Non-invasive diagnostics of the maxillary and frontal sinuses based on diode laser gas spectroscopy*

Märta Lewander¹, Sven Lindberg², Tomas Svensson¹, Roger Siemund³, Katarina Svanberg⁴ and Sune Svanberg¹

¹ Division of Atomic Physics, Department of Physics, Lund University, Lund, Sweden
² Department of Oto-Rhino-Laryngology, University Hospital, Lund University, Lund, Sweden
³ Department of Diagnostic Radiology, University Hospital, Lund University, Lund, Sweden
⁴ Department of Oncology, University Hospital, Lund University, Lund, Sweden

Summary
Problem: Suspected, but objectively absent, rhinosinusitis constitutes a major cause of visits to the doctor, high health care costs, and the over-prescription of antibiotics, contributing to the serious problem of resistant bacteria. This situation is largely due to a lack of reliable and widely applicable diagnostic methods.

Method: A novel method for the diagnosis of rhinosinusitis based on non-intrusive diode laser gas spectroscopy is presented. The technique is based on light absorption by free gas (oxygen and water vapour) inside the sinuses, and has the potential to be a complementary diagnostic tool in primary health care. The method was evaluated on 40 patients with suspected sinus problems, referred to the diagnostic radiology clinic for low-dose computed tomography (CT), which was used as the reference technique.

Main results: The data obtained with the new laser-based method correlated well with the grading of opacification and ventilation using CT. The sensitivity and specificity were estimated to be 93% and 61%, respectively, for the maxillary sinuses, and 94% and 86%, respectively, for the frontal sinuses. Good reproducibility was shown.

Principal conclusion: The laser-based technique presents real-time clinical data that correlate well to CT findings, while being non-intrusive and avoiding the use of ionizing radiation.

Key words: paranasal sinus disease, diagnostic techniques and procedures, optical device, laser, absorption

Introduction
Rhinosinusitis, i.e. inflammation of the nose and paranasal sinuses due to e.g. bacterial, viral or allergic reaction, constitutes a major socio-economic and health problem. Costs include the consultation of a physician, tests and medical treatment. In 2002, over one fifth of the prescriptions for antibiotics in the USA were for the treatment of rhinosinusitis. Most diagnoses of rhinosinusitis, approximately 87%, are performed in primary health care clinics and are based on symptoms and clinical signs (1-3).

The definition of rhinosinusitis is symptom-based: the major factors being facial pain, facial fullness/congestion, nasal obstruction/blockage, purulence in the nasal cavity, and discharge/purulence/discoloured postnasal drainage together with fever (4,5). In 2007, the European Position Paper on Rhinosinusitis (6) expanded the recommendations for the diagnosis of this complaint to include objective signs, such as mucosal obstruction, revealed by endoscopy or computed tomography (CT). It has been found that when objective measurements are included in the diagnostic process, a large proportion (approximately 73%) of rhinosinusitis patients diagnosed as having...
rhinosinusitis according to the symptoms mentioned above, showed no evidence of the disease according to CT and/or endoscopic evaluation (7). This group of patients thus does not require treatment with antibiotics, a fact that is important in restricting the use of antibiotics.

The cost of chronic rhinosinusitis to society is high due to the large number of outpatient visits and the medication prescribed. The major contributors to the high outpatient cost are CT and nasal endoscopy. CT is especially costly, and considerable savings could be made if this could be replaced by an inexpensive non-invasive method for the diagnosis of rhinosinusitis. We present a potential diagnostic method, and the instrumentation required, that can be used by both ENT specialists and physicians in primary health care, which allows easy, non-invasive, repeated measurements. This newly developed technique provides information on the gas volume and composition of the gas in the maxillary and frontal sinuses. The method is based on a laser spectroscopic technique called GASMAS (GAs in Scattering Media Absorption Spectroscopy), which probes the free gas inside solids and liquids. GASMAS has already shown promising results in volunteers for both the frontal and maxillary sinuses. The present paper presents the results from the first clinical trial carried out to study the diagnostic accuracy of GASMAS by comparing the findings to those from CT scans.

Method

Participants

The inclusion criterion for the study population was referral to the Lund University Hospital for low-dose CT due to suspected sinus disease. The patients received information on the clinical trial in the CT examination referral letter. Forty patients who gave their written consent were examined with the index test, i.e. the developed laser-based technique (GASMAS), directly after undergoing the CT scan. Both examinations were performed at the Diagnostic Radiology Clinic at the Lund University Hospital. No treatment was administered between the tests. The study was approved by the Regional Ethics Committee in Southern Sweden (No. 210-2007).

Reference standard – Computed tomography

The diagnostic accuracy of the laser-based technique GASMAS was evaluated against CT as a reference standard. CT was chosen as the reference technique because it provides objective confirmation of sinus disease.

The CT examinations were performed using a Philips Brilliance 64-slice scanner (Philips Healthcare, Best, the Netherlands) with the patient in the supine position. The acquisition parameters were: spiral mode, 80 kVp, CTDI\textsubscript{vol} 0.7 mGy, 1 mm slice thickness reconstructed with 50% overlap. From these primary slices, multiplanar reformations with 2 mm coronal and axial slices including the maxillary and frontal sinuses were reconstructed.

Index test – Laser-based technique

The examinations with the laser-based technique GASMAS were performed with an instrument developed at the Division of Atomic Physics at Lund University, depicted in Figure 1. The instrument measures the amount of free gas, oxygen (O\textsubscript{2}), and water vapour (H\textsubscript{2}O), inside the sinus cavity, by measuring light absorption. The absorption by H\textsubscript{2}O is closely related to the volume of the gas probed, due to humidity saturation. The ratio of the absorption by O\textsubscript{2} and H\textsubscript{2}O provides a measure of the O\textsubscript{2} concentration inside the sinus. Both the absorption by H\textsubscript{2}O and the O\textsubscript{2}/H\textsubscript{2}O absorption ratio were analysed.

A detailed technical description of the instrument and data processing has been presented previously. The instrument interfaces with the patient through two probes: a fibre probe (10 cm long, 1 cm in diameter), and a detector probe (a large-area photodiode, measuring 18 × 18 mm\textsuperscript{2}), which are held against the patient’s face. The fibre probe transmits light (total energy of about 1.5 mW) from two diode lasers, emitting at 760 nm and 935 nm for the detection of O\textsubscript{2} and H\textsubscript{2}O, respectively. Laser drivers, an optical breadboard and a computer with a data acquisition card (DAQ) are assembled on a rolling cart. A long optical fibre to the light transducer (fibre probe) and a long signal cable to the light detector (detector probe) enable easy access to the sinuses of the patient during examination.

The index tests were performed on the maxillary and frontal sinuses on both sides of the patient. The measurement geometries are depicted in Figure 1. For examination of the maxillary sinus, the fibre probe is placed inside the mouth on the palate, on the right or left side pointing towards the sinus to be examined. The detector probe is placed on the cheekbone anteriorly under the eye, in direct contact with the skin. For frontal sinus examinations the fibre probe is placed at the location of the frontal bone directed towards the sinus. The detector is positioned on the frontal bone above the eyebrow. Each sinus was examined four times, the probes being repositioned between measurements. A measuring time of 20 seconds, corresponding to 200 averages of the signal, was used. During measurements the lights in the room were turned off, and the patient wore laser-light protective goggles.

Data evaluation

The diagnostic accuracy of the laser-based technique was investigated by comparing the results obtained from CT scans (i.e. the reference standard). The status of the examined
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The status of the study population, according to the reference standard evaluation (CT scans).

<table>
<thead>
<tr>
<th>Maxillary sinus</th>
<th>Frontal sinus</th>
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<td>OMC 80 sinuses</td>
<td>OMC 74 sinuses - 6 absent</td>
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<table>
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The results of the reference standard and the index test were obtained under blinded conditions. The investigators were blinded regarding the results of the CT scan during the index test (laser measurements) and during the subsequent data processing. Likewise, the radiologist analysing the CT scans was unaware of the results obtained with the laser-based method.

Statistical methods

The sensitivity and specificity of the index test were estimated by applying threshold values of the test parameters, calculated from the healthy patients. The examined sinuses were divided into two subgroups: healthy sinuses and sinuses with pathology. The results obtained from the healthy sinuses with the index test, i.e. classified as showing no opacification or obstruction of the OMC in the CT evaluation, were analysed assuming a normal distribution. Thresholds of the index test parameters, H₂O absorption and ratio of O₂ and H₂O absorption, were determined from the 95% confidence intervals. The sensitivity and specificity were estimated using the thresholds obtained for the frontal and maxillary sinus, individually.

The reproducibility of the index test was evaluated from the four consecutive individual gas absorption measurements. The deviation between the index test parameters obtained from the same sinus was calculated and used for reproducibility analysis.

Results

In the present study, 40 patients underwent examinations with the reference standard and the index tests. The examinations took place between October 21, 2008 and February 29, 2009. The study population consisted of 10 men, aged 22 - 83 years (57 ± 16 years), and 30 women aged 30 - 84 years (54 ± 14 years). The patients did not report any discomfort during the laser-based examination, which lasted approximately 15 minutes.

The distribution of the sinus conditions of the study population shows that the majority of the sinuses were classified as healthy according to the results of the reference standard (Table 1). Only 25% of the maxillary sinuses examined were opacified (Grade 1 or 2) or showed obstructed ventilation (OMC 2). The corresponding proportion for the frontal sinuses was 12%, including 7.5% with an absent sinus.

Data from 25 measurements using the index test are missing due to technical problems or limited light transmission.
through the examined sinus. The missing data correspond to 3.9% of the total 640 index test measurements performed (4 sinuses in each patient, and 4 measurements on each sinus). Technical problems, such as incorrectly saved files and saturated amplification, resulted in the loss of 9 measurements. Four of the maxillary sinuses examined (16 measurements) could not be evaluated due to the low level of laser light detected. Of the 25 missing measurements, five are a single measurement among the four consecutive measurements performed on a particular sinus. The 3 values obtained for these sinuses were used for the evaluation of the diagnostic accuracy of the method. The 5 sinuses for which no index test data were available were not included in the comparison of the reference standard and index test. The CT evaluation of these sinuses showed four healthy sinuses (Grade 0 and OMC 0), and one partially patent sinus (Grade 1 and OMC 0).

The results of the index test parameters, absorption by H₂O and the ratio of O₂ and H₂O absorption are shown in Figure 2, in relation to their classification from the CT images. Sinuses with H₂O absorption below the detection limit of the index test instrument, i.e. 2 mm [12], were excluded from the evaluation of the ratio. It can be seen that the sinuses with no gas-containing cavity, Grade 2 or absent sinuses, exhibit no H₂O absorption. The maxillary sinuses with a partially gas-filled cavity (Grade 1) show a lower H₂O absorption than aerated sinuses (Grade 0). This clear differentiation between Grades 1 and 0 was not seen in the data obtained from the frontal sinuses. Gas-filled, but unventilated maxillary and frontal sinuses (Grade 0 and OMC 2) show no anomaly in H₂O absorption, but 3 out of 5 exhibited a reduced oxygen concentration. Low H₂O absorption in the index test identifies obstructed and absent sinuses well, and a low O₂/H₂O absorption ratio identifies unventilated sinuses.

The H₂O absorption (which is a measure of the volume of gas) and the O₂/H₂O absorption ratio (which provides a measure of the oxygen concentration) provide complementary information. The clinical applicability of the index test should thus be evaluated using both these parameters together. Presenting the measured absorption of O₂ and H₂O of a particular sinus together, illustrates a way to expose both index test parameters with retained sinus identity. Figure 3 shows the index test data obtained in this way demonstrating the suitability of this approach. The sinuses classified as healthy (Grade 0 and OMC 0) exhibit strongly positively correlated values of O₂ and H₂O absorption that lie around the line y = x. This indicates that the gas volumes of O₂ and H₂O probed were similar, and that the O₂ concentration is similar to the ambient concentration. A clear deviation from the line y = x indicates an abnormal oxygen concentration. Low values of H₂O and O₂ absorption, i.e. low gas volume, are located in the lower left corner of each plot. The data presented in Figure 3 provide information on the gas volume and oxygen concentration in a particular sinus, in contrast to the data presented in Figure 2.

The data and thresholds presented in Figure 3 yield a sensitivity and specificity of 93% and 61% for the maxillary sinuses,
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and 94% and 86% for the frontal sinuses (Table 2). Sinuses with any kind of pathological finding according to the reference standard classification (CT scans) are treated as one group. Both parameters of the index test, H\textsubscript{2}O absorption and O\textsubscript{2}/H\textsubscript{2}O absorption, were used to determine the diagnostic accuracy. The estimated sensitivity and specificity provide information about the possibility of distinguishing sinuses with pathological conditions from healthy sinuses (Grade 0 and OMC 0).

The index test showed good reproducibility. The variation between the four consecutive measurements on each sinus with the laser-based technique was small. For the maxillary sinus, the average deviation between the measurements on the same sinus was 2.0 mm for the H\textsubscript{2}O absorption and 0.09 for the O\textsubscript{2}/H\textsubscript{2}O ratio. The corresponding values for the frontal sinus were 1.2 mm and 0.09.

**Discussion**

The results of this study show that there is good correlation between the reference standard (CT scans) and the index test (laser-based technique GASMAS). They also demonstrate that diagnostically valuable information can be obtained non-invasively, with a simple instrument not requiring ionizing radiation. The laser-based system consists of low-cost components that can be incorporated into a small device with good usability, as well as in instruments with additional sensors. The price of the research instrument was about €20,000, and exhibits large flexibility and alteration possibilities. The cost of a commercially produced instrument can be expected to be considerably lower.

The non-intrusive and harmless nature of the technique allows repeated examinations, making it possible to study the course of the disease, as well as recurring problems. The diagnostic parameters provided by the technique are the absorption of light by H\textsubscript{2}O (which is correlated to the gas volume probed), and the ratio of the absorption by O\textsubscript{2} and H\textsubscript{2}O (which provides a measure of the oxygen concentration). As can be seen in Figure 2, the former parameter is related to the obstruction of the sinus, while the latter is related to the ventilation of the sinus. A useful way of combining the parameters in the analysis is to plot the O\textsubscript{2} absorption vs. the H\textsubscript{2}O absorption in a scatter plot, as shown in Figure 3. Here, healthy sinuses lie close to the line y = x above the H\textsubscript{2}O absorption threshold. Distinct deviation from the line indicates a low or high (high being physiologically unlikely) oxygen concentration. Small values close to the origin of the plot indicate low gas absorption, i.e. opacified, absent or small sinuses.

This study on the clinical applicability of the laser-based method demonstrated good sensitivity and specificity between sinuses with and without pathological findings according to the CT classification scheme. Due to the low occurrence of sinuses with pathological conditions in the study population, no statistical differentiation could be made between the CT classification subgroups, however, particular groups, such as those with opacified sinuses (Grade 2 and absent sinuses) are well distinguished with the laser-based technique. Further studies on a larger study population are needed to determine the sensitivity and specificity for specific subgroups. The large proportion of healthy sinuses in the present study population provides a good basis for threshold calculations, whereas the statistical data are not conclusive with respect to sensitivity and specificity.

Since the laser-based method provides information on the gas volume probed by the laser light, there is no intrinsic way to distinguish between small healthy sinuses and partially obstructed larger sinuses, or between an absent or a completely obstructed sinus. Using an additional method, such as acoustic techniques, together with the laser-based method could allow such distinctions to be made. Further investigations into this are required. Additional information that could be incorpora-
Figure 3. Absorption by O$_2$ vs. H$_2$O together with 95% confidence interval threshold values of H$_2$O absorption and ratio of O$_2$ to H$_2$O absorption (dashed lines), for the maxillary sinuses (left) and the frontal sinuses (right).

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Author contributions
SS, KS, SL and ML designed the trial and SS and ML developed the technique. The collection of clinical data was performed with ML as the main responsible. The index test instrument including software was constructed by ML and TS. CT classification was conducted by RS. The analysis of the data obtained with the index test and the diagnostic correlations were performed by ML. The report was drafted by ML in collaboration with the other authors. All authors approved the final text.

Conflict of interest statement
ML, KS and SS are shareholders in a spin-off company, GASPOROX AB.
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References

Märta Lewander
Division of Atomic Physics
Department of Physics
Lund University
Lund
Sweden

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