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FILED AS OF DATE: 20190923
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FILER:

COMPANY DATA:

COMPANY CONFORMED NAME: Anchiano Therapeutics Ltd.
CENTRAL INDEX KEY: 0001534248
STANDARD INDUSTRIAL CLASSIFICATION: PHARMACEUTICAL PREPARATIONS [2834]
IRS NUMBER: 000000000
STATE OF INCORPORATION: L3
FISCAL YEAR END: 1231

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SEC ACT: 1934 Act
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BUSINESS ADDRESS:

STREET 1: 1/3 HIGH-TECH VILLAGE
STREET 2: GIVAT RAM, P.0. BOX 39264
CITY: JERUSALEM
STATE: L3
ZIP: 9139102
BUSINESS PHONE: 972-2-5486555

MAIL ADDRESS:

STREET 1: 1/3 HIGH-TECH VILLAGE
STREET 2: GIVAT RAM, P.0. BOX 39264
CITY: JERUSALEM
STATE: L3
ZIP: 9139102

FORMER COMPANY:

FORMER CONFORMED NAME: BioCancell Ltd.
DATE OF NAME CHANGE: 20111104

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<P STYLE="font: 14pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">SECURITIES AND EXCHANGE COMMISSION </P>

<P STYLE="font: 14pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">WASHINGTON, D.C. 20549</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">REPORT OF FOREIGN PRIVATE ISSUER PURSUANT
TO RULE 13a-16 or 15d-16 UNDER THE </P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">SECURITIES EXCHANGE ACT OF 1934</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">For the month of September 2019</P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center"> </P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">Commission File Number: 001-38807</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">ANCHIANO THERAPEUTICS LTD.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">(Translation of registrant's name into English)</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">1/3 High-Tech Village, Givat Ram, P.O. Box 39264</P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">Jerusalem, 9139102 Israel</P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">(Address of principal executive office)</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">Form 20-F x ¨ </P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1): ¨ </P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-indent: 24.5pt; background-color: white">On September 20, 2019, Anchiano Therapeutics, Inc. (“Anchiano”), a Delaware corporation and wholly-owned subsidiary of Anchiano Therapeutics Ltd. (the “Company”), and ADT Pharmaceuticals, LLC, a Delaware corporation (“ADT”), entered into a collaboration and license agreement with an effective date as of September 13, 2019 (the “Collaboration Agreement”). Pursuant to the terms and conditions set forth in the Collaboration Agreement, the parties agreed to use commercially reasonable efforts to conduct research and development activities of novel small-molecule inhibitors (RAS and PDE10/β-catenin) (“Compounds”) under the oversight of a joint steering committee established by the parties. As part of the arrangement, Anchiano will be primarily responsible for the research, development,

manufacturing and regulatory activities and ADT will assist with the research activities as necessary in exchange for a quarterly fee from Anchiano. In connection with the Collaboration Agreement, ADT also granted Anchiano an exclusive option to research, develop, manufacture and commercialize Compounds relating to patents owned by ADT and any products containing such Compounds worldwide (the "Option"). In consideration for the rights granted under the Collaboration Agreement, Anchiano will pay ADT (i) a \$3 million upfront fee; (ii) a fee upon exercise of the Option; and (iii) milestone payments with respect to the development and commercialization of any products containing the Compounds. In addition, Anchiano will pay ADT royalties ranging in the low- to mid-single digit percentage on sales of any products containing the Compounds. Anchiano may terminate the Collaboration Agreement at any time in its entirety or on a compound-by-compound basis after providing 90 days written notice to ADT.

The foregoing summary of the Collaboration Agreement is qualified in its entirety by reference to the complete text of the Collaboration Agreement, a copy of which is filed as Exhibit 10.1 and is incorporated herein by reference.

On September 23, 2019, the Company issued a press release announcing that Anchiano has entered into the Collaboration Agreement with ADT, as described above, and providing an update to its ongoing pivotal Phase 2 Codex Trial and the timing of the 35-patient interim analysis. The Company currently has 31 patients enrolled or in active screening, of which 23 have been dosed, and estimates that data collection and interim analysis will be completed by the first quarter of 2020.

A copy of this press release is attached hereto as Exhibit 99.1 and is incorporated herein by reference.

Exhibit Index

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<p>No.</p>

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Description

10.1	Collaboration and License Agreement, dated as of September 13, 2019, by and between Anchiano Therapeutics Inc. and ADT Pharmaceuticals, LLC.
99.1	Press Release dated September 23, 2019.

* Portions of this exhibit have been omitted in accordance with Regulation S-K Item 601(b)(2). The registrant hereby undertakes to furnish a unredacted copy of the exhibit to the Securities and Exchange Commission upon its request.

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SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

Date: September 23, 2019

Anchiano Therapeutics Ltd.	

	By:</TD>
/s/ Dr. Frank G. Haluska</TD>	
	Name: Dr. Frank G. Haluska</TD>
	Title: Chief Executive Officer</TD>

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<P STYLE="margin: 0; text-align: right">Exhibit 10.1</P>

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<P STYLE="margin: 0">CERTAIN PORTIONS OF THIS EXHIBIT, MARKED BY BRACKETS AND ASTERISKS [***], WERE OMITTED BECAUSE THOSE PORTIONS ARE NOT MATERIAL AND WOULD BE COMPETITIVELY HARMFUL TO THE COMPANY IF PUBLICLY DISCLOSED.</P>

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<P STYLE="margin: 0"></P>

<P STYLE="font: bold 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">COLLABORATION AND LICENSE AGREEMENT</P>

<P STYLE="font: bold 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center"> </P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; text-indent: 0.5in">THIS COLLABORATION AND LICENSE AGREEMENT (this “Agreement”) is entered into as of September 13, 2019 (the “Effective Date”) by and between ADT PHARMACEUTICALS, LLC, a company organized and existing under the laws of Delaware and having offices at 31691 Shoal Water Dr. Orange Beach, Alabama 36561 (“ADT”) and ANCHIANO THERAPEUTICS, INC., a company organized under the laws of Delaware and having offices at One Kendall Square, Building 500, Suite 6-106, Cambridge, MA (“Anchiano”). ADT and Anchiano are sometimes referred to herein individually as a “Party” and collectively as the “Parties”</P>

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<P STYLE="font: bold 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">BACKGROUND</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; text-indent: 0.5in">WHEREAS, ADT owns or otherwise controls certain patents, patent applications, technology, know-how, scientific and technical information and other proprietary rights and information relating to certain compounds, pharmaceutical compositions thereof and methods of using the same;</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; text-indent: 0.5in">WHEREAS, Anchiano is a biopharmaceutical company focused on the research, discovery, development, and commercialization of pharmaceutical products; and</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; text-indent: 0.5in">WHEREAS, Anchiano and ADT desire to collaborate together to research, develop and commercialize Compounds and Products

Development means, together with all correlative meanings, pre-clinical and clinical drug development activities, conducted before or after obtaining Regulatory Approval that are reasonably related to or leading to the development, preparation, and submission of data and information to a Regulatory Authority for the purpose of obtaining, supporting or expanding Regulatory Approval or to the appropriate body for obtaining, supporting or expanding pricing and reimbursement approval, including without limitation, all activities related to preclinical testing, assay development and validation, in vivo testing, biomarker development and validation, toxicology, pharmacokinetic profiling, design and conduct of Clinical Trials and any other clinical trials or studies, regulatory affairs, statistical analysis, report writing, and regulatory filing creation and submission (including the services of outside advisors and consultants in connection therewith). Development expressly excludes (a) Research, (b) Commercialization and (c) the Manufacture and accumulation of commercial inventory of a product.

1.32 **Effective Date** has the meaning set forth in the preamble to this Agreement.

1.33 **EMA** means the European Medicines Agency or its successor.

1.34 **EU** means all of the European Union member states as of the applicable time during the Term.

1.35 **Executive Officer** means (a) in the case of Anchiano, Frank G. Haluska, and (b) in the case of ADT, Michael R. Boyd.

surveillance studies and clinical or other research studies.

1.82 Product means any product

containing a Compound as an active ingredient, in any form, presentation, dosage, or formulation. For clarity, a Product includes a Combination Product.

1.83 Publishing Party has the meaning set forth in Section 10.4.

1.84 RAS Inhibitor Compound means (a) any compound, or prodrug thereof, claimed or disclosed specifically or generically by the Optioned IP or Licensed IP, wherein said compound is a RAS inhibitor [***], and (b) any RAS inhibitor compounds, or prodrug thereof, derived in one or more steps by either Party starting from any of the foregoing compounds, prodrugs, precursors or functional sub-units thereof.

1.85 RAS Inhibitor Product means any Product containing a RAS Inhibitor Compound as an active ingredient.

1.86 Registrational Cohort or Study means a pivotal cohort of a study or a pivotal study that is designed and executed to get statistically significant evidence of efficacy and safety as required by Regulatory Authorities for approval of an NDA or MAA.

1.87

and Sublicensees to keep (as applicable), adequate books and records of accounting for the purpose of calculating all amounts due to ADT hereunder. For three (3) years next following the end of the calendar year to which each will pertain, such books and records of accounting (including those of Anchiano's Affiliates and Sublicensees, as applicable) will be made available for inspection at reasonable times and upon reasonable notice by an independent certified accountant selected by ADT, and which is reasonably acceptable to Anchiano, for the sole purpose of inspecting the amounts due to ADT under this Agreement. In no event will such inspections be conducted hereunder more frequently than once every twelve (12) months or cover more than thirty-six (36) months prior to the date of request for inspection. Such accountant must have executed and delivered to Anchiano and its Affiliates and Sublicensees, as applicable, a confidentiality agreement as reasonably requested by Anchiano, which will include provisions limiting such accountant's disclosure to ADT to only whether the royalty reports are correct or incorrect and the amount of any discrepancy. The results of such inspection, if any, will be binding on both Parties if not disputed within thirty (30) days following receipt by the Parties of the inspection report. Any such dispute over an inspection report shall be subject to the dispute resolution procedure of Article 12, and no payment shall be required until the dispute is resolved. If it is determined that Anchiano underpaid, Anchiano shall pay to ADT such amount it was determined to have within thirty (30) days of such determination. If it is determined that Anchiano overpaid, ADT shall pay to Anchiano such amount it was determined to have been overpaid within thirty (30) days following such determination. Any undisputed underpayments will be paid by Anchiano within thirty (30) days of notification of the results of such inspection. Any undisputed overpayments will be fully creditable against amounts payable in subsequent payment periods. ADT will pay for such inspections, except that in the event there is any upward adjustment in amounts payable for any calendar year shown by such inspection of more than [***], Anchiano will reimburse ADT for any reasonable out-of-pocket costs of such accountant.

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(b) <u>Taxes</u>. Where required by Applicable Law, Anchiano shall have the right to withhold applicable taxes from any payments to be made by Anchiano to ADT pursuant to this Agreement; provided that, to the extent allowed by Applicable Law, prior to such withholding, Anchiano shall give written notice of its

intention to withhold and allow ADT sufficient time to furnish any documentation or forms to the applicable Governmental Authority to minimize or eliminate such withholding. Anchiano shall provide ADT with receipts from the appropriate taxing authority for all payments of taxes withheld and paid by Anchiano to such authorities on behalf of ADT. ADT shall have the right to appeal to the appropriate taxing authority any such withholding and payment of such taxes.

(c) Interest Due. Anchiano will pay ADT interest on any payments that are not paid on or before the date such payments are due under this Agreement at a rate of [***] or the maximum applicable legal rate, if less, calculated on the total number of days payment is delinquent.

(d) No Other Compensation. Other than as explicitly set forth (and as applicable) in this Agreement or the CCRSA, Anchiano will not be obligated to pay any additional fees, milestone payments, royalties or other payments of any kind to or on behalf of ADT or its Affiliates under this Agreement.

(e) Right to Set-off. Either Party will have the right to deduct from amounts otherwise payable hereunder any amounts payable to such Party (or its Affiliates) from the other Party (or its Affiliates) under this Agreement that have been determined by a final, non-appealable judgment by a court of competent jurisdiction or otherwise agreed to by the Parties.

Article
INTELLECTUAL PROPERTY

7.1

be provided to the other Party within five (5) days after receipt of such notification and will be sent to the address set forth in Section 13.3.

Article

**REPRESENTATIONS,
WARRANTIES AND COVENANTS**

8.1 Mutual Representations, Warranties and Covenants. Each Party hereby represents and warrants to the other Party as of the Effective Date, and covenants, as applicable, as a material inducement for such other Party's entry into this Agreement, as follows:

(a) Corporate Existence and Power. It is a company or corporation duly organized, validly existing, and in good standing under the laws of the jurisdiction in which it is incorporated, and has full corporate power and authority and the legal right to own and operate its property and assets and to carry on its business as it is now being conducted and as contemplated in this Agreement, including the right to grant the licenses granted by it hereunder.

(b) Authority and Binding Agreement. (i) It has the corporate power and authority and the legal right to enter into this Agreement and perform its obligations hereunder; (ii) it has taken all necessary corporate action on its part required to authorize the execution and delivery of this Agreement and the performance of its obligations hereunder; and (iii) this Agreement has been duly executed and delivered on behalf of such Party, and constitutes a legal, valid, and binding obligation of such Party that is enforceable against it in accordance with its terms.

by ADT to Anchiano, neither ADT nor any of its Affiliates have entered into any agreements with Third Parties with respect to the Compounds or Products.

(f) Disclosure of Information. All Information and data provided by or on behalf of ADT to Anchiano or its agents or representatives prior to or on the Effective Date with respect to this Agreement was and is true, accurate and complete in all material respects, and ADT has not disclosed, failed to disclose or caused to be disclosed any Information or data that could reasonably be expected to be misleading in any material respect.

(g) Employee Assignment. As of the Effective Date, ADT has secured from all employees, consultants, contractors and other Persons who have contributed to the development, creation, conception or invention of any of the Licensed IP a written agreement assigning to ADT or its Affiliates all rights to such developments, creations, conceptions or inventions, or Licensed IP and such Affiliates have assigned such rights to ADT, and neither ADT nor any of its Affiliates has received any written communication challenging ADT's ownership or right to the Licensed IP, unless such an agreement with the inventor is not required under Applicable Law for ownership in such Licensed IP to vest in ADT.

(h) All Material Information Furnished. As of the Effective Date, ADT has furnished or made available to Anchiano or its agents or representatives all material information that is in ADT's or its Affiliates' possession concerning the Compounds, the Products (in each case in the form being developed by ADT as of the Effective Date) and the Licensed IP, including relevant to the safety or efficacy of such Compounds and Products, and all material regulatory filings and other material correspondence with Regulatory Authorities relating to any such Compound or Product, and such information is accurate, complete and true in all material respects.

(i)

as such to be considered Confidential Information. The terms of this Agreement shall be deemed to be the Confidential Information of both Parties. During the Term, information relating to the Compounds or Products shall be deemed to be the Confidential Information of Anchiano. Notwithstanding the foregoing, Confidential Information will not include any information to the extent that it can be established by written documentation by the receiving Party that such information:

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(a) was already known to the receiving Party, other than under an obligation of confidentiality (except to the extent such obligation has expired or an exception is applicable under the relevant agreement pursuant to which such obligation was established), at the time of disclosure;</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; text-indent: 1.5in">
(b) was generally available to the public or otherwise part of the public domain at the time of its disclosure to the receiving Party;</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; text-indent: 1.5in">
(c) became generally available to the public or otherwise part of the public domain after its disclosure and other than through any act or omission of the receiving Party in breach of this Agreement;</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; text-indent: 1.5in">
(d) was independently developed by the receiving Party as demonstrated by written documentation prepared contemporaneously with such independent development; or</P>

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copies of the disclosure to the other Party as far in advance of such filing or other disclosure as is reasonably practicable under the circumstances, (B) such Party has promptly notified the other Party in writing of such requirement and any respective timing constraints, and (C) such Party has given the other Party a reasonable time under the circumstances from the date of notice by such Party of the required disclosure to comment upon, request confidential treatment or approve such disclosure, then such Party will have the right to make such public disclosure at the time and in the manner reasonably determined by its counsel to be required by Applicable Law. Notwithstanding anything to the contrary herein, it is hereby understood and agreed that if a Party seeking to make a disclosure to the SEC as set forth in this Section 10.2, and the other Party provides comments within the respective time periods or constraints specified herein or within the respective notice, the Party seeking to make such disclosure or its counsel, as the case may be, will in good faith (1) consider incorporating such comments and (2) use reasonable efforts to incorporate such comments, limit disclosure or obtain confidential treatment to the extent reasonably requested by the other Party. Each Party will have the right to issue additional press releases or to make public disclosures with the prior written agreement of the other Party.

with Anchiano or ADT prior to any disclosure of patentable subject matter, (c) that all such publications and presentations are consistent with good scientific practice and accurately reflect work done and the contributions of the Parties, and (d) that no such publication or presentation be made except to the extent approved by the JSC in advance in writing. Unless otherwise mutually agreed upon by the Parties, (i) the Party desiring to publish or present any publication or presentation concerning the activities to be conducted hereunder (the "Publishing Party") will transmit to the other Party (the "Reviewing Party") for review and comment a copy of the proposed publication or presentation, at least thirty (30) days prior to the proposed submission of the publication or presentation to a Third Party; and (ii) the Publishing Party will postpone the publication or presentation upon request by the Reviewing Party in order to allow the consideration of appropriate patent applications or other protection on information contained in the publication or presentation.

10.5
Attorney-Client Privilege. Neither Party is waiving, nor will be deemed to have waived or diminished, any of its attorney work product protections, attorney-client privileges or similar protections and privileges as a result of disclosing information pursuant to this Agreement, or any of its Confidential Information (including Confidential Information related to pending or threatened litigation) to the receiving Party, regardless of whether the disclosing Party has asserted, or is or may be entitled to assert, such privileges and protections. The Parties: (a) share a common legal and commercial interest in such disclosure that is subject to such privileges and protections; (b) are or may become joint defendants in proceedings to which the information covered by such protections and privileges relates; (c) intend that such privileges and protections remain intact should either Party become subject to any actual or threatened proceeding to which the disclosing Party's Confidential Information covered by such protections and privileges relates; and (d) intend that after the Effective Date both the receiving Party and the disclosing Party will have the right to assert such protections and privileges.

Article 11
TERM AND TERMINATION

11.1
Term. This Agreement will commence on the Effective Date and, unless earlier terminated pursuant to this Article 11, will expire on a country-by-country basis and Product-by-Product basis at the end of the applicable Royalty Term (the "Term"). Following the end of the Term for any such

part, any corporate name
or logo of Anchiano or its Affiliates or Sublicensees).

(c) In consideration of the licenses to be granted by Anchiano to ADT under Section 11.5(a) and Section 11.5(b), ADT shall pay Anchiano a royalty on annual worldwide Net Sales of Reversion Compounds or Reversion Products at the royalty rates set forth in the below table only to the extent the sale of such Reversion Compounds or Reversion Products, as applicable, would otherwise infringe on the Reversion IP:

Timing of Termination	Royalty Rate

Field: Page; Sequence: 29; Value: 21

Field: Sequence; Type: Arabic; Name: PageNo -->29

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11.6

Rights in Bankruptcy. All rights and licenses granted under or pursuant to this Agreement by ADT and Anchiano are, and will otherwise be deemed to be, for purposes of Section 365(n) of the U.S. Bankruptcy Code, licenses of right to intellectual property as defined under Section 101 of the U.S. Bankruptcy Code. The Parties agree that each Party, as licensee of certain rights under this Agreement, will retain and may fully exercise all of its rights and elections under the U.S. Bankruptcy Code. The Parties further agree that, in the event of the commencement of a bankruptcy proceeding by or against a Party (such Party, the "Bankrupt Party") under the U.S. Bankruptcy Code, the other Party will be entitled to a complete duplicate of (or complete access to, as appropriate) any intellectual property licensed to such other Party and all embodiments of such intellectual property, which, if not already in such other Party's possession, will be promptly delivered to it (a) upon any such commencement of a bankruptcy proceeding upon such other Party's written request therefor, unless the Bankrupt Party elects to continue to perform all of its obligations under this Agreement or (b) if not delivered under clause (a), following the rejection of this Agreement by the Bankrupt Party upon written request therefor by the other Party. The Parties acknowledge and agree that of the milestones and royalties to be paid pursuant to Article 6, only the royalties contained in Section 6.4 will constitute royalties within the meaning of Bankruptcy Code section 365(n) with respect to the licenses of intellectual property hereunder.

11.7

Other Remedies. Termination or expiration of this Agreement for any reason will not release either Party from any liability or obligation that already has accrued prior to such expiration or termination, nor affect the survival of any provision hereof to the extent it is expressly stated to survive such termination. Termination or expiration of this Agreement for any reason will not constitute a waiver or release of, or otherwise be deemed to prejudice or adversely affect, any rights, remedies or claims, whether for damages or otherwise, that a Party may have hereunder or that may arise out of or in connection with such termination or expiration.

11.8

Survival. Termination or expiration of this Agreement will not affect rights or obligations of the Parties under this Agreement that have accrued prior to the date of termination or expiration of this Agreement. Notwithstanding anything to the contrary, the following provisions will survive and apply after expiration or termination of this Agreement in its entirety: Sections 7.1, 7.2(b), 8.3, 11.4, 11.5, 11.6, 11.7 and 11.8 and Article 1, Article 9, Article 10, Article 12 and Article 13. In addition, the other applicable provisions of Article 6 will survive such expiration or termination of this Agreement in its entirety to the extent required to make final reimbursements, reconciliations or other payments incurred or accrued prior to the date of termination or expiration. For any surviving provisions requiring action or

decision by the JSC or an Executive Officer, each Party will appoint representatives to act as its JSC members or Executive Officer, as applicable. All provisions not surviving in accordance with the foregoing will terminate upon expiration or termination of this Agreement and be of no further force and effect.

Article 12

DISPUTE RESOLUTION

12.1 **Dispute Resolution**. If an unresolved dispute arises out of or relates to this Agreement, or the breach thereof, either Party may refer such dispute to the Executive Officers of Anchiano and ADT, who shall meet in person or by telephone within thirty (30) days after such referral to attempt in good faith to resolve such dispute. If such matter cannot be resolved by discussion of such officers within such thirty (30) days period (as may be extended by mutual agreement), either Party shall be entitled to seek resolution of such dispute pursuant to Section 12.2 below.

12.2 **Arbitration**. If the parties are unable to resolve a dispute on an issue of interpretation, breach or enforcement of this Agreement, the parties shall refer such dispute to be finally resolved by binding arbitration under the terms of this Section 12.2, except that all disputes with respect to the validity or infringement of Patents shall be subject to applicable federal court jurisdiction and not subject to the terms of this Section 12.2. Whenever a party shall decide to institute arbitration proceedings, it shall give written notice to that effect to the other party. Any such arbitration shall be conducted under the Commercial Arbitration Rules and Mediation Procedures (including procedures for large, complex commercial disputes) of the American Arbitration Association by a panel of three (3) arbitrators in Boston, Massachusetts. Each party shall select one (1) arbitrator who is not employed by, or otherwise affiliated with, such party within fifteen (15) days after the institution of arbitration proceedings, and the two (2) arbitrators so selected shall designate the third arbitrator. The parties shall use their commercially reasonable efforts to conclude the arbitration hearings within six (6) months following the confirmation of the third and presiding arbitrator.

limitation" (or "includes without limitations"). "Herein," "hereby," "hereunder," "hereof" and other equivalent words refer to this Agreement as an entirety and not solely to the particular portion of this Agreement in which any such word is used. The term "or" means "and/or" hereunder. All definitions set forth herein will be deemed applicable whether the words defined are used herein in the singular or the plural. Unless otherwise provided, all references to Sections and Exhibits in this Agreement are to Sections and Exhibits of this Agreement. References to any Sections include Sections and subsections that are part of the related Section (e.g., a section numbered "Section 3.2" would be part of "Section 3"; and references to "Section 3.2" would also refer to material contained in the subsection described as "Section 3.2(a)"). Unless otherwise stated, dollar amounts set forth in this Agreement are U.S. dollars.

13.6 Assignment. Neither Party may assign or transfer (whether by operation of Applicable Law or otherwise) this Agreement or any rights or obligations hereunder without the prior written consent of the other, except that a Party may make such an assignment without the other Party's consent to an Affiliate or to a successor to substantially all of the business to which this Agreement relates, whether in a merger, sale of stock, sale of assets, reorganization or other transaction. Any permitted successor or assignee of rights or obligations hereunder will expressly assume performance of such rights or obligations (and in any event, any Party assigning this Agreement to an Affiliate will remain bound by the terms and conditions hereof). Any permitted assignment will be binding on and inure to the benefit of the successors of the assigning Party. Any assignment or attempted assignment by either Party in violation of the terms of this Section 13.6 will be null, void and of no legal effect.

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13.7 Further Actions. Each Party agrees to execute, acknowledge and deliver such further instruments, and to do all such other acts, as may be necessary or appropriate in order to

carry out the purposes and intent of this Agreement.

13.8

Severability. If any one or more of the provisions of this Agreement is held to be invalid or unenforceable by an arbitrator or by any court of competent jurisdiction from which no appeal can be or is taken, the provision will be considered severed from this Agreement and will not serve to invalidate any remaining provisions hereof. The Parties will make a good faith effort to replace any invalid or unenforceable provision with a valid and enforceable one such that the objectives contemplated by the Parties when entering into this Agreement may be realized.

13.9

No Waiver. Any delay in enforcing a Party's rights under this Agreement or any waiver as to a particular default or other matter will not constitute a waiver of such Party's rights to the future enforcement of its rights under this Agreement, except with respect to an express written and signed waiver relating to a particular matter for a particular period of time.

13.10

Independent Contractors. Each Party will act solely as an independent contractor, and nothing in this Agreement will be construed to give either Party the power or authority to act for, bind, or commit the other Party in any way. Nothing herein will be construed to create the relationship of partners, principal and agent, or joint-venture partners between the Parties.

13.11

Counterparts. This Agreement may be executed in one (1) or more counterparts, by facsimile, pdf or other electronic format, each of which will be deemed an original, but all of which together will constitute one and the same instrument.

13.12

	ANCHIANO:
	ANCHIANO THERAPEUTICS, INC.
	Name:
	Title:

</P>

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<P STYLE="font: bold 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">SCHEDULE 1.72</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">EXHIBIT A</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">CCRSA Terms</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center"><I>Consulting and Collaboration Research Support Agreement</I></P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">Description and Purpose:</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify"><I>Under the CCRSA, the CCRSA Fee from Anchiano to ADT will enable ADT to provide dedicated, consultative and hands-on research support for a) continued strengthening and broadening of the Optioned Patents or Licensed Patents and b) facilitation of tactical and strategic decision-making in Anchiano's preclinical and clinical Research and Development of the Compounds and/or Products. If Anchiano exercises the Option, the CCRSA Fee would continue for the first forty-eight months of the Term of the Collaboration and License Agreement.</I></P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">Functions supported:</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify"><I>ADT's functions in support of Anchiano's drug development program for ADT's RAS and PDE10 inhibitors may include, but are not limited to, the following examples:</I></P>

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<TD STYLE="width: 0.25in"><TD STYLE="width: 0.25in">·

</TD><TD STYLE="text-align: left">Consulting, and hands-on small-scale synthesis and biological investigations of novel compounds and methods of use encompassed in as well as extending from the Optioned Patents or Licensed Patents;</TD></TR></TABLE>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">CCRSA Fee:</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-indent: 0in"><I>Starting with execution of the CCRSA, Anchiano will pay ADT <U>[***]</U>. The base quarterly fee will be allocated internally by ADT to partially cover ADT's direct and indirect costs, including but not limited to: (a) salary support for ADT employee(s); (b) materials, supplies and minor equipment; (c) lab lease space; (d) insurance; (e) accounting services; (f) attorney fees; (g) consultant costs, (h) business travel and meeting participation; (i) other necessary expenses, all as more fully set forth in the CCRSA. </I></P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-indent: 0in"><I>In addition to the base CCRSA Fee, supplemental equipment and supply funds may be provided by Anchiano to ADT for major equipment and/or supply purchases (single item above \$5,000), requested and justified by ADT, subject to approval by Anchiano. Supplemental equipment and supply funds will be used only for major equipment and/or supply purchases specifically approved and funded by Anchiano. </I></P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-indent: 0in"><I>In addition to the base CCRSA Fee and any supplemental equipment and supply funds, supplemental subcontract funds may be provided by Anchiano to ADT for subcontracted services or facilities access provided to ADT by the University of South Alabama (USA). Supplemental subcontract funds may be requested and justified by ADT on an as-needed basis, subject to approval and funding by Anchiano; such funds will be used only for services or facilities access provided by USA to ADT solely for support of Anchiano's drug development program for the Compounds and/or Products. </I></P>

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<P STYLE="margin: 0; text-align: right">Exhibit 99.1</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">FOR IMMEDIATE RELEASE </P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0"> </P>

<P STYLE="font: 14pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center">Anchiano Therapeutics Enters into An Exclusive Option To License Agreement for Novel Pan-RAS Inhibitor and PDE10/β-catenin Inhibitor Programs </P>

<P STYLE="font: 14pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center"> </P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center"><I>Anchiano will make an upfront payment in exchange for the option to in-license at any time through obtaining an Investigational New Drug (IND) designation</I></P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center"><I>Company also provides update on accrual to its ongoing pivotal Phase 2 Codex trial </I></P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center"><I>Anchiano to hold a call on Tuesday, September 24, 2019 at 8:00 am Eastern Time</I></P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: center"> </P>

<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">CAMBRIDGE, Mass., September 23, 2019 – Anchiano Therapeutics Ltd. (Nasdaq: ANCN) (“Anchiano” or the “Company”), a pivotal-stage biopharmaceutical company focused on the discovery and development of targeted therapies to treat cancer, today reported that Anchiano Therapeutics, Inc., the Company’s U.S. subsidiary, has entered into an exclusive worldwide collaboration and option to license agreement with ADT Pharmaceuticals, LLC (“ADT”) to develop novel small-molecule inhibitors of RAS and PDE10/β-catenin. This collaboration reflects Anchiano’s ongoing strategy to grow a pipeline beyond its pivotal-stage asset, inodiftagene vixteplasmid, with programs that have the potential to address significant clinical needs, while leveraging Anchiano’s expertise in small-molecule oncology development.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">“We are excited to bring both of these important programs aimed at difficult-to-treat genetically-defined cancers into our portfolio. They complement our pivotal program and Codex trial and underscore our commitment to develop therapies with targeted approaches,” said Frank Haluska M.D., Ph.D., President and Chief Executive Officer of Anchiano. “Mutations in RAS are found in approximately one-third of all cancers. Recent exciting advances have been made in treating a subset of these cancers, but the successful development of a RAS-targeted therapy with broad inhibitory activity, irrespective of RAS isoform or mutation, has the potential for great clinical impact. Likewise, APC or β-catenin alterations are found almost uniformly in colorectal cancer and polyposis syndromes, and are observed in other tumor types as well, but effective targeted approaches to these lesions have been lacking. We are enthusiastic about the opportunity to develop PDE10 inhibition to target the Wnt/APC/β-catenin pathway where it is an oncogenic driver.”</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">Michael Boyd M.D., Ph.D., President and Chief Executive Officer of ADT, added, “We are thrilled to partner with Anchiano, a company with a well-respected management team with a track record of success in development of targeted cancer therapies in their prior experience. We have confidence that this team has the knowledge, capability, and commitment to fully develop these two programs, and a shared vision of bringing new therapies to patients in need.”</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">Under the terms of the collaboration and license agreement, Anchiano will be granted an exclusive option to license the RAS and PDE10/β-catenin inhibitors in exchange for a \$3 million upfront payment to ADT and will fund certain research activities. At any time through obtaining an Investigational New Drug (IND) designation, Anchiano will have the option to exclusively license the compounds worldwide and will be responsible

for all aspects of pre-clinical and clinical development and global commercialization. If Anchiano exercises its option, it will be responsible for development and commercialization and will incur additional payment obligations, including milestone and royalty payments to ADT.

LifeSci Advisors, LLC acted as exclusive transaction advisor to Anchiano.

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Dr. Haluska added, "In addition to the news of our option to license agreement, we are also providing an enrollment update on our pivotal Phase 2 Codex trial of inodiftagene vixteplasmid in BCG-unresponsive non-muscle invasive bladder cancer patients. We currently have 31 patients enrolled or in active screening, of which 23 have been dosed. We had previously estimated that enrollment for the 35-patient interim analysis would be complete by the end of September, allowing for the 12-week readout to take place in the fourth quarter of this year. While at this time we are close to that projection, our conservative estimate is that data collection and interim analysis will be completed by the first quarter of 2020."

About the Pan-RAS Program¹²

Oncogenic mutations in the RAS family of genes (KRAS, HRAS, and NRAS) are present in approximately 30% of cancer. RAS plays a pivotal role in signal transduction pathways leading to tumor cell proliferation and survival. ADT's program has identified novel small molecules that exhibit potent and selective inhibition of activated RAS signaling regardless of isoform or mutation, or pan-RAS inhibition.

Historically, direct inhibition of RAS has been challenging. However, investigational compounds that selectively target the KRAS G12C mutation

recently have shown antitumor activity in the clinic, clinically validating RAS as a therapeutic target. These current investigational drugs are mutation specific—with G12C representing approximately 11% of KRAS mutations in cancer. A broadly active pan-RAS inhibitor with the potential to treat RAS-driven cancers regardless of RAS isoform or mutation would be clinically useful.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; background-color: white">The RAS inhibitor program is comprised of a novel series of indene derivatives that potently, selectively and reversibly inhibit growth of tumor cells harboring mutant RAS, while having greater than 100-fold selectivity over cells with normal RAS activity. Inhibitory activity has been observed with low nanomolar potency in KRAS-, HRAS-, and NRAS-driven tumor cell models with a variety of mutations across a variety of tumor types. These compounds inhibit downstream signaling through RAF and PI3K pathways, initiate cell-cycle arrest and induce apoptosis, demonstrated blockade of GTP loading of RAS in the nucleotide-free state in cell-free biochemical assays, and have exhibited in vivo activity in RAS mutant tumor models. They have potential for RAS inhibition in a broad variety of clinical settings. </P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; background-color: white">About the PDE10/β-catenin Program⁴⁵⁶⁷⁸</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; background-color: white">Genetic alterations in components that make up the Wnt signaling pathway, which includes APC (adenomatous polyposis coli) and β-catenin, are prevalent in a number of cancer types, occurring in upwards of 80% of colorectal cancers. Additionally, germline mutations of APC lead to the hereditary cancer syndrome Familial Adenomatous Polyposis (FAP). Wnt signaling controls the level of intracellular activated β-catenin, a key effector of oncogenic signal transduction, and oncogenic alterations in Wnt, APC, or β-catenin all result in elevated and uncontrolled levels of β-catenin. Recent studies have shown that the cyclic nucleotide phosphodiesterase 10A (PDE10) is overexpressed during early stages of tumorigenesis and is essential for tumor cell growth. PDE10 inhibition activates protein kinase G and leads to the degradation of the oncogenic pool of β-catenin to suppress critical proteins essential for tumor cell proliferation and survival. Thus, targeting PDE10 provides a novel approach to selectively suppress β-catenin-mediated transcriptional activity.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">ADT’s program has identified small molecules that selectively and potently inhibit PDE10 and suppress Wnt/β-catenin signaling in preclinical models. PDE10 inhibition has been shown to downregulate β-catenin expression, and inhibits polyp and tumor growth. It has potential for application

in the treatment of cancer as well as spontaneous and familial polyposis syndromes.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">Conference Call and Webcast Information</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify">Company management will discuss the licensing agreement and corporate update on a call scheduled for Tuesday, September 24, 2019 at 8:00 am Eastern Time. To participate in the call, dial 1-877-451-6152 (domestic) or 1-201-389-0879 (international) fifteen minutes before the conference call begins and reference the conference passcode 13694843. The live conference call and replay can be accessed via audio webcast at <U><http://public.viavid.com/index.php?id=136227></U>.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">About Anchiano</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; background-color: white">Anchiano is a pivotal-stage biopharmaceutical company dedicated to the discovery, development, and commercialization of novel targeted therapies to treat cancer in areas of significant clinical need, with offices in Cambridge, MA, and Jerusalem, Israel. Anchiano's most advanced product candidate, inodiftagene vixteplasmid, is in development as a treatment for non-muscle-invasive bladder cancer. For more information on Anchiano, please visit <U>www.anchiano.com</U>.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; background-color: white">About ADT</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0; text-align: justify; background-color: white">ADT Pharmaceuticals is a private company focused on discovering, developing and securing patent protection for novel molecules that inhibit constitutively activated RAS- or Wnt-mediated signaling pathways that drive the growth of many human cancers. ADT's technology currently comprises a broad, novel proprietary small-molecule class, encompassing at least two distinct mechanistic sub-classes that share a common chemical core; one subclass targets RAS and the other subclass inhibits PDE10 to activate cGMP/PKG signaling and induce degradation of the oncogenic pool of β -catenin. For more information on ADT, please visit

www.ADT-Pharma.com.

Forward-Looking Statements

This press release contains "forward-looking statements" that are subject to risks and uncertainties. Words such as "believes," "intends," "expects," "projects," "anticipates" and "future" or similar expressions are intended to identify forward-looking statements. These forward-looking statements are subject to the inherent uncertainties in predicting future results and conditions, many of which are beyond the control of Anchiano, including, without limitation, the risk factors and other matters set forth in its filings with the Securities and Exchange Commission, including its Annual Report on Form 20-F for the year ended December 31, 2018. Anchiano undertakes no obligation to publicly update or revise any forward-looking statements, whether as a result of new information, future events or otherwise, except as may be required by law.

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">References</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">¹ NCI RAS initiative website <U><https://www.cancer.gov/research/key-initiatives/ras></U>.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">² Fakhri M, et al. Phase 1 study evaluating the safety, tolerability, pharmacokinetics (PK), and efficacy of AMG 510, a novel small molecule KRASG12C inhibitor, in advanced solid tumors. <I>J Clin Oncol</I> 37, 2019 (suppl; abstr 3003).</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">³ O'Connell B, et al. Pharmacological Targeting of RAS: Recent Success with Direct Inhibitors. <I>Pharmacol Res</I>, 2019, 139:503-511.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">⁴ Kinzler KW, Vogelstein B. Lessons from hereditary colorectal cancer. <I>Cell</I>, 1996, 87(2): 159–170.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">⁵ Li N, et al. Phosphodiesterase 10A: a novel target for selective inhibition of colon tumor cell growth and β-catenin-dependent TCF transcriptional activity. <I>Oncogene</I> 2015 Mar 19;34(12):1499-509.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">⁶ Li N, et al. Suppression of β-catenin/TCF transcriptional activity and colon tumor cell growth by dual inhibition of PDE5 and 10.

<I>Oncotarget.</I> 2015 Sep 29;6(29):27403-15.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">⁷ Lee K, et al. β-catenin nuclear translocation in colorectal cancer cells is suppressed by PDE10A inhibition, cGMP elevation, and activation of PKG. <I>Oncotarget. </I>2016 Feb 2;7(5):5353-65.</P>

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<P STYLE="font: 10pt Times New Roman, Times, Serif; margin: 0pt 0">⁸ Zhu B, et al. Phosphodiesterase 10A is overexpressed in lung tumor cells and inhibitors selectively suppress growth by blocking β-catenin and MAPK signaling. <I>Oncotarget.</I> 2017 Aug 27;8(41):69264-69280.</P>

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