Determining the presence of IgE antibodies with ImmunoCAP Phadiatop in appropriate evaluation of atopic patients

ImmunoCAP Phadiatop is a blood test designed to differentiate between atopic and non-atopic patients. This test demonstrates the presence of IgE antibodies to common inhalant allergens and acts as an objective and reliable first step when testing for allergy.

**Clinical utility**
- An assay for graded determination of atopy with semi-quantitative or qualitative results
- The result provided demonstrates either a high or low probability for atopy
- A positive test result allows the physician to proceed with more specific testing to identify the causative allergen(s)
- A negative test result indicates that symptoms are not caused by common environmental allergens and the physician can then explore other allergens or causes of disease

**Outstanding performance**
- Over 90% probability of correctly classifying atopic vs. non-atopic patients
- High sensitivity and specificity
- Clinical performance documented in trials with more than 900 patients
- Recommended by the National Institute of Health (NIH)

*ImmunoCAP Phadiatop is a reliable and accurate first step when testing for allergy.*
Phadia® Laboratory
Systems provide optimal allergy testing solutions using advanced, state-of-the-art technology.

ImmunoCAP tests give reliable results that support primary care physicians as well as specialists in providing optimal patient management. Through fully automated Phadia Laboratory Systems you can increase your operational efficiency and shorten the lead times – whether being a small local clinic or a large commercial laboratory.

A family to grow with

When your allergy testing grows you can simply add new Phadia instrumentation without having to abandon your previous system. The unique Phadia Information Data Manager software allows you to integrate several Phadia instruments into one network without having to learn new software.

Technical features ImmunoCAP Phadiatop

- Semi quantitative or qualitative results
- Reliable and reproducible test results
- 40 µl serum or plasma needed per test