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# **EDITORIAL**

Welcome to the 2018 first issue of INSOAP times, your source to what's happening in the aptamer world presented by the INSOAP team. Happy New Year everyone. I hope you had a great start to 2018. New questions to be asked, new answers to be discovered, new topics to be discussed and old topics to be updated this year. We are currently gearing up for the 5th Aptamer Symposium to be held at



Oxford, UK on April 11-12th 2018. Registration is open and we are looking forward to catching up with you all in April. If you've been keeping an eye on the program schedule, you'll be aware that Emeritus Professor Uli Hahn is receiving a life-time achievement award. I can't tell you how happy I'll be to see Uli receive this. He has been a member of INSOAP since the beginning and has brought his outstanding humour to the conference each year. You'll also remember that we featured Uli in our 'Interview with a researcher' section in our first newsletter issue last year. I'm also excited about our new feature at the conference this year which will see poster presenters giving flash talks.

You will see a few updates in this newsletter as we try to stay ahead of changes. We have updated the companies offering aptamer based services globally, and provided an update on clinical trials of various aptamers. We also have a new section on new aptamers and in this newsletter we covered the aptamers generated in 2017. In each issue, we will be providing updates on which new aptamers are appearing in the literature. If you'd like to ensure your aptamers make the list, please get in touch with us. We've also, keeping with our intention of introducing you to as many researchers as possible, interviewed Professor Dr Beatrix Suess and I hope you enjoy her answers to the questions as much as we have! I would like to take this opportunity to thank the entire INSOAP team for their efforts, and generation donation of time and talent. Things wouldn't have been possible without you and I am greatly appreciative of all the assistance of the team for ensuring we produce timely updates for INSOAP. If you would like to contribute to the newsletter, please get in touch! As always, the INSOAP welcomes your ideas and suggestions.

As a final note, have you liked our Facebook page? We are currently providing links to new aptamer research papers on a daily basis. Don't have time to keep up to date on current literature? Get our daily updates in your morning newsfeed at <a href="https://www.facebook.com/AptaSoc/">https://www.facebook.com/AptaSoc/</a>. Please don't forget to also follow us on twitter (@Aptamer Society, @Japtamers).

May you all have a great few months, and we'll see you in Oxford in April!

Dr Sarah Shigdar President



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# From the Editor

If you have anything you would like to see in the next issue of the INSOAP newsletter, send it directly to

<u>sarah.shiqdar@deakin.edu.au</u>. Please ensure that your articles and information are in by close of business on 31<sup>st</sup> August 2017

#### **Aptamers Journal**

We announced the official journal of INSOAP at Aptamers 2017. Please email us at

aptasoc@gmail.com to express your interest in joining the editorial or reviewer team. Please see http://JAptamers.co.uk to submit your article.

# Aptamers 2018 Symposium

Web: http://libpubmedia.co.uk/aptamers-2018 Twitter: @AptamerSociety; @JAptamers; #AptaOx18 Email: AptamersOxford@gmail.com

I'm so excited reading our list of confirmed speakers for the 5<sup>th</sup> symposium. There are so many familiar names, both from previous attendees at the conference and from papers that I've read. As we are all focussed on developing our aptamers for clinical applications, I'm also pleased to announce that we will have a presentation from a patent attorney. I'm so happy to announce additional sponsors, and we have secured a sponsor for best poster award. Which leads me to our new feature at the conference this year – we will be introducing 3 minute student/ECR presentations for all poster presenters. This will provide students/ECRs a chance to practice their scientific 'elevator pitch' and also provides a great opportunity to network later at the conference dinner. As I mentioned in the editorial, we are also awarding our inaugural life-time achievement award to Emeritus Professor Ulrich Hahn and I can't wait to hear his speech. While we wait for the conference, check out our report on last year's conference in the Aptamers Journal. See you all in April in Oxford! And if you can't make it, check out our conference report in the Aptamers Journal later this year.

We would like to thank our sponsors (so far) for their generous support, making this meeting possible.



# **Aptamers Journal**

The Aptamers journal is the official journal of the International Society on Aptamers and will publish studies on all aspects of aptamer research. The journal has a strong belief that both positive and negative data can have a large impact on scientific research so we encourage the submission of both. Do you have a troubling troubleshooting issue that you want to share? A protocol that you are proud of and want to share? Or even some R & D news or an Editorial you want to contribute? We would like to hear from you. We will also be accepting full research articles, research reports, reviews and mini reviews, as well as meeting reports. We're hoping to publish the first meeting report of 4th Aptamers Symposium soon. So if you'd like to publish your work in the first Aptamers journal, please follow this link http://JAptamers.co.uk.



# Aptamers generated in 2017 – list of newly published aptamers

It has been an ongoing challenge to maintain and update a resource to reliably search for aptamers and their targets. There have been several efforts towards storing aptamer information in a database format. For example, Ponomarenko et al. developed the first aptamer database, SELEX\_DB, in 2002 that focused on sequences from SELEX to help define natural DNA and RNA recognition sites for proteins 1. In 2004, the Ellington group created a more complete database, the Aptamer Database, which encompassed information from all in vitro experiments 2. This database was intended for identifying aptamers and unnatural ribozymes that already existed as well as for collecting information about SELEX experiments to gain an understanding of the distribution of functional nucleic acids in the sequence space. In 2006, RiboaptDB was created and contained the same information as the Aptamer Database with a greater emphasis on artificial ribozymes 3. More recently, a collaborative database was introduced (Aptamer Base), using an openly licensed community-built resource 4. Ironically, the goal of this effort was to use this platform to keep the database readily available and updated. Unfortunately, all of these and other databases are either no longer publicly available, or are considerably out-of-date.

Perhaps one of the most currently functional databases is run through Aptagen (<u>https://www.aptagen.com/aptamer-index/aptamer-list.aspx</u>). Here, you can search a list of over 500 aptamers. However, this list is certainly far from complete! In fact, each month, there are several publications reporting new aptamers. In spite of this, the research community continues to make use of a small subset of common aptamers (e.g., thrombin, theophylline) 5. Therefore, our goal is to bring newly-reported aptamers to your attention! Specifically, in each issue we will provide a list of new aptamer targets that have been reported in the literature since the last Newsletter. We hope this will be a valuable resource.

Since this is a new effort, here we first provide a list of reported aptamers that have been characterized with a dissociation constant throughout 2017 (Table 1). We will be using the Pubmed search engine (keywords "aptamer" and "SELEX"). If we have missed any newly reported aptamers, please let us know (<u>mmckeague@gmail.com</u>)! Readers should consult the literature (link provided) for verification and further information.

Target	Link	Nucleic acid type
Human fatty acid binding protein		
(FABP3)	http://www.ncbi.nlm.nih.gov/pubmed/27545084	DNA
Proteasome-Associated		
Deubiquitylating Enzyme UCH37	https://www.ncbi.nlm.nih.gov/pubmed/27930845	RNA
		DNA + one unnatural
von Willebrand Factor A1-Domain	https://www.ncbi.nlm.nih.gov/pubmed/27966933	hydrophobic base
Human epidermal growth factor		
receptor 2 (HER2)	https://www.ncbi.nlm.nih.gov/pubmed/28122449	DNA
Group A Streptococcus serotype M3	https://www.ncbi.nlm.nih.gov/pubmed/28121169	DNA
Bacterial sepsis agents	https://www.ncbi.nlm.nih.gov/pubmed/28119514	DNA
Clenbuterol Hydrochloride	https://www.ncbi.nlm.nih.gov/pubmed/28161951	DNA
Respiratory syncytial virus	https://www.ncbi.nlm.nih.gov/pubmed/28220811	DNA
Vaccine antigen in the human		
papillomavirus (HPV) vaccine Gardasil	https://www.ncbi.nlm.nih.gov/pubmed/28233502	Somamer
Geniposide	https://www.ncbi.nlm.nih.gov/pubmed/28264528	DNA
Carbendazim	https://www.ncbi.nlm.nih.gov/pubmed/28264568	DNA
Proprotein convertase subtilisin/kexin		
type 9	https://www.ncbi.nlm.nih.gov/pubmed/28265062	Somamer
E. coli, E. aerogenes, K. pneumoniae,		
C. freundii, B. subtilis, and S.		
epidermidis	https://www.ncbi.nlm.nih.gov/pubmed/28272554	DNA
AMPA and kainate receptors	https://www.ncbi.nlm.nih.gov/pubmed/28325839	RNA
Dimethylindole red	https://www.ncbi.nlm.nih.gov/pubmed/28391845	DNA
Filament vimentin	https://www.ncbi.nlm.nih.gov/pubmed/28396463	RNA
Bifidobacterium bifidum	https://www.ncbi.nlm.nih.gov/pubmed/28441340	DNA
Gremlin-1	https://www.ncbi.nlm.nih.gov/pubmed/28452949	DNA
Mycobacterium tuberculosis	https://www.ncbi.nlm.nih.gov/pubmed/28454652	DNA
Benzylpenicillin	https://www.ncbi.nlm.nih.gov/pubmed/28522308	DNA
Coagulation factor Xia	https://www.ncbi.nlm.nih.gov/pubmed/28522812	DNA
VEGF and Doxycyline	https://www.ncbi.nlm.nih.gov/pubmed/28634758	DNA and RNA
ATP	https://www.ncbi.nlm.nih.gov/pubmed/28661647	RNA
Mycobacterium tuberculosis H37Rv	https://www.ncbi.nlm.nih.gov/pubmed/28689112	DNA
Cholangiocarcinoma (CCA) cells	https://www.ncbi.nlm.nih.gov/pubmed/28713479	DNA

#### Table 1: Newly-reported aptamers published in 2017.



Glucagon receptor (GCGR)	https://www.ncbi.nlm.nih.gov/pubmed/28775305	DNA
Amanita phalloides	https://www.ncbi.nlm.nih.gov/pubmed/28787470	DNA
E. coli O157	https://www.ncbi.nlm.nih.gov/pubmed/28818557	DNA
Insulin like growth factor II receptor	https://www.ncbi.nlm.nih.gov/pubmed/28839458	RNA
renal cell carcinoma	https://www.ncbi.nlm.nih.gov/pubmed/28841985	DNA
CD123	https://www.ncbi.nlm.nih.gov/pubmed/28845698	DNA
Posaconazole	https://www.ncbi.nlm.nih.gov/pubmed/28861519	DNA
Human lung cancer cell line PC-9	https://www.ncbi.nlm.nih.gov/pubmed/2887519	DNA
PD-1 and PD-L1	https://www.ncbi.nlm.nih.gov/pubmed/28912094	XA Library
Muscovy duck parvovirus	https://www.ncbi.nlm.nih.gov/pubmed/28917743	DNA
Lysosomal-associated membrane		
protein 1	https://www.ncbi.nlm.nih.gov/pubmed/28918021	DNA
Cytotoxic T lymphocyte antigen-4	https://www.ncbi.nlm.nih.gov/pubmed/28918052	DNA
LAG3 (CD223)	https://www.ncbi.nlm.nih.gov/pubmed/28934318	RNA
		Diversely
Thrombin	https://www.ncbi.nlm.nih.gov/pubmed/28938065	functionalized DNA
Pseudomonas aeruginosa 692 (PA692)	https://www.ncbi.nlm.nih.gov/pubmed/28937998	DNA
Proteolytic Amyloidogenic Fragment		
of β2 m	https://www.ncbi.nlm.nih.gov/pubmed/28960840	DNA
Sickle Hemoglobin	https://www.ncbi.nlm.nih.gov/pubmed/29039727	2'F-RNA
Interleukin 2 receptor alpha	https://www.ncbi.nlm.nih.gov/pubmed/29055191	DNA
Breast cancer cells	https://www.ncbi.nlm.nih.gov/pubmed/29054799	DNA
Mammaglobin B (MGB2) and		
mammaglobin A (MGB1)	https://www.ncbi.nlm.nih.gov/pubmed/29101327	DNA
Escherichia coli O157:H7	https://www.ncbi.nlm.nih.gov/pubmed/29242148	DNA
EpCAM	https://www.ncbi.nlm.nih.gov/pubmed/29245156	DNA
Polymorphonuclear myeloid-derived		
suppressor cells	https://www.ncbi.nlm.nih.gov/pubmed/29290791	DNA
Lambda cl repressor	https://www.ncbi.nlm.nih.gov/pubmed/29284756	RNA
Fipronil	https://www.ncbi.nlm.nih.gov/pubmed/29283416	DNA

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2. Lee, J. F.; Hesselberth, J. R.; Meyers, L. A.; Ellington, A. D., Aptamer database. Nucleic acids research 2004, 32 (Database issue), D95-100.

3. Thodima, V.; Pirooznia, M.; Deng, Y., RiboaptDB: a comprehensive database of ribozymes and aptamers. BMC Bioinformatics 2006, 7 Suppl 2, S6.

4. Cruz-Toledo, J.; McKeague, M.; Zhang, X.; Giamberardino, A.; McConnell, E.; Francis, T.; DeRosa, M. C.; Dumontier, M., Aptamer Base: a collaborative knowledge base to describe aptamers and SELEX experiments. Database (Oxford) 2012, 2012, bas006.

5. Mascini, M.; Palchetti, I.; Tombelli, S., Nucleic acid and peptide aptamers: fundamentals and bioanalytical aspects. Angew Chem Int Ed Engl 2012, 51 (6), 1316-32.

# **Aptamers in clinical trials**

Aptamers embrace refined properties as therapeutic molecules that are functionally comparable to antibodies such as the high binding affinity and specificity. However, aptamers have superior advantages compared to antibodies including low toxicity, lack of immunogenicity, can be chemically modified and functionalized with stealth stabilizing motifs, fast chemical and enzymatic production, small size which allow deep tissue penetration, can be selected to bind wide range of targets and can distinguish very closely related molecules, highly stabile and have flexible structure. For therapeutic approaches, aptamers can work as agonist, antagonist, and targeting ligand for drug delivery. Although, aptamers were discovered 28 years ago, there is only one federal approved aptamer-based drug, Pegaptanib (Macugen), which is anti-VEGF used for the treatment of AMD. Moreover, there are several aptamers involved in clinical trials (Table 1), either as direct therapeutic molecules or as targeting ligands for selective delivery of other therapeutic molecules. Such clinical trials offer valuable information for more understanding of aptamers behaviour in human. Despite the tremendous success and the future promises of using of aptamers as therapeutics, there are crucial challenges that need to be solved. The physiochemical properties are complicated and need more investigation and development, the lack of safety data, the pharmacokinetics and pharmacodynamics are not well understood and need more deep exploration. Moreover, the protection by intellectual properties portfolio limit the commercial use and distribution of aptamers. In addition, the promising clinical applications of aptamers need more education and expansion into clinical practitioners. There are more and more researchers dedicated to develop aptamers for clinical applications and the believe that such efforts will provide more understanding to the behaviour aptamers as therapeutic molecules and provide class of therapeutic aptamers that meet the market demands.

4

Table 2: Clinical trials on aptamers

#### Aptamer/ Condition or Location Phase Status **Trial Name** Drug disease Xijing Hospital Nuclear Medicine Department. The Clinical Application of 68Ga Xi'an, Shaanxi, Farly Labeled ssDNA Aptamer Sgc8 in China 68Ga-Sgc8 Recruiting Colorectal cancer phase 1 Healthy Volunteers and Xijing Hospital, **Colorectal Patients** Xi'an, Shaanxi, China (2017-No Contacts or Arrhythmia No Atrial Fibrillation Research In AtrialFibrillation Locations No informati Active Provided information Hypertension CATalonia (AFRICAT) on **Diabetes Mellitus** (2016-Fresno, California, United States Fullerton, Zimura in Subjects With California, United Geographic Atrophy Geographic Atrophy Secondary t States Zimura Recruiting Phase 2 macular Dry Age-Related Macular Mountain View, degeneration Degeneration California, United States and other 31 more locations (2016-University of California Irvine, No Urinary Bladder Molecular Biosensors for Orange, No informati Recruiting **Detection of Bladder Cancer** Neoplasms California, United information on States (2015 Establish the Safety and Idiopathic Polypoidal Pheonix, Arizona, Tolerability of Zimura<sup>®</sup> (AntiC5 Choroidal United States Zimura Phase 2 Completed Aptamer) in Combination With Vasculopathy Anti-VEGF Therapy in Subjects (2015-2017) (IPCV) With (IPCV) Pheonix, Arizona, Anti-PDGF **United States** BB An 18 Month Phase 2a Open Beverly Hills, Pegylated A Label, Randomized Study of California, United ptamer/ Fo States Age related Avastin<sup>®</sup>, Lucentis<sup>®</sup>, or Eylea<sup>®</sup> vista®: Sacramento, Phase 2 Maculare (Anti-VEGF Therapy) bevacizuma Terminated California, United degeneration Administered in Combination h: States and other With Fovista<sup>®</sup> (Anti-PDGF BB ranibizuma Pegylated Aptamer) 21 more b : locations aflibercept (2015-2017) Fovista™(A nti-PDGF-B An Open-Label Investigator Sponsored Trial to Investigate th pegylated a Retinal ptamer) Safety, Tolerability and Consultants of Drugs: **Development of Subfoveal** Neovascular Age-Arizona, Phoenix, Lucentis® Unknown Fibrosis By Intravitreal Phase 1 Related Macular Arizona, United Anti-VEGF Status Administration of Altering Degeneration States \ Avastin® Regimens of Fovista and Anti-(2014-2015) Anti-VEGF VEGF Therapy in Subjects With \ Evlea® Neovascular Age-Related Macula Anti-VEGF Degeneration E10030, Gilbert, Arizona, A Phase 3 Safety and Efficacy **United States** Drug bevaci Study of Fovista® (E10030) Phoenix, Arizona, zumab or Age related maculare Intravitreous Administration in Phase 3 Terminated aflibercept\ Combination With Either Avastin United States degeneration E10030 or Eylea® Compared to Avastin® Tucson, Arizona, United States or Eylea<sup>®</sup> Monotherapy sham



	-	-	-	-	
and other 204					
more locations					
(2014-2017) Germany/ Italy/ United kingdom (2014-2015)	Lexaptepid pegol (NOX-H94) (anti- hepcidin L- RNA- aptamer)	Phase 1,2	Completed	Anemia End Stage Renal Disease	Lexaptepid Pegol (NOX-H94) in ESA-hyporesponsive Anemia in Dialysis Patients
Phoenix, Arizona, United States Bakersfield, California, United States Beverly Hills, California, United States and other 112 more locations (2013-2017)	E10030, Drugs: ranibizuma b \E10030 sham	Phase 3	Terminated	Age related maculare degeneration	A Phase 3 Safety and Efficacy Study of Fovista® (E10030) Intravitreous Administration in Combination With Lucentis® Compared to Lucentis® Monotherapy
Bankstown Lidcome Hospital Bankstown, New South Wales, Australia Royal North Shore Hospital Sydney, New South Wales, Australia Barwon Health Geelong, Victoria, Australia and other 2 more locations (2013-last update 2017)	Nab- Paclitaxel	Phase 2	Active	Breast cancer	IST Neoadjuvant Abraxane in Newly Diagnosed Breast Cancer (Neonab)
Retina Institute of Hawaii, Honolulu, Hawaii, United States, (2011-	pegaptanib sodium (Macugen)	Phase1	Completed	Proliferative Diabetic Retinopathy (PDR)	A Single-Center Trial of Intravitreous Injections of Macugen (Pegaptanib Sodium) Given at Least 7 Days Before Vitrectomy Secondary To Tractional Retinal Detachment in Proliferative Diabetic Retinopathy
Retina Institute of Hawaii, Honolulu, Hawaii, United States (2011-	pegaptanib sodium (Macugen)	No informati on	Available	Diabetic Macular Edema	A Single-Center Trial of High Frequency Pegaptanib for Rapid Restoration of VEGF Levels in Diabetic Retinal Edema (GUARDIAN)
Palmetto Retinal Center, West Columbia, South Carolina, United States (2010-2017)	E10030 plus Lucentis Drug	Phase 2	Completed	Age related macular degeneration (AMD)	E10030 (Anti-PDGF Pegylated Aptamer) Plus Lucentis for Neovascular Age-Related Macular Degeneration
Scope Life Sciences GmbH Hamburg, Germany (2009-2014)	NOX-A12	Phase 1	Completed	Autologous Stem Cell Transplantation	NOX-A12 First-in-human (FIH) Study
Ophthotech, New York, New York, United States (2009-2017)	ARC 1905	Phase 1	Completed	Age related maculare degeneration	ARC1905 (Anti-C5 Aptamer) in Subjects With Dry Age-related Macular Degeneration
No Contacts or Locations Provided (2009-2013)	NOX-E36	Phase1	Completed	Chronic Inflammatory Diseases Type 2 Diabetes Mellitus	NOX-E36 First-in-Human (FIH) Study



				Systemic Lupus	
Ophthotech Corp, New York, New York, United States (2008-2012)	ARC1905	Phase 1	Completed	Erythematosus Age related macular degeneration (AMD)	ARC1905 (ANTI-C5 APTAMER) Given Either In Combination Therapy With Lucentis® 0.5 mg/Eye In Subjects With Neovascular Age-Related Macula Degeneration
No Contacts or Locations Provided (2008-2009)	ARC1779	Phase 2	Withdrawn (sponsor decided not to go forwaed with the study)	Von Willebrand Disease	A Study of the Pharmacokinetics, Pharmacodynamics, and Safety of ARC1779 Injection in Patients With Von Willebrand Disease Type 2B
Archemix Investigational Site, Vienna, Austria (2008-2009)	ARC1779	Phase 2	Completed	Purpura, Thrombotic ThrombocytopenicV on Willebrand Disease Type-2b	ARC1779 Injection in Patients With Von Willebrand Factor- Related Platelet Function Disorders
Denise Teuber, New York, New York, United States (2007-2010)	E10030	Phase 1	Completed	Age related macular degeneration (AMD)	A Phase 1, Safety, Tolerability an Pharmacokinetic Profile of Intravitreous Injections of E1003 (Anti-PDGF Pegylated Aptamer) i Subjects With Neovascular Age Related Macular Degeneration
Charlotte Eye, Ear, Nose and Throat Associates, P.A., Charlotte, North Carolina, United States (2006-2007)	pegaptanib sodium	Phase4	Terminated	Macular degeneration	A Clinical Trial to Explore the Safety and Efficacy of Three Different Doses of Pegaptanib Sodium in Patients With Wet Age Related Macular Degeneration (AMD)
National Heart, Lung and Blood Institute (NHLBI), Bethesda, Maryland, United States ( 2005-2008)	REG1	Phase 1	Completed	Health\ anticoagulant treatment	Safety and Dosing Evaluation of REG1 Anticoagulation System
National Eye Institute (NEI), Bethesda, Maryland, United States (2003-2008)	EYE001 (Anti-VEGF Pegylated aptamer)	Phase 1	Completed	Hippel-Lindau disease	EYE001 to Treat Retinal Tumors i Patients With Von Hippel-Lindau Syndrome
No Contacts or Locations Provided (2002-2006)	pegaptanib sodium (Macugen)	Phase 2	Completed	Diabetic macular edema	Pegaptanib Sodium Compared to Sham Injection in Patients With DME Involving the Center of the Macula
No Contacts or Locations Provided (2002-2006)	pegaptanib sodium (Macugen)	Phase 2,3	Completed	Age related macular degeneration	Study of the Safety, Tolerability and Pharmacokinetics of 1 Mg/Eye and 3 Mg/Eye Pegaptani Sodium in Patients With Exudative Age-Related Macular Degeneration (AMD)
No Contacts or Locations Provided (2001-2006)	pegaptanib sodium (Macugen)	Phase 2,3	Completed	Age related macular degeneration	A Clinical Trial to Explore Safety and Efficacy of Different Doses o Pegaptanib Sodium, Compared t Sham, in Patients With Wet AME
Foundation for Fighting Blindness, Baltimore, Maryland, United States (2001 2005)	EYE001 (anti- VEGF apta mer)	Phase 2,3	Completed	Macular Degeneration Choroidal Neovascularization	Phase II/III Study of Anti-VEGF in Neovascular AMD

# **Aptamer Companies- Updated!!**



Given the growing market in the field of aptamers in recent years and few lingering questions in my mind, I thought of performing extensive web search about aptamer companies and updating the list of companies offering aptamer based services or involved in aptamer research. Are there more companies involved in aptamer research than we know off? Are the companies focused in therapeutics only? Are they all clustered in the US? Quite unexpectedly, I found companies that focus on aptamer development are present worldwide and not limited to the US. As a matter of fact, there are more companies in Europe and UK compared to the US and more and more companies emerging in the Asia Pacific region. Another interesting observation I found during my search is aptamer development in the industry is not restricted to therapeutic applications. Several companies are working towards translating basic aptamer science into development of customizable diagnostic assays and detection products. Furthermore, these companies are developing and using diverse and innovative ways of aptamer selection technology starting from basic chemical synthesis of aptamers through SELEX to use of exclusive technologies such as High Throughput Screening of Aptamers (HTSA) approach and Rapid Isolation of DNA Aptamers (RIDA) method for aptamer discovery. I hope to see all these promising companies growing tremendously in the near future and more new companies coming up and bringing aptamer based products and services to the customers all around the globe.

Table 3. Commercial entities working on aptamers

Name of the	Location	Focus and exclusive technology		
aptamer company				
2bind, GmbH	Germany	The company's <u>customers</u> and collaborators mainly work in the		
		fields of drug development, aptamer generation, and antibody		
		discovery.		
AM Biotechnologies	USA	The company uses a proprietary bead-based technology to select		
		X-Aptamers.		
AMSBIO'S	UK/USA/Deutschland/	The company provides a full range of high quality custom service		
	Switzerland	for a number of key areas including aptamers.		
Aptabharat	India	The company aim to address all the diagnostic and research need		
		by creating specific and high affinity aptamers and by developing		
		aptamer based innovative diagnostic assays, research and teachi		
		products		
Aptagen	USA	The company offers aptamer (synthetic antibody) products and		
10.		services as research reagents, diagnostic and biomarker discover		
		tools, as well as for use in drug discovery and targeted delivery for		
		therapeutics, and bioindustrial applications.		
Aptahem	Sweden	The company develops aptamer-based pharmaceuticals for the		
Aptanem	Sweden	treatment of life-threatening conditions in which a combination		
		coagulation and inflammation are involved.		
Aptamer Group	UK	The Aptamer Group of companies focuses on the development of		
Aptailler Group	UK			
		aptamer technologies. The company develop nucleic acid aptam		
		for use in research & development, biomarker discovery,		
		diagnostics or therapeutic developments.		
AptaIT GmbH	Germany	The company is providing innovative solutions to exploit the full		
		potential of next-generation sequencing data analysis and SELEX		
		Custom NGS data analysis services and software platform		
		COMPAS.		
Aptamer Sciences, Inc.	South Korea	The company is focused on commercializing cutting-edge		
		technologies for analysis of proteins, based on its proprietary		
		aptamer technology platforms.		
Apterna	UK	The company focuses on developing chemically synthesized RNA		
		aptamers that bind to target molecules with outstanding specific		
		and affinity. They are developing internalizing RNA aptamers tha		
		enable targeted delivery of various payloads including RNA, toxir		
		enzymes, chemotherapy agents, photodynamic molecules,		
		radionuclides and nanocarriers.		
Aptitude Medical	USA	The company is revolutionizing molecular recognition – they are		
Systems		enabling the interaction with key molecules which have been out		
,		reach. Their products allow detection and manipulation of protein		
		for cancer, autoimmune, heart disease, and personalized drug		
		response. The company's systems are integrated in devices whic		
		are deployable outside of central facilities, directly at the point o		
		care.		
Aptus Biotech S.L	Spain	APTUS activities are focused on:		
riptus Dioteen s.E	Spain			
		<ul> <li>Aptamer selection services.</li> </ul>		



		<ul> <li>Supply of customized aptamers for different biotechnological applications.</li> </ul>
		R&D projects in collaboration with other research
		centers and companies, where aptamers provide a competitive advantage.
Apta Biosciences	UK/Singapore	The company's focus is on the development of next generation
		affinity molecules for bespoke diagnostic applications.
AptaMatrix	USA	The company's focus is to accelerate the rate of aptamer discove using its patent pending High Throughput Screening of Aptamers
		(HTSA) approach in addition to developing its novel AlloSwitch™
		sensor technology capable of
		a) creating rapid diagnostic tools for detection of chemical and
		biological targets, and b) leveraging this diagnostic platform for drug discovery applications.
AptaTargets	Spain	The company is focused in developing therapeutic applications
17501		based on aptamer technology.
ATDBio Ltd	UK	The company supplies a wide range of custom-made unmodified and chemically modified oligonucleotides for small and large scal
		applications.
AuramerBio	New Zealand	The company specializes in the translation of aptamer science int
		analytical tools and diagnostic assays. It provides customizable sensing solutions for medical and environmental research and
		beyond.
Base Pair Technologies	USA	The company is the provider of highly customized aptamer
PPI Croup		discovery and development services. No response from the company yet
BBI Group Centauri Therapeutics	UK UK	The company's Alphamer™ technology is based on "programmab
Ltd		immunity" in which chemically synthesised molecules redirect
		naturally occurring antibodies to selected pathogens to fight the
		infection. The molecules have two distinct parts: one end binds a cell-surface target on the pathogen using an aptamer whereas th
		other end presents specific epitopes that attach to the circulating
		antibodies.
DSM Biotechnology Dhewa Biotech Private	Netherlands India	No online information available. The company provides services of aptamer, aptamer library,
Limited	illula	monoclonal antibodies, antibody & diagnostics and biosensors
Firefly Bioworks		Acquired by Abcam
Iba GmbH	Germany	The company is dedicated to providing high quality product and
		service solutions for life science research in industry and academ The comprehensive portfolio ranges from nucleic acid custom
		services to products and services for cloning, transfection,
Internets of DNIA	Global(USA/Europe/As	recombinant protein production and cell isolation.
Integrated DNA Technologies (IDT)	(JODAILUNA/FURODE/AS	IDT synthesize nucleic acid aptamers and aptamer libraries, and
		collaborate with outside research groups on aptamer design and
	ia- Pacific/Australia/Japa	collaborate with outside research groups on aptamer design and aptamer applications.
	ia- Pacific/Australia/Japa n)	aptamer applications.
Izon Science	ia- Pacific/Australia/Japa n) Global	aptamer applications. PhD student in the company are working on developing new
Izon Science	ia- Pacific/Australia/Japa n)	aptamer applications.
Izon Science	ia- Pacific/Australia/Japa n) Global (USA/Europe/Asia-	aptamer applications. PhD student in the company are working on developing new platform for sensitively detecting small molecules like environmental pollutants using aptamer functionalized nanoparticles in conjunction with Izon's resistive pulse sensing
	ia- Pacific/Australia/Japa n) Global (USA/Europe/Asia- Pacific)	aptamer applications. PhD student in the company are working on developing new platform for sensitively detecting small molecules like environmental pollutants using aptamer functionalized nanoparticles in conjunction with Izon's resistive pulse sensing technology.
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		expertise is focused on the identification and the characterization of aptamers for biotechnological development of analytical and diagnostic aptamer projects.		
Opthotech corp.	USA	The company's focus is developing novel therapeutics to treat ophthalmic diseases, with a focus on orphan and age-related retinal diseases.		
OTC Biotech	USA	Company's website not available		
Pure Biologics	Poland	The company is developing an innovative, chemically-modified aptamers discovery platform, called PureApta.		
Ribomic	Japan	Ribomic Inc. aims to create artificial RNA aptamers to disease- causing proteins based on the aptamer's superior potential of strong and specific target capturing compared with antibody, a promotes their therapeutic development.		
SomaLogic	USA	The company monitors health and disease through the analysis or protein concentration changes in biological samples. They developed the SOMAscan platform, which allows scientists and researchers to identify protein biomarkers for diseases and conditions, and apply them to drug and diagnostic research and development.		
Тадсух	Japan	TagCyx has established an innovative nucleic acid-based drug discovery platform, the "Xenoligo™ system". It enables screening of drug candidates from diverse oligonucleotide libraries containing the highly functional "fifth base", and stabilizing molecules using our proprietary technology.		
Tocris Bioscience	USA	The company develops aptamers for CD133 and EpCam target.		
TriLink Biotechnologies, Inc	USA	The company provides services for educators and academic researchers in gene therapy, nucleoside chemotherapy, oligonucleotide therapy, and diagnostics. It manufactures custon oligonucleotides, modified nucleoside triphosphates, and CleanAmp PCR products for research, diagnostic, therapeutic, an OEM markets; modified nucleic acid products, including phosphoramidites and other small molecules; and oligonucleotides, including custom synthesis and stocked oligonucleotides.		
Veraptus	China	The company is focused on the research, development, and commercialization of Veraptus diagnostic and therapeutic aptamers.		
Vivonics, Inc.	USA	The company has developed and patented a one-step rapid technique for developing Aptamers called RIDA. Rapid Isolation of DNA Aptamers (RIDA) is a technique for the isolation of high affinity and high selectivity Aptamers, which can be completed in days as compared to months and can produce better performing Aptamers.		



# Interview with a researcher: Beatrix Suess

Professor Dr Beatrix Suess studied Biology at the Ernst-Moritz-Arndt-University Greifswald and the Friedrich-Alexander-University Erlangen-Nürnberg. In Erlangen, she completed her PhD in the lab of Wolfgang Hillen studying structure-function relationship of a bacterial repressor protein TetR. It was the time when scientists discovered that RNA is more than simply the blueprint of our genome but that it has important cellular functions on its own. Therefore she decided to study this interesting biomolecule, with a special focus on the use of *in vitro* selected RNA molecules, so called aptamers, as molecular switches. During her time as junior group leader, she



spent some time as a research fellow in the lab of Ron Breaker (Yale University) and Renee Schroeder (Vienna University). In 2007, she was appointed associated professor for Chemical Biology, and in 2013, full professor for Synthetic Biology at the Technical University, Darmstadt. Still, RNA is her main interest, and Beatrix wants to understand how RNA molecules can exert regulation.

## Q1) How did you become interested in the field of aptamers?

To be honest – it was a fortunate coincidence. The topic was suggested by my former supervisor, right after finishing my PhD and I didn't really know what was coming, I was eager to learn something new, I never regret it ... ©

I was also fascinated by the interdisciplinarity of the field.

## Q2) From your point of view, what is unique about aptamers?

Their perfect mixture of physics, chemistry, biology and how they interact with gene regulation. Also their widespread applicability (due to stability, size, inherent structure, specificity and affinity...) to a range of problems in vivo and in vitro.

# Q3) What do you think is the future of aptamers?

Better and diversified SELEX will increase the ligand range, efficiency and open up new fields of research and application.

### Q4) What are the major challenges that need to be solved?

Improved speed and reproducibility of SELEX, tackling difficult targets (maybe develop a universal SELEX procedure?), understanding the black box of the SELEX process better.

## Q5) What we should do for the aptamer science?

Improve networks between people who need aptamers (specific or random) and people who have expertise in generating them. This would solve plenty of communications problems, speed up the development of real-world applications and further the standing of the field (e.g. as an alternative to antibodies).

## Q6) Tell us about your research.

We specialize in developing novel, synthetic riboswitches based on small molecule binding aptamers. The goal is to establish a comprehensive and reliable toolbox for synthetic biology.

## Q7) How did you know about the INSOAP?

From the very beginning, I was attending the Oxford Aptamer conferences – I was fascinated by the fact that there is a conference dedicated to the molecules I am mostly interested in <sup>(2)</sup>.

## Q8) How will you support the INSOAP?

I will encourage all members and guests of my lab to become member the society. Q9) What kind of advice can give to the young researchers about aptamers? Aptamers are way more cool than proteins! Also, there are so many different methods you can apply to them, that studying aptamers teaches you a lot of the tools necessary for all of synthetic biology.

## Q10) What is your personal philosophy on life and science?

Make science repeatable, scalable and reduce the "magic" that seems to go into some experiments.



Q11) What was your favourite part about research? Exchange and discussion with my students and colleagues.

#### Q12) What do you like to do in your free time?

Spend time with family and friends, relax by doing sports or arts and crafts.

## Q13) Any other fun facts/tidbits you'd like us to know!

If you do a PCR with an aptamer-pool, containing (for secret reasons) glucose, your PCR cycler starts to smell like caramel. We do not take responsibility of the downstream application, though.

(answers also contain input from the Suess lab members <sup>(2)</sup>)

# **Nominations for INSOAP committee**

We are currently asking for expressions of interest for membership of the management committee of INSOAP. If you would like to be an integral part of our Society as it moves forward, please contact me at sarah.shigdar@deakin.edu.au.

# Updates to the website

We have been working on updating the website for INSOAP and you will now see that we have a listing of all aptamer companies throughout the world, as well as a listing of all the aptamer laboratories to date. If we haven't got you listed, please get in touch and we will add you to our growing list. We are also providing a careers page so please get in touch with any vacancies you wish to be listed. Finally, if there are any suggestions for improvements to the website, please contact us and we will make the changes.

## Keep in touch

http://aptamersociety.org http://www.linkedin.com/groups/8282517 www.facebook.com/AptaSoc https://twitter.com/AptamerSociety https://twitter.com/aptamer\_connect.