



Press Release
Montrouge, France, June 4, 2020

DBV Technologies to Present Data on Epicutaneous Immunotherapy and Food Allergy Quality of Life at EAACI Digital Congress 2020

Virtual presentations will include data about children with and without multiple food allergies from the pivotal Phase III peanut allergy immunotherapy trial PEPITES

Biologics License Application for investigational Viaskin Peanut under review with U.S. Food and Drug Administration

DBV Technologies (Euronext: DBV – ISIN: FR0010417345 – Nasdaq Stock Market: DBVT), a clinical-stage biopharmaceutical company, today announced that seven abstracts, including four late breakers, highlighting new data from the Company's pre-clinical and clinical food allergy programs were accepted at the European Academy of Allergy & Clinical Immunology (EAACI) Digital Congress, June 6-8, 2020. The virtual oral and poster presentations will be available on the EAACI website to conference registrants starting Saturday, June 6 at 9:00am CEST / 3:00am EDT through December of this year.

“The data we are presenting at this year’s digital EAACI congress highlight potential applications of epicutaneous immunotherapy across patient populations as well as new insights into promising cellular pathways for biomarker identification,” said Pharis Mohideen, M.D., Chief Medical Officer of DBV Technologies. *“These presentations highlight the breadth of research DBV is conducting on the science and impact of food allergy as part of our commitment to improving the lives of patients through innovative potential treatments like epicutaneous immunotherapy. Our lead candidate, Viaskin Peanut, is currently under FDA review, and we look forward to making this investigational therapy available for patients as soon as possible, if approved.”*

Viaskin™ Peanut is an investigational therapy which aims to deliver biologically active compounds to the immune system through the skin to potentially safeguard peanut-allergic children in the event of accidental exposure to peanut. The Viaskin Peanut Biologics License Application, which received Breakthrough (2015) and Fast



Track (2012) designations by U.S. Food and Drug Administration (FDA), is currently under review with a target action date of August 5.

Abstracts of Interest:

Oral Presentations:

“Efficacy and Safety of Epicutaneous Immunotherapy (EPIT) for Peanut Allergy in Subjects With and Without Concomitant Food Allergies” will be presented by Philippe Bégin, M.D., PhD, Université de Montréal, Allergy and clinical immunology section, CHU Sainte-Justine

- *Abstract Number: 1362*
- *Session Title: Novel Perspectives on Diagnosis and Management of Food Allergy*

Late Breaker: “Differences in Epitome Response in Peanut-Allergic Subjects Treated with Different Immunotherapy Preparations” will be presented by Dianne Campbell, M.D., University of Sydney, Department of Allergy and Immunology, and Vice President of Clinical Development & Medical Affairs at DBV Technologies (joint submission with AllerGenis, Stanford University and Imperial College London)

- *Abstract Number: 1776*
- *Session Title: Immunotherapy: From Bench to Bedside*

Late Breaker: “Epicutaneous Immunotherapy in Murine Model Modulates Humoral Immunity Through Regulation of IL-13+ T Follicular Helper Cells” will be presented by Vincent Dioszeghy, PhD, DBV Technologies

- *Abstract Number: 1739*
- *Session Title: Immunotherapy: From Bench to Bedside*

Late Breaker: “Distinct Contribution of Skin Dendritic Cell Subsets to the Efficacy of Epicutaneous Immunotherapy in Murine Models of Food Allergy” will be presented by Leo Laoubi, PhD Fellow, DBV Technologies (Oral Presentation)

- *Abstract Number: 1747*
- *Session Title: Immunotherapy: From Bench to Bedside*



Poster Presentations:

“Evaluation of Psychometric Parameters of Food Allergy Quality-of-Life Questionnaires With Item Response Theory for the Assessment of Health-Related Quality of Life during Food Allergy Treatments” will be presented by Audrey Dunn Galvin, M.D., University College Cork, Department of Pediatrics & Child Health

- *Abstract Number: 1247*
- *Session Title: Food Allergy*

Late Breaker: “Quality of Life of Children and Adolescents with Food Allergy: Mapping FAQLQ-PF onto Paediatric-Specific Health State Utility Scores” will be presented by Gang Chen, PhD, Monash University

- *Abstract Number: 1719*
- *Session Title: Food Allergy*

“Safety Benefits of an Increased Threshold in Milk-Allergic Patients: A Quantitative Risk Assessment Study” will be presented by Benjamin C. Remington, PhD, University of Nebraska-Lincoln Food Allergy Research and Resource Program, and the Remington Consulting Group B.V. (Affiliation during study was TNO, Netherlands)

- *Abstract Number: 1192*
- *Session Title: Food Allergy*

About DBV Technologies

DBV Technologies is developing Viaskin[®], an investigational proprietary technology platform with broad potential applications in immunotherapy. Viaskin is based on epicutaneous immunotherapy, or EPIT[™], DBV’s method of delivering biologically active compounds to the immune system through intact skin. With this new class of non-invasive product candidates, the Company is dedicated to safely transforming the care of food allergic patients. DBV’s food allergies programs include ongoing clinical trials of Viaskin Peanut and Viaskin Milk, and preclinical development of Viaskin Egg. DBV is also pursuing a human proof-of-concept clinical trial of Viaskin Milk for the treatment of Eosinophilic Esophagitis, and exploring potential applications of its platform in vaccines and other immune diseases. DBV Technologies has global headquarters in Montrouge, France and offices in Bagneux, France, and North American operations in Summit, NJ and New York, NY. The Company’s ordinary shares are traded on segment B of Euronext Paris (Ticker: DBV, ISIN code: FR0010417345), part of the SBF120 index, and the Company’s ADSs (each representing one-half of one ordinary share) are traded on the Nasdaq Global Select Market (Ticker: DBVT).



Forward Looking Statements

This press release may contain forward-looking statements and estimates, including statements regarding the potential of Viaskin™ Peanut as a treatment for peanut-allergic children, Viaskin Milk as a treatment for cow's milk protein allergy and Viaskin Egg as a treatment for hen's egg allergy, the conduct and timing of the Company's clinical trials of Viaskin Peanut and Viaskin Milk and the Company's research, development and regulatory plans for its product candidates and preclinical pipeline, including the Company's planned interactions with the FDA regarding Viaskin Peanut and the target action date for the Company's BLA. These forward-looking statements and estimates are not promises or guarantees and involve substantial risks and uncertainties. At this stage, the products of the Company have not been authorized for sale in any country. Among the factors that could cause actual results to differ materially from those described or projected herein include uncertainties associated generally with research and development, clinical trials and related regulatory reviews and approvals. Furthermore, the timing of any action by the FDA and possible regulatory paths forward cannot be guaranteed. A further list and description of these risks, uncertainties and other risks can be found in the Company's regulatory filings with the French Autorité des Marchés Financiers, the Company's Securities and Exchange Commission filings and reports, including in the Company's Annual Report on Form 20-F for the year ended December 31, 2019, and future filings and reports by the Company. Existing and prospective investors are cautioned not to place undue reliance on these forward-looking statements and estimates, which speak only as of the date hereof. Other than as required by applicable law, DBV Technologies undertakes no obligation to update or revise the information contained in this Press Release.

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