Data from Incyte's Dermatology Portfolio Accepted for Presentation at the 2022 European Academy of Dermatology and Venereology (EADV) Congress

August 24, 2022

WILMINGTON, Del.--(BUSINESS WIRE)--Aug. 24, 2022-- Incyte (Nasdaq:INCY) today announced that multiple abstracts featuring data from its dermatology portfolio will be presented at the upcoming 2022 European Academy of Dermatology and Venereology (EADV) Congress, held September 7-10, 2022, in Milan and virtually.

"We are excited for the opportunity to present research from our growing dermatology portfolio at this year's EADV Congress, including data on ruxolitinib cream in vitiligo, on the heels of our recent U.S. Food and Drug Administration (FDA) approval in this indication," said Jim Lee, M.D., Ph.D., Group Vice President, Inflammation & Autoimmunity, Incyte. "We additionally look forward to several presentations of data on our oral JAK1 inhibitor, povorcitinib (formerly INCB054707), which we are advancing into Phase 3 development for hidradenitis suppurativa."

Key abstracts include:

Oral Presentation

Vitiligo

Efficacy and Safety of Ruxolitinib Cream for the treatment of Vitiligo: Week 52 Pooled Analysis of the TRuE-V Phase 3 Studies (Abstract #FC01.04. Session: Free Communications in Miscellaneous. Thursday, September 8, 3:00 a.m. EST)

Poster Presentations

Vitiligo

Efficacy and Safety of Ruxolitinib Cream for the Treatment of Vitiligo by Patient Demographics and Baseline Clinical Characteristics: Week 52 Pooled Subgroup Analysis from Two Randomized Phase 3 Studies (Abstract #P1383. Session: Pigmentary Disorders)

Mental Health and Psychosocial Burden Among Patients Living with Vitiligo In Europe: Findings from the Global VALIANT Study (Abstract #P1384. Session: Pigmentary Disorders)

Exploring the Natural and Treatment History of Vitiligo in Europe: Findings from the Global VALIANT Study (Abstract #P1401. Session: Pigmentary Disorders)

Treatment Burden Among Patients with Vitiligo: Findings from the Global VALIANT Study (Abstract #P1402. Session: Pigmentary Disorders)

Quality of Life and Disease Severity in Patients with Vitiligo: Findings from the Global VALIANT Study (Abstract #P1403. Session: Pigmentary Disorders)

Atopic Dermatitis

Ruxolitinib Cream Provided Progressive Improvement in Patients with Atopic Dermatitis who did not Achieve Investigator's Global Assessment Treatment Success at Week 8: Pooled Results from Two Phase 3 Studies (Abstract #P0247. Session: Atopic Dermatitis/Eczema)

Efficacy of Ruxolitinib Cream for the Treatment of Atopic Dermatitis by Anatomic Region: Pooled Analysis from Two Randomized Phase 3 Studies (Abstract #P0248. Session: Atopic Dermatitis/Eczema)

Hidradenitis Suppurativa

Efficacy and Safety of the Janus Kinase (JAK) 1 Inhibitor INCB054707 in Patients (Pts) with Hidradenitis Suppurativa (HS): Results from a Randomized, Placebo-Controlled, Phase 2 Dose-Ranging Study (Abstract #P0004. Session: Acne and Related Disorders, Hidradenitis Suppurativa)

Effects of the Janus Kinase (JAK) 1 Inhibitor INCB054707 on Patient (Pt)-Reported Quality of Life (QoL) in Hidradenitis Suppurativa (HS): Results from a Randomized, Placebo-Controlled, Phase 2 Dose-Ranging Study (Abstract #P0005. Session: Acne and Related Disorders, Hidradenitis Suppurativa)

Effects of the Janus Kinase (JAK) 1 Inhibitor INCB054707 on Patient (Pt)-Reported Skin Pain, Analgesic Use, and Itch in Hidradenitis Suppurativa (HS): Results from a Randomized, Placebo-Controlled, Phase 2 Dose-Ranging Study (Abstract #P0006. Session: Acne and Related Disorders, Hidradenitis Suppurativa)

More information regarding the congress is available on the EADV website: https://eadvcongress2022.org/. Congress content will be available on the EADV Virtual platform for a period of three months following the congress.

About Povorcitinib (INCB054707)
Povorcitinib (INCB054707) is an oral small-molecule JAK1 inhibitor currently in Phase 2 clinical trials for hidradenitis suppurativa (HS), vitiligo and prurigo nodularis. A Phase 3 study in HS is expected to begin by end of 2022.

About Opzelura™ (ruxolitinib) Cream 1.5%
Opzelura, a novel cream formulation of Incyte's selective JAK1/JAK2 inhibitor ruxolitinib approved by the U.S. Food & Drug Administration for the

topical treatment of nonsegmental vitiligo in patients 12 years of age and older, is the first and only treatment for repigmentation approved for use in the United States. Opzelura is also approved in the U.S. for the topical short-term and non-continuous chronic treatment of mild to moderate atopic dermatitis (AD) in non-immunocompromised patients 12 years of age and older whose disease is not adequately controlled with topical prescription therapies, or when those therapies are not advisable. Use of Opzelura in combination with therapeutic biologics, other JAK inhibitors, or potent immunosuppressants, such as azathioprine or cyclosporine, is not recommended.

The marketing authorization application (MAA) for ruxolitinib cream as a potential treatment for adolescents and adults (age >12 years) with nonsegmental vitiligo with facial involvement is under review at the European Medicines Agency (EMA).

Incyte has worldwide rights for the development and commercialization of ruxolitinib cream, marketed in the United States as Opzelura. In April 2022, Incyte entered into a strategic alliance agreement with Maruho Co., Ltd. for the development, manufacturing and exclusive commercialization of ruxolitinib cream for treatment of autoimmune and inflammatory dermatology indications in Japan.

Opzelura is a trademark of Incyte.

IMPORTANT SAFETY INFORMATION

OPZELURA is for use on the skin only. Do not use OPZELURA in your eyes, mouth, or vagina.

OPZELURA may cause serious side effects, including:

Serious Infections: OPZELURA contains ruxolitinib. Ruxolitinib belongs to a class of medicines called Janus kinase (JAK) inhibitors. JAK inhibitors are medicines that affect your immune system. JAK inhibitors can lower the ability of your immune system to fight infections. Some people have had serious infections while taking JAK inhibitors by mouth, including tuberculosis (TB), and infections caused by bacteria, fungi, or viruses that can spread throughout the body. Some people have been hospitalized or died from these infections. Some people have had serious infections of their lungs while taking OPZELURA. Your healthcare provider should watch you closely for signs and symptoms of TB during treatment with OPZELURA.

OPZELURA should not be used in people with an active, serious infection, including localized infections. You should not start using OPZELURA if you have any kind of infection unless your healthcare provider tells you it is okay. You may be at a higher risk of developing shingles (herpes zoster) while using OPZELURA.

Increased risk of death due to any reason (all causes): Increased risk of death has happened in people 50 years of age and older who have at least 1 heart disease (cardiovascular) risk factor and are taking a medicine in the class of medicines called JAK inhibitors by mouth.

Cancer and immune system problems: OPZELURA may increase your risk of certain cancers by changing the way your immune system works. Lymphoma and other cancers have happened in people taking a medicine in the class of medicines called JAK inhibitors by mouth. People taking JAK inhibitors by mouth have a higher risk of certain cancers including lymphoma and lung cancer, especially if they are a current or past smoker. Some people have had skin cancers while using OPZELURA. Your healthcare provider will regularly check your skin during your treatment with OPZELURA. Limit the amount of time you spend in the sunlight. Wear protective clothing when you are in the sun and use a broad-spectrum sunscreen.

Increased risk of major cardiovascular events: Increased risk of major cardiovascular events such as heart attack, stroke, or death have happened in people 50 years of age and older who have at least 1 heart disease (cardiovascular) risk factor and taking a medicine in the class of medicines called JAK inhibitors by mouth, especially in current or past smokers.

Blood clots: Blood clots in the veins of your legs (deep vein thrombosis, DVT) or lungs (pulmonary embolism, PE) can happen in some people taking OPZELURA. This may be life-threatening. Blood clots in the vein of the legs (deep vein thrombosis, DVT) and lungs (pulmonary embolism, PE) have happened more often in people who are 50 years of age and older and with at least 1 heart disease (cardiovascular) risk factor taking a medicine in the class of medicines called JAK inhibitors by mouth.

Low blood cell counts: OPZELURA may cause low platelet counts (thrombocytopenia), low red blood cell counts (anemia), and low white blood cell counts (neutropenia). If needed, your healthcare provider will do a blood test to check your blood cell counts during your treatment with OPZELURA and may stop your treatment if signs or symptoms of low blood cell counts happen.

Cholesterol increases: Cholesterol increase has happened in people when ruxolitinib is taken by mouth. Tell your healthcare provider if you have high cholesterol or triglycerides.

Before starting OPZELURA, tell your healthcare provider if you:

- have an infection, are being treated for one, or have had an infection that does not go away or keeps coming back
- have diabetes, chronic lung disease, HIV, or a weak immune system
- have TB or have been in close contact with someone with TB
- have had shingles (herpes zoster)
- have or have had hepatitis B or C
- live, have lived in, or have traveled to certain parts of the country (such as the Ohio and Mississippi River valleys and the Southwest) where there is an increased chance for getting certain kinds of fungal infections. These infections may happen or become more severe if you use OPZELURA. Ask your healthcare provider if you do not know if you have lived in an area where these infections are common.
- think you have an infection or have symptoms of an infection such as: fever, sweating, or chills, muscle aches, cough or shortness of breath, blood in your phlegm, weight loss, warm, red, or painful skin or sores on your body, diarrhea or stomach pain, burning when you urinate or urinating more often than usual, feeling very tired
- have ever had any type of cancer, or are a current or past smoker
- have had a heart attack, other heart problems, or a stroke

- have had blood clots in the veins of your legs or lungs in the past
- have high cholesterol or triglycerides
- have or have had low white or red blood cell counts
- are pregnant or plan to become pregnant. It is not known if OPZELURA will harm your unborn baby. There is a pregnancy exposure registry for individuals who use OPZELURA during pregnancy. The purpose of this registry is to collect information about the health of you and your baby. If you become exposed to OPZELURA during pregnancy, you and your healthcare provider should report exposure to Incyte Corporation at 1-855-463-3463.
- are breastfeeding or plan to breastfeed. It is not known if OPZELURA passes into your breast milk. Do not breastfeed during treatment with OPZELURA and for about 4 weeks after the last dose.

After starting OPZELURA:

- Call your healthcare provider right away if you have any symptoms of an infection. OPZELURA can make you more likely to get infections or make worse any infections that you have.
- Get emergency help right away if you have any symptoms of a heart attack or stroke while using OPZELURA, including:
    - discomfort in the center of your chest that lasts for more than a few minutes, or that goes away and comes back
    - severe tightness, pain, pressure, or heaviness in your chest, throat, neck, or jaw
    - pain or discomfort in your arms, back, neck, jaw, or stomach
    - shortness of breath with or without chest discomfort
    - breaking out in a cold sweat
    - nausea or vomiting
    - feeling lightheaded
    - weakness in one part or on one side of your body
    - slurred speech

- Tell your healthcare provider right away if you have any signs and symptoms of blood clots during treatment with OPZELURA, including: swelling, pain, or tenderness in one or both legs, sudden, unexplained chest or upper back pain, or shortness of breath or difficulty breathing.
- Tell your healthcare provider right away if you develop or have worsening of any symptoms of low blood cell counts, such as: unusual bleeding, bruising, tiredness, shortness of breath, or fever.

Tell your healthcare provider about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements.

The most common side effects of OPZELURA in people treated for atopic dermatitis include: common cold (nasopharyngitis), diarrhea, bronchitis, ear infection, increase in a type of white blood cell (eosinophil) count, hives, inflamed hair pores (folliculitis), swelling of the tonsils (tonsillitis), and runny nose (rhinorrhea).

The most common side effects of OPZELURA in people treated for nonsegmental vitiligo include: acne at the application site, itching at the application site, common cold (nasopharyngitis), headache, urinary tract infection, redness at the application site, and fever.

These are not all of the possible side effects of OPZELURA. Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088. You may also report side effects to Incyte Corporation at 1-855-463-3463.

Please see the Full Prescribing Information, including Boxed Warning, and Medication Guide for OPZELURA.

INDICATIONS AND USAGE

OPZELURA is a prescription medicine used on the skin (topical) for:

- short-term and non-continuous chronic treatment of mild to moderate eczema (atopic dermatitis) in non-immunocompromised adults and children 12 years of age and older whose disease is not well controlled with topical prescription therapies or when those therapies are not recommended
- the treatment of a type of vitiligo called nonsegmental vitiligo in adults and children 12 years of age and older

The use of OPZELURA along with therapeutic biologics, other JAK inhibitors, or strong immunosuppressants such as azathioprine or cyclosporine is not recommended.

It is not known if OPZELURA is safe and effective in children less than 12 years of age with atopic dermatitis or nonsegmental vitiligo.

About Incyte Dermatology

Incyte's science-first approach and expertise in immunology has formed the foundation of the company. In Dermatology, the Company's research and development efforts are focused on leveraging our knowledge of the JAK-STAT pathway to identify and develop topical and oral therapies with the potential to modulate immune pathways driving uncontrolled inflammation and help restore normal immune function.

Currently, Incyte is exploring the potential of JAK inhibition for additional immune-mediated dermatologic conditions with a high unmet medical need, including hidradenitis suppurativa. To learn more, visit the Dermatology section of Incyte.com.

About Incyte

Incyte is a Wilmington, Delaware-based, global biopharmaceutical company focused on finding solutions for serious unmet medical needs through the discovery, development and commercialization of proprietary therapeutics. For additional information on Incyte, please visit Incyte.com and follow @Incyte.

Forward-Looking Statements

Except for the historical information set forth herein, the matters set forth in this press release, including statements regarding Incyte's dermatology portfolio; its plans to present data from this portfolio at EADV; whether or when any development compounds or combinations from that portfolio will be approved or commercially available for use in humans (beyond the specific geographic regions and indications for which Opzelura already has been approved); and whether and when Opzelura and/or any of the other compounds in Incyte's clinical development pipeline, if approved, might provide a successful treatment option for patients with atopic dermatitis, vitiligo or other additional immune-mediated dermatologic conditions with a high unmet medical need contain predictions, estimates and other forward-looking statements.

These forward-looking statements are based on Incyte's current expectations and subject to risks and uncertainties that may cause actual results to differ materially, including unanticipated developments in and risks related to: unanticipated delays; further research and development and the results of clinical trials possibly being unsuccessful or insufficient to meet applicable regulatory standards or warrant continued development; the ability to enroll sufficient numbers of subjects in clinical trials; the effects of the COVID-19 pandemic and measures to address the pandemic on Incyte's clinical trials, supply chain, other third-party providers and development and discovery operations; determinations made by the U.S. FDA and other regulatory authorities outside of the United States; the efficacy or safety of Incyte's products; the acceptance of Incyte's products in the marketplace; market competition; sales, marketing, manufacturing and distribution requirements; and other risks detailed from time to time in Incyte's reports filed with the Securities and Exchange Commission, including its annual report and its quarterly report on Form 10-Q for the quarter ended June 30, 2022. Incyte disclaims any intent or obligation to update these forward-looking statements.

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Media
Catalina Loveman
+1 302 498 6171
cloveman@incyte.com

Investors
Christine Chiou
+1 302 274 4773
cchiou@incyte.com

Source: Incyte