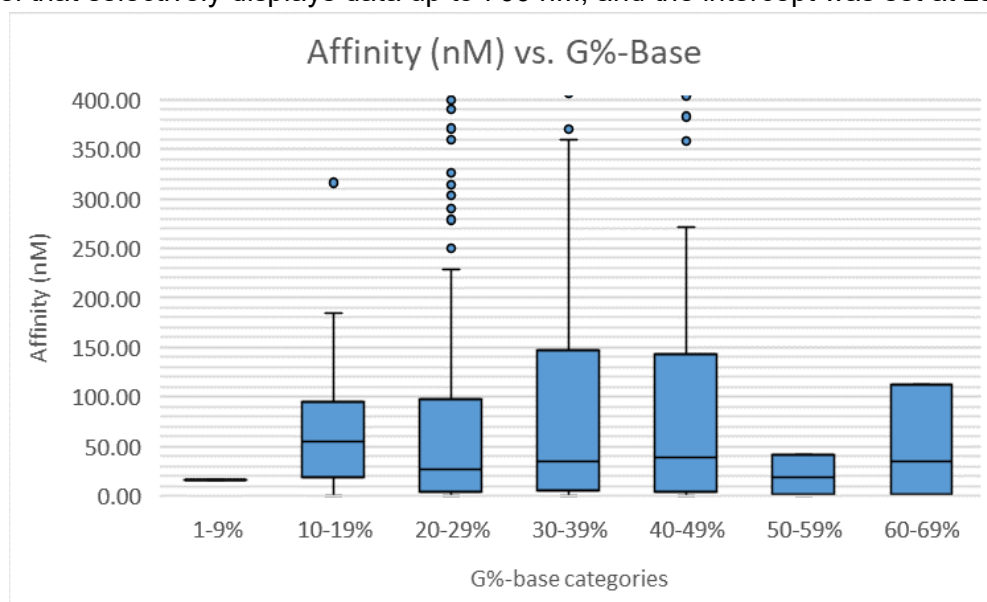


Figure S13: Aptamer binding affinity in nM vs. G% (boxplot). There are six G% categories to assess whether the specific range of G% affects affinity. Categories 1-9%, 50-59%, and 60-69% have less than 20 sample sizes. The y-axis was adjusted to a zoom level that selectively displays data up to 400 nM.

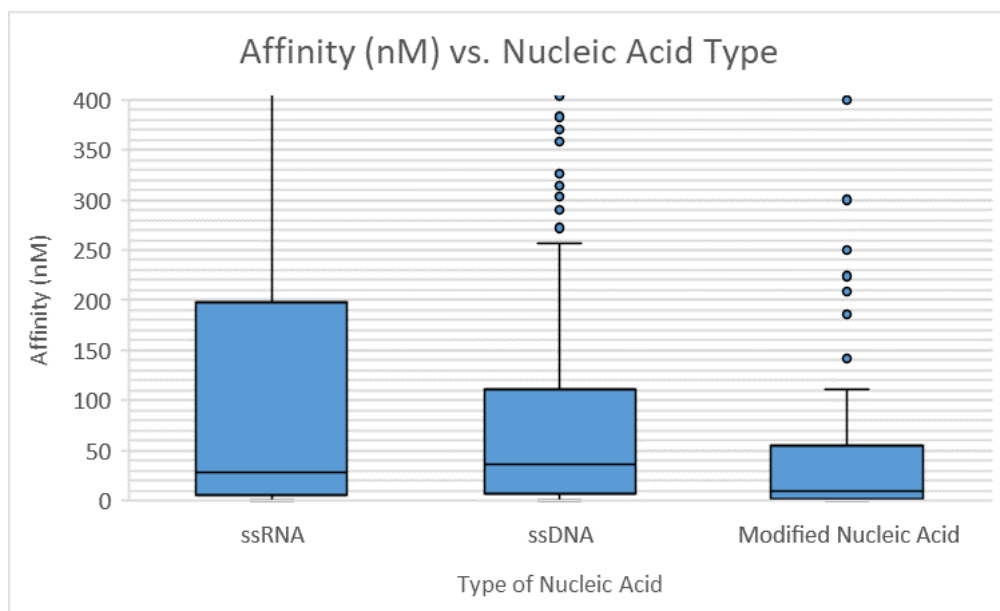


Figure S14: Nucleic acid type versus aptamer affinity (boxplot).
 The y-axis was adjusted to a zoom level that selectively displays data up to 400 nM.

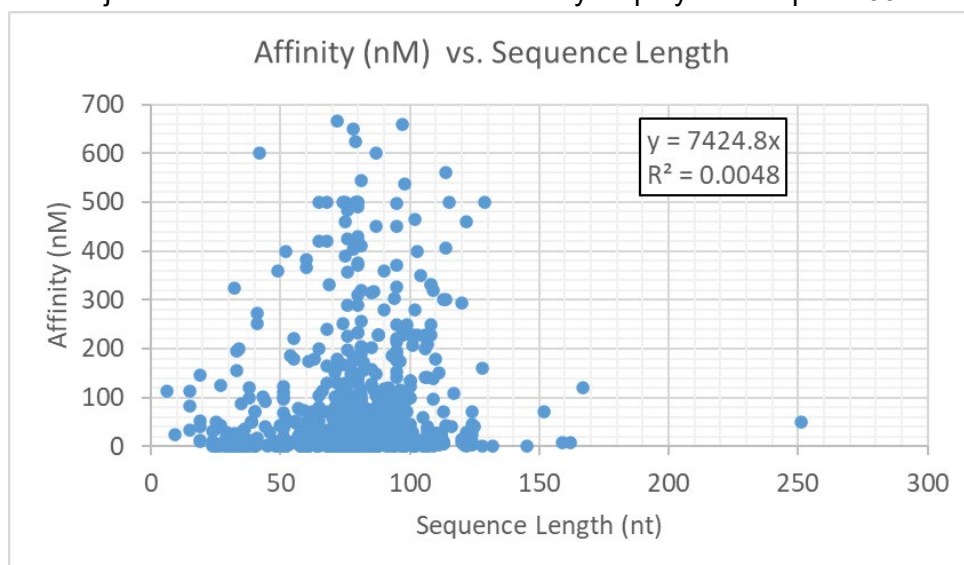


Figure S15: Aptamer binding affinity versus sequence length.
 The relationship between these two variables is assessed using the line of best fit. With an R^2 value of 0.0048, there is no correlation between the two variables. The y-axis was adjusted to a zoom level that selectively displays data up to 700 nM, and the intercept was set at zero.

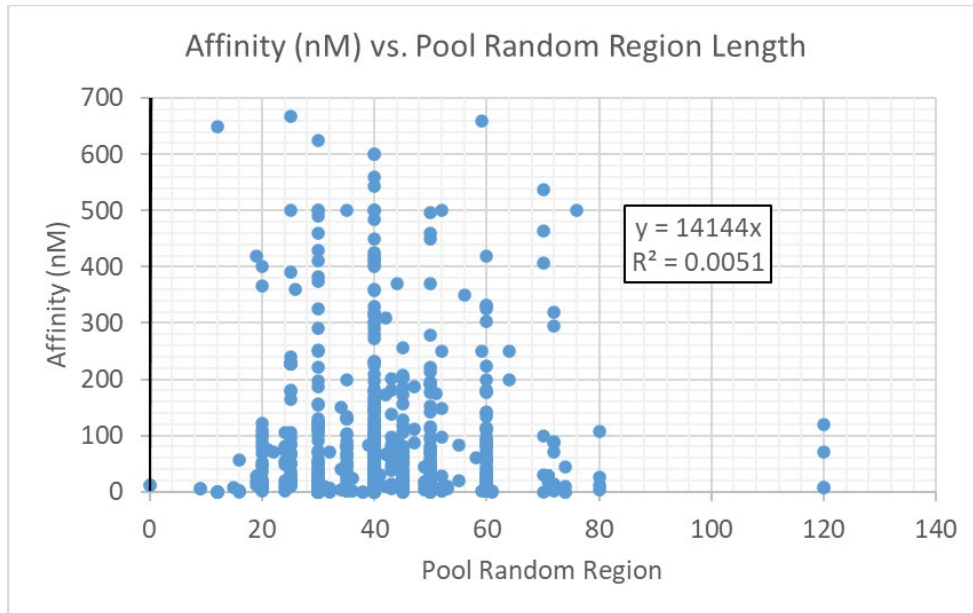


Figure S16: Aptamer binding affinity versus pool random region.

The relationship between these two variables is assessed using the line of best fit. With an R^2 value of 0.0051, there is no correlation between the two variables. The y-axis was adjusted to a zoom level that selectively displays data up to 700 nM, and the intercept was set at zero.

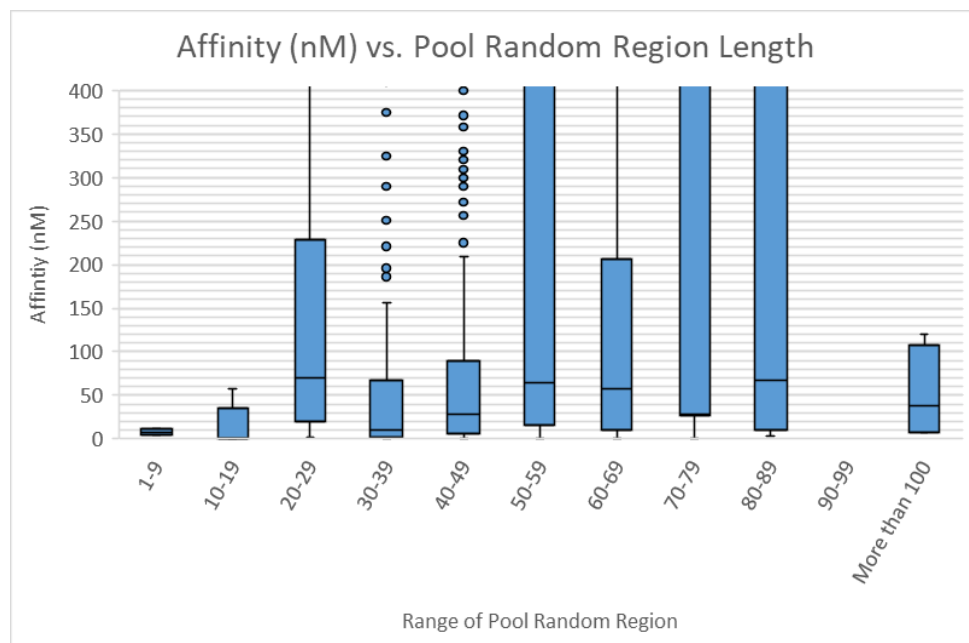


Figure S17: Length of pool random region versus aptamer affinity (boxplot).

Categories 1-9, 10-19, 80-89, 90-99, and more than 100 have less than 20 sample sizes. The y-axis was adjusted to zoom-in on the area around the boxplots, data up to 400 nM displayed.

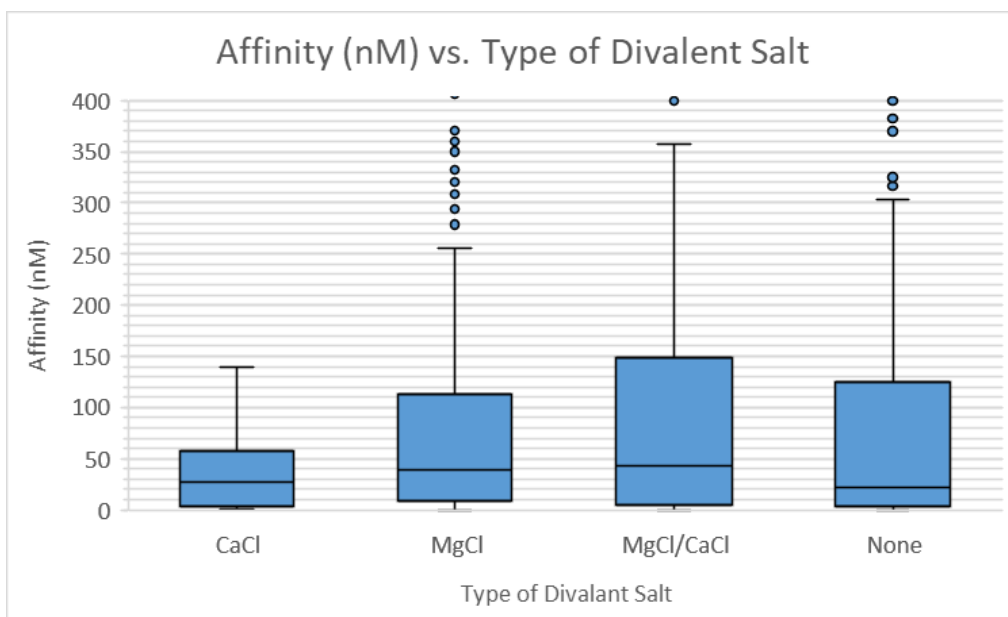


Figure S18: Divalent salt included in binding buffer versus aptamer affinity (boxplot). The y-axis was adjusted to a zoom level that selectively displays data up to 400 nM.

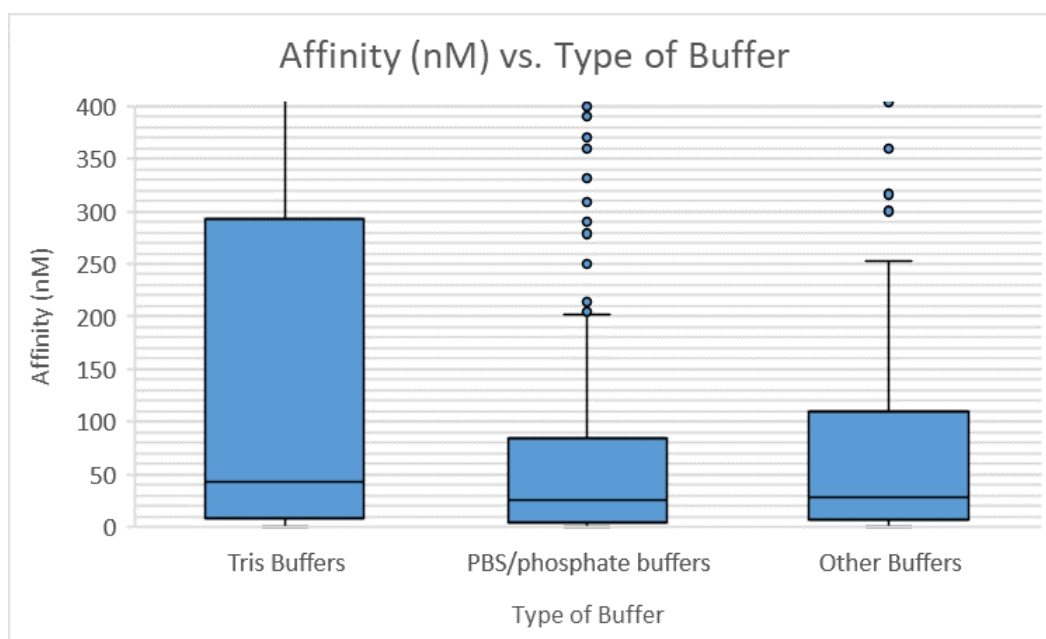


Figure S19: Buffer type versus aptamer binding affinity (boxplot). The y-axis was adjusted to zoom-in on the area around the boxplot, data up to 400 nM displayed.

UTexas DB Serial Number and Citation	Aptamer Name, Target Sequence Comparison	Source of Error
# 10000063 Lorsch, J. R., & Szostak, J. W. (1994). Biochemistry, 33(4), 973- 982.	35-mer aptamer, anti-Cobinamide dicyanide UTexas: 5'GGAACCGGUGCGCAUAACCACCUCAGUGCGAGCAA3' Apta-Index: 5'-----CCGGUGCGCAUAACCACCUCAGUGCGAGCAAGGAA3'	Apta-Index added GGAA to wrong end
# 10001000 Kwon, H. M., et al. (2014). A PloS one, 9(5), e97574.	HA12-16, anti-gHA1 UTexas: 5'GGGUUCACUGCAGACUUGACGAAGCUUGCUUGACGGAGAUCAAGGGCGAGUCGCAUACCAAGUUGAUGGGGAAUGGAUCCACAUCUACGAAUUC3' Apta-Index: 5'GACAAGGATAAATCCTTCAATGAAGTGGGTCACTCATCTGTGA3'	Apta-Index reported an entirely different sequence.
# 10001063 Dong, L., et al. (2015). Scientific reports, 5(1), 15552.	AP273, anti-Alpha-fetoprotein (AFP) UTexas: 5'GTGACGCTCCTAACGCTGACGTGACGCTCCTAACGCTGACTCAGGTGCAGTTCTCGACTCGGTCTTGATGTGGGTCTGTCCGTCCGAACCAATCCCTGTCCGTCCGAACCAATC3' Apta-Index: 5'GTGACGCTCCTAACGCTGACTCAGGTGCAGTTCTCGACTCGGTCTTGATGTGGGTCTGTCCGTCCGAACCAATC3'	Both had errors: UTexas duplicated primer regions and Apta- Index deleted an internal C.
#10001073 Li, P., et al. (2015). BMC Veterinary Research, 11(1), 1-11.	QA-36, anti-Soft-shelled turtle iridovirus (STIV) UTexas: 5'GACGCTTACTCAGGTGTGACTCGTGTGCGGGGAGGGGAGTGGCGCTGTTGGTGCGGGTATAGCGCGTGGTGTGCAAGGACGCAGAGAAGTCTC3' 5'-----TGTGCGGGGAGGGGAGTGGCGCTGTTGGTGCGGGTATAGCGCGTGGTGT-----3' ^ Apta-Index:	Apta-Index deleted the primer regions.

<p># 10001268</p> <p>Xiang, J., et al. (2019). Molecular Therapy- Nucleic Acids, 16, 302-312.</p>	<p>BI-1, anti-BACE-1</p> <p>UTexas: 5' ATCCAGAGTGACGCAGCAAGCGATACTGCGTGGCTGGAGGCGGGTAGGGCCAGAGTTC TGGACACGGTGGCTTAGT3'</p> <p>Apta-Index: 5' AGCGATACTGCGTGGCTGGAGGCGGGTAGGGCCAGAGTTC3'</p>	<p>UTexas added primer regions.</p>
<p># 10001297</p> <p>Diaz-Fernandez, A., et al. (2019). Biosensors and Bioelectronics, 128, 83-90.</p>	<p>PSA-1, anti-Prostate-specific antigen</p> <p>UTexas: 5' AGGGTTGATAGGTTAAGAGCGGACGGTTGCGCTATATTTAACCAAAAGTCTGGATTAACA CGATGTCAACTAGCTGTTGGG3'</p> <p>Apta-Index: 5' GGACGGTTGCGCTATATTTAACCAAAAGTCTGGATTAACA3'</p>	<p>UTexas added primer regions.</p>
<p># 10001334</p> <p>Song, Z., et al. (2020). Molecules, 25(23), 5585.</p>	<p>CD63-1, anti- CD63 positive cells</p> <p>UTexas: 5' TAGGGAAGAGAAGGCATATGATTAACACGACAGACGTTCCGGAGTTCGAACCCCTGACAGCGTGGGCTTGACTAGTACATGACCACTTGA3'</p> <p>Apta-Index: 5' TAACACGACAGACGTTCCGGAGTTCGAACCCCTGACAGCGTGGG3'</p>	<p>UTexas added primer regions.</p>
<p># 10001339</p> <p>Kohlberger, M., et al. (2020). PLoS One, 15(11), e0241560.</p>	<p>C7, anti-Rituximab, anti-CD20 IgG1 antibody</p> <p>UTexas: 5' ATACCAGCTTATTCAATTGGCCATTGTGGACTTCTTTGGGTAATTCAGGGGCTCGATTAGATAGTAAGTGAATCT3'</p> <p>Apta-Index: 5' GGCCATTGTGGACTTCTTTGGGTAATTCAGGGGCTCGATT3'</p>	<p>UTexas added primer regions.</p>
<p># 10001343</p> <p>Kohlberger, M., et al. (2020). PloS one, 15(11), e0241560.</p>	<p>C10, anti-Rituximab, anti-CD20 IgG1 antibody</p> <p>UTexas: 5' ATACCAGCTTATTCAACTACTTCGGCTAGTTAGGGGTAGTTTAGATCGTCTCTACATAGATAGTAAGTGAATCT3'</p> <p>Apta-Index: 5' ACTTCGGCTAGTTAGGGGTAGTTTAGATCGTCTCTACAT3'</p>	<p>UTexas added primer regions.</p>

<p># 10001348</p> <p>Chinnappan, R., et al. (2021). Talanta, 224, 121818.</p>	<p>BC1, anti- Beta-crosslap (BC)</p> <p>UTexas: 5' ATACCAGCTTATTCAATTATGACGGGGTCTAGGCAAGTAATAACGGGGCAAGCTTTTCTATCTCGTTCTAGGGTAAGATAGTAAGTGCAATCT3'</p> <p>Apta-Index: 5' ATGACGGGGTCTAGGCAAGTAATAACGGGGCAAGCTTTTCTATCTCGTTCTAGGGTA3'</p>	<p>UTexas added primer regions.</p>
<p># 10001349</p> <p>Zhu, C., et al. (2021). Talanta, 223(Pt 1), 121690.</p>	<p>T-2, anti- Thyroglobulin (Tg)</p> <p>UTexas: 5' CCTAACCGATATCACACTCACCGGTGAGCGGGGAGGCGATGCCCAGGCTAACTTGACTCAGTTGGTCGTCATTGGAGTATC3'</p> <p>Apta-Index: 5' -----CGCGTGAGCGGGGAGGCGATGCCCAGGCTAACTTGACTCA-----3'</p>	<p>Apta-Index deleted the primer regions.</p>
<p># 10001358</p> <p>Yu, Q., et al. (2021). Journal of fish diseases, 44(1), 33–44.</p>	<p>GVI-7, anti-Grass carp reovirus (GCRV)-infected CIK cells</p> <p>UTexas: 5' GACGCTTACTCAGGTGTGACTCGTGAACCCACCTCAGGGCATCTTACATTTCTTCTAAGTTGTTACCATGTTTCGAAGGACGCAGATGAAGTCTC3'</p> <p>Apta-Index: 5' GTCTGAAGTAGACCGCAGGAGTGAACCCACCTCAGGGCATCTTACATTTCTTCTAAGTTGTTACCATGTTTAGTCACACCTGAGTAAGCGT3'</p>	<p>Apta-Index added incorrect primer regions (and paper has entirely different forward primer in folded aptamer, Fig 2).</p>
<p># 10001372</p> <p>Wu, H., et al. (2021). ACS omega, 6(5), 3771–3779.</p>	<p>HPA-2, anti-Helicobacter pylori</p> <p>UTexas: 5' AAGGAGCAGCGTGGAGTTACCAGGAGGACCCATTCTCGTGTATCGACGAGATCCAGTGACCACGACGACACCCCTAA3'</p> <p>Apta-Index: 5' -----CCAGGAGGACCCATTCTCGTGTATCGACGAGATCCAGTG-----3'</p>	<p>Apta-Index deleted the primer regions.</p>
<p># 10001395</p> <p>Lorenzo-Gómez, R., et al. (2022). Analytica chimica acta,</p>	<p>D1, anti- 16-amino acid peptide from collagen XIa1</p> <p>UTexas: 5' ATACCAGCTTATTCAATTTTTGGTTGACGGCAGTCGGCGGTATGCGCATATCGTGTGGTAACAATCGTAATCAGTTAG3'</p> <p>Apta-Index: 5' GGTTGACGGCAGTCGGCGGTATGCGCATATCGTGTGGTA3'</p>	<p>UTexas added primer regions.</p>

1189, 339206.		
# 10000117 Xu, W., & Ellington, A. D. (1996). PNAS 93(15), 7475–748	C2, anti-HIV-1 Rev UTexas: 5' UCUAAUACGACUCACUUAUGGGAGAGACAAGCUUGGGUC UCGACCUCGCGCAGGAGGGUGGAGGGUCGUAGAGCGCGUA AGAAGAGAAAAGAGAAGUAAUUAAGGAUCCUCAC 3' Apta-Index: 5' GGAGGUCGACCUCGCGCAGGAGGGUGGAGGGUCGUAGAGCGCGUAGGAGG3'	UTexas added primer regions rather than motif regions.
# 10000358 Proske, D., et al. (2002). The Journal of biological chemistry, 277(13), 11416– 11422.	DP3, anti-Neuropeptide Y UTexas: 5' UCGGAGAAAGGGAAGCUUGAGCAGCAGGAGGGCCGGCGUUAGGGUUAGCGAGCCGAUUGAAAGAAGAAGGAACGAGCGUACGGAUCCGAUC3' Apta-Index: 5' -GGGAGAAAGGGAAGCUUGAGCAGCAGGAGGGCCGGCGUUAGGGUUAGCGAGCCGAUUGAAAGAAGAAGGAACGAGCGUACGGAUCCGAUC3'	Apta-Index deleted one nt and inserted an extra G.
# 10000392 Pileur, F., et al. (2003). Nucleic acids research, 31(19), 5776–5788.	VI-2, anti-Ribonuclease H1 UTexas: 5' GCCTGTTGTGAGCCTCCTCTCGAA CGGTCGCTCCGTGTGGCTTGGGTTGGGTGTGGCAGTGACT TGAGCGTTTATTCTTGTCTCCC 3' Apta-Index: 5' CGGTCGCTCCGTGTGGCTTGGGTTGGGTGTGGCAGTGAC3'	UTexas added primer regions.
# 10000645 Tang, Z., et al. (2007). Analytical chemistry, 79(13), 4900–4907.	TD05, anti-Ramos cells UTexas: 5' AAGGAGCAGCGTGGAGGATA AACACCGTGGAGGATAGTTCGGTGGCTGTT CAGGGTCTCCTCCC GGT TAGGGTGTGTCTCGTGGT 3' Apta-Index: 5' AACACCGTGGAGGATAGTTCGGTGGCTGTT CAGGGTCTCCTCCC GGT3'	UTexas added primer regions.

<p># 1000697</p> <p>Li, M., et al. (2008). Journal of the American Chemical Society, 130(38), 12636–12638.</p>	<p>85A, anti-Fibrinogen</p> <p>UTexas: 5' CCTTCGTTGTCTGCCTTCGTAGCGGATCGAATTACGCGTTAACGGCAACCGATAACGGGACCGATTGCACACCCTTCAGAATTCGCACCA3' Apta-Index: 5' CCTTCGTTGTCTGCCTTCGTAGGACCCGACACATCGACGCAGGGAAATTCGGCAAGTCCAGCCAAATGCCACCCTTCAGAATTCGCACCA3'</p>	<p>Apta-Index reported a different random region and the correct primer regions.</p>
<p># 1000800</p> <p>Savory, N., et al. (2010). Biosensors & bioelectronics, 26(4), 1386–1391.</p>	<p>PSap4#5, anti-Prostate specific antigen (PSA)</p> <p>UTexas: 5' CATGCTTACCTATAGTGAACTTTATTAGCCTCCCGGAAGAGCACCTCTTTCATGCTTACCTATAGTGAAC3' Apta-Index: 5' TTTTAATTAAAGCTCGCCATCAAATAGCTTT3'</p>	<p>Apta-Index reported the ΔPSap4#5 aptamer as PSap4#5</p>
<p># 1000885</p> <p>Song, KM, Jet al. Anal Bioanal Chem. 2012;402(6): 2153-2161.</p>	<p>AMP18, anti-Ampicillin</p> <p>UTexas: 5' CACCTAATACGACTCACTATAGCGGATCCGA-TTAGTTGGGGTTCAGTTGCTGGCTCGAACAAGCTTGC3' Apta-Index: 5' TTTAGTTGGGGTTCAGTTG3'</p>	<p>UTexas added primer regions and deleted a T nucleotide.</p>
<p># 1000899</p> <p>Woo, H. M., et al. (2013). Antiviral research, 100(2), 337–345.</p>	<p>NS1 aptamer, anti-Influenza virus non-structural protein 1 (NS1) protein</p> <p>UTexas: 5' GCAATGGTACGGTACTTCCGCGGTCCGGGGTGGGTGGGTGGGGGTGCGGG-----CAAAGTGCACGCTACTTTGCTAA3' Apta-Index: 5' GCAATGGTACGGTACTTCCGCGGTCCGGGGTGGGTGGGTGGGGGTGCGGGGGGCGGCCGCAAAGTGCACGCTACTTTGCTAA3'</p>	<p>UTexas deleted 10 nucleotides.</p>

<p># 10000918</p> <p>Eissa, S., et al. (2013). Analytical chemistry, 85(24), 11794–11801.</p>	<p>OA34, anti-Okadaic acid (OA)</p> <p>UTexas: 5' ATACCAGCTTATTC AATTGGTCACCAACAACAGGGAGCGCTACGCGAAGGGTCAATGTGACGTCATGCGGATGTGTGGAGATAGTAAGTGAATCT3'</p> <p>Apta-Index: 5' GGTACCAACAACAGGGAGCGCTACGCGAAGGGTCAATGTGACGTCATGCGGATGTGTGG3'</p>	<p>UTexas added primer regions.</p>
<p># 10000968</p> <p>Yang, M., et al. (2013). Sensors (Basel, Switzerland), 13(5), 6865–6881.</p>	<p>Apt22, anti- Salmonella Paratyphi A</p> <p>UTexas: 5' GAATTCAGTCGGACAGCGATGGACGAATATCGTCTCCAGTGAATTCAGTCGGACAGCGGATGGACGAATATCGTCTCCC3'</p> <p>Apta-Index: 5' ATGGACGAATATCGTCTCCAGTGAATTCAGTCGGACAGCG3'</p>	<p>Sequence from the paper is unclear (i.e., unclear if primer regions are added or not).</p>
<p># 10000981</p> <p>Xu, D., et al. (2014). Nucleic acid therapeutics, 24(3), 226–238.</p>	<p>AptER-1, anti-Estrogen receptor alpha (ERα)</p> <p>UTexas: 5' GGGCAGAGGCACCGCGAACAACGCAAGACAGAGUGCCGACAAGAGCACAAGCUUCUGCCC3'</p> <p>Apta-Index: 5' GGGCAGACCGCACC CGCAACAACGCAAGACAGAGUGCCGACAAGAGCACAAGCUUCUGCCC3'</p>	<p>Apta-Index introduced a mutation (changed C to G)</p>
<p># 10000795</p> <p>Chang, T. W., et al. (2010). Biochemical and biophysical research communications, 396(4), 854–860.</p>	<p>S132B-C22, anti-Light chain of type A botulinum neurotoxin (BoNT/A) (LCA)</p> <p>UTexas: 5' GGGAGGAGGAGAGAUGUGAACUUGACAGCGUGCCUAGAAGUCCAAGCUUAAAUAACCACGCUAGCAGAAACUCUACACUGGACUGGCG3'</p> <p>Apta-Index: 5' GGGAGGAGGAGAGATGTGAAC TTGACAGCGUGCCUAGAAGUCCAAGCUUAAAUAACCACGCUAGCAGAAACTCTACACTGGACTGGCG3'</p>	<p>Apta-Index included T's, instead of U's for an RNA Aptamer</p>

<p># 10001357</p> <p>Yu, Q., et al. (2021). Journal of fish diseases, 44(1), 33–44.</p>	<p>GVI-1, anti-Grass carp reovirus (GCRV)-infected CIK cells</p> <p>UTexas: 5'GACGCTTACTCAGGTGTGACTCGGGGTGTAGCTCGTTATGATTTCGGACAAGACTTACCTTGCGCCTCTGGGATCGAAGGACGCAGATGAAGTCTC3'</p> <p>Apta-Index: 5'GTCTGAAGTAGACCGCAGGAGGGGTGTAGCTCGTTATGATTTCGGACAAGACTTACCTTGCGCCTCTGGGATAGTCACACCTGAGTAAGCGT3'</p>	<p>Apta-Index recorded incorrect primer regions.</p>
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Table S3: Comparison of Aptamer Databases: Examination of those sequences that differed.

When comparing the established Apta-Index to the UTexas Aptamer Database, we found 90 aptamer sequences in common between the two. Of the 27 that were different, we examined the source of the differences to build our internal training practices when extracting sequence information from the literature.