Persbericht
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Gereguleerde informatie

Celyad benoemt Filippo Petti tot Chief Financial Officer

Mont-Saint-Guibert, België - Celyad (Euronext Brussel en Parijs, en NASDAQ: CYAD), een klinisch biofarmaceutisch bedrijf dat zich richt op ontwikkeling van celgebaseerde CAR-T behandelingen, kondigt vandaag de benoeming aan van Filippo Petti tot Chief Financial Officer. Hij volgt hierbij Patrick Jeanmart op. Op 3 september 2018 neemt de heer Pettizijn functie op, en de heer Jeanmart blijft tot 31 december 2018 aan als raadgever om een vlotte en efficiënte overgang te verzekeren.

"De komst van Filippo in het team is essentieel voor onze ontwikkeling op de Amerikaanse kapitaal- en financiële markten. Zijn grote kennis van de sector, zijn ervaring in de oncologie en zijn connectiviteit met de Amerikaanse beleggerswereld vormen een grote troef voor Celyad nu het bedrijf voor belangrijke mijlpalen staat. In naam van het hele bedrijf en van de raad van bestuur wil ik Patrick Jeanmart bedanken voor zijn toewijding en belangrijke bijdrage aan het succes van Celyad", **verkondigde dr. Christian Homsy, CEO van Celyad**.

De heer Petti is momenteel vicevoorzitter voor healthcare investment banking bij Wells Fargo Securities. Vóór zijn overstap naar Wells Fargo Securities was hij vicevoorzitter voor healthcare investment banking bij William Blair & Company. Vooraleer hij functies in investment banking vervulde, hield de heer Petti zich zowel bij William Blair & Company als bij Wedbush Securities bezig met aandelenresearch. Hij begon zijn carrière als onderzoekswetenschapper bij OSI Pharmaceuticals, Inc. en stapte daarna binnen dat bedrijf over naar bedrijfsontwikkeling. De heer Petti heeft een diploma van Master of Business Administration van de Cornell University, van Master of Science van de St. John's University en van Bachelor of Science van de Syracuse University.

Filippo Petti verklaarde: "*De gedifferentieerde benadering van behandeling met CAR-T-cellen van Celyad kan een belangrijke impact hebben op het leven van allerlei types kankerpatiënten. Ik vind het een eer om deel uit te maken van dit bedrijf in deze opwindende fase van innovatie op het vlak van autologe en allogene behandelingen die moeten leiden tot vernieuwende behandelingsopties om patiënten te helpen. Ik kijk uit naar mijn samenwerking met het Celyad-team om onze aanwezigheid op de Amerikaanse financiële en investeringsmarkten uit te breiden".*

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About Celyad

Celyad is a clinical-stage biopharmaceutical company focused on the development of specialized CAR-T cell-based therapies. Celyad utilizes its expertise in cell engineering to target cancer. Celyad's CAR-T cell platform has the potential to treat a broad range of solid and hematologic tumors. Its lead oncology candidate, CYAD-01 (CAR-T NKG2D), is currently evaluated in a Phase I dose escalation clinical trial to assess the safety and clinical activity of multiple administrations of autologous CYAD-01 cells in seven refractory cancers including five solid tumors (colorectal, ovarian, bladder, triple-negative breast and pancreatic cancers) and two hematological tumors (acute myeloid leukemia and multiple myeloma). The safety and clinical activity of the CYAD-01 therapy concurrently administered with standard-of-care treatments or preconditioning chemotherapy is also assessed in a full clinical development program focused on acute myeloid leukemia and colorectal cancer. Celyad was founded in 2007 and is based in Mont-Saint-Guibert, Belgium, and New York, NY. Celyad's ordinary shares are listed on the Euronext Brussels and Euronext Paris exchanges, and its American Depository Shares are listed on the NASDAQ Global Market, all under the ticker symbol CYAD.

For more information, please contact:

Celyad

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Forward-looking statements

This release may contain forward-looking statements, including statements regarding the safety and efficacy of CYAD-01 and the new mAb manufacturing method used to manufacture this drug product candidate; statements concerning the ongoing and planned clinical development of CYAD-01. Forward-looking statements may involve known and unknown risks, uncertainties and other factors which might cause actual results, financial condition and liquidity, performance or achievements of Celyad, or industry results, to differ materially from those expressed or implied by such forward-looking statements. In particular it should be noted that the interim data summarized above are preliminary in nature. There is limited data concerning safety and clinical activity following treatment with the CYAD-01 drug product candidate. These results may not be repeated or observed in ongoing or future studies involving the CYAD-01 drug product candidate. These forward-looking statements are further qualified by important factors and risks, which could cause actual results to differ materially from those in the forward-looking statements, including statements about: the initiation, timing, progress and results of our preclinical studies and clinical trials, and our research and development programs; our ability to advance drug product candidates into, and successfully complete, clinical trials; our ability to successfully manufacture drug product for our clinical trials, including with our new mAb manufacturing process and with respect to manufacturing drug product with the desired number of T cells under our clinical trial protocols; our reliance on the success of our drug product candidates, including our dependence on the regulatory approval of CYAD-01 in the United States and Europe and subsequent commercial success of CYAD-01, both of which may never occur; the timing or likelihood of regulatory filings and approvals; our ability to develop sales and marketing capabilities; the commercialization of our drug product candidates, if approved; the

pricing and reimbursement of our drug product candidates, if approved; the implementation of our business model, strategic plans for our business, drug product candidates and technology; the scope of protection we are able to establish and maintain for intellectual property rights covering our drug product candidates and technology; our ability to operate our business without infringing, misappropriating or otherwise violating the intellectual property rights and proprietary technology of third parties; cost associated with enforcing or defending intellectual property infringement, misappropriation or violation; product liability; and other claims; regulatory development in the United States, the European Union, and other jurisdictions; estimates of our expenses, future revenues, capital requirements and our needs for additional financing; the potential benefits of strategic collaboration agreements and our ability to enter into strategic arrangements; our ability to maintain and establish collaborations or obtain additional grant funding; the rate and degree of market acceptance of our drug product candidates, if approved; our financial performance; developments relating to our competitors and our industry, including competing therapies and statements regarding future revenue, hiring plans, expenses, capital expenditures, capital requirements and share performance. A further list and description of these risks, uncertainties and other risks can be found in Celyad's U.S. Securities and Exchange Commission (SEC) filings and reports, including in its Annual Report on Form 20-F filed with the SEC on April 6, 2018 and subsequent filings and reports by Celyad. Given these uncertainties, the reader is advised not to place any undue reliance on such forward-looking statements. These forward-looking statements speak only as of the date of publication of this document and Celyad's actual results may differ materially from those expressed or implied by these forward-looking statements. Celyad expressly disclaims any obligation to update any such forward-looking statements in this document to reflect any change in its expectations with regard thereto or any change in events, conditions or circumstances on which any such statement is based, unless required by law or regulation.