
**UNITED STATES
SECURITIES AND EXCHANGE COMMISSION**

Washington, D.C. 20549

FORM 8-K

**Current Report
Pursuant to Section 13 or 15(d) of
the Securities Exchange Act of 1934**

Date of Report (Date of earliest event reported): **March 2, 2023**

ADAPT IMMUNE THERAPEUTICS PLC

(Exact name of registrant as specified in its charter)

England and Wales
(State or other jurisdiction of
incorporation)

1-37368
(Commission File Number)

Not Applicable
(IRS Employer Identification No.)

**60 Jubilee Avenue, Milton Park
Abingdon, Oxfordshire OX14 4RX
United Kingdom**
(Address of principal executive offices, including zip code)

(44) 1235 430000
(Registrant's telephone number, including area code)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol	Name of each exchange on which registered
American Depositary Shares, each representing 6 Ordinary Shares, par value £0.001 per share	ADAP	The Nasdaq Global Select Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act.

Item 1.02 Termination of a Material Definitive Agreement

On March 2, 2023, Adaptimmune Limited and Universal Cells, Inc (“Universal Cells”) mutually agreed to the termination of the Collaboration and License Agreement between Adaptimmune Limited and Universal Cells dated January 13, 2020 (the “Collaboration Agreement”), pursuant to which Universal Cells and Adaptimmune Limited had agreed to collaborate to research, develop, and commercialize certain cellular therapy products directed to certain targets. Termination of the Collaboration Agreement became effective March 6, 2023 (the “Effective Date”). The parties previously terminated an Amended and Restated Research Collaboration and License Agreement, dated January 13, 2020, effective February 26, 2023.

In connection with the termination, all licenses and sublicenses granted to either party pursuant to the Collaboration Agreement ceased, and each party is required to return all confidential information of the other party within 30 days of the Effective Date. Each party also agreed to destroy all cell lines and other materials of the other party in its possession within 30 days of the Effective Date. There were no termination penalties in connection with the termination.

Item 2.02 Results of Operations and Financial Conditions.

On March 6, 2023, Adaptimmune Therapeutics plc (the “Company”) announced its financial results for the fourth quarter and year ended December 31, 2022 and provided a corporate update. A copy of the press release is being furnished as Exhibit 99.1 hereto and is incorporated by reference herein.

The information in Item 2.02 of this Current Report on Form 8-K, including the Exhibit 99.1 attached hereto, shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, (the "Exchange Act"), or incorporated by reference in any filing made by the Company under the Securities Act of 1933, as amended, or the Exchange Act, except as expressly set forth by the Company by specific reference in such a filing.

Item 5.02 Departure of Directors or Certain Officers; Election of Directors; Appointment of Certain Officers; Compensatory Arrangements of Certain Officers.

On March 3, 2023, Adaptimmune, LLC (“Adaptimmune”), a wholly-owned subsidiary of Adaptimmune Therapeutics plc (the “Company”), entered into a separation and consulting agreement (the “Agreement”) with Cintia Piccina who served as the Company’s Chief Commercial Officer since January 31, 2022. Ms. Piccina also received notice pursuant to the Worker Adjustment and Retraining Notification Act of 1988 (WARN Act). The Agreement is effective as of March 5, 2023 and provides that Ms. Piccina’s employment with Adaptimmune ends on that date and she will provide consulting services to the Company for a period from March 6, 2023 until September 6, 2023 unless the consulting arrangement is terminated earlier or extended (the “Consulting Period”).

The Agreement provides that Adaptimmune will pay Ms. Piccina a severance payment equal to nine months base pay, in the amount of \$333,750, less all applicable deductions and withholdings and a payment equal to the gross value of nine months of health care coverage under the Consolidated Omnibus Budget Reconciliation Act of 1985 (COBRA) totaling \$5,730.84. These payments will be made in lump-sum form in the next available month-end pay date following the effective date of the Agreement. Ms. Piccina acknowledged and agreed that the payments are in full satisfaction of Adaptimmune’s obligations under its Executive Severance Policy dated March 10, 2017.

An option covering 3,376,992 ordinary shares (the “Market Value Options”) granted to Ms. Piccina on January 31, 2022 pursuant to the Adaptimmune Therapeutics plc 2016 Employee Share Option Scheme and related plan documents (collectively, the “Plan”) will continue to vest during the Consulting Period, subject to the relevant Plan rules and in accordance with the respective vesting schedule. The Agreement provides that Ms. Piccina will not receive any additional compensation for the consulting services. Provided that the Agreement is not terminated by the Company for cause, Ms. Piccina will be permitted a period of 12 months from the date that she ceases to be Connected (as defined in the Plan) to exercise the Market Value Options that shall have vested by the date that she ceases to be Connected.

The foregoing summary of the Agreement is qualified in its entirety by reference to the complete text of the Agreement, a copy of which is filed as Exhibit 10.1 to this Current Report on Form 8-K and is incorporated herein by reference.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits.

<u>Exhibit No.</u>	<u>Description of Exhibit</u>
10.1	<u>Separation and Consulting Agreement dated as of March 3, 2023 by and between Adaptimmune, LLC and Cintia Piccina and effective as of March 5, 2023.</u>
99.1	<u>Press release dated March 6, 2023.</u>
104	Cover Page Interactive Date File (embedded within the Inline XBRL document)

SIGNATURES

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

ADAPTIMMUNE THERAPEUTICS PLC

Date: March 6, 2023

By: /s/ Margaret Henry

Name: Margaret Henry

Title: Corporate Secretary

SEPARATION AND CONSULTING AGREEMENT

THIS SEPARATION AND CONSULTING AGREEMENT (together with Exhibit A, the “ **Agreement**”) is made and entered into effective as of March 5, 2023 (“**Effective Date**”) by and between Adaptimmune, LLC (“**Adaptimmune**”) and Cintia Piccina (“**Piccina**”).

WHEREAS, Piccina and Adaptimmune previously entered into an Employment Agreement, dated as of January 26, 2022 (the “**Employment Agreement**”).

WHEREAS, Piccina’s employment with Adaptimmune shall end, and Adaptimmune and Piccina wish to conclude their employment relationship on mutually satisfactory terms and to settle fully and finally all matters and potential disputes that Piccina may have with Adaptimmune and certain others.

WHEREAS, Piccina and Adaptimmune have agreed that following the Effective Date, Piccina shall continue to provide services to Adaptimmune under the terms of this Agreement in order to facilitate a smooth transition for both parties.

NOW, THEREFORE, in consideration of the mutual promises contained herein and intending to be legally bound, Adaptimmune and Piccina hereby agree as follows:

1. **Separation from Employment.** Piccina’s employment with Adaptimmune will end permanently and irrevocably effective March 5, 2023 (“**Separation Date**”).

2. **Payment.** If Piccina timely executes and is and remains in compliance with this Agreement, Adaptimmune shall pay Piccina the following amounts, collectively defined as the “**Payment**”:

- i. A payment equal to gross value of nine (9) months’ base pay, in the amount of \$333,750, less all applicable deductions and withholdings; and
- ii. A payment equal to the gross value to cover nine (9) months of health care coverage through COBRA totaling a net of \$5,730.84.

The Payment will be made in lump-sum form in the next available Company month-end pay date following the Effective Date of this Agreement. Piccina acknowledges and agrees that the Payment is in full satisfaction of Adaptimmune’s obligations to Piccina under the Executive Severance Policy dated March 10, 2017 (the “**Severance Policy**”). Piccina will be entitled to any accrued and unused vacation time, as communicated separately, regardless of whether she elects to sign this Agreement.

Regardless of whether Piccina enters into this Agreement, Piccina will remain eligible to receive group health benefits (medical and dental) through the end of the month of the Separation Date in accordance with plan terms and enrollment status, and will receive a notice of rights under the Consolidated Omnibus Budget Reconciliation Act (“**COBRA**”), pursuant to which Piccina can, if eligible, elect to extend health benefits on a self-pay basis. If Piccina enters into this Agreement, Piccina will receive the taxable lump sum cash payment identified in Section 2(ii) above intended

for use to extend health benefits under COBRA. However, this amount is not legally restricted as to use and it will be paid regardless of whether Piccina actually enrolls for COBRA coverage. *To receive COBRA coverage, Piccina must fulfill all enrollment requirements and pay all applicable premiums in a timely manner. Adaptimmune will not enroll Piccina for COBRA coverage or pay any COBRA premiums on Piccina's behalf.*

3. **No Further Employment-Related Payments, Benefits or Rights.** Piccina acknowledges entering into the Employment Agreement in exchange for adequate consideration, and Piccina hereby reaffirms Piccina's commitments and obligations under the Employment Agreement that remain in effect during and after the Separation Date and Consulting Period, including but not limited to Piccina's obligations under Sections 5 through 24 of the Employment Agreement (the "**Continuing Obligations**"). Piccina further acknowledges that Piccina has a copy of the Employment Agreement, that Piccina has read the Employment Agreement again before signing this Agreement, and that the consideration Piccina received in exchange for signing the Employment Agreement was adequate and reasonable. Piccina further acknowledges that, other than the Payment described in Section 2 above, Piccina has received payment in full of all of the compensation, benefits and/or payments of any kind due to Piccina from Adaptimmune and any other Released Parties (as defined below) related to Piccina's employment and under the Employment Agreement and the Severance Policy, including all compensation (including both straight time and overtime), bonuses, expense reimbursements, payments to or from benefit plans, unused accrued vacation time, personal time, severance, sick pay or any other payment under a plan, program, practice or promise of Adaptimmune or that of any other Released Parties (as defined below). Piccina further acknowledges that Piccina is not, and shall not be, entitled to receive from Adaptimmune or any other Released Parties any payments, benefits or perquisites (whether monetary and non-monetary) other than those expressly described in this Agreement.

4. **Consulting Period.**

a. **Duration of Consulting Period.** Piccina will provide services as described in this Section 4 beginning on March 6, 2023, and continuing until September 6, 2023, unless the consulting arrangement is earlier terminated or extended as provided in this Section 4.a (such period, the "**Consulting Period**").

i. **Termination for Cause.** The Consulting Period may be terminated by Adaptimmune for Cause upon written notice to Piccina. "**Cause**" shall mean with respect to Piccina one or more of the following: (i) acts or omissions constituting gross negligence, recklessness or willful misconduct on the part of Piccina with respect to Piccina's obligations or otherwise relating to the business of Adaptimmune; (ii) Piccina's material breach of Adaptimmune rules, policies and/or procedures; (iii) Piccina's material insubordination or material non-performance or willful neglect of assigned duties; (iv) acts or omissions which bring the reputation of Adaptimmune into material disrepute; (v) except as specifically approved in advance in writing by the Chief Executive Officer or General Counsel of Adaptimmune, any act or omission by Piccina aiding or abetting a competitor, supplier or customer of Adaptimmune and/or any of its subsidiaries or affiliates to the material disadvantage or detriment of Adaptimmune and/or any of its subsidiaries or affiliates; (vi) Piccina's commission of fraud, misappropriation, embezzlement or theft; or (vii) Piccina's material breach of this Agreement, including, but not limited to, violation of any of the Continuing Obligations.

ii. **Termination for Convenience.** Either party may terminate the Consulting Period upon two weeks' written notice to the other party.

iii. Extension by Mutual Agreement. The parties may extend the Consulting Period for up to an additional three (3) months (through and including December 6, 2023) by mutual agreement.

b. **Services.** During the Consulting Period, Piccina will provide her expertise and knowledge to Adaptimmune in an advisory role from time to time and at such locations as mutually agreed upon by Piccina and Adaptimmune from time to time (the “**Services**”). The Services will include, but not be limited to: (i) advising on commercial planning and strategy; (ii) specific projects related to patient treatment/site operational execution, and (iii) internal and external positioning/communication/culture. Piccina will have primary control over the means and manner of performing the Services under this Agreement but will perform the Services in a quality and efficient manner in accordance with the reasonable requirements of Adaptimmune. Adaptimmune understands that Piccina may provide services to other entities during the Consulting Period provided Piccina complies with this Agreement and the Continuing Obligations. Unless otherwise agreed by the parties, Piccina will provide all materials, equipment and supplies necessary to perform the Services.

c. **Continued Vesting of Certain Equity; No Additional Compensation.** Piccina was granted certain stock options under Award No: 4182 dated January 31, 2022 (the “**Award**”) pursuant to the Adaptimmune Therapeutics PLC 2016 Employee Share Option Scheme and related plan documents (collectively, the “**Plan**”). Piccina agrees and acknowledges that she will remain Connected (as defined in the Plan) and her share options granted under the Award will continue to vest during the Consulting Period, subject to the relevant Plan rules and in accordance with the respective vesting schedule. Piccina understands and agrees she will not receive any additional compensation for the Services provided under this Section 4.

d. **Expenses.** During the Consulting Period, Adaptimmune will reimburse Piccina for all reasonable expenses reasonably incurred by her in connection with the performance of the Services, provided such expenses are approved in advance in writing by Adaptimmune. Piccina shall provide Adaptimmune with such receipts or other evidence of actual payment of the expenses as shall be requested by Adaptimmune.

e. **Independent Contractor Status.** Piccina confirms that during the Consulting Period she will be an independent contractor and that her performance of the Services under this Agreement will not entitle her to any rights to any pension, insurance, car or other fringe benefits from Adaptimmune. Piccina represents, confirms and warrants that she is free to perform the Services and there are no third party obligations or restrictions (including any restrictions imposed by a third party employer) including obligations or restrictions concerning providing services to others, confidentiality of proprietary information and assignment of inventions, ideas, patents or copyrights, and Piccina agrees that she will not do anything in the performance of Services hereunder that would violate any such duty. Piccina will notify Adaptimmune immediately if she becomes aware of any circumstances which might lead to a conflict of interest or breach of any professional conduct rules.

f. **Obligations During and After the Consulting Period.**

i. Piccina agrees that the Continuing Obligations (including but not limited to Piccina's obligations regarding Confidential Information and Work Product and Intellectual Property, Inventions and Patents as set forth in the Employment Agreement) will apply during the Consulting Period to Piccina's performance of the Services.

ii. All materials, equipment, documents, data compilations (in whatever form), software programs, electronic materials and other Confidential Information ("**Adaptimmune Property**") that Piccina receives or makes during the Consulting Period are and shall remain the property of Adaptimmune, and Piccina shall immediately return Adaptimmune Property (including any copies thereof) to Adaptimmune upon Adaptimmune's request and/or upon the termination or expiration of the Consulting Period. Piccina shall not remove from Adaptimmune's offices any Adaptimmune Property except as authorized in writing by Adaptimmune. The obligation to return such property extends to anything received or made as a result of performing the Services for Adaptimmune.

iii. All forms of intellectual property including, without limitation, patents, rights in know-how and confidential information, copyrights, designs, trademarks and any applications for the same ("**Intellectual Property**") shall be owned by Adaptimmune where first created or reduced to practice in the performance of the Services. Piccina hereby assigns and agrees to assign all of her right and title to such Intellectual Property to Adaptimmune. Where necessary, Piccina agrees to execute and provide such further documentation as may be reasonably required by Adaptimmune to vest title to Intellectual Property in Adaptimmune in accordance with this clause. This clause shall survive any termination or expiration of the Consulting Period.

g. **Reaffirmation of Agreement Required at End of Consulting Period; Additional Consideration.** Within forty-five (45) days after the end of the Consulting Period, Piccina will sign the Reaffirmation of Agreement attached as Exhibit A to this Agreement (the "**Reaffirmation**") for additional consideration as follows:

i. Provided the Consulting Period is not terminated by Adaptimmune for Cause, and if Piccina signs and does not revoke the Reaffirmation within the revocation period described therein, Piccina shall be eligible for the following benefit as additional consideration: she will be permitted a period of 12 months from the date that she ceases to be Connected (as defined in the Plan) to exercise any share options under the Award that shall have vested by the date that she ceases to be Connected, notwithstanding the language of the Plan, and such benefit shall be the "**Additional Consideration**" as that term is used in this Agreement, including the Reaffirmation. All other relevant Plan rules shall apply to Piccina's exercise of any share options under the Award.

ii. In the event of a termination of the Consulting Period by Adaptimmune for Cause: (x) the period for Piccina to exercise any share options under the Award shall expire on the date of termination of the Consulting Period, subject always to the relevant Plan rules; and (y) Adaptimmune shall pay Piccina the lump sum of [\$5,000] within thirty (30) days after Piccina signs the Reaffirmation, provided that she does not revoke her acceptance of the Reaffirmation within the revocation period described therein, and such payment shall be the "**Additional Consideration**" as that term is used in this Agreement, including the Reaffirmation.

5. **General Release.** In consideration of the promises contained herein and intending to be legally bound, Piccina, for Piccina, Piccina's heirs, executors, administrators, successors,

assigns, and legal and personal representatives, hereby unconditionally and irrevocably remises, releases, and forever discharges Adaptimmune and each and every one of its subsidiaries and related or affiliated entities (together, the “**Entities**”) and each of the Entities’ current and former directors, members, officers, shareholders, employees, agents, and attorneys (collectively, the “**Released Parties**”) of and from any and all claims, causes of action, liabilities, obligations, controversies, damages, lawsuits, debts, demands, costs, charges and/or expenses (including attorneys’ fees and costs) of any nature whatsoever, asserted or unasserted, known or unknown, suspected or unsuspected, that Piccina ever had, now has or hereafter may have against Adaptimmune or any of the other Released Parties that arose at any time regarding any matter up to and including the date of this Agreement. Without in any way limiting the generality of the foregoing, Piccina specifically acknowledges and agrees that the claims released herein include, to the fullest extent permitted by law, (a) all claims arising under any federal, state or local statute, ordinance, or regulation, including but not limited to the Americans with Disabilities Act, Title VII of the Civil Rights Act of 1964, the Age Discrimination in Employment Act, the Older Workers Benefit Protection Act, the Worker Adjustment and Retraining Notification Act (WARN), in each case as amended, (b) all claims arising under any common law principle, including claims for breach of any implied or express contract or quasi-contract, wrongful discharge, constructive discharge, defamation, unjust enrichment, or negligent or intentional infliction of emotional distress, (c) all claims arising out of or relating to Piccina’s employment with Adaptimmune or any of the other Released Parties or the termination of that employment, including any claims under Piccina’s employment, and (d) all claims for any attorneys’ fees and costs. Notwithstanding the foregoing, Piccina does not release the Released Parties from any claims that Piccina may have (w) under this Agreement, (x) for unemployment insurance benefits, (y) arising out of facts occurring after the date of Piccina’s execution of this Agreement, or (z) that as a matter of federal and/or state law may not be waived, and this release is subject to Section 13 below. Further, Piccina is not waiving any claim to benefits under retirement benefits or savings and investment plans Adaptimmune may have, subject to their terms, or to file a claim for benefits under Section 502(a)(1)(B) of ERISA, to the extent applicable, although Piccina does waive any rights to claim penalties, any claim under Section 510 or 511 of ERISA, or relief for any alleged breach of fiduciary duties under any ERISA-governed plans.

6. **Covenant Not To Sue.** Piccina agrees that neither Piccina nor any person or entity on Piccina’s behalf shall commence, maintain or prosecute any lawsuit or court complaint against Adaptimmune or any of the other Released Parties with respect to any act, omission or other matter that is released by the provisions of the preceding Section. This Section shall not operate to waive any rights that may not legally be waived, nor shall it preclude Piccina from bringing an action under this Agreement. Piccina affirms that, as of this date, Piccina has not taken or initiated any action encompassed by this Section.

7. **Mutual Non-Disparagement.** Subject to Section 13 below, Piccina shall not communicate or publish, directly or indirectly, any disparaging comments or information about Adaptimmune or any of the other Released Parties or make any comments that would in any way place any of these entities and individuals in a negative light. Adaptimmune shall not communicate or publish, directly or indirectly, any disparaging comments or information about Piccina or make any comments that would in any way place Piccina in a negative light. Nothing in this Section, however, prohibits either Piccina or Adaptimmune from making any communication that Piccina or Adaptimmune, respectively, is required or entitled to make by nonwaivable law.

8. **Employment Reference.** Any professional reference concerning Piccina’s

employment with Adaptimmune shall be limited only to disclosure of Piccina's job title and dates of employment subject to all such inquiries being made to Adaptimmune's Global Head of Human Resources.

9. **Knowing and Voluntary Agreement.** Piccina acknowledges that Piccina has carefully read and reviewed this Agreement and fully understands that Piccina enters into it knowingly and voluntarily. Piccina acknowledges that in compliance with the Older Workers Benefit Protection Act (OWBPA), Adaptimmune has informed Piccina of the group of individuals who were considered and who were selected for separation as part of the same action resulting in Piccina's separation from employment by providing Piccina a disclosure document showing the job titles and ages of all such employees (the "**Disclosure**"). Piccina understands and acknowledges that the release provided in this Agreement is in exchange for consideration that is in addition to anything to which Piccina is already entitled and that, by this Section, Adaptimmune has advised Piccina to consult with an attorney of Piccina's choosing prior to executing this Agreement and Piccina hereby warrants and represents that Piccina has either consulted with Piccina's counsel or knowingly opted not to seek such consultation. Piccina acknowledges that neither Adaptimmune nor any of its employees, representatives or attorneys have made any representations or promises concerning the terms or effects of this Agreement other than those contained herein.

10. **Enforcement.** Piccina acknowledges that any compensation (including the Payment) conditioned on timely execution, nonrevocation and noncancellation, and adherence to the terms, of this Agreement shall be subject, to the extent permitted by law, to return or reimbursement (if already paid) to Adaptimmune or cancelled and forever discharged by Adaptimmune (if not yet paid), with the remaining terms of the Agreement remaining in full force and effect.

11. **Governing Law.** This Agreement shall in all respects be interpreted, enforced, and governed under the laws of the Commonwealth of Pennsylvania, without reference to the principles of conflicts of law otherwise applicable therein. The language of all parts of this Agreement shall in all cases be construed as a whole, according to its fair meaning, and not strictly for or against either party.

12. **Good Faith Settlement and Non-Admission of Liability.** Piccina agrees that the Payment made pursuant to this Agreement is a good faith settlement of claims and is not to be construed as an admission of legal liability by Adaptimmune and the other Released Parties and that no person or entity shall utilize this Agreement, or the consideration received pursuant to this Agreement, as evidence of any admission of liability. Piccina agrees not to assert that this Agreement is an admission of guilt or wrongdoing and acknowledges that Adaptimmune and the other Released Parties does not believe or admit that it has done anything wrong or engaged in any conduct for which it is liable to Piccina.

13. **Non-Interference.** For clarity, Adaptimmune confirms that nothing in this Agreement – including in the Confidentiality, Non-Disparagement, General Release, and Covenant Not to Sue provisions – is intended to prohibit Piccina from filing a charge with any agency which enforces anti-discrimination or other employment laws, or from cooperating with or providing truthful information to any governmental agency. However, Piccina understands that by signing this Agreement and not revoking it, Piccina is waiving the right to recover any money from Adaptimmune or any other Released Parties, other than the Payment. Further, nothing in this Agreement shall prevent either party from disclosing facts related to claims of discrimination,

retaliation or harassment. Nothing in this Agreement requires confidentiality of discrimination, retaliation or harassment allegations.

14. **Section 409A Compliance.** All payments or benefits under this Agreement are subject to any applicable employment or tax withholdings or deductions. In addition, the parties hereby agree that it is their intention that all payments or benefits provided under this Agreement comply with an exemption from Section 409A of the Internal Revenue Code of 1986, as amended (the “Code”) as a short-term deferral and this Agreement shall be interpreted accordingly. Piccina is hereby advised to seek independent advice from her tax advisor(s) with respect to the application of the exemption from Section 409A of the Code to any payments or benefits under this Agreement. Notwithstanding the foregoing, Adaptimmune does not guarantee the tax treatment of any payments or benefits under this Agreement, including without limitation under the Code, federal, state, or local laws.

15. **Entire Agreement.** This Agreement, together with the Disclosure, sets forth the entire agreement between the parties with respect to the subject matter hereof and fully supersedes any and all written or oral contracts, agreements or understandings between the parties pertaining to the subject matter hereof; provided, however, that, notwithstanding the foregoing, Piccina re-affirms and shall remain bound by the post-separation obligations set forth in any document between Piccina and Adaptimmune, including but not limited to the Continuing Obligations under the Employment Agreement. Piccina agrees to notify Adaptimmune in writing prior to accepting any offer of employment that may conflict in any way with any such post-separation obligations.

BY SIGNING THIS AGREEMENT, EMPLOYEE ACKNOWLEDGES DOING SO VOLUNTARILY AFTER CAREFULLY READING AND FULLY UNDERSTANDING EACH PROVISION AND ALL OF THE EFFECTS OF THIS AGREEMENT, WHICH INCLUDES A RELEASE OF KNOWN AND UNKNOWN CLAIMS AND A RESTRICTION ON FUTURE LEGAL ACTION AGAINST ADAPT IMMUNE AND OTHER RELEASED PARTIES.

IN WITNESS WHEREOF, and intending to be legally bound hereby, the parties execute this Separation and Consulting Agreement.

Employee:

For Adaptimmune, LLC:

By: /s/ Cintia Piccina
Printed Name: Cintia Piccina
Date: March 3, 2023

By: /s/ Adrian Rawcliffe
Printed Name: Adrian Rawcliffe
Title: CEO
Date: March 3, 2023

EXHIBIT A
REAFFIRMATION OF SEPARATION AND CONSULTING AGREEMENT

I, Cintia Piccina, through this Reaffirmation of Separation and Consulting Agreement (this “**Reaffirmation**”) hereby reaffirm and commit to the terms and conditions of the Separation and Consulting Agreement effective March 5, 2023 (the “**Separation Agreement**”), that I entered into with Adaptimmune, LLC (“**Adaptimmune**”).

In particular, I reaffirm the General Release and Covenant Not to Sue provisions set forth in Sections 5 and 6 of the Separation Agreement, as well as Sections 7 through 15 of the Separation Agreement. I understand and acknowledge that the General Release is intended to be as broad as legally permissible and applies to both employment-related and non-employment-related claims against any and all Released Parties (as defined in the Separation Agreement) up to the time that I execute this Reaffirmation.

Notwithstanding the foregoing, I understand that the following are not included in this reaffirmation of the General Release: (i) any rights or claims that arise after the date I sign this Reaffirmation; (ii) any rights or claims that are not waivable as a matter of law; or (iii) any claims arising from the breach of the Separation Agreement or this Reaffirmation. In addition, I understand that nothing in this Reaffirmation prevents me from filing, cooperating with, or participating in any investigation or proceeding before the Equal Employment Opportunity Commission, or any other government agency.

I further acknowledge and confirm that I have been given a period of at least forty-five (45) calendar days within which to consider the Agreement and whether to execute this Reaffirmation, and the parties agree that any changes to this Agreement, whether material or immaterial, have not re-started the running of this period. For my signature on this Reaffirmation to be effective, it must be dated *no earlier than the last day of the Consulting Period and no more than forty-five (45) days following the last day of the Consulting Period*.

I may revoke or cancel my acceptance of this Reaffirmation (but not the Separation Agreement) within seven (7) calendar days after execution of it by notifying Adaptimmune of my desire to do so in writing delivered to Adaptimmune at: (i) 351 Rouse Boulevard, Philadelphia, PA 19112, or (ii) legal@adaptimmune.com. To be effective, Adaptimmune must receive such notice of revocation or cancellation before the close of business on the seventh (7th) calendar day following my execution of this Reaffirmation. I understand and agree that I will not be entitled to the Additional Consideration (as defined in the Separation Agreement) if I revoke this Reaffirmation in the time and manner described above.

I understand that my receipt of the Additional Consideration described in Section 4 of the Separation Agreement is conditioned upon my returning this Reaffirmation to Adaptimmune within forty-five (45) days after the end of the Consulting Period. By signing this Reaffirmation, I hereby reaffirm and agree that I will continue to be bound by the terms of the Separation Agreement.

Dated: _____

Signed: _____



Adaptimmune Reports Fourth-Quarter and Full Year Financial Results and Business Update

BLA submission initiated for afami-cel, with aim to complete in mid-2023; afami-cel has the potential to be the first marketed engineered TCR T-cell therapy for a solid tumor

52% (13/25) response rate in ovarian, bladder and head & neck cancers in the Phase 1 SURPASS trial with next-generation MAGE-A4 product; 75% (9/12) response rate amongst patients with these tumor types who received ≤ 3 prior lines of therapy

Adaptimmune will advance its wholly owned optimized PRAME TCR to be IND-ready in 2023

Strategic combination to create a preeminent cell therapy company for solid tumors announced earlier today

Webcast to be held today, March 6, 2023, at 8:00 a.m. EST (1:00 p.m. GMT)

PHILADELPHIA, PA. and OXFORD, UK, March 6, 2023 – Adaptimmune Therapeutics pl (Nasdaq: ADAP), a leader in cell therapy to treat cancer, today reported financial results for the fourth quarter and full year ended December 31, 2022 and provided a business update.

Adrian Rawcliffe, Adaptimmune’s Chief Executive Officer: “The last twelve months have seen immense progress in autologous cell therapies for people with cancer. CAR-T therapies have established autologous T-cell therapy as viable businesses within the broader cell and gene therapy market, which has annual sales exceeding three billion dollars. Our progress with T-cell therapies in solid tumors is truly exciting, as solid tumors account for nearly 90% of all adult cancers. I think 2023 will be a breakout year for T-cell therapies to address the broader cancer market, with Adaptimmune at the forefront.”

Adaptimmune’s first potential commercial product, afami-cel, for the treatment of synovial sarcoma

Adaptimmune initiated its BLA submission to the U.S. Food and Drug Administration (FDA) in the fourth-quarter 2022 and is on track to complete the BLA in mid-2023. This BLA is supported by data from Cohort 1 of the pivotal trial SPEARHEAD-1, which met its primary endpoint for efficacy. The Company has Regenerative Medicine Advanced Therapy (RMAT) designation from the FDA for afami-cel for the treatment of synovial sarcoma.

As reported in November 2022, data presented at the Connective Tissue Oncology Society (CTOS) annual meeting indicate continued clinical responses with an acceptable safety profile in heavily pre-treated patients with late-stage synovial sarcoma after a single dose of afami-cel.

- Overall response rate was 39% in heavily pre-treated patients with synovial sarcoma, with a median duration of response of ~12 months
- Afami-cel shown to drive tumor infiltration of activated and proliferative cytotoxic ("killer") T-cells into tumors - which likely contributes to antitumor responses.
- Benefit:risk profile of afami-cel has been favorable, to date

Potential of next-gen MAGE-A4 TCR T-cell therapy (ADP-A2M4CD8) in multiple solid tumors

- The following results were reported from 43 evaluable patients the Phase 1 SURPASS trial at the beginning of the year in heavily pre-treated patients with late-stage cancers after a single dose of ADP-A2M4CD8

- 37% overall response rate across multiple solid tumors
- 52% response rate in the focus indications of ovarian, bladder and head & neck cancers
- 75% response rate in these focus indications amongst patients who received ≤ 3 prior lines of therapy
- The Company is initiating a Phase 2 trial, SURPASS-3, in 1H 2023 for people with ovarian cancer.
 - SURPASS-3 will be conducted in patients with platinum resistant ovarian cancer who have received ≤ 4 prior lines of therapy; ADP-A2M4CD8 will be investigated as monotherapy and also combined with the checkpoint inhibitor nivolumab.
 - SURPASS-3, which could become registrational, is supported by RMAT designation from the FDA and is being developed in collaboration with The GOG Foundation, Inc.
- Adaptimmune is initiating two new cohorts in the Phase 1 SURPASS trial, combining ADP-A2M4CD8 with the checkpoint inhibitor pembrolizumab in 1) the second line treatment setting for bladder cancer and 2) in the first line treatment setting for head & neck cancer.
- As announced last year, the Company has closed the SURPASS-2 trial in gastroesophageal cancers to further enrollment

Preclinical pipeline update

- Last year, the Company announced that it will gain full control of the late-stage preclinical optimized PRAME TCR as well as the NY-ESO cell therapy program; discussions with GSK to finalize termination and transfer terms remain ongoing
- The Company aims for the PRAME program to be IND-ready in 2023
- Adaptimmune will continue to focus on its MAGE-A4 franchise while determining the optimal development path for complementary PRAME and NY-ESO programs
- Partnered programs with Genentech continue with the allogeneic pipeline and the Company is also advancing its own wholly owned allogeneic programs
- Last year, the Company took the decision to change the cell line being used to develop its MAGE-A4 allogeneic cell therapy. This change was due to the presence of a chromosomal abnormality in the original cell line and will delay the timing of the first allogeneic IND submission to 2025. This original cell line is not used in any of the Company's partnered programs.

Corporate and other news

- Adaptimmune announced a strategic combination with TCR² Therapeutics Inc. earlier today (please refer to separate press release dated March 6, 2023). As a result, and following the closing of the transaction, it is anticipated that the combined company's cash runway will extend into 2026.
- Adaptimmune and Universal Cells have agreed to terminate the Collaboration and License Agreement dated January 13, 2020 under which the parties agreed to co-develop certain allogeneic cell therapies. Termination is effective as of March 6, 2023. Termination does not impact the development of our allogeneic cell lines for our internal allogeneic programs or for our collaboration with Genentech Inc.
- Completed the majority of the expenditure on two capital projects to prepare manufacturing network for the next phase of growth.
 - Additional cleanroom space in the manufacturing facility at the Navy Yard in Philadelphia, PA for future commercial launch of afami-cel
 - Construction of a dedicated allogeneic manufacturing facility in the United Kingdom (co-located with its UK research headquarters) to supply future allogeneic products.
- Completed restructuring with a reduction in headcount of approximately 25%.

- In connection with the Company's restructuring, Cintia Piccina separated from the Company as its Chief Commercial Officer effective March 5, 2023. Ms. Piccina remains engaged with Adaptimmune on a consultancy basis.

Financial Results for the fourth quarter and year ended December 31, 2022

- **Cash / liquidity position:** As of December 31, 2022, Adaptimmune had cash and cash equivalents of \$108.0 million and Total Liquidity¹ of \$204.6 million, compared to \$149.9 million and \$369.6 million, respectively, as of December 31, 2021.
- **Revenue:** Revenue for the fourth quarter and year ended December 31, 2022 was \$11.0 million and \$27.1 million, respectively, compared to \$1.4 million and \$6.1 million for the same periods in 2021. Revenue has increased primarily due to an increase in development activities under our collaboration arrangements, in particular due to development activities under the Genentech Strategic Collaboration and License Agreement, which became effective in October 2021. Revenue also increased due to a \$6 million payment receivable from GSK as a result of the termination and amendment to the GSK Collaboration and License Agreement.
- **Research and development (R&D) expenses:** R&D expenses for the fourth quarter and year ended December 31, 2022 were \$23.1 million and \$127.7 million, respectively, compared to \$29.5 million and \$111.1 million for the same periods in 2021. R&D expenses increased due to an increase in the average number of employees engaged in research and development, increases in subcontracted expenditures and a decrease in offsetting reimbursements receivable for research and development tax and expenditure credits.
- **General and administrative (G&A) expenses:** G&A expenses for the fourth quarter and year ended December 31, 2022 were \$15.2 million and \$63.4 million, respectively, compared to \$14.8 million and \$57.3 million for the same periods in 2021 due to increases in employee-related costs and other corporate costs and restructuring charges.
- **Net loss:** Net loss attributable to holders of the Company's ordinary shares for the fourth quarter and year ended December 31, 2022 was \$29.3 million and \$165.5 million, respectively (\$0.03 and \$(0.17) per ordinary share), compared to \$38.9 million and \$158.1 million, respectively (\$(0.04) and \$(0.17) per ordinary share), for the same periods in 2021.

Financial Guidance

The Company believes that its existing cash, cash equivalents and marketable securities, together with the additional payments under the Strategic Collaboration and License Agreement with Genentech and reductions in the Company's operating costs as a result of the restructuring of the Company that is expected to be completed in the first quarter of 2023, will fund the Company's current operations into early 2025, as further detailed in the Company's Quarterly Report on Form 10-K for the fourth quarter and year ended December 31, 2022, to be filed with the Securities and Exchange Commission following this earnings release.

On March 6, 2023 the Company announced entry into a merger agreement under which the Company will combine with TCR² Therapeutics Inc in an all-stock transaction. Following the closing of the transaction, we currently estimate that the cash runway of the combined company will extend into early 2026.

¹ Total liquidity is a non-GAAP financial measure, which is explained and reconciled to the most directly comparable financial measures prepared in accordance with GAAP below

Webcast Information

The Company will host a live webcast to provide additional details at 8:00 a.m. EDT (1:00 p.m. GMT) today, March 6, 2023. A live webcast of the conference call and replay can be accessed at <https://api.newsfilecorp.com/redirect/e4WKKsxwna>. Call in information is as follows: (800)-319-4610 (US or Canada) or +1 (416)-915-3239 (International and additional options available [HERE](#)). Callers should dial in 5-10 minutes prior to the scheduled start time and simply ask to join the Adaptimmune call.

About Adaptimmune

Adaptimmune is a clinical-stage biopharmaceutical company focused on the development of novel cancer immunotherapy products for people with cancer. The Company's unique SPEAR (Specific Peptide Enhanced Affinity Receptor) T-cell platform enables the engineering of T-cells to target and destroy cancer across multiple solid tumors.

Forward-Looking Statements

This release contains "forward-looking statements" within the meaning of the Private Securities Litigation Reform Act of 1995 (PSLRA). These forward-looking statements involve certain risks and uncertainties. Such risks and uncertainties could cause our actual results to differ materially from those indicated by such forward-looking statements, and include, without limitation: the success, cost and timing of our product development activities and clinical trials and our ability to successfully advance our TCR therapeutic candidates through the regulatory and commercialization processes. For a further description of the risks and uncertainties that could cause our actual results to differ materially from those expressed in these forward-looking statements, as well as risks relating to our business in general, we refer you to our Annual Report on Form 10-K filed with the Securities and Exchange Commission for the year ended December 31, 2021 filed on March 14, 2022, Current Reports on Form 8-K, and our other filings with the Securities and Exchange Commission. The forward-looking statements contained in this press release speak only as of the date the statements were made and we do not undertake any obligation to update such forward-looking statements to reflect subsequent events or circumstances.

Total Liquidity (a non-GAAP financial measure)

Total Liquidity (a non-GAAP financial measure) is the total of cash and cash equivalents and marketable securities (available-for-sale debt securities). Each of these components appears separately in the condensed consolidated balance sheet. The U.S. GAAP financial measure most directly comparable to Total Liquidity is cash and cash equivalents as reported in the condensed consolidated financial statements, which reconciles to Total Liquidity as follows (in millions):

	December 31, 2022	December 31, 2021
Cash and cash equivalents	\$ 108,033	\$ 149,948
Marketable securities - available-for-sale debt securities	96,572	219,632
Total Liquidity	\$ 204,605	\$ 369,580

The Company believes that the presentation of Total Liquidity provides useful information to investors because management reviews Total Liquidity as part of its assessment of overall solvency and liquidity, financial flexibility, capital position and leverage.

Condensed Consolidated Statement of Operations
(unaudited, in thousands, except per share data)

	Three months ended December 31,		Year ended December 31,	
	2022	2021	2022	2021
Development revenue	11,028	1,417	27,148	6,149
Revenue	\$ 11,028	\$ 1,417	\$ 27,148	\$ 6,149
Operating expenses				
Research and development	(23,052)	(29,505)	(127,726)	(111,090)
General and administrative	(15,218)	(14,776)	(63,387)	(57,305)
Total operating expenses	(38,270)	(44,281)	(191,113)	(168,395)
Operating loss	(27,242)	(42,864)	(163,965)	(162,246)
Interest income	523	179	1,542	1,095
Other (expense) income, net	(1,537)	4,036	(536)	3,852
Loss before income tax expense	(28,256)	(38,649)	(162,959)	(157,299)
Income tax expense	(994)	(210)	(2,497)	(791)
Net loss attributable to ordinary shareholders	\$ (29,250)	\$ (38,859)	\$ (165,456)	\$ (158,090)
Net loss per ordinary share				
Basic and diluted	\$ (0.03)	\$ (0.04)	\$ (0.17)	\$ (0.17)
Weighted average shares outstanding:				
Basic and diluted	984,715,238	937,328,712	967,242,403	934,833,017

Condensed Consolidated Balance Sheets
(unaudited, in thousands, except share data)

	December 31, 2022	December 31, 2021
Assets		
Current assets		
Cash and cash equivalents	\$ 108,033	\$ 149,948
Marketable securities - available-for-sale debt securities	96,572	219,632
Accounts receivable, net of allowance for doubtful accounts of \$0 and \$0	7,435	752
Other current assets and prepaid expenses	43,330	45,126
Total current assets	255,370	415,458
Restricted cash	1,569	1,718
Operating lease right-of-use assets, net of accumulated amortization of \$9,470 and \$7,253	18,019	20,875
Property, plant and equipment, net of accumulated depreciation of \$38,588 and \$36,253	53,516	30,494
Intangible assets, net of accumulated amortization of \$4,676 and \$4,051	442	1,000
Total assets	\$ 328,916	\$ 469,545
Liabilities and stockholders' equity		
Current liabilities		
Accounts payable	\$ 4,753	\$ 8,113
Operating lease liabilities, current	2,728	2,320
Accrued expenses and other current liabilities	31,215	29,909
Restructuring provision	2,285	—
Deferred revenue, current	23,520	22,199
Total current liabilities	64,501	62,541
Operating lease liabilities, non-current	20,349	23,148
Deferred revenue, non-current	160,892	177,223
Other liabilities, non-current	1,296	673
Total liabilities	247,038	263,585
Stockholders' equity		
Common stock - Ordinary shares par value £0.001, 1,282,773,750 authorized and 987,109,890 issued and outstanding (2021: 1,240,853,520 authorized and 937,547,934 issued and outstanding)	1,399	1,337
Additional paid in capital	990,656	959,611
Accumulated other comprehensive loss	(875)	(11,142)
Accumulated deficit	(909,302)	(743,846)
Total stockholders' equity	81,878	205,960
Total liabilities and stockholders' equity	\$ 328,916	\$ 469,545

Condensed Consolidated Cash Flow Statement
(unaudited, in thousands)

	Year ended December 31, 2022	Year ended December 31, 2021
Cash flows from operating activities		
Net loss	\$ (165,456)	\$ (158,090)
<i>Adjustments to reconcile net loss to net cash used in operating activities:</i>		
Depreciation	5,266	5,630
Amortization	809	937
Share-based compensation expense	18,240	20,629
Unrealized foreign exchange (gains)/losses	(2,438)	540
Amortization on available-for-sale debt securities	2,525	5,276
Other	816	1,173
<i>Changes in operating assets and liabilities:</i>		
(Increase)/decrease in receivables and other operating assets	(9,813)	(19,358)
Increase in payables and other current liabilities	4,408	4,207
Increase in deferred revenue	3,874	149,785
Net cash (used in)/provided by operating activities	(141,769)	10,729
Cash flows from investing activities		
Acquisition of property, plant and equipment	(29,496)	(8,574)
Acquisition of intangible assets	(244)	(207)
Maturity or redemption of marketable securities	166,994	224,343
Investment in marketable securities	(48,117)	(139,762)
Net cash provided by/(used in) investing activities	89,137	75,800
Cash flows from financing activities		
Proceeds from issuance of common stock from offerings, net of commissions and issuance costs	12,817	2,529
Proceeds from exercise of stock options	50	759
Net cash provided by financing activities	12,867	3,288
Effect of currency exchange rate changes on cash, cash equivalents and restricted cash	(2,299)	365
Net (decrease)/increase in cash, cash equivalents and restricted cash	(42,064)	90,182
Cash, cash equivalents and restricted cash at start of period	151,666	61,484
Cash, cash equivalents and restricted cash at end of period	\$ 109,602	\$ 151,666
Supplemental cash flow information		
Interest received	\$ 5,149	\$ 7,765
Amortization on available-for-sale debt securities	(2,525)	(5,276)
Income taxes paid	630	535

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