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ASIT BIOTECH ANNOUNCES ITS INTENTION TO LAUNCH AN INITIAL PUBLIC OFFERING AND TO LIST ITS SHARES ON Euronext BRUSSELS AND Euronext PARIS

Brussels, Belgium, 14 April, 2016 – ASIT biotech SA (“ASIT biotech” or the “Company”), a Belgian clinical stage biopharmaceutical company focused on the development and future commercialisation of a range of breakthrough immunotherapy products for the treatment of allergies, announces today its intention to raise new funds through an Initial Public Offering with admission to trading of all shares on Euronext Brussels and Euronext Paris.

Thierry Legon, Chief Executive Officer of ASIT biotech, commented: *“ASIT biotech has the ambition to change the face of the current allergy immunotherapy (AIT) market by overcoming the main drawbacks of the existing AIT treatments. We believe this initial public offering will support the strong momentum established by our achievements and will allow us to bring our gp-ASIT+™ lead product candidate for grass pollen allergy through Phase III, currently underway in Europe, and to move towards a first application for commercialisation in Germany.”*

Company highlights:

- **Innovative ASIT+™ immunotherapy:** the Company's ASIT+™ technology platform (Allergen Specific ImmunoTherapy) allows for the production, characterisation and quality control of innovative active ingredients consisting of a unique mixture of highly purified natural

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allergen fragments in an optimal size selection. With this innovative technology the Company believes it can address both the risks and limitations of current allergy immunotherapy treatments (“AIT”). The ASIT+™ allergen fragments rapidly trigger a protective immune response without the need of an adjuvant.

This very rapid onset of action results in a 4 doctor visit treatment for gp-ASIT+™ (over 3 weeks vs. 40 visits over 3 years for classical subcutaneous AIT). The shorter course of treatment is expected to improve patient acceptance and compliance leading to a higher real-life effectiveness. The absence of adjuvant improves the safety profile and is expected to increase the acceptance of ASIT+™ products.

The Company believes that its innovative ASIT+™ technology platform is applicable to a range of allergies.

- **Addressing significant unmet medical need in a \$10bn respiratory allergies market¹:** allergic rhinitis, for which the Company is developing its two lead product candidates, affects a significant part of the population in Europe and the United States. Most of the current medications treat the symptoms but not the root cause of the disease. The only current treatment available on the market tackling the cause of allergic rhinitis is whole allergen immunotherapy. However, the drawbacks of this type of treatment limit its acceptance and the compliance to it. The Company targets allergic patients with poor disease control and who are looking for a more effective treatment.
- **1st product candidate in Phase III clinical trial:** the Company’s first lead product candidate for grass pollen allergy (*gp-ASIT+™*) has successfully completed Phase IIa and IIb clinical studies generating compelling and positive statistically significant proof of concept results. The Company has launched a Phase III clinical study with *gp-ASIT+™* in 6 European countries and has finalised patient screening and enrolment for the same.
- **2nd product candidate to enter clinical development:** the Company has completed the first phase of required regulatory preclinical development of its second lead product candidate for house dust mite allergy (*hdm-ASIT+™*) and intends to start the first clinical study in Europe in 2016.
- **Large market potential:** based on the results of the completed clinical studies to date, the Company believes that its lead product candidates have the potential to become best-in-class immunotherapy products for grass pollen and house dust mite rhinitis overcoming the limitations of current AIT treatments. As such, the Company believes that its lead product candidates could significantly expand the current immunotherapy market.
- **Manufacturing arrangements in place:** the Company has a framework service agreement in place with a contract manufacturing organisation (CMO) for the production of its novel active pharmaceutical ingredients for the European market.

¹ Visiongain, allergic rhinitis drugs market forecast 2015-2025.

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Béatrice De Vos, Chairwoman of ASIT biotech, added: *"With the development of the ASIT+™ technology platform we hope to allow a large group of allergic patients to access effective, safe and short-course immunotherapy treatments. The intended initial public offering demonstrates the significant progress the ASIT biotech team made and the advanced phase the Company is in with respect to its lead product candidate for grass pollen which we are determined to further develop towards commercialisation. The company is targeting Germany as the first country to launch this short course novel treatment of grass pollen allergic rhinitis with gp-ASIT."*

Clinical development plan

- A Phase III clinical study with *gp-ASIT+™* was launched in Europe with patient screening and enrolment completed, and the Company expects to initiate the clinical development in the United States end of 2016.
- Subject to the results of the ongoing Phase III trial with *gp-ASIT+™*, which are expected early 2017, the Company intends to file a marketing authorisation application (MAA) in Germany in order to launch *gp-ASIT+™* on the German market in the course of 2018.
- The clinical development of *gp-ASIT+™* in the United States is expected to result in filing of a Biologics Licence Application (BLA) in the United States and a MAA in the other key European countries from 2019 onwards.
- The Company's second lead product candidate *hdm-ASIT+™* is expected to enter clinical development in 2016.

The Offering

Subject to the approval of the prospectus by the FSMA, ASIT biotech intends to offer new ordinary shares in the IPO with admission to trading of all shares on Euronext Brussels and Paris. The Company intends to use the net proceeds of such offering to support the clinical development of its grass pollen rhinitis product candidate (*gp-ASIT+™*), the early stage clinical development of its second product candidate in house dust mite rhinitis, (*hdm-ASIT+™*), the working capital needs, general corporate purposes, and the early preclinical development of new ASIT+™ platform based product candidates.

KBC Securities and Société Générale Corporate & Investment Banking are acting as Joint Global Coordinators and Joint Bookrunners.

About ASIT biotech

ASIT biotech is a Belgian clinical stage biopharmaceutical company focused on the development and future commercialisation of a range of breakthrough immunotherapy products for the treatment of allergies. Thanks to its innovative ASIT+™ technology platform, ASIT biotech is currently the only developer of AIT product candidates consisting of a unique mixture of highly purified natural allergen fragments in an optimal size selection. This innovation results in a short treatment, expected to improve patient compliance and

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real-life effectiveness. ASIT biotech's product pipeline entails two novel ASIT+™ product candidates targeting respiratory allergy with the highest prevalence (i.e. grass pollen: *gp-ASIT+™* and house dust mite: *hdm-ASIT+™*), that could significantly expand the current immunotherapy market. The Company believes that its innovative ASIT+™ platform is flexible and would be applicable across a range of allergies.

ASIT biotech has a headcount of 22 staff members, at its headquarters in Brussels and its laboratory in Liège, Belgium.

The Company's shareholding includes private and public investors such as SFPI, Mr. Rodolphe de Spoelberch, Société de Développement et de Participation du Bassin de Liège SA (Meusinvest), Synergies WALLONIE SA (previously SRIW Techno SA), Finance.brussels/SRIB Group, Start-It SA and Epimède SA.

Further information can be found at: www.asitbiotech.com

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as of the date of this announcement and, without prejudice to the Company's obligations under applicable law in relation to disclosure and ongoing information, the Company does not intend, and does not assume any obligation, to update the forward-looking statements set forth in this announcement.

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