

Improved Inverse Method for Radiative Characteristics of Closed-Cell Absorbing Porous Media

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Radiative characteristics such as the extinction coefficient, the scattering albedo and the scattering phase function of fused quartz containing closed cells are determined by using an inverse method based on theoretical and experimental bi-directional transmittances. The theoretical transmittances are obtained by solving the Radiative Transfer Equation with the Discrete Ordinate Method. Improvements have been made over previously reported experimental determination of porous fused quartz radiative characteristics by using a more accurate phase function and an adaptive quadrature to compute more precisely the intensities in the measurement directions. In addition, a two step inverse method to compute accurately and simultaneously the radiative parameters has been developed. The results are shown to be independent of samples thickness. Exhaustive comparison between experimental measurements of hemispherical transmittance and reflectance and computational results using the retrieved radiative characteristics shows good agreement. The retrieved absorption coefficient of porous fused quartz appears to be more realistic than that reported in our earlier publication.

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Nomenclature

a	bubble radius, m
b	corrective factor used in Eq. (10)
c_{ij}	matrix elements of the sensitivity coefficients J
CN	condition number of the sensitivity matrix J
e	sample thickness, m
f_1, f_2	spectral weights of the Henyey-Greenstein phase function Φ_{HG}
g_1, g_2	spectral parameters of the Henyey-Greenstein phase function Φ_{HG}
g	spectral asymmetry factor
I	spectral radiation intensity, $Wm^{-2}sr^{-1}$
IN	iteration number
J	matrix of the sensitivity coefficients
k	volumetric absorption coefficient, m^{-1}
m	fused quartz refractive index
Mb	quadrature order of the discrete ordinate method
MN	measurement number
n	number of unknown parameters including ω , β , f_1 , g_1 , and/or g_2
Nb	number of measurement directions
p	unknown parameter such as ω , β , f_1 , g_1 , or g_2
Q	ratio of the measured scattered to the incident radiation fluxes
r	interface reflectivity
S	minimization function
T	spectral transmittance or reflectance, sr^{-1}
\bar{T}	average spectral transmittance or reflectance, sr^{-1}
x	bubble size parameter
y	spatial coordinate along the sample thickness, m
w	angular weight of the discrete ordinate method, sr
α	angle between incident radiation and measurement directions, rad
β	volumetric extinction coefficient, m^{-1}

χ	experimental error, %
δ	Kronecker delta function
$\varepsilon_0, \varepsilon_1, \varepsilon_2, \varepsilon_3$	coefficients of the third order polynomial estimating T_{sca} in Eq. (17)
Φ	spectral phase function
γ	relaxation factor used in Eq. (5)
η	cosine of the angle α
φ	azimuthal angle, <i>rad</i>
κ	fused quartz absorption index
λ	radiation wavelength, <i>m</i>
μ	cosine of the angle θ
ν_i	weight associated to measurement in the direction $i=1$ to N_b
Π	dimensionless sensitivity coefficient
θ	angle between incident radiation direction and radiation inside the porous medium, <i>rad</i>
Θ	scattering angle defined in Eq. (21), <i>rad</i>
$\Delta\theta$	divergence angle of the incident radiation, <i>rad</i>
σ	standard deviation
τ_0	optical thickness
ξ	random number defined between 0 and 1
ω	volumetric scattering albedo
$\Delta\Omega$	solid angle, <i>sr</i>

Superscripts

+	refers to hemispherical transmittance
-	refers to hemispherical reflectance

Subscripts

<i>bulk</i>	refers to the continuous phase (quartz)
<i>coll</i>	refers to collimated radiation

<i>d</i>	refers to detection
<i>e</i>	refers to experimentally measured value
<i>exact</i>	refers to the exact radiative parameters
<i>HG</i>	Henyey-Greenstein phase function
<i>max, min</i>	refers to the higher and lower integration bound in Eq. (30), respectively
<i>NC</i>	refers to the Nicolau phase function
<i>sca</i>	refers to the scattered
<i>t</i>	refers to the theoretical value
<i>TPF</i>	refers to the truncated phase function
λ	refers to spectral value
<i>0</i>	refers to incident radiation
<i>12</i>	refers to radiation from the air to the air-glass interface
<i>21</i>	refers to radiation from the glass to the glass-air interface

I. Introduction

FOAM and cellular materials have practical importance in many applications. Examples range from food processes, where foam can disrupt the process, to space and building applications where they are used as insulating materials. Thermal radiation in cellular materials is a significant mode of energy transfer in most of these applications. Thus, the modeling of radiative transfer in cellular materials has a primary importance for optimizing performance in engineering applications. An extensive review of radiative transfer in dispersed media was carried out by Viskanta and Mengüç,¹ and by Baillis and Sacadura.² A porous medium is often treated as a continuous, homogeneous, absorbing, and scattering medium. In order to evaluate the radiative heat transfer, radiative characteristics such as the extinction coefficient, the scattering albedo, and the scattering phase function are required. They can be determined by different approaches:

1) radiative characteristics can be predicted from porosity and bubble size distribution by considering a random arrangement of particles by using for example the Mie theory or the geometric optics laws assuming independent scattering;³⁻⁸

2) other methods consist of determining the radiative characteristics from a Monte Carlo approach at the microscopic scale, taking into account the complex morphology of the porous medium;⁹⁻¹⁴

3) finally, other approaches are based on the experimental measurement of reflectance and transmittance of the medium on a macroscopic scale combined with an inverse method.¹⁵⁻²⁰ Hemispherical emittance measurements have also been exploited to retrieve the radiative characteristics of porous media.^{21,22}

The present study focuses on radiative characteristics of fused quartz containing bubbles or closed cells as illustrated in Fig. 1. Few studies on such media have been reported. Pilon and Viskanta⁶ have studied the effects of volumetric void fraction and bubble size distribution on the radiative characteristics of semitransparent media containing gas bubbles. They used the model proposed by Fedorov and Viskanta⁷ which is based on the anomalous-diffraction approximation. Wong and Mengüç¹² used a ray-tracing method in a porous medium composed of spherical air pockets embedded in a non-absorbing matrix to study the depolarization of the incident radiation. More recently, experimental determination of radiative characteristics of fused quartz containing bubbles¹⁸ based on an inverse method has been carried out. The Henyey-Greenstein phase function model was adopted and theoretical transmittance in the experimental directions was interpolated from the solution of the Radiative Transfer Equation. The retrieved absorption coefficient of the porous fused quartz was found to be greater than that of dense fused quartz which a priori, seems to contradict physical intuition since bubbles entrapped in the glass matrix are transparent.^{23, 24} On the other hand, the larger absorption coefficient could be attributed to trapping of radiation by successive inter-reflections within the bubbles¹⁸ or to the increases optical path within the glass matrix due to reflections at the surface of the bubbles. These apparent contradictions are due to the choice of the phase function model used in the calculations of the theoretical bi-directional transmittance and reflectance and will be clarified in this paper.

The present study aims at completing the previous one¹⁸ by investigating (1) the influences of the phase function model on the retrieved radiative characteristics, (2) the best way to calculate the transmitted intensity in the specific measurement directions, and (3) the development of a more efficient identification technique. First, the inverse method using experimental and theoretical transmittances is described, including details regarding (i) different forms of the minimization and (ii) scattering phase functions, and as well as (iii) the direct computation of the extinction coefficient. Then, the experimental setup and the measurements are briefly presented while the theoretical model for calculating the bi-directional transmittance and reflectance is explained in detail. Finally, the results are presented and discussed.

II. Parameter identification method

A. Description

The spectral radiative characteristics of semitransparent media are the single scattering albedo ω_λ , the extinction coefficient β_λ and the scattering phase function Φ_λ which depends on n-2 parameters that will be here denoted by $(p_l)_{l=3,n}$. As a result, the n unknown parameters are $(p_l)_{l=1,\dots,n}=(\omega, \beta, p_{l=3,n})$. For a given sample, the parameter identification method is based on:

- 1) the experimental measurements of the bi-directional reflectance and transmittance (T_e) obtained for several directions (i),
- 2) the theoretical bi-directional reflectance and transmittance (T_i) calculated for the same directions.

For each wavelength λ , the goal is to determine the radiative parameters $(p_l)_{l=1,\dots,n}$, which minimize a function S characterized by the quadratic differences between the experimentally measured bi-directional transmittances $T_{e,i}$ and the corresponding numerically calculated value $T_{i,i}$ for Nb measurement directions:

$$S(p_1, \dots, p_n) = \sum_{i=1}^{Nb} v_i^2 [T_{ii}(p_1, \dots, p_n) - T_{ei}]^2 \quad (1)$$

The bi-directional transmittance or reflectance T_i for normal incident intensity are defined by the following expression:

$$T_i = \frac{I_i}{I_0 \Delta\Omega_0} \quad (2)$$

where I_i is the transmitted or reflected intensity in the direction i and I_0 is the intensity of the collimated beam normally incident on the sample within the incident solid angle $\Delta\Omega_0$. The weight v_i , associated with the direction i , is introduced to decrease the importance of inaccurate measurements.

The optimization method adopted is the Gauss linearization method²⁵ which minimizes S by setting to zero the derivatives of Eq. (1) with respect to each of the unknown parameters. As the system is non-linear, an iterative procedure is performed over j iterations:

$$\begin{bmatrix} \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_1} \right)^2 & \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_1} \right) \left(\frac{\partial T_{ii}}{\partial p_2} \right) & \dots & \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_1} \right) \left(\frac{\partial T_{ii}}{\partial p_n} \right) \\ \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_1} \right) \left(\frac{\partial T_{ii}}{\partial p_2} \right) & \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_2} \right)^2 & \dots & \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_2} \right) \left(\frac{\partial T_{ii}}{\partial p_n} \right) \\ \dots & \dots & \dots & \dots \\ \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_1} \right) \left(\frac{\partial T_{ii}}{\partial p_n} \right) & \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_2} \right) \left(\frac{\partial T_{ii}}{\partial p_n} \right) & \dots & \sum_{i=1}^{Nb} v_i^2 \left(\frac{\partial T_{ii}}{\partial p_n} \right)^2 \end{bmatrix}^j \begin{bmatrix} \Delta p_1 \\ \Delta p_2 \\ \dots \\ \Delta p_n \end{bmatrix} = \begin{bmatrix} \sum_{i=1}^{Nb} v_i^2 (T_{ii} - T_{ei}) \frac{\partial T_{ii}}{\partial p_1} \\ \sum_{i=1}^{Nb} v_i^2 (T_{ii} - T_{ei}) \frac{\partial T_{ii}}{\partial p_2} \\ \dots \\ \sum_{i=1}^{Nb} v_i^2 (T_{ii} - T_{ei}) \frac{\partial T_{ii}}{\partial p_n} \end{bmatrix} \quad (3)$$

Solutions of the system of Eq. (3) gives the variation parameter Δp_l^j added to the value of each parameter p_l^j at the j^{th} iteration, i.e.,

$$p_l^{j+1} = p_l^j + \Delta p_l^j \quad \text{with } l=1,2,\dots,n \quad (4)$$

The use of Eq. (4) limits the convergence of the inverse process due to large values of Δp_l^j during the first few iterations. In this study, we propose to weight the parameter Δp_l^j with a relaxation factor γ ($1 \geq \gamma > 0$):

$$p_l^{j+1} = p_l^j + \gamma \Delta p_l^j \quad \text{with } l=1,2,\dots,n \quad (5)$$

The converged solution is estimated to be reached when $\Delta p_l^j / p_l^j < 10^{-3}$. The matrix on the left-hand side of Eq. (3) will be termed J. It is composed of the sensitivity coefficients $\partial T_{ii} / \partial p_l$ calculated from the theoretical model. The condition number (CN) of the matrix J can be calculated from the following relation:

$$CN(J) = \| J^{-1} \| \cdot \| J \| \quad (6)$$

where $\| J \|$ is the norm of the matrix, calculated from the matrix elements c_{ij} as follows:

$$\| J \| = \max_{i=1,n} \sum_{j=1}^n c_{ij} \quad (7)$$

Note that the condition number CN is always larger than unity. The larger CN is, the more ill-conditioned the system. Thus, small uncertainties in the measurements can result in very large changes in the Δp vector. A large condition number occurs when at least two of the sensitivity coefficients are quasi-linearly interdependent and/or when at least one is very small or very large compared to the others. The analysis of the sensitivity coefficients and

the condition number is an efficient tool for understanding the physical behavior of the problem and for studying the feasibility of simultaneous determination of the unknown parameters.²⁶

B. Measurements weights

Two common expressions for the weights can be used:¹⁷⁻¹⁹

$$v_i = 1 \quad \text{for } i=1 \text{ to } Nb \quad (8)$$

$$v_i = \frac{1}{T_{ei}} \quad \text{for } i=1 \text{ to } Nb \quad (9)$$

When using Eq. (8), the minimization gives more importance to the highest measurement values. This is inconvenient in the situation where physical information is “hidden” behind small data values. Thus, using Eq. (9) enables one to give the same importance to each measurement and appears to be more appropriate.

Usually, some measurements feature large experimental uncertainties as detailed hereafter in section III. B. Such data can potentially contain information about the porous material. It is therefore preferable to decrease the importance of that data in the identification procedure rather than to discard them completely. This can be done by modifying the weight v_i by a corrective factor b_i as follows:

$$v_i = \frac{b_i}{T_{ei}} \quad \text{for } i=1 \text{ to } Nb \quad (10)$$

The value of the parameter b_i depends on the accuracy of the measurements estimated through the experimental errors χ discussed in section V.C. In this study, the following values were used for different measurement error ranges:

$$\begin{cases} \chi_i < 10 \% & \Rightarrow b_i = 1 \\ 10 \% < \chi_i < 20 \% & \Rightarrow b_i = 0.1 \\ \chi_i > 20 \% & \Rightarrow b_i = 0.01 \end{cases} \quad \text{for } i=1 \text{ to } Nb \quad (11)$$

The weights given by Eqs. (10) and (11) for different experimental uncertainties are adopted in this study.

C. Models for the scattering phase function

The expression for the phase function plays an important role in describing the appropriate directional scattering behavior. In practice, the representation of the scattering phase function as an expansion in Legendre polynomial²⁷ is not suitable in the case of highly anisotropic material due to the larger number of unknown parameters.²⁰ Among the useful models, the Henyey-Greenstein (HG) approximation²⁸ is the most popular with only one unknown parameter g_λ :

$$\Phi_{HG}(\Theta, g_\lambda) = \frac{1 - g_\lambda^2}{(1 + g_\lambda^2 - 2g_\lambda \cos \Theta)^{3/2}} \quad (12)$$

where Θ is the angle between the directions of the incident and scattered radiation intensities on a scattering point.

However, some materials having a more complex anisotropic scattering pattern require the use of a more complex scattering function to properly describe the directional scattering behavior. For example, Nicolau et al.¹⁹ proposed a combination of Henyey-Greenstein functions for fibrous media:

$$\Phi_{NC}(\Theta, g_{1\lambda}, g_{2\lambda}, f_{1\lambda}, f_{2\lambda}) = f_{2\lambda} [f_{1\lambda} \Phi_{HG}(\Theta, g_{1\lambda}) + (1 - f_{1\lambda}) \Phi_{HG}(\Theta, g_{2\lambda})] + (1 - f_{2\lambda}) \quad (13)$$

where $f_{1\lambda}$ and $f_{2\lambda}$ are the weights associated to the scattering functions $\Phi_{HG}(\Theta, g_{1\lambda})$ and $[f_{1\lambda} \Phi_{HG}(\Theta, g_{1\lambda}) + (1 - f_{1\lambda}) \Phi_{HG}(\Theta, g_{2\lambda})]$, respectively.

According to the previous study¹⁹, the simultaneous computation of these parameters combined with the extinction coefficient and the scattering albedo remains critical. The reduction of the number of unknown parameters (ω , β , f_1 , f_2 , g_1 and g_2) to be simultaneously identified is required.

This study proposes a new combination of scattering functions called the Truncated Phase Function (TPF) depending only on three parameters (f_1 , g_1 and g_2):

$$\begin{cases} \Phi_\lambda(\Theta, f_1, g_1, g_2) = \Phi_{1\lambda}(\Theta, f_1, g_1, g_2) = f_1 \Phi_{HG\lambda}(\Theta, g_1) + (1 - f_1) \Phi_{HG\lambda}(\Theta, g_2) & \text{for } 0^\circ \leq \Theta \leq 90^\circ \\ \Phi_\lambda(\Theta, f_1, g_1, g_2) = 0.03 \Phi_{1\lambda}(\Theta, f_1, g_1, g_2) & \text{elsewhere} \end{cases} \quad (14)$$

Figure 2 compares the TPF to the exact Mie scattering phase function²⁹ in the case of an optically large bubble ($x=2\pi a/\lambda=2000$) located in a refracting medium ($m_\lambda=1.44$). The corresponding TPF parameters are $f_1=0.2$, $g_1=0.98$, and $g_2=0.45$. One can note that the TPF function can properly approximate the exact phase function.

The method previously adopted by Nicolau et al.¹⁹ is used to normalize this function. The resulting function satisfies:

$$\frac{1}{2} \int_0^\pi \Phi_\lambda(\Theta, f_1, g_1, g_2) \sin\Theta d\Theta = 1 \quad (15)$$

The asymmetry factor corresponding to this phase function can be determined by:²⁹

$$g_\lambda = \frac{1}{2} \int_0^\pi \Phi_\lambda(\Theta, f_1, g_1, g_2) \cos\Theta \sin\Theta d\Theta \quad (16)$$

D. Direct estimation of the extinction coefficient

The collimated transmittance T_{coll} in the incident radiation direction which is attenuated only by extinction and reflection at interfaces can be written as:²⁸

$$T_{coll} = T_{e1} - T_{sca1} = \frac{(1-r_{12})^2 \exp(-\beta e)}{1-r_{12}^2 \exp(-2\beta e)} \quad (17)$$

where T_{e1} refers to the measured transmittance in the incident direction while T_{sca1} refers to the scattered transmittance toward the same direction. The subscript 1 refers to the first direction of measurement ($i=1$) which is, in this study, the same as the incident radiation direction. The Fresnel reflectivity is denoted r_{12} at the air-glass interface for normal incidence and e is the sample thickness.

Here, T_{sca1} is estimated by a third order polynomial in terms of the measurement direction cosine η ($=\cos\alpha$) i.e., $T_{scai} = \varepsilon_3 \eta_i^3 + \varepsilon_2 \eta_i^2 + \varepsilon_1 \eta_i + \varepsilon_0$ with $i=1$ and $\eta_1=1$. The four coefficients ε_0 , ε_1 , ε_2 , and ε_3 are obtained by matching the conditions: $T_{scai} = T_{ei}$ for the four directions $i=2$ to 5.

After some manipulations of Eq. (17), the extinction coefficient can be expressed as:

$$\beta = -\frac{1}{e} \ln \left\{ \frac{\sqrt{(1-r_{12})^4 + 4T_{coll}^2 r_{12}^2} - (1-r_{12})^2}{2T_{coll} r_{12}^2} \right\} \quad (18)$$

The extinction coefficient can be directly calculated from Eq. (18). Thus, the set of unknown parameters to be simultaneously identified is reduced to ω , f_1 , g_1 and g_2 .

III. Experimental Measurements

A. Experimental Setup

The experimental data of spectral bi-directional transmittance and reflectance are obtained from an experimental setup including a Fourier-transform infrared spectrometer (FTS 60 A, Bio-Rad Inc) associated with a detector (liquid nitrogen cooled MCT detector) mounted on a goniometric system.¹⁷⁻¹⁹ The incident radiation emitted by the source is modulated after which the resulting spectrum range varies from 1.67 to 14 μm . The collimated beam is perpendicularly incident on the sample with a divergence half-angle of $\Delta\theta_0=2.21 \cdot 10^{-2}$ rad and a beam size d equal to 40 mm. We assume that the samples are thin enough (see Section V.A) to guarantee one-dimensional radiation transfer. Unfortunately, to the best of our knowledge, no criteria for the ratio of sample thickness to beam diameter is available in the literature. Note that measurements were performed with and without a gold coating deposited on the edges of the slab. No noticeable effects were recorded indicating that the side boundary conditions have no effect on the directional transmittance and reflectance. Therefore, the problem can be treated as one-dimensional.

The intensity transmitted or reflected by the sample is collected by a spherical mirror which focuses it on the detector. The corresponding detection solid angle ($\Delta\Omega_d$) is characterized by a detection half-angle equal to $\Delta\theta_d=0.33 \cdot 10^{-2}$ rad.

Then, the measured bi-directional transmittance in the direction i can be computed from.³⁰

$$T_{ei} = \frac{I(\eta_i)}{I_0 \Delta\Omega_0} = \frac{Q_i}{\eta_i \max(\Delta\Omega_d, \Delta\Omega_0)} \quad (19)$$

where Q_i is the ratio of the radiation flux transmitted or reflected by the sample to that incident on the sample, directly estimated from the FTIR measurement in the direction i . The incident and detection solid angles can be expressed as $\Delta\Omega_0=2\pi(1-\cos\Delta\theta_0)$ and $\Delta\Omega_d=2\pi(1-\cos\Delta\theta_d)$, respectively.³⁰

The bi-directional measurements are carried out over $N_b=24$ directions as shown in Fig. 3: 12 directions in the forward hemisphere (transmittances) and 12 directions in the backward hemisphere (reflectances). These directions are chosen by combining two Gaussian quadratures aimed at increasing the number of measurements around the direction of the incident radiation and suitable for forward and/or backward scattering media.¹⁹ Note that N_b must be a positive even number.

B. Measurement uncertainties

There are two major sources of experimental uncertainties involved in the bi-directional FTIR measurements, namely noise and misalignment. Indeed, the measurements become erroneous when the signal to noise ratio is too

small. It is the case for our measurements (i) at wavelengths beyond 4.04 μm for which the fused quartz becomes optically thick and (ii) far from the incident radiation direction (from 48 ° to 90 ° for transmittances and from 90 ° to 148 ° for reflectances) where the magnitude of the scattered flux is small.

Moreover, for the measurement in the 2nd direction (the closest to the incident radiation direction), the signal decreases sharply and a slight overestimation of the measured signal occurs due to the diffraction of the incident beam from the aperture inside the FTIR spectrometer. Also, the same problem occurs for measurement close to the backward specular direction (i.e., 23rd direction). Measuring precisely the specular reflectance is also difficult with our goniometric system due to the optical misalignments.

In order to reduce these uncertainties, the measurements are repeated five times (each one corresponding to a new goniometric system alignment) and the resulting average bi-directional transmittances and reflectances are used in the inversion procedure. Moreover, only the essential measurements which are required for identification are retained. These measurements are presented and analyzed in sections V.B and V.C.

IV. Theoretical model

The theoretical spectral bi-directional transmittance and reflectance are computed by solving the Radiative Transfer Equation (RTE) based on the assumptions that (1) radiative transfer is one-dimensional, (2) a steady-state regime is established, (3) azimuthal symmetry prevails, and (4) medium emission can be disregarded thanks to the radiation modulation and the phase sensitive detection.

A. RTE and boundary conditions

Under the above assumptions, the RTE can be written as follows:¹⁸

$$\mu \frac{\partial I_{\lambda}(y, \mu)}{\beta_{\lambda} \partial y} + I_{\lambda}(y, \mu) = \frac{\omega_{\lambda}}{2} \int_{-1}^1 I_{\lambda}(y, \mu') \Phi_{\lambda}(\Theta) d\mu' \quad (20)$$

where y indicates the spatial coordinate along the sample thickness and μ is the direction cosine of the intensity with respect to the y axis.

The scattering angle Θ can be expressed in terms of the direction cosines μ and μ' as:³¹

$$\Theta = \cos^{-1} \left(\mu \mu' + \left[(1 - \mu^2)(1 - \mu'^2) \right]^{1/2} \cos \varphi \right) \quad (21)$$

where φ is the azimuthal angle which can be taken any arbitrary value in case of azimuthal symmetry.³¹

The boundary conditions are obtained by assuming that the interfaces are optically smooth i.e., the surface roughness is smaller than the wavelength of the incident radiation and the area associated with the open bubbles at the sample surfaces is negligible due to the small void fraction. In fact, the sample surfaces are mechanically polished as described in the previous work¹⁸ and the open bubbles occupy only 7.5 % of the total sample surface exposed to the incident radiation. Then, the boundary conditions associated with the RTE for normal incident radiation are:¹⁸

$$I_{\lambda}(0,\mu) = r_{21}I_{\lambda}(0,-\mu) + m_{\lambda}^2(1-r_{12})\delta_{\mu_0,\mu}I_{\lambda}(0,\mu_0) \quad \mu > 0 \quad (22)$$

$$I_{\lambda}(e,\mu) = r_{21}I_{\lambda}(e,-\mu) \quad \mu < 0 \quad (23)$$

where r_{12} and r_{21} are the interfacial reflectivities at the air-glass and glass-air interfaces, respectively. The Kronecker delta function is defined as $\delta_{\mu_0,\mu} = 1$ if $\mu = \mu_0$ and $\delta_{\mu_0,\mu} = 0$ otherwise where $\mu_0 = 1$ in the case of normal incident radiation.

When the absorption index κ_{λ} is small such as $\kappa_{\lambda} < 10^{-3}$ which is the case for fused quartz in the spectral range of interest from 1.67 to 4.04 μm as one can see in Fig. 4,^{23, 32-34} the reflectivities r_{12} and r_{21} are determined entirely from the refractive index m_{λ} by means of the Fresnel's equations.^{28, 31} The air-glass reflectivity r_{12} for normal incident radiation can be calculated as:³¹

$$r_{12} = \frac{(m_{\lambda} - 1)^2}{(m_{\lambda} + 1)^2} \quad (24)$$

Due to scattering by bubbles embedded in the continuous phase, the radiation field inside the porous slab does not reach the back face of the slab perpendicularly. Thus, the glass-air reflectivity r_{21} is given by:³¹

$$r_{21} = \frac{1}{2} \left[\frac{\sin^2(\theta - \alpha)}{\sin^2(\theta + \alpha)} + \frac{\text{tg}^2(\theta - \alpha)}{\text{tg}^2(\theta + \alpha)} \right] \quad (25)$$

where $\theta = \cos^{-1}\mu$ is the angle between the internal radiation towards the surface of the slab and the y axis, and $\alpha = \cos^{-1}\eta$ is the angle between the refracted radiation leaving the slab and the incident radiation direction. The angles θ and α are related by Snell's law³¹ expressed as:

$$m_\lambda \sin\theta = \sin\alpha \quad (26)$$

B. Method of solution of the RTE

The Discrete Ordinate Method (DOM)^{28, 31} is applied to solve the RTE [Eq. (20)]. It consists of replacing the integral term in the RTE by a sum over Mb directions namely “quadrature”. Several standard quadratures such as the Gaussian, Radau, and Fiveland quadratures³⁵ can be used in the integral calculation. Then, a system of partial differential equations is obtained. Previously reported studies on the experimental determination of radiative characteristics of open-cell porous media^{17, 19} neglected the reflection at the interfaces due to the large porosity of the medium. Then, a simpler system of equations could be solved analytically by separating the collimated and scattered radiation. In the present study however, the system is more complicated due to the reflection at the front and back interfaces. Thus, the space is discretized along the y-direction in order to solve numerically the system of partial differential equations with the associated boundary conditions [Eqs. (22) and (23)] by using the control volume method.³⁶ A linear scheme (diamond) is employed to evaluate the radiative intensity in the middle of the control volume knowing the radiative intensities on the control volume boundaries.³⁶ For a number of control volumes larger than 190, the numerical results were found to be independent of the number of control volumes,¹⁸ i.e., numerical convergence was reached.

C. Transmitted and reflected intensity calculations

The intensities leaving the sample with smooth interfaces can be written as:¹⁸

$$I_\lambda(0,\eta) = r_{12}\delta_{\mu_0,-\eta}I_\lambda(0,-\eta) + \left(\frac{1}{m_\lambda}\right)^2(1-r_{21}) I_\lambda(0,\mu) \quad \eta > 0 \quad (27)$$

$$I_\lambda(e,\eta) = \left(\frac{1}{m_\lambda}\right)^2(1-r_{21}) I_\lambda(e,\mu) \quad \eta < 0 \quad (28)$$

where the direction cosine $\eta = \cos\alpha$ with α is the measurement angle while $\mu = \cos\theta$ is related to η through Eq. (26).

In general, the measurement angle α is different from one of the quadrature angles θ_i selected for the numerical calculations due to refraction at the interfaces of the slab except for the direct transmission and backscattering directions (i.e., for $\eta=\pm 1$). To circumvent this difficulty, an interpolation can be used to evaluate the intensity in the measurement angle α using the computed intensity in the quadrature directions.¹⁸ Different interpolation laws such as linear (LL), exponential (EL) and combined exponential-linear laws (ELL) can be used.

D. Adaptive Composite Quadrature

Another way of avoiding the above mentioned difficulty related to interpolation is to use an Adaptive Composite Quadrature (ACQ)³⁷ to solve Eq. (20). In the present study, the ACQ quadrature depends on the radiation wavelength and consists of $Mb/2$ directions in each hemisphere such that $Mb > Nb$ (Fig. 3). Here also, Mb must be a positive even number.

In the forward hemisphere, the first $Nb/2$ directions among the $Mb/2$ directions are related directly to the experimental direction through Snell's Law. Rearranging Eq. (26) yields:

$$\mu_i = \cos \left\{ \sin^{-1} \left(m_\lambda^{-1} \sin(\cos^{-1} \eta_i) \right) \right\} \quad \text{for } i=1 \text{ to } Nb/2 \quad (29)$$

The weight w_i associated to direction i from 1 to $Nb/2$ can be geometrically interpreted as the solid angle $\Delta\Omega_i$ around each direction divided by 2π , i.e.,

$$w_i = \frac{\Delta\Omega_i}{2\pi} = \int_{\theta_{i,min}}^{\theta_{i,max}} \sin\theta \, d\theta = - \int_{\mu_{i,min}}^{\mu_{i,max}} d\mu = \mu_{i,min} - \mu_{i,max} \quad \text{for } i=1 \text{ to } Nb/2 \quad (30)$$

where $\theta_{i,min}$ and $\theta_{i,max}$ are the minimum and maximum polar angles around direction i , respectively. For $i=1$, $\mu_{1,min}=1$ and $w_1=\Delta\Omega_1/2\pi=\Delta\Omega_0/2\pi$. Thus, we can calculate $\mu_{i,min}$, $\mu_{i,max}$, and w_i for values of i between 2 and $Nb/2$ thanks to Eqs. (29) and (30).

Due to the refraction mismatch between the sample and the surrounding medium, these $Nb/2$ directions are confined under a critical angle defined by:

$$\mu_{\frac{Nb}{2},max} = \cos \left\{ \sin^{-1} (1/m_\lambda) \right\} \quad (31)$$

Then, the $Mb/2-Nb/2$ remaining directions between $\mu_{Nb/2,max}$ and $\mu_{Nb/2}=0$ can be defined by using the standard quadrature rules. However, since the intensity variation outside the critical angle is quasi-linear, a regular discretization is sufficient.

Therefore, the remaining direction cosines can be expressed as:

$$\begin{cases} \mu_{Nb/2+1} = \cos(\theta_{Nb/2,max} + \Delta\theta/2) \\ \mu_i = \cos(\theta_{i-1} + \Delta\theta) \end{cases} \quad \text{for } i = Nb/2+2 \text{ to } Mb/2 \quad (32)$$

with $\Delta\theta = \frac{\pi - 2\theta_{Nb/2,max}}{Mb - Nb}$. The associated weights are given by:

$$w_i = \cos(\theta_i - \Delta\theta) - \cos(\theta_i + \Delta\theta) \quad \text{for } i = Nb/2+1 \text{ to } Mb/2 \quad (33)$$

The $Mb/2$ directions in the backward hemisphere and those in the forward hemisphere are symmetrical with respect to sample surface, i.e.,

$$\begin{cases} \mu_i = -\mu_j \\ w_i = w_j \end{cases} \quad \text{for } i = 1 \text{ to } Mb/2 \text{ and } j = Mb/2+1 \text{ to } Mb \quad (34)$$

The theoretical bi-directional transmittance obtained by using the interpolation methods with a Gaussian quadrature and the ACQ are compared. The quadrature order considered are $Mb=30$ and 40 . Two cases of refracting, absorbing, and scattering but non-emitting medium assuming the HG phase function are considered: (1) $m_\lambda=1.41$, $\beta e=\tau_0=1$, $\omega=0.9$, $g=0.75$; and (2) $m_\lambda=1.40$, $\beta e=\tau_0=2$, $\omega=0.3$, $g=0.5$.

Figure 5.a shows the computed transmittances in the experimental directions using three different interpolation laws, namely linear (LL), exponential (EL) and combined exponential-linear laws (ELL), and the ACQ for quadrature orders 30 and 40. Figures 5b and 5c depict the relative differences in directional transmittance obtained for each interpolation law with respect to the ACQ quadrature order 40. One can note that the same conclusions can be drawn for the two media considered. When the added directions ($Mb-Nb$) in the ACQ exceed 6, corresponding to $Mb=30$, the solution of the RTE converges numerically, i.e., it is independent of the number of directions and control volumes, the relative difference in transmittance fall below 1 %. The linear interpolation (LL) with quadrature order $Mb=30$ overestimates significantly the intensities near the incident direction, the errors exceed 1000 % and decrease to 100 % for a quadrature order $Mb=40$. The exponential interpolation (EL) with quadrature

order Mb=30 gives errors reaching 100 % in directions far from the incident direction and from the backward direction. These errors decrease down to 20 % for quadrature order Mb=40. The combined exponential-linear law (ELL) gives better results. The maximum deviation appears only in the directions around 90 ° and does not exceed 10 % for quadrature order Mb=40. The results for each interpolation laws can be improved by increasing the quadrature order but this approach increases the CPU time and is not convenient for the inverse method. Thus, the ACQ quadrature order Mb=30 is adopted in the present study.

V. Results

A. Sample characteristics

Three samples of different thickness ($e=5, 6$ and 9.9 mm) are studied; all of them feature an average void fraction of 4 % and an average bubble radius a equal to 0.64 mm. As one can see in Fig. 1, the bubbles are spherical in shape and randomly distributed. The sample thickness e and the fused quartz refractive index m_λ are used as input data in the identification process. Different correlations for m_λ have been suggested in the literature for different spectral regions.³⁸⁻⁴¹ The most widely accepted is the Malitson's correlation which is valid over the spectral range from 0.21 to 3.71 μm at 20 °C:

$$m_\lambda^2 = 1 + \frac{0.696\lambda^2}{\lambda^2 - (0.068)^2} + \frac{0.408\lambda^2}{\lambda^2 - (0.116)^2} + \frac{0.897\lambda^2}{\lambda^2 - (9.896)^2} \quad (35)$$

The validity of Eq. (35) was also confirmed by Tan⁴¹ up to 6.7 μm . Therefore, due to its wide range of validity at room temperature, Eq. (35) is used in the present study. The identification of parameters has been performed for more than 100 different wavelengths in the spectral region from 1.67 to 4.04 μm .

B. Sensitivity Coefficients

In order to investigate the influence of each measurement direction on the inverse method, the sensitivity coefficients of the theoretical model based on the TPF phase function [Eq. (14)] are investigated. For illustration purposes, we consider three cases of semitransparent media with Fresnel interfaces characterized by: (1) for $\lambda=1.89$ μm , $m_\lambda=1.44$, $\beta e=\tau_0=0.5$, $\omega=0.90$, $f_1=0.22$, $g_1=0.98$, and $g_2=0.50$; (2) for $\lambda=3.20$ μm , $m_\lambda=1.40$, $\beta e=\tau_0=1.0$, $\omega=0.70$, $f_1=0.21$, $g_1=0.98$, and $g_2=0.45$; and (3) for $\lambda=3.96$ μm , $m_\lambda=1.39$, $\beta e=\tau_0=2.5$, $\omega=0.35$, $f_1=0.17$, $g_1=0.96$, and $g_2=0.35$. The variations of the absolute dimensionless sensitivity coefficients defined as $\Pi = p_l/T_{ei}(\partial T_{ii}/\partial p_l)$ for $l=\beta, \omega, f_1, g_1$, and g_2 and $i=1$ to N_b , are depicted in Figs. 6a to 6c versus the measurement angle α . The sensitivity of the theoretical

bi-directional transmittance and reflectance to β increases when the optical thickness increases particularly in the incident direction. The sensitivity to ω tends to 0 near the incident direction and increases with the measurement angle. This sensitivity decreases slightly as the optical thickness increases but remains significant and close to unity. For optically thin medium, the sensitivity to ω has similar trend to that of β . As far as the parameter f_1 is concerned, the model is sensitive only to directions near the forward and backward directions. The parameter g_2 has the lowest sensitivity for optically thin medium. On the contrary, the sensitivity coefficient is the largest for the parameter g_1 , especially around the forward and backward directions.

It is clear that the first direction is essential to determine the extinction coefficient of media with moderate optical thickness ($\tau_0=1$ and 2.5). In addition, measurements in the scattering directions are required to identify ω and β for $\tau_0=0.5$. In the case of optically thin media ($\tau_0 < 1$), ω and β may be linearly dependent resulting in a high condition number i.e., an ill-conditioned system, such that their simultaneous estimation appears difficult. Some directions near the forward ($i=2$ to 6) and backward directions ($i=Nb-4$ to Nb) are certainly sufficient to determine the parameters f_1 and g_1 . As for ω , the scattering directions are required to identify the parameter g_2 .

C. Influence of the number of measurements and the experimental uncertainties

To investigate the influence of the number of measurements on the parameters identification, a parametric study was performed by considering different combinations of forward and backward measurements such as 12/12, 12/5, 7/12, 9/7, and 9/9. The first number refers to the number of forward measurements counted from the incident direction ($\eta=\mu_0=1$) and the second one corresponds to the number of backward measurements counted from the specular direction ($\eta=-1$). The relaxation factor γ used in Eq. (5) is chosen equal to 0.5. Instead of using experimental measurements where the exact solutions are unknown, we used simulated measurements based on the solutions of Eq. (20) by using the TPF model and the radiative characteristics obtained from the identification results (Section V.G): (1) $\lambda=1.89 \mu\text{m}$, $m_\lambda=1.44$, $\beta e=\tau_0=0.5$, $\omega=0.90$, $f_1=0.22$, $g_1=0.98$, and $g_2=0.50$; (2) $\lambda=3.20 \mu\text{m}$, $m_\lambda=1.40$, $\beta e=\tau_0=1.0$, $\omega=0.70$, $f_1=0.21$, $g_1=0.98$, and $g_2=0.45$; and (3) $\lambda=3.96 \mu\text{m}$, $m_\lambda=1.39$, $\beta e=\tau_0=2.5$, $\omega=0.35$, $f_1=0.17$, $g_1=0.96$, and $g_2=0.35$. To take into account the experimental uncertainties, the simulated measurements (T_i) are corrupted by adding a normally distributed random errors:⁴²

$$T_{ei}=T_{ti}+\xi\sigma_i \quad \text{for } i=1 \text{ to } Nb \quad (36)$$

where $0 < \xi < 1$ is a normally distributed random number and σ_i is the standard deviation for an experimental error χ_i corresponding to an confidence interval of 99 %. The standard deviation σ_i for direction i is expressed as:

$$\sigma_i = \frac{\chi_i \bar{T}_i}{2.58} \quad (37)$$

where \bar{T}_i is the average transmittance defined as:

$$\left\{ \begin{array}{ll} \bar{T}_i = T_{ti} & \text{for } i=1, 2, Nb-1, \text{ and } Nb \\ \bar{T}_i = \frac{1}{10} \sum_{j=3}^{Nb/2} T_{tj}, & \text{for } i=3, Nb/2 \\ \bar{T}_i = \frac{1}{10} \sum_{Nb/2+1}^{Nb-2} T_{tj}, & \text{for } i=Nb/2+1, Nb-2 \end{array} \right. \quad (38)$$

The relative error χ_i associated with the direction i is chosen to be equal to the largest experimental error whose origins have been discussed in section III.B. After several measurements, they are estimated as:

$$\left\{ \begin{array}{ll} \chi_1 = 10 \% \\ \chi_2 = 20 \% \\ \chi_i = 10 \% & \text{for } i=3, Nb-2 \\ \chi_{Nb-1} = 20 \% \\ \chi_{Nb} = 40 \% \end{array} \right. \quad (39)$$

The exact and corrupted bi-directional transmittances are shown in Figs. 7a to 7c as a function of the measurement angle α . As one can see, the largest absolute errors correspond to the directions far from the direction of the incident radiation.

Moreover, two identification approaches are tested: (A1) approach A1 consists of direct computation of β from Eq. (18) and simultaneous estimation of ω , f_1 , g_1 , and g_2 parameters; (A2) approach A2 consists of estimating simultaneously β , ω , f_1 , g_1 , and g_2 . The initial values of these parameters are taken identical in both cases. The difference between the identified parameters and the exact solutions, the number of iterations (IN), and the condition number (CN) are summarized in Tables 1a to 1c. The following conclusions can be drawn:

1) The parameters ω , f_1 , and g_2 are the most sensitive to the experimental uncertainties. Their deviations from the exact solutions increase as the optical thickness increases. On the contrary, the parameter g_1 is not influenced by the experimental uncertainty and is in good agreement with the exact solution (the deviation is less than 4 % in all cases).

2) When an insufficient number of measurements is used such as 12/5, 7/12, and/or 9/7, either the inverse process does not converge or it converges toward a wrong solution (the errors can reach 300 %). However, if too many noisy measurements are used such as 12/12, the computation leads to large errors that may reach up to 20 % and 60 % depending on the optical thickness. In our case, the 9/9 combination gives a good compromise between convergence and accuracy for every wavelength, and will be used in the identification using experimental data.

3) The condition number obtained by simultaneous estimation of all parameters (approach A2), is very large (10^{+6} to 10^{+13}) for all the optical thicknesses studied. When the CN is greater than 10^{+10} , the inversion procedure converges toward erroneous solutions. On the other hand, the independent computation of the extinction coefficient (approach A1) leads to an acceptable condition number ($\sim 10^{+3}$).

4) The directly computed extinction coefficient (approach A1) is less precise than that retrieved from the approach A2 for small optical thickness ($\tau_0=0.5$). This deviation is due to the approximation of the intensity variation as a third order polynomial function.

D. Influence of the parameters' initial values

The parameters identification is performed by using the simulated experimental transmittance and reflectance described in section V.C with 9 forward and 9 backward measurements. To investigate the effect of the parameters' initial value on the identification results three combinations of initial parameters are considered as reported in Tables 2a to 2c. In addition, the two identification approaches A1 and A2 previously described are compared. The relative difference between the computed and exact parameters, the number of iterations (IN), and the condition number (CN) are summarized in Tables 2a to 2c. One can note that the initial guesses for the unknown parameters do not affect the results of approach A1 but only the number of iterations, i.e., the CPU time. However, the approach A2 is influenced by the parameters' initial values especially for absorbing materials ($\tau_0=2.5$). Therefore, Approach A1 is more stable than A2 with respect to the initial guesses for the radiation characteristics to be identified.

E. The two step inverse process

The analysis in Sections V.C and V.D show that 9 forward and 9 backward measurements are sufficient for the current identification of parameters. The approach A1 using the direct computation of β is more robust than the simultaneous parameter estimation (approach A2). It avoids dealing with a very ill-conditioned system which reduces the efficiency of the method. However, it is less precise for small optical thickness. The approach A2 is always characterized by a large CN and requires a better knowledge of the initial parameters.

Moreover, one can note that these two identification schemes are complementary. Thus, this suggests performing two successive inversion procedures to optimize the final results. First, a preliminary calculation using approach A1 is carried out which provides an approximate value of the unknown parameters. Then, a second identification step based on approach A2 is performed using the results from approach A1 as initial guesses. This second step can be done without estimating g_1 since it can be determined precisely from the first step (see section V.D). By using this two step inverse procedure, there are no restrictions either on the parameters' initial value or on the optical thickness range. This two step inversion scheme is applied in the present study.

F. Influence of the phase function model

To investigate the effects of the scattering phase function model, the radiative parameters are identified by using measurements corresponding to the 6 mm thick sample. The two phase function models considered are the HG and the TPF. The identified parameters are summarized in Table 3 for typical wavelengths. The absorption coefficient of the fused quartz $k_{\text{bulk}} = 4\pi\kappa_{\text{bulk}}/\lambda$ is also reported on the same table where the absorption index κ_{bulk} is taken from Dombrovsky et al.²³ The resulting bi-directional transmittances are shown in Figs. 8a and 8b.

Table 3 indicates that the phase function model has significant influence on the retrieved radiative characteristics especially in the spectral region where fused quartz is transparent (from 1.67 to 2.7 μm and 2.9 to 3.5 μm in this study). As for the bi-directional transmittance and reflectance reported in Fig. 8, one can note that the theoretical results obtained using the TPF function gives good agreement between the measured and retrieved bi-directional transmittance and reflectance while results from the HG function agrees only for the 5 forward ($i=1$ to 5) and 3 backward directions ($i=N_b-4$ to N_b-1). Overall, the HG function fails to properly describe the directional scattering behavior of the studied material.

G. Identified radiative characteristics

The radiative characteristics (Figs. 9a to 9f) are determined from the three samples of different thickness previously described. For comparison, the bulk quartz radiative characteristic and the average results obtained by Baillis et al.¹⁸ using the HG phase function are also reported.

First, one can conclude that the retrieved parameters are independent of samples thickness. The observed dispersion of data is mainly attributed to the measurement uncertainties. The standard deviation is especially large for the parameter g_2 reaching about 12 %.

Overall, the retrieved radiation characteristics tend to disagree with previously reported data¹⁸ except for the extinction coefficient which can be estimated properly using the HG function. This is mainly due to the phase function model adopted in the theoretical formulations since it was shown to have little effect on the extinction coefficient but a significant one on the absorption and scattering coefficients (see section V.F). However, in all cases, results using the HG model gives a small scattering albedo associated with a highly forward anisotropic phase function and an absorption coefficient 8 to 10 times higher than that of bulk quartz (k_{bulk}). On the contrary, the retrieved absorption coefficient using the TPF model is of the same order of magnitude as that of bulk quartz.

Moreover, the scattering behavior of the porous quartz is generally forward anisotropic with g equal to about 0.78, its absorption coefficient is slightly smaller than that of the bulk material, and the scattering dominates the extinction ($\omega \sim 0.8-0.9$) in the transparency bands of fused quartz. These mean that the bubbles are non-absorbing but scatter only radiation and the quartz matrix is the only absorbing substance.

H. Comparison with hemispherical measurements and influences of experimental uncertainties

To verify that the retrieved parameters represent accurately the radiative characteristics of the material studied, the calculated hemispherical transmittance termed T_t^+ and reflectance termed T_t^- based on (1) the current identified parameters and (2) the results of Baillis et al.¹⁸ are compared with those measured experimentally and denoted T_e^+ and T_e^- . The Fourier-Transform InfraRed spectrometer is used in combination with a gold coated integrating sphere (CSTM RSA-DI-40D) to measure the spectral hemispherical transmittance T_e^+ and reflectance T_e^- . The experimental errors are evaluated for each sample from five different measurements. Depending on the wavelength, errors range from 3 to 8 % for transmittance and from 9 to 16 % for reflectance. The average radiative characteristics are introduced in the RTE [Eq. (20)] to compute the hemispherical transmittance and reflectance (after integration of the bi-directional transmittance over each hemisphere). The computational errors are evaluated as follows: first, Eq. (20) is solved for each sample thickness by using the associated radiative parameters, then the standard deviation are determined from the computed hemispherical transmittances and reflectances of different thicknesses. The comparisons are reported in Figs. 10a to 10c. Overall, good agreement is observed between the measured hemispherical transmittances and reflectances, and their values computed using the radiative characteristics retrieved with the TPF function. On the other hand, the numerical results obtained using the radiative parameters reported by Baillis et al.¹⁸ are always smaller than the hemispherical transmittance T_e^+ and reflectance

T_e^- measurements. This can be attributed to the underestimation of the scattering albedo and overestimation of the absorption coefficient obtained when the HG phase function is adopted.

Finally, the comparison of numerical results with hemispherical measurements not only enables one to quantify the experimental errors but also offers an efficient validation tool.

VI. Conclusions

Recently, the experimental determination of radiative characteristics of fused quartz containing bubbles was performed using an inverse method based on bi-directional transmittance measurements. The Henyey-Greenstein phase function was used in the theoretical model based on the RTE. Due to refraction at both interfaces of the slab, the theoretical transmittance and reflectance in the measurement directions were evaluated by using an interpolation law. In the present study, several improvements are proposed particularly for (i) the scattering phase function model, (ii) the identification procedure, and (iii) the quadrature used in the theoretical calculations. From the above discussion, the following conclusions can be drawn:

- 1) The importance of the phase function model on the inverse method based on bi-directional transmittance measurements has been demonstrated. The use of a common scattering phase function such as the Henyey-Greenstein function underestimates the scattering coefficient and overestimates the absorption coefficient while it properly estimates the extinction coefficient. The present study proposed a more elaborated phase function, the so-called Truncated Phase Function (TPF), which depends on three parameters and enables one to take into account the complex scattering behavior of the samples. The hemispherical transmittance and reflectance computed using these newly retrieved coefficients gives better agreement with the experimental measurements than those obtained using the previously reported radiative characteristics.
- 2) The interpolation methods used to evaluate the theoretical transmittance in the measurement directions were shown to be less accurate than the proposed Adaptive Composite Quadrature (ACQ) unless a high quadrature order was used. The use of an adaptive quadrature has been proposed and found to be advantageous in terms of both computational time and precision.
- 3) The importance of the choice of the measurement directions on the identification results has been highlighted. It is recommended to perform the sensitivity coefficients analysis for similar study using bi-directional transmittance and reflectance measurements.

- 4) A new inverse method based on a two step inversion procedure is proposed. It uses a preliminary parameters estimation step. This technique enables one (i) to avoid the errors induced by the direct computation of the extinction coefficient from the collimated transmittance and (ii) to accelerate the convergence.
- 5) The radiative characteristics of the porous fused quartz were then identified and shown to be independent of the sample thickness. Unlike the results obtained using the Henyey-Greenstein function, the absorption coefficient of porous samples of porosity equal to 4 % obtained using the TPF phase function is slightly smaller than that of the dense matrix. Compared to the previously published data, the current radiative characteristics appear to be in better agreement with the physical intuition.
- 6) The limitations of the experimental setup to measure the bi-directional measurements have been pointed out. In particular, the influence of the measurement number, the measurement noises, and the alignment uncertainties on the results of the inverse method have been observed. In order to obtain more reliable measurements, a more sensitive detector is required to improve accuracy.
- 7) Finally, the same experimental methodology can be used for other semitransparent materials.

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Table captions

Table 1 Influence of measurement number on the identified parameters for different wavelengths.

Table 2 Influence of the parameters initial values on the identification results for different wavelengths.

Table 3 Influence of the phase function model on the radiative characteristics for 6 mm sample thickness at different wavelengths.

Table 1a Influence of bi-directional measurement number (MN) on the identified parameters for $\lambda=1.89 \mu\text{m}$, $m_\lambda=1.44$ with the exact parameters: $\beta_{\text{exact}}=50 \text{ m}^{-1}$, $\omega_{\text{exact}}=0.9$, $f_{1\text{exact}}=0.22$, $g_{1\text{exact}}=0.98$, and $g_{2\text{exact}}=0.5$ and the initial parameters $\beta_0=70 \text{ m}^{-1}$, $\omega_0=0.5$, $f_{10}=0.5$, $g_{10}=0.7$, and $g_{20}=0.2$.

MN	$\Delta\beta/\beta_{\text{exact}}, \%$		$\Delta\omega/\omega_{\text{exact}}, \%$		$\Delta f_1/f_{\text{exact}}, \%$		$\Delta g_1/g_{1\text{exact}}, \%$		$\Delta g_2/g_{2\text{exact}}, \%$		IN		CN	
	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1
12/12	4.52	13.92	-9.09	-10.41	5.12	5.56	-0.32	-0.16	-5.29	-7.72	24	15	$3.1 \cdot 10^7$	$1.1 \cdot 10^3$
12/5	-2.58	13.92	-7.02	-8.88	-0.47	2.48	-0.46	-0.10	-4.76	-7.27	27	15	$9.3 \cdot 10^7$	$1.2 \cdot 10^3$
7/12	-	13.92	-	-10.56	-	6.83	-	-0.12	-	-12.53	-	35	-	$1.5 \cdot 10^3$
9/7	-	13.92	-	-8.32	-	4.66	-	0.07	-	0.37	-	17	-	$1.4 \cdot 10^3$
9/9	2.12	13.92	-6.63	-8.68	3.42	7.57	-0.21	0.15	-2.44	2.38	23	20	$2.2 \cdot 10^7$	$1.0 \cdot 10^3$

Table 1b Influence of bi-directional measurement number (MN) on the identified parameters: for $\lambda=3.20 \mu\text{m}$, $m_\lambda=1.41$ with the exact parameters: $\beta_{\text{exact}}=100 \text{ m}^{-1}$, $\omega_{\text{exact}}=0.7$, $f_{1\text{exact}}=0.21$, $g_{1\text{exact}}=0.98$, $g_{2\text{exact}}=0.45$ and the initial parameters $\beta_0=70 \text{ m}^{-1}$, $\omega_0=0.5$, $f_{10}=0.5$, $g_{10}=0.7$, and $g_{20}=0.2$.

MN	$\Delta\beta/\beta_{\text{exact}}, \%$		$\Delta\omega/\omega_{\text{exact}}, \%$		$\Delta f_1/f_{\text{exact}}, \%$		$\Delta g_1/g_{1\text{exact}}, \%$		$\Delta g_2/g_{2\text{exact}}, \%$		IN		CN	
	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1
12/12	1.47	1.98	-13.78	-13.80	21.83	22.19	0.27	0.29	22.32	22.38	25	16	$7.3 \cdot 10^6$	$7.2 \cdot 10^2$
12/5	-132.89	1.98	-30.15	-13.24	-61.10	19.51	-1.50	0.22	-7.23	21.24	31	20	$2.8 \cdot 10^{11}$	$1.0 \cdot 10^3$
7/12	1.41	-	-16.45	-	21.14	-	0.10	-	-0.75	-	34	-	$1.1 \cdot 10^7$	-
9/7	2.61	1.98	-10.57	-10.67	14.92	14.59	0.25	0.20	14.76	14.48	21	16	$7.4 \cdot 10^6$	$9.5 \cdot 10^2$
9/9	2.19	1.98	-11.92	-11.91	19.14	19.21	0.36	0.35	19.04	19.01	24	14	$5.9 \cdot 10^6$	$8.0 \cdot 10^2$

Table 1c Influence of bi-directional measurement number (MN) on the identified parameters: for $\lambda=3.96 \mu\text{m}$, $m_\lambda=1.39$ with the exact parameters: $\beta_{\text{exact}}=250 \text{ m}^{-1}$, $\omega_{\text{exact}}=0.35$, $f_{1\text{exact}}=0.17$, $g_{1\text{exact}}=0.96$, $g_{2\text{exact}}=0.35$ and the initial parameters $\beta_0=70 \text{ m}^{-1}$, $\omega_0=0.5$, $f_{10}=0.5$, $g_{10}=0.7$, and $g_{20}=0.2$.

MN	$\Delta\beta/\beta_{\text{exact}}, \%$		$\Delta\omega/\omega_{\text{exact}}, \%$		$\Delta f_1/f_{\text{exact}}, \%$		$\Delta g_1/g_{1\text{exact}}, \%$		$\Delta g_2/g_{2\text{exact}}, \%$		IN		CN	
	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1
12/12	1.21	1.27	-35.41	-35.33	40.25	40.23	-0.30	-0.26	59.60	59.62	24	14	$1.8 \cdot 10^6$	$1.6 \cdot 10^2$
12/5	-318.91	1.27	-149.57	-32.84	-206.36	36.32	-3.93	-0.36	-75.51	50.59	31	15	$2.8 \cdot 10^{13}$	$1.8 \cdot 10^2$
7/12	1.10	1.27	-39.78	58.47	39.50	-102.13	-0.41	-3.46	37.63	-171.85	26	15	$1.6 \cdot 10^6$	$2.5 \cdot 10^5$
9/7	1.40	1.27	-31.94	-32.18	39.18	39.01	-0.14	-0.11	68.78	68.64	38	15	$2.0 \cdot 10^6$	$4.4 \cdot 10^2$
9/9	1.45	1.27	-31.77	-32.04	38.90	38.75	-0.15	-0.12	68.07	68.06	24	15	$2.0 \cdot 10^6$	$4.3 \cdot 10^2$

Table 2a Influence of the parameters initial values (piv) on the identification results for $\lambda=1.89 \mu\text{m}$, $m_\lambda=1.44$ with 9 forward and 9 backward measurements and the exact parameters: $\beta_{\text{exact}}=50 \text{ m}^{-1}$, $\omega_{\text{exact}}=0.90$, $f_{1\text{exact}}=0.22$, $g_{1\text{exact}}=0.98$, and $g_{2\text{exact}}=0.5$. The initial parameters are: piv 1: $\beta_0=70 \text{ m}^{-1}$, $\omega_0=0.5$, $f_{10}=0.5$, $g_{10}=0.7$, and $g_{20}=0.2$; piv 2: $\beta_0=500 \text{ m}^{-1}$, $\omega_0=0.1$, $f_{10}=0.8$, $g_{10}=0.2$, and $g_{20}=0.1$; and piv 3: $\beta_0=100 \text{ m}^{-1}$, $\omega_0=0.8$, $f_{10}=0.1$, $g_{10}=0.7$, and $g_{20}=0.2$.

piv	$\Delta\beta/\beta_{\text{exact}}, \%$		$\Delta\omega/\omega_{\text{exact}}, \%$		$\Delta f_1/f_{\text{exact}}, \%$		$\Delta g_1/g_{1\text{exact}}, \%$		$\Delta g_2/g_{2\text{exact}}, \%$		IN		CN	
	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1
1	2.12	13.92	-6.63	-8.32	3.42	4.66	-0.21	0.07	-2.44	0.37	23	17	$2.2 \cdot 10^7$	$1.4 \cdot 10^3$
2	1.22	13.92	-7.47	-8.32	7.80	4.65	-0.04	0.07	4.42	0.37	62	32	$1.2 \cdot 10^7$	$1.1 \cdot 10^3$
3	1.18	13.92	-7.47	-8.30	7.78	4.73	-0.04	0.07	4.40	0.49	25	15	$1.2 \cdot 10^7$	$1.2 \cdot 10^3$

Table 2b Influence of the parameters initial values (piv) on the identification results for $\lambda=3.20 \mu\text{m}$, $m_\lambda=1.41$ with 9 forward and 9 backward measurements and the exact parameters: $\beta_{\text{exact}}=100 \text{ m}^{-1}$, $\omega_{\text{exact}}=0.7$, $f_{1\text{exact}}=0.21$, $g_{1\text{exact}}=0.98$, $g_{2\text{exact}}=0.45$. The initial parameters are: piv 1: $\beta_0=70 \text{ m}^{-1}$, $\omega_0=0.5$, $f_{10}=0.5$, $g_{10}=0.7$, and $g_{20}=0.2$; piv 2: $\beta_0=500 \text{ m}^{-1}$, $\omega_0=0.1$, $f_{10}=0.8$, $g_{10}=0.2$, and $g_{20}=0.1$; and piv 3: $\beta_0=100 \text{ m}^{-1}$, $\omega_0=0.8$, $f_{10}=0.1$, $g_{10}=0.7$, and $g_{20}=0.2$.

piv	$\Delta\beta/\beta_{\text{exact}}, \%$		$\Delta\omega/\omega_{\text{exact}}, \%$		$\Delta f_1/f_{\text{exact}}, \%$		$\Delta g_1/g_{1\text{exact}}, \%$		$\Delta g_2/g_{2\text{exact}}, \%$		IN		CN	
	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1
1	2.19	1.98	-11.92	-10.67	19.14	14.59	0.36	0.20	19.04	14.48	24	16	$5.9 \cdot 10^6$	$9.4 \cdot 10^2$
2	0.04	1.52	-15.06	-15.46	10.88	15.75	-0.19	-0.01	12.95	17.00	50	25	$3.3 \cdot 10^7$	$1.4 \cdot 10^3$
3	-0.04	1.52	-15.10	-15.46	10.76	15.81	-0.20	0.00	12.86	16.99	37	16	$3.7 \cdot 10^7$	$1.4 \cdot 10^3$

Table 2c Influence of the parameters initial values (piv) on the identification results for $\lambda=3.96 \mu\text{m}$, $m_\lambda=1.39$ with 9 forward and 9 backward measurements and the exact parameters: $\beta_{\text{exact}}=250 \text{ m}^{-1}$, $\omega_{\text{exact}}=0.35$, $f_{1\text{exact}}=0.17$, $g_{\text{exact}}=0.96$, and $g_{2\text{exact}}=0.35$. The initial parameters are: piv 1: $\beta_0=70 \text{ m}^{-1}$, $\omega_0=0.5$, $f_{10}=0.5$, $g_{10}=0.7$, and $g_{20}=0.2$; piv 2: $\beta_0=500 \text{ m}^{-1}$, $\omega_0=0.1$, $f_{10}=0.8$, $g_{10}=0.2$, and $g_{20}=0.1$; and piv 3: $\beta_0=100 \text{ m}^{-1}$, $\omega_0=0.8$, $f_{10}=0.1$, $g_{10}=0.7$, and $g_{20}=0.2$.

piv	$\Delta\beta/\beta_{\text{exact}}, \%$		$\Delta\omega/\omega_{\text{exact}}, \%$		$\Delta f_1/f_{\text{exact}}, \%$		$\Delta g_1/g_{1\text{exact}}, \%$		$\Delta g_2/g_{2\text{exact}}, \%$		IN		CN	
	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1	a.2	a.1
1	2.19	1.98	-11.92	-10.67	19.14	14.59	0.36	0.20	19.04	14.48	24	16	$5.9 \cdot 10^6$	$9.4 \cdot 10^2$
2	0.04	1.52	-15.06	-15.46	10.88	15.75	-0.19	-0.01	12.95	17.00	50	25	$3.3 \cdot 10^7$	$1.4 \cdot 10^3$
3	-0.04	1.52	-15.10	-15.46	10.76	15.81	-0.20	0.00	12.86	16.99	37	16	$3.7 \cdot 10^7$	$1.4 \cdot 10^3$

Table 3 Influence of the phase function model on the radiative characteristics for 6 mm sample thickness at different wavelengths.

$\lambda, \mu\text{m}$	$k_{\text{bulk}}, \text{m}^{-1}$	$\beta_{\text{HG}}, \text{m}^{-1}$	$\beta_{\text{TPF}}, \text{m}^{-1}$	ω_{HG}	ω_{TPF}	$k_{\text{HG}}, \text{m}^{-1}$	$k_{\text{TPF}}, \text{m}^{-1}$	g_{HG}	g_{TPF}
1.89	5.94	75.2	78.23	0.47	0.94	39.85	4.06	0.95	0.70
2.76	151.61	170.74	176.48	0.17	0.27	141.71	128.28	0.93	0.84
3.96	204.04	262.21	264.18	0.14	0.23	225.50	201.97	0.94	0.81

Figure captions

Fig. 1 Photo of the studied samples.

Fig. 2 Mie scattering function of an optically large bubble of size parameter $x=2000$ and TPF function characterized by $f_1=0.2$, $g_1=0.98$, and $g_2=0.45$.

Fig. 3 Measurement and ACQ directions

Fig. 4 Index of absorption of fused quartz from literature.

Fig. 5a Comparison between transmittances from the interpolation laws (linear law (LL), exponential law (EL), and exponential linear law (ELL)) and the ACQ.

Fig. 5b Relative transmittance deviation of the different interpolation laws (linear law (LL), exponential law (EL), and exponential linear law (ELL)) compared to the ACQ for $\lambda=3.20 \mu\text{m}$.

Fig. 5c Relative transmittance deviation of the different interpolation laws (linear law (LL), exponential law (EL), and exponential linear law (ELL)) compared to the ACQ for $\lambda=2.74 \mu\text{m}$.

Fig. 6a Dimensionless sensitivity coefficients for $\lambda=1.89 \mu\text{m}$.

Fig. 6b Dimensionless sensitivity coefficients for $\lambda=3.20 \mu\text{m}$.

Fig. 6c Dimensionless sensitivity coefficients for $\lambda=3.96 \mu\text{m}$.

Fig. 7a Simulated exact and corrupted bi-directional transmittances for $\lambda=1.89 \mu\text{m}$.

Fig. 7b Simulated exact and corrupted bi-directional transmittances for $\lambda=3.20 \mu\text{m}$.

Fig. 7c Simulated exact and corrupted bi-directional transmittances for $\lambda=3.96 \mu\text{m}$.

Fig. 8a Influences of the phase function model on the transmittances for the sample $e=6 \text{ mm}$ at $\lambda=1.89 \mu\text{m}$.

Fig. 8b Influences of the phase function model on the transmittances for the sample $e=6 \text{ mm}$ at $\lambda=3.96 \mu\text{m}$.

Fig. 9a Identified extinction (β) and absorption (k) coefficients.

Fig. 9b Identified scattering albedo (ω).

Fig. 9c Identified scattering parameter f_1 .

Fig. 9d Identified scattering parameter g_1 .

Fig. 9e Identified scattering parameter g_2 .

Fig. 9f corresponding asymmetry factor ($\langle g \rangle$).

Fig. 10a Comparison between the computed and measured hemispherical transmittance and reflectance for the sample $e=5$ mm.

Fig. 10b Comparison between the computed and measured hemispherical transmittance and reflectance for the sample $e=6$ mm.

Fig. 10c Comparison between the computed and measured hemispherical transmittance and reflectance for the sample $e=9.9$ mm.

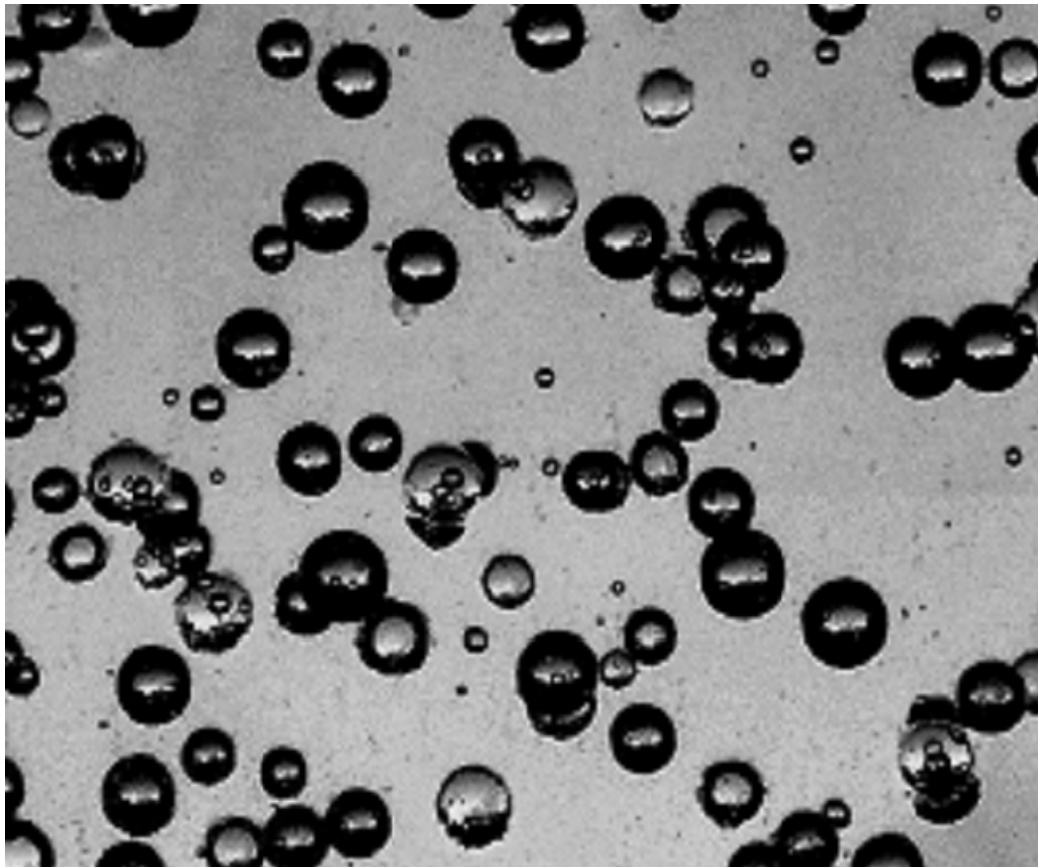


Fig. 1, Randrianalisoa et al.

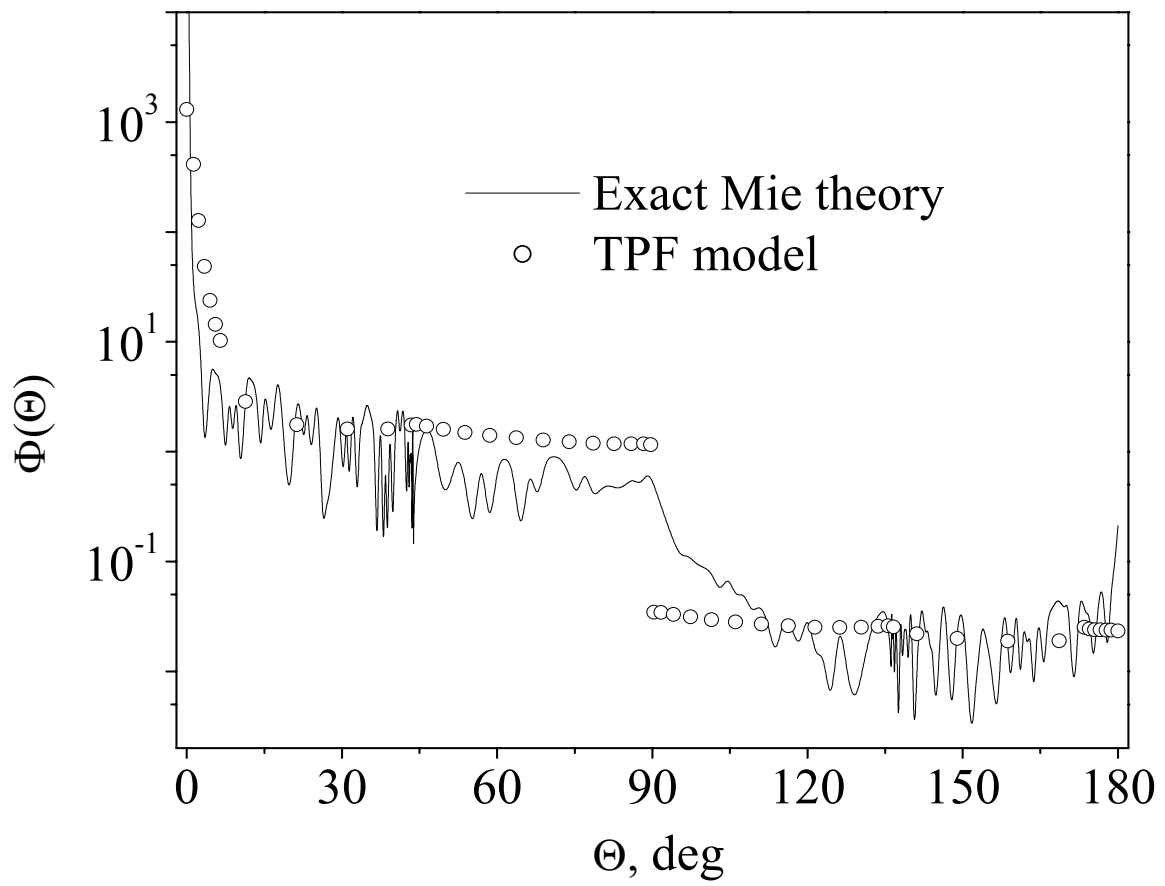


Fig. 2, Randrianalisoa et al.

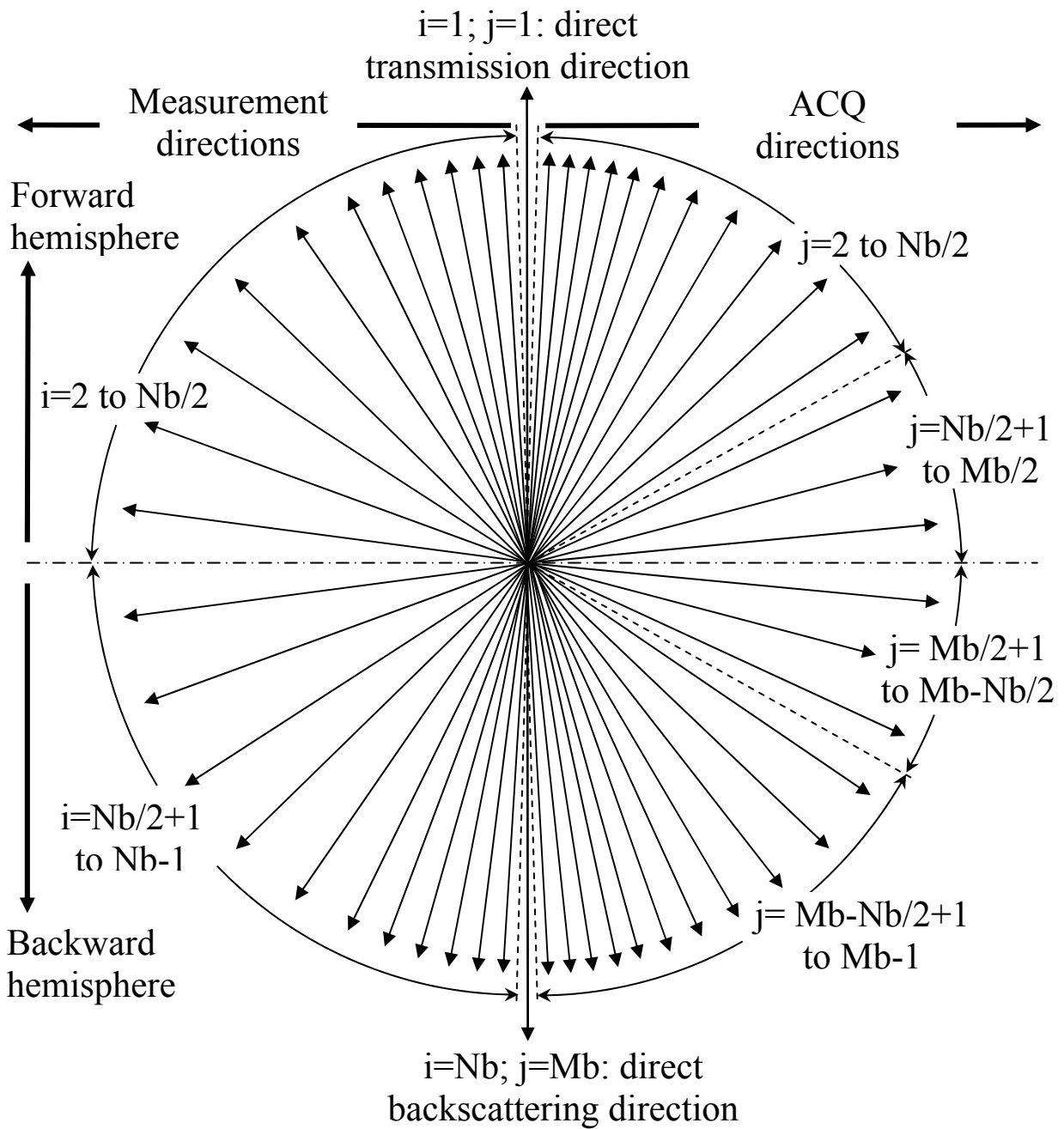


Fig. 3, Randrianalisoa et al.

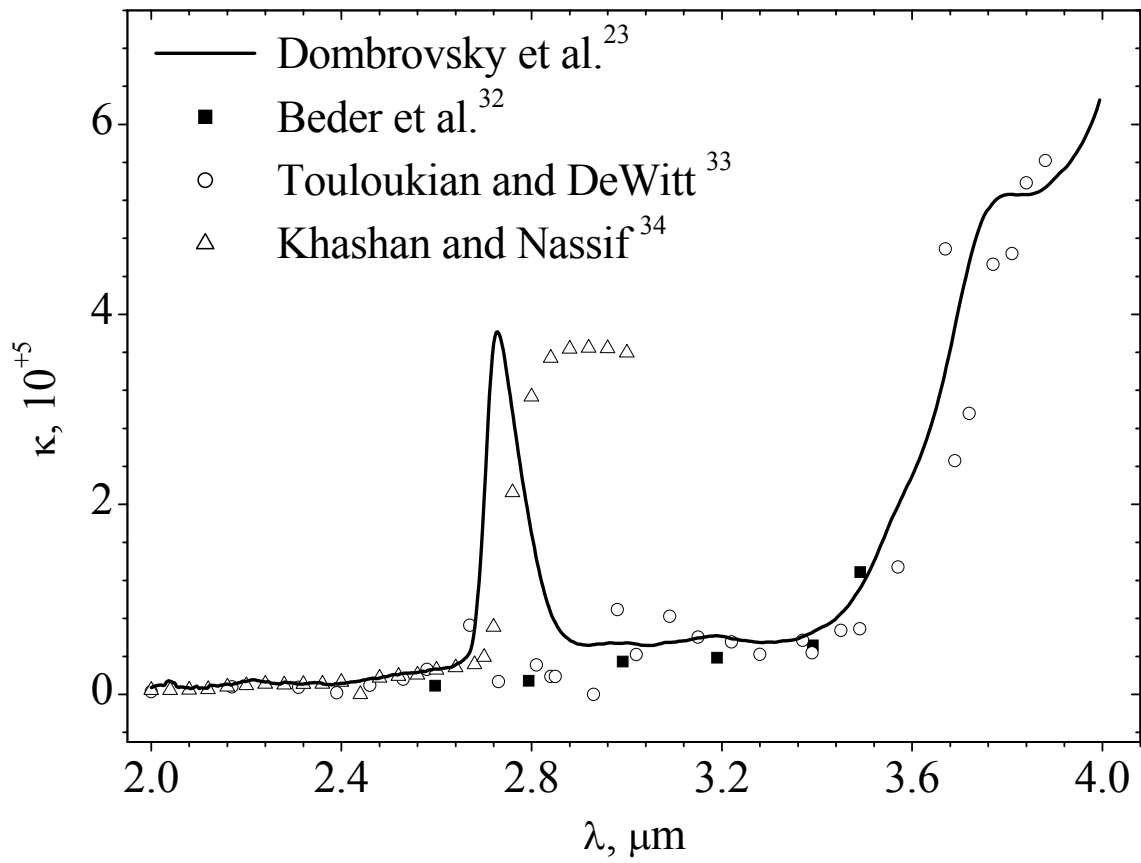


Fig. 4, Randrianalisoa et al

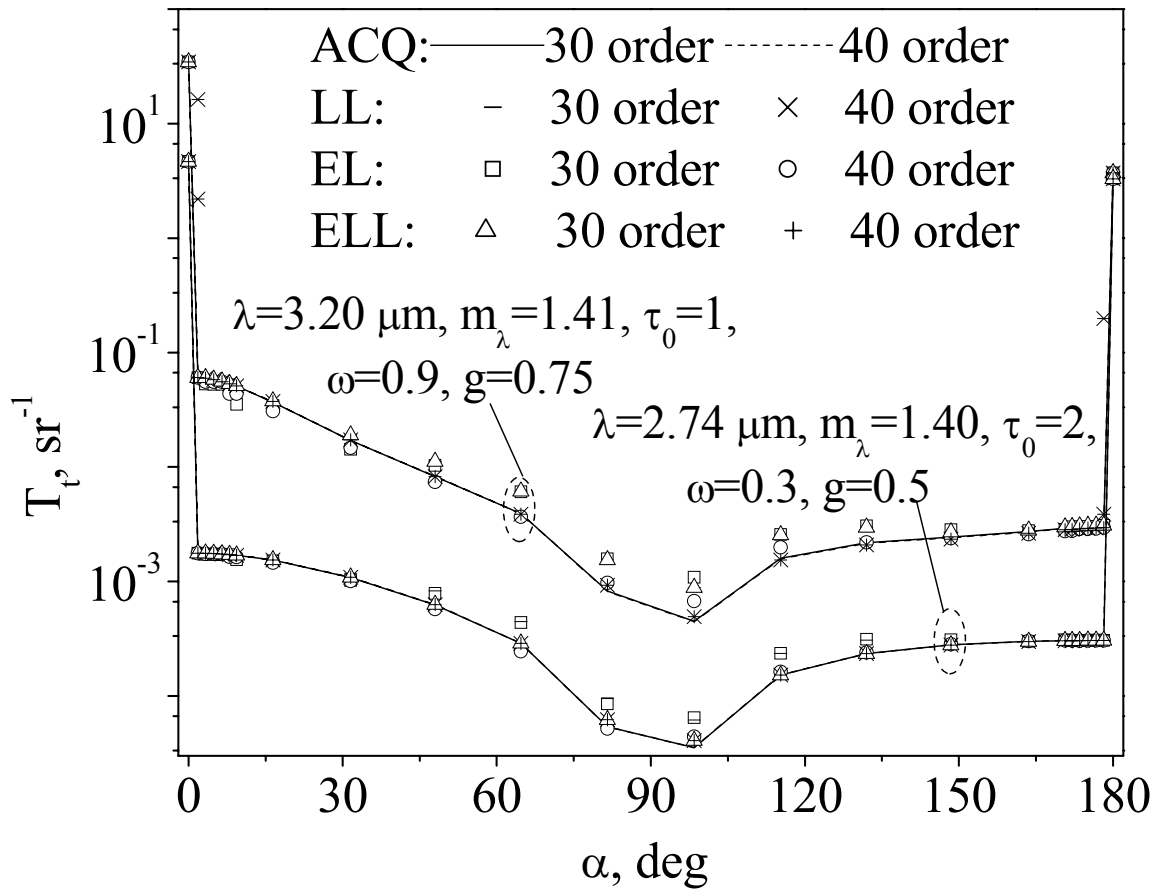


Fig. 5a, Randrianalisoa et al.

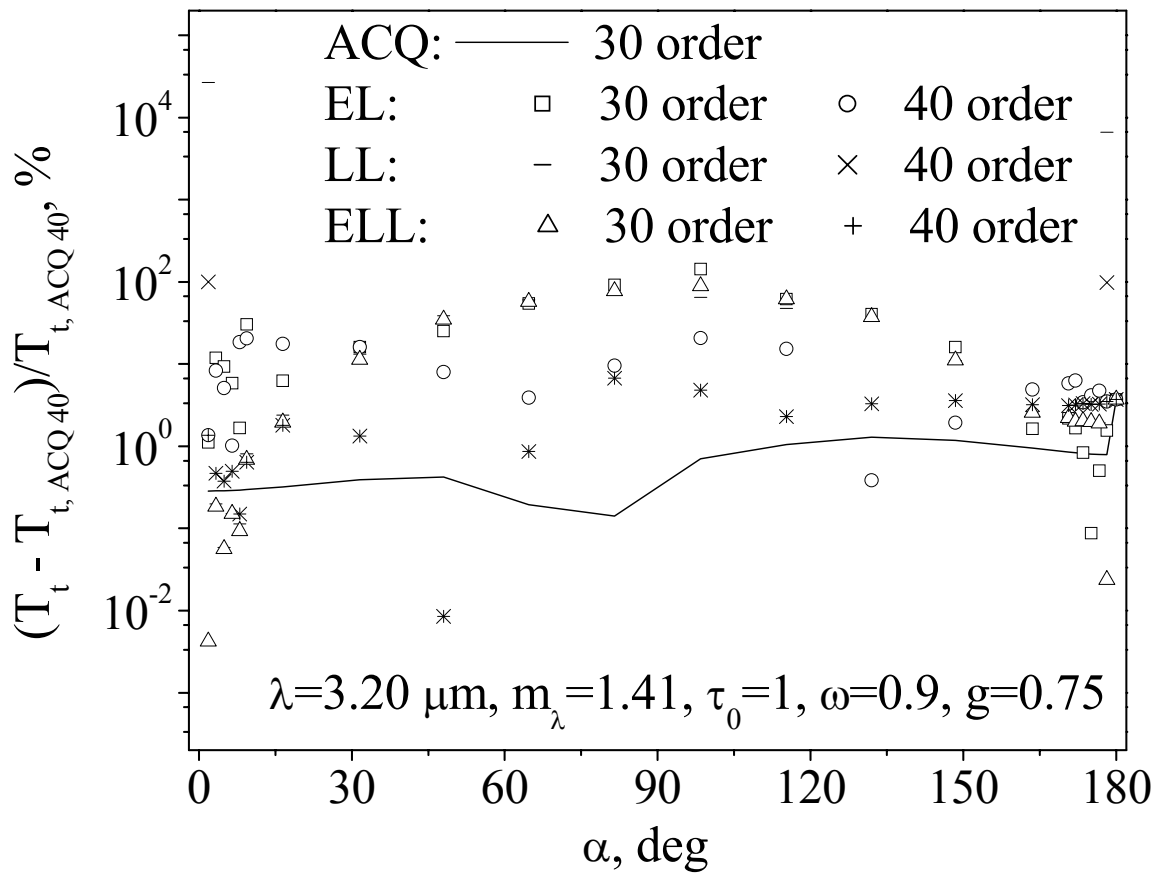


Fig. 5b, Randrianalisoa et al.

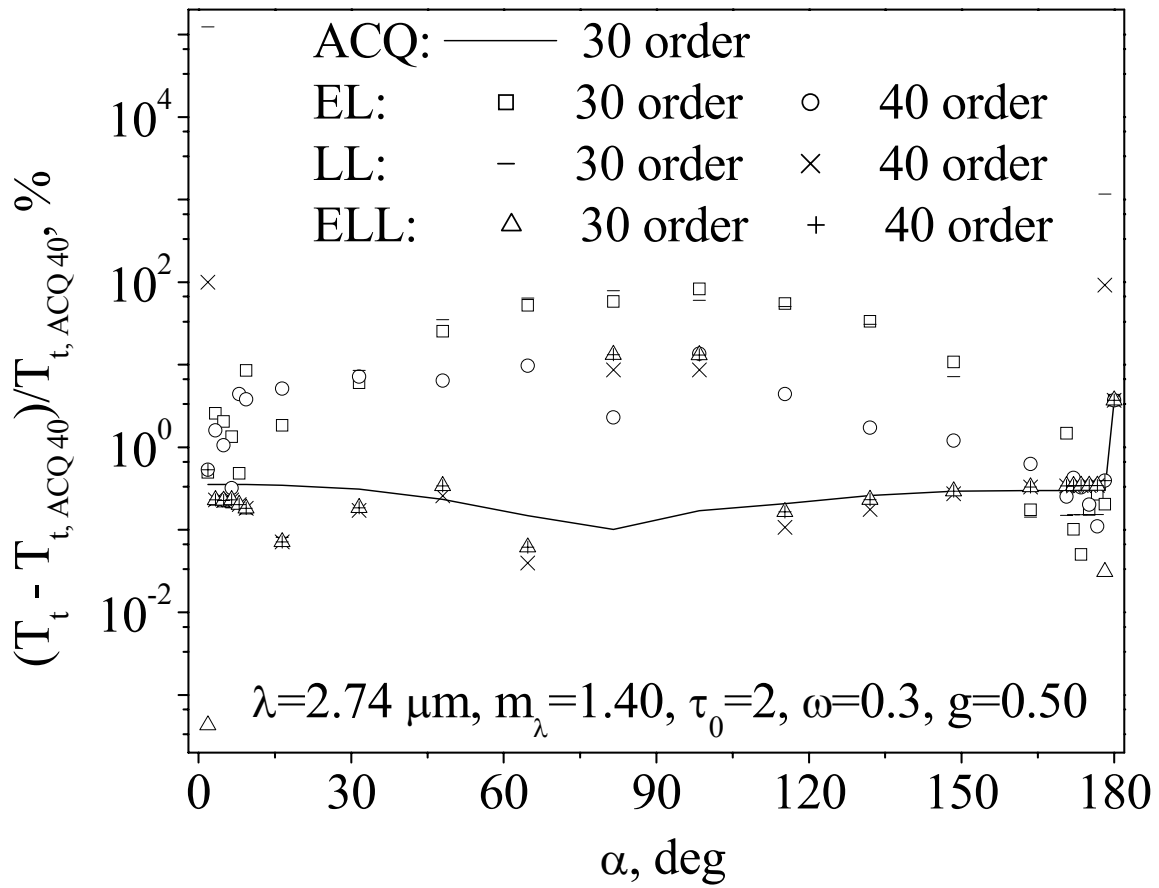


Fig. 5c, Randrianalisoa et al.

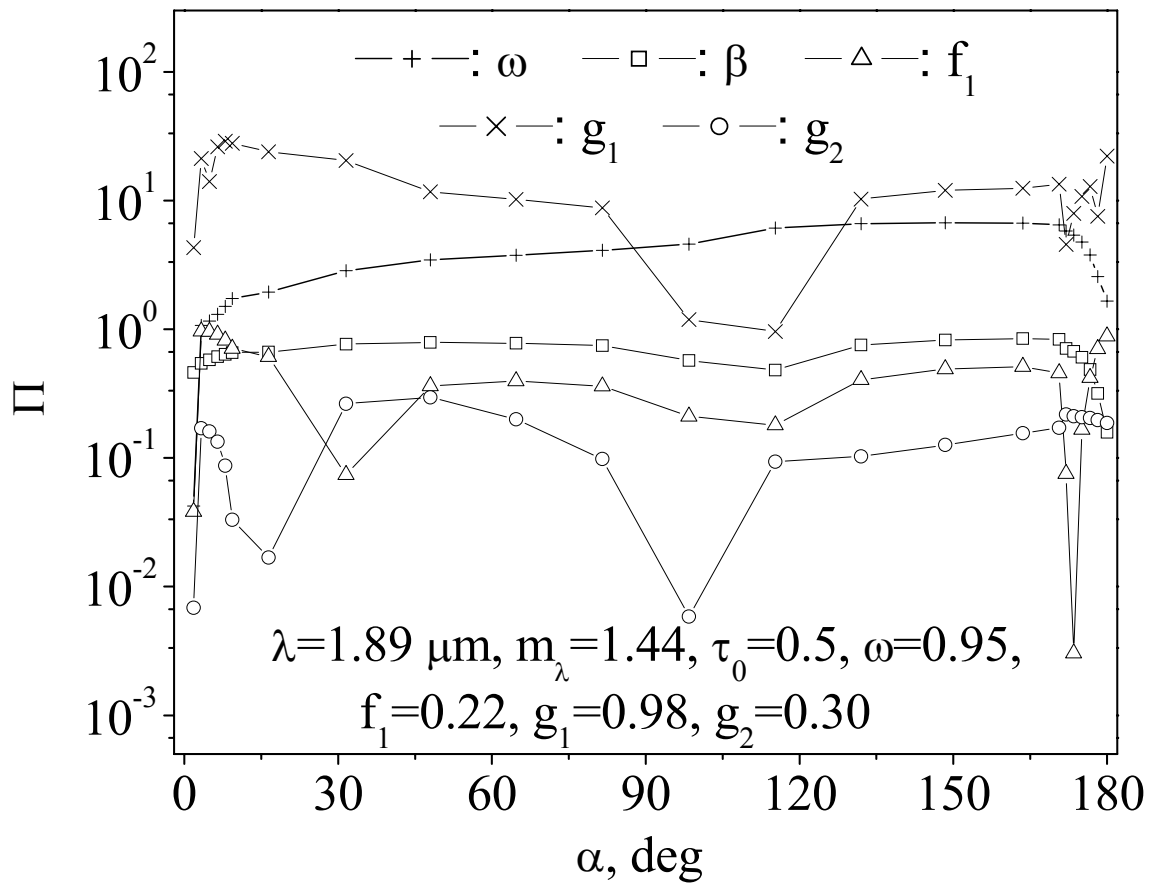


Fig. 6a, Randrianalisoa et al.

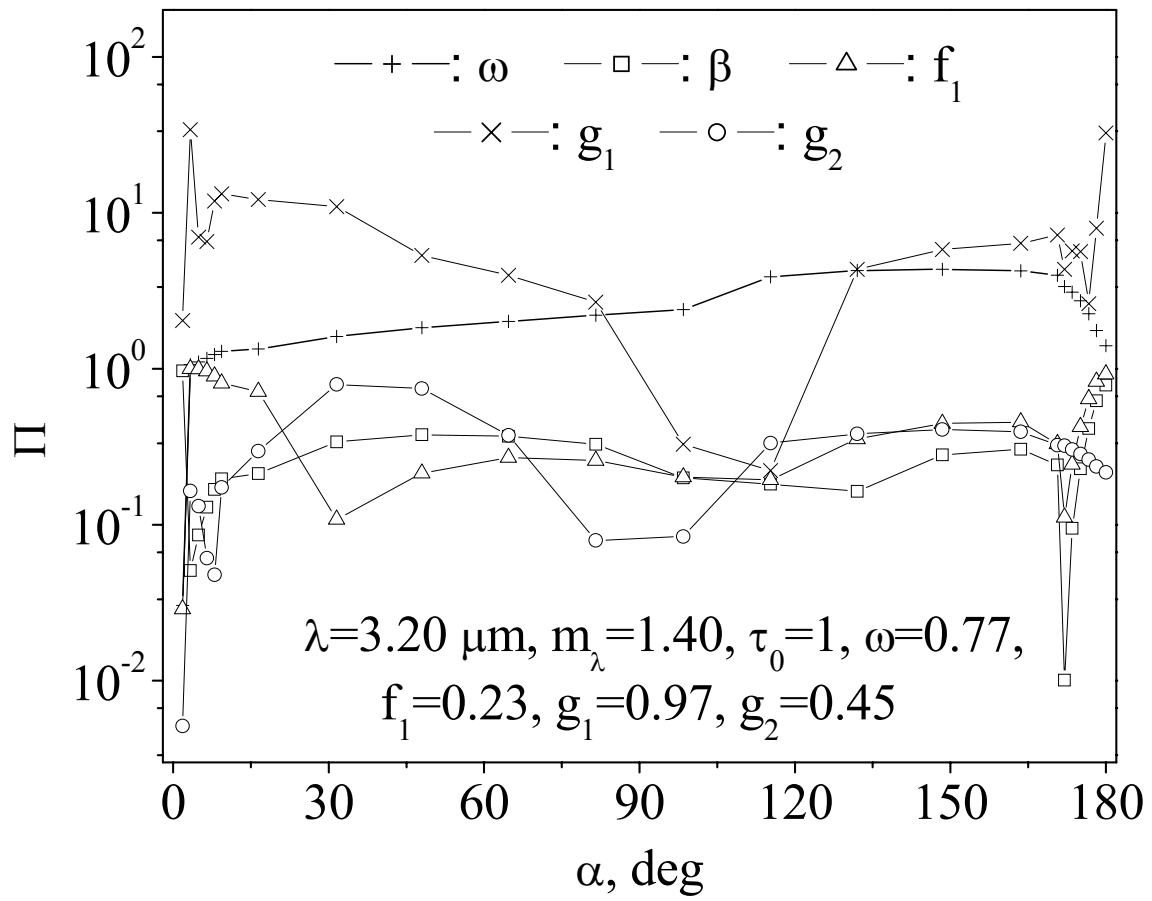


Fig. 6b, Randrianalisoa et al.

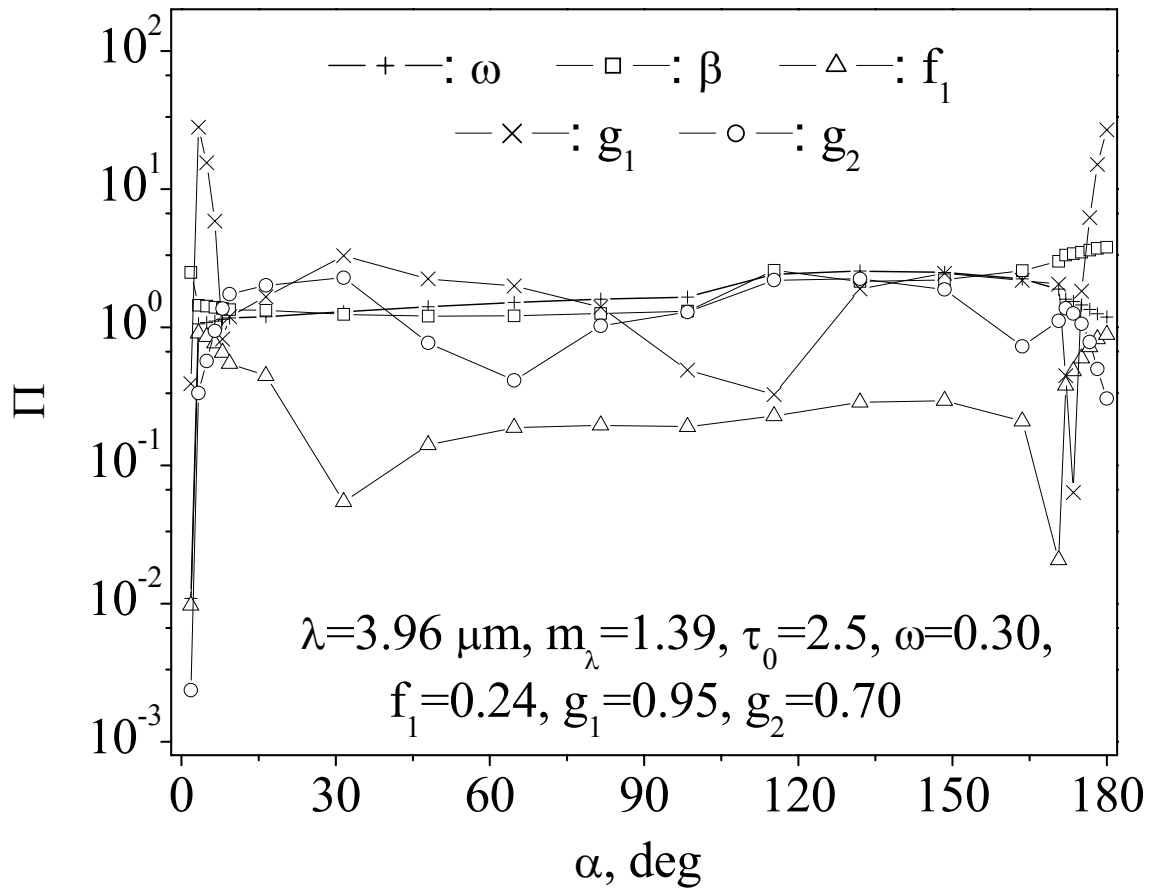


Fig. 6c, Randrianalisoa et al.

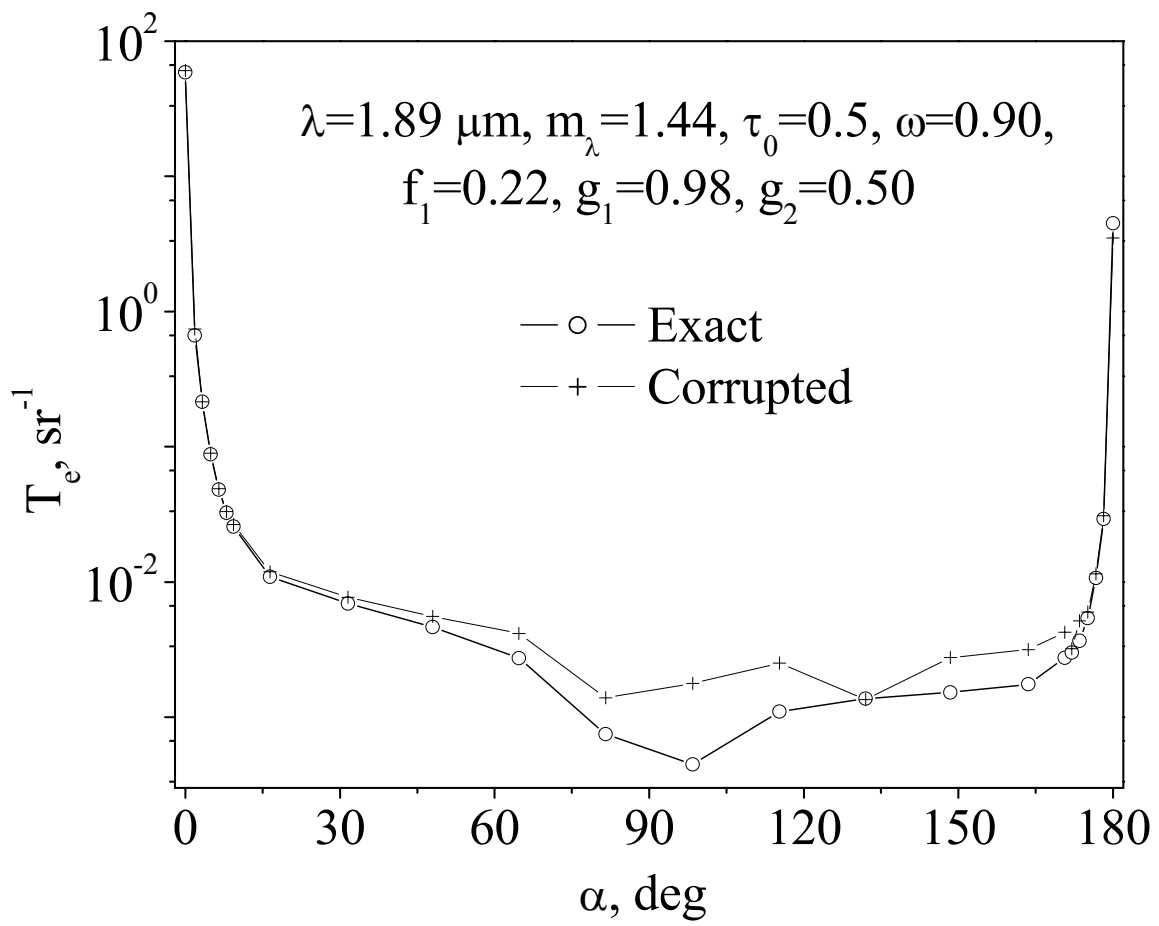


Fig. 7a, Randrianalisoa et al.

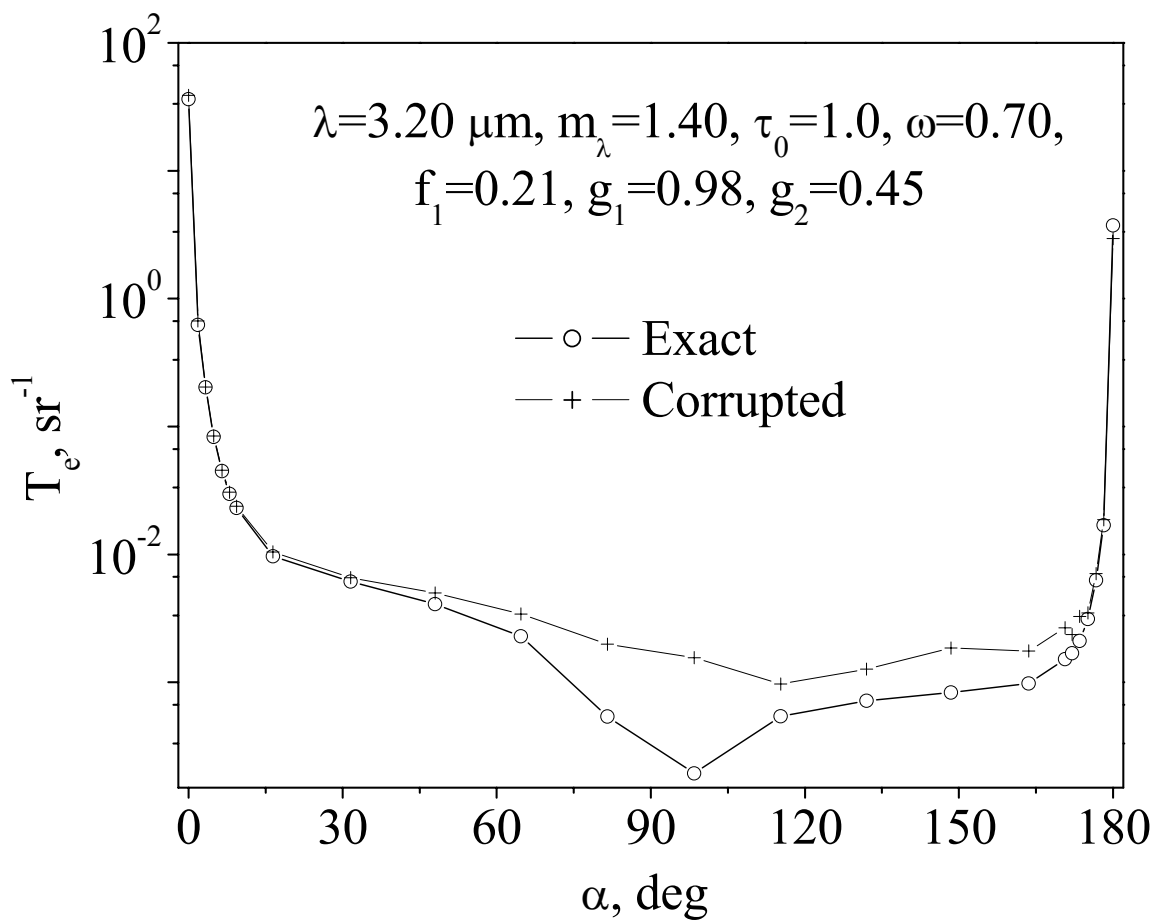


Fig. 7b, Randrianalisoa et al.

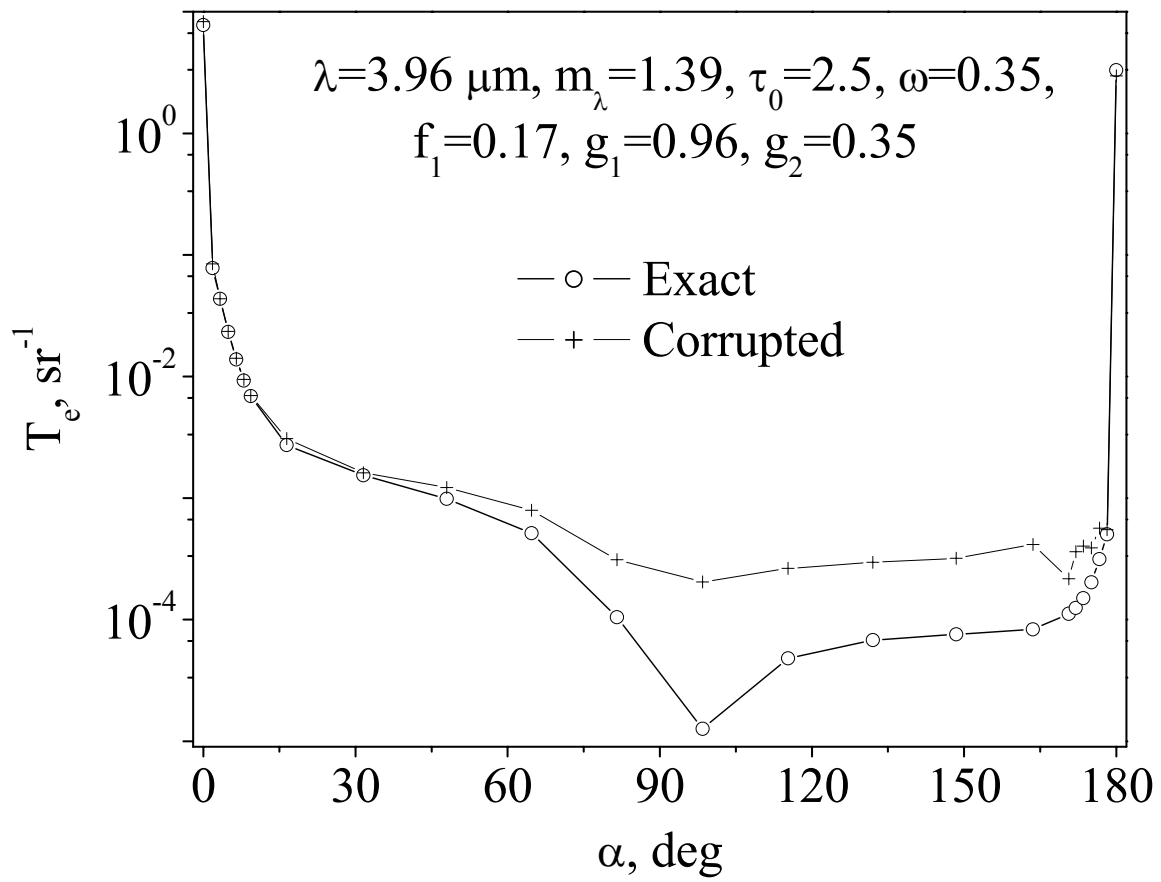


Fig. 7c, Randrianalisoa et al.

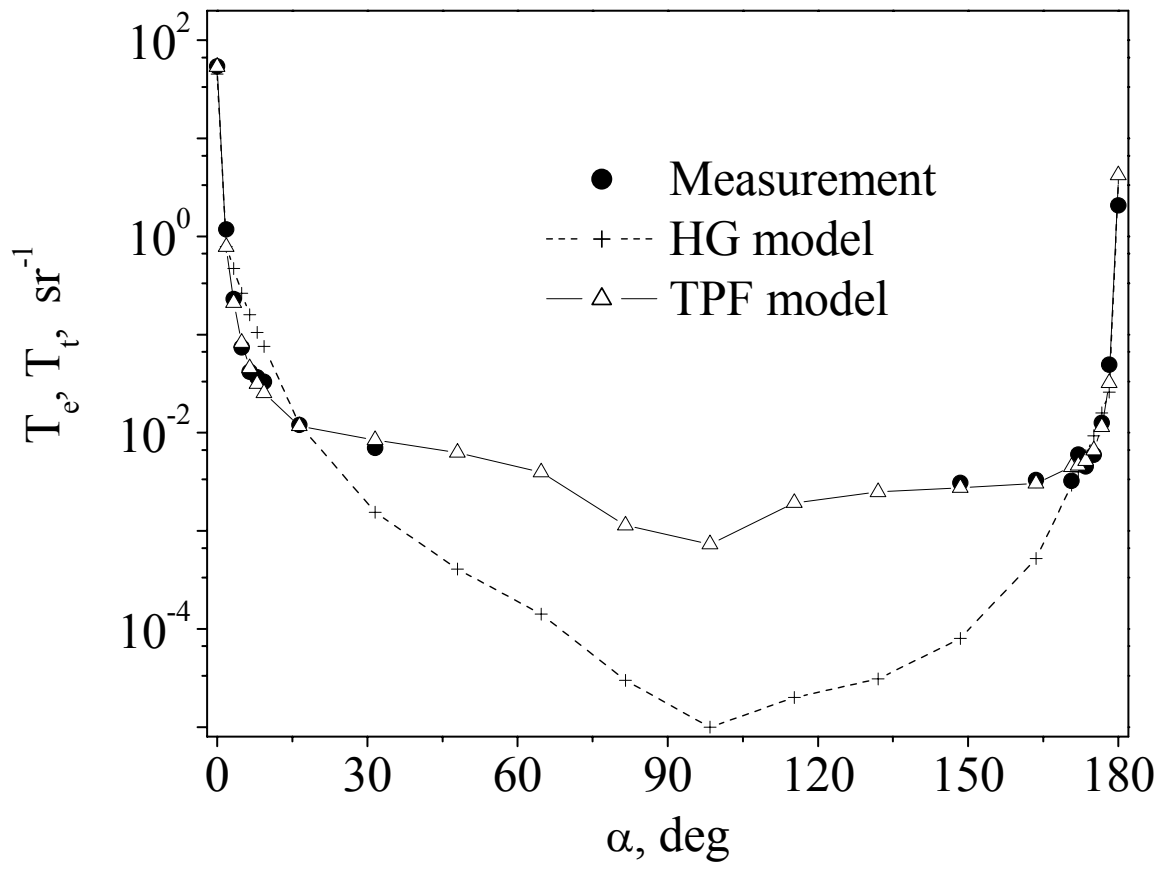


Fig. 8a, Randrianalisoa et al.

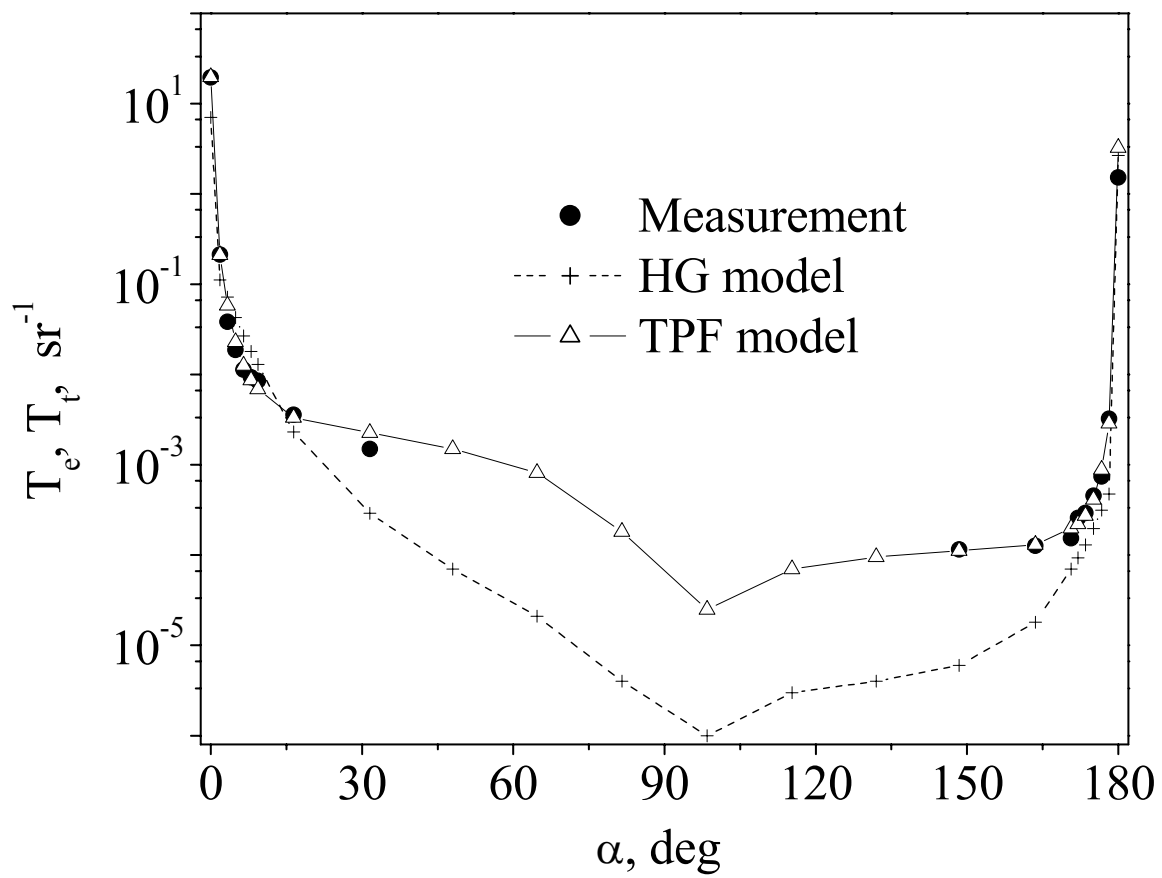


Fig. 8b, Randrianalisoa et al.

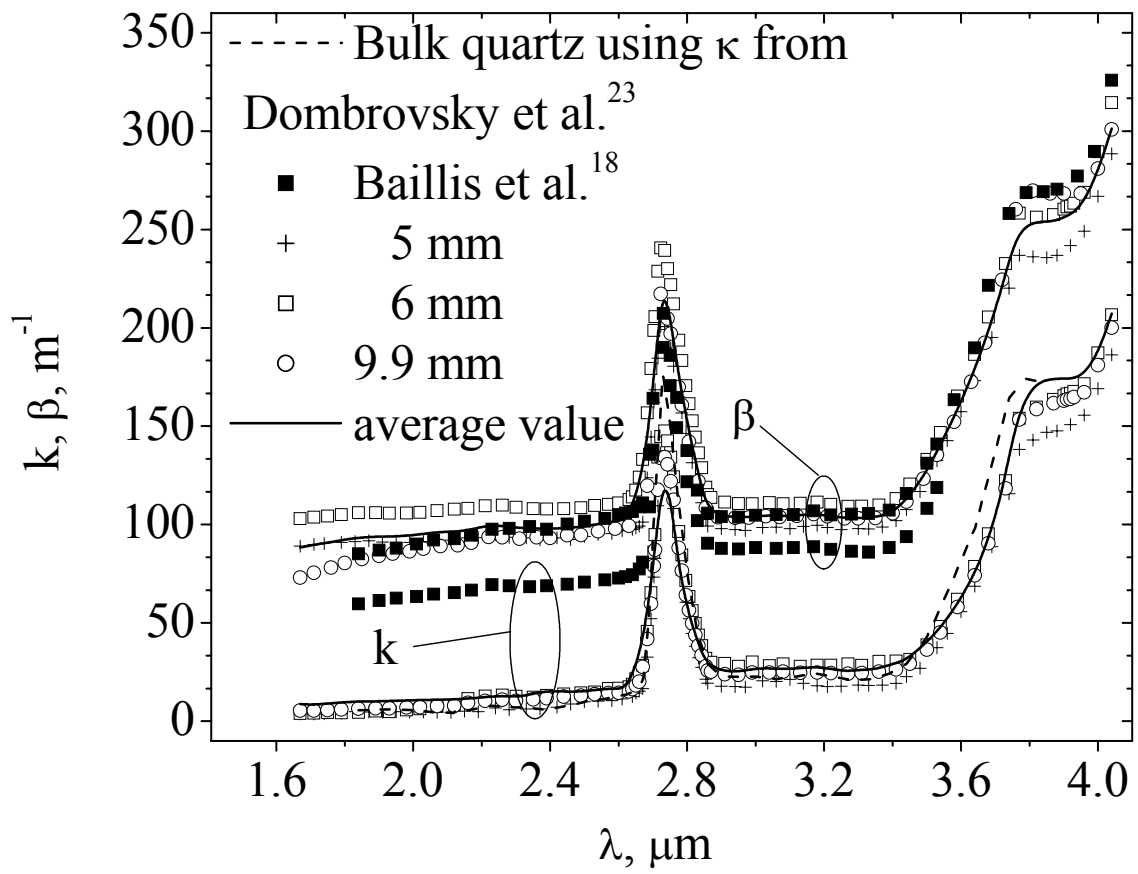


Fig. 9a, Randrianalisoa et al.

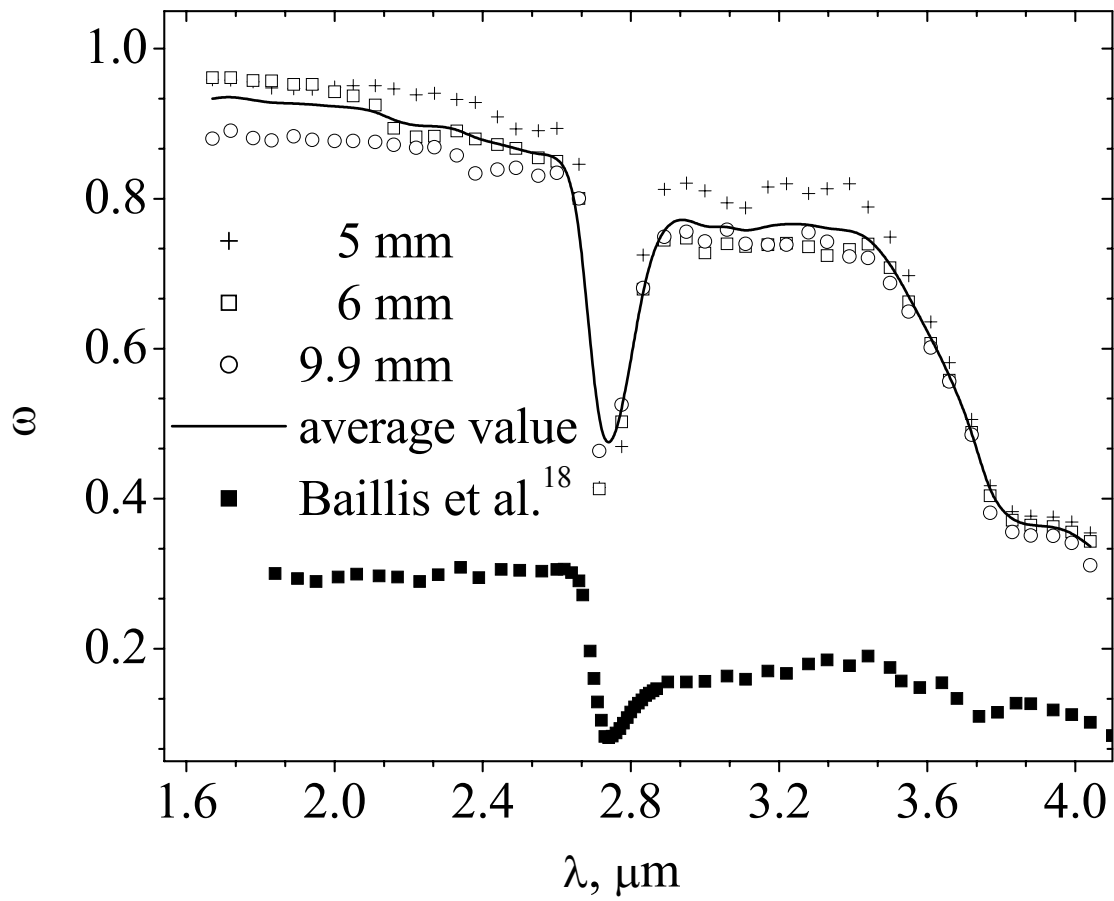


Fig. 9b, Randrianalisoa et al.

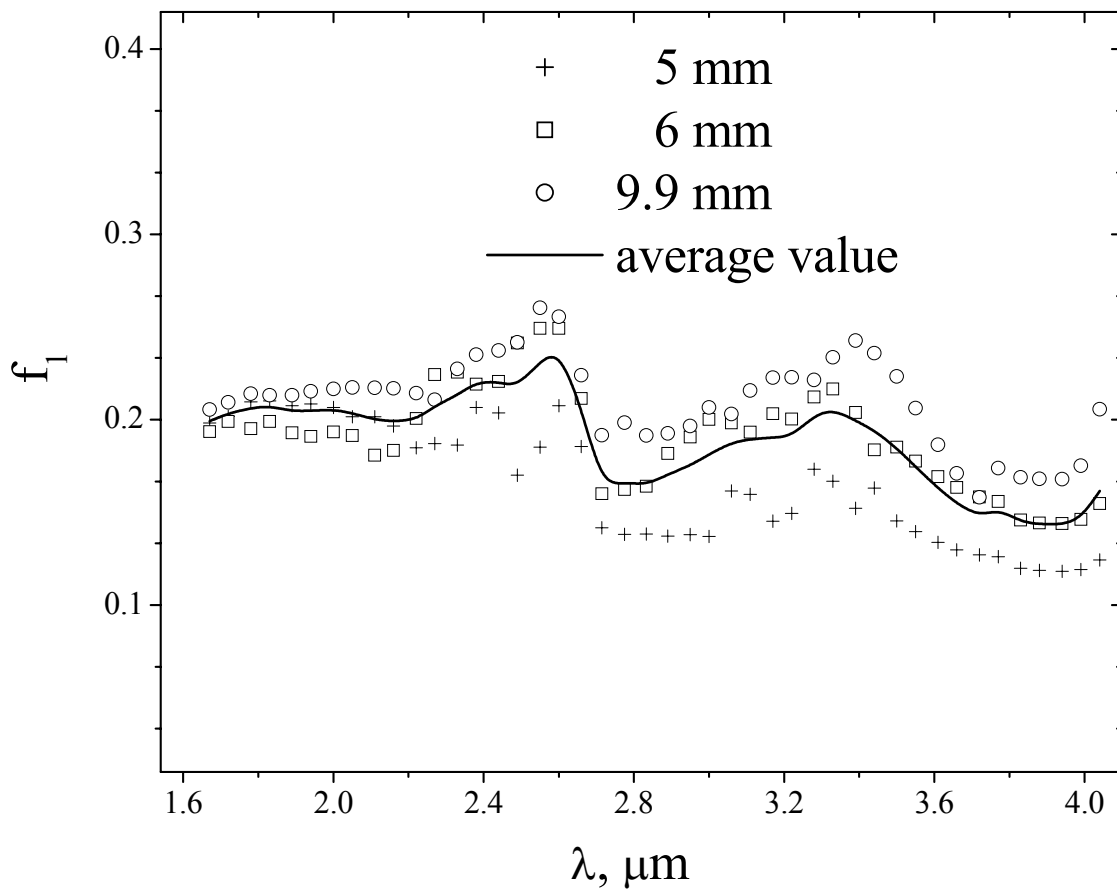


Fig. 9c, Randrianalisoa et al.

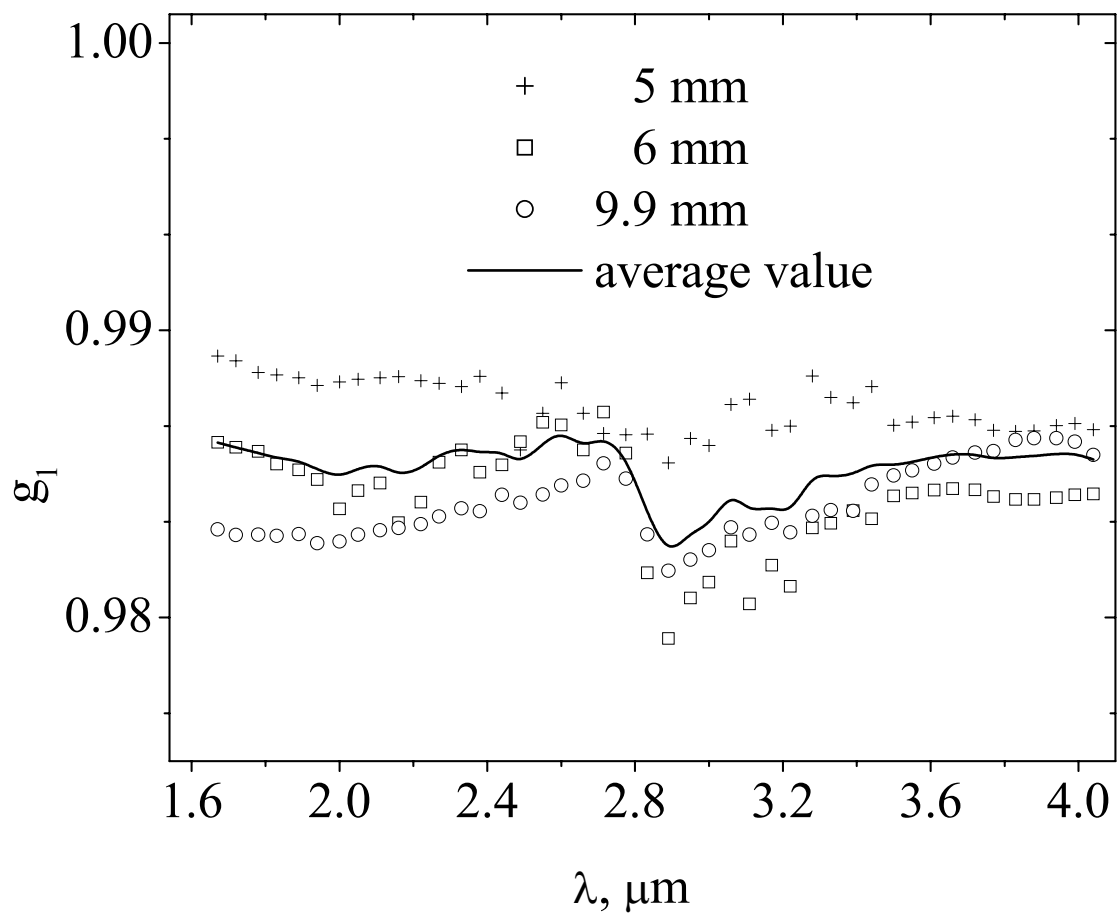


Fig. 9d, Randrianalisoa et al.

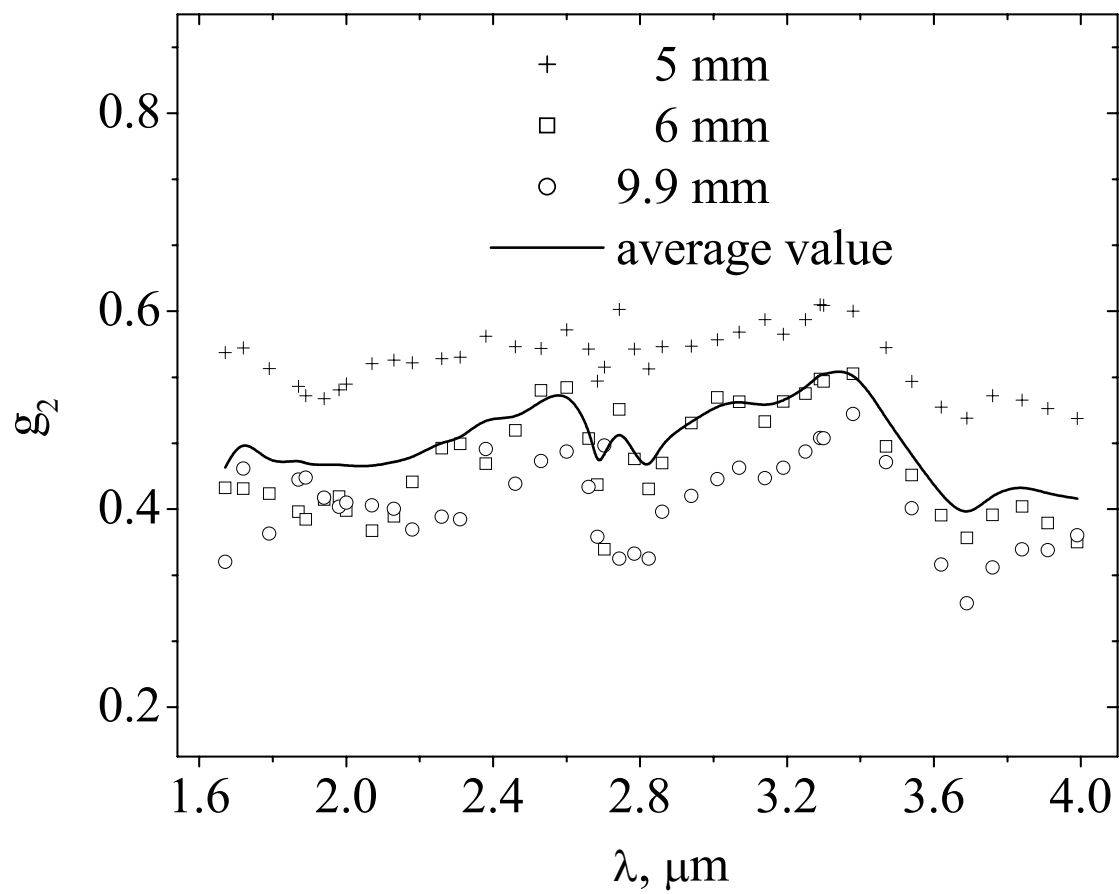


Fig. 9e, Randrianalisoa et al.

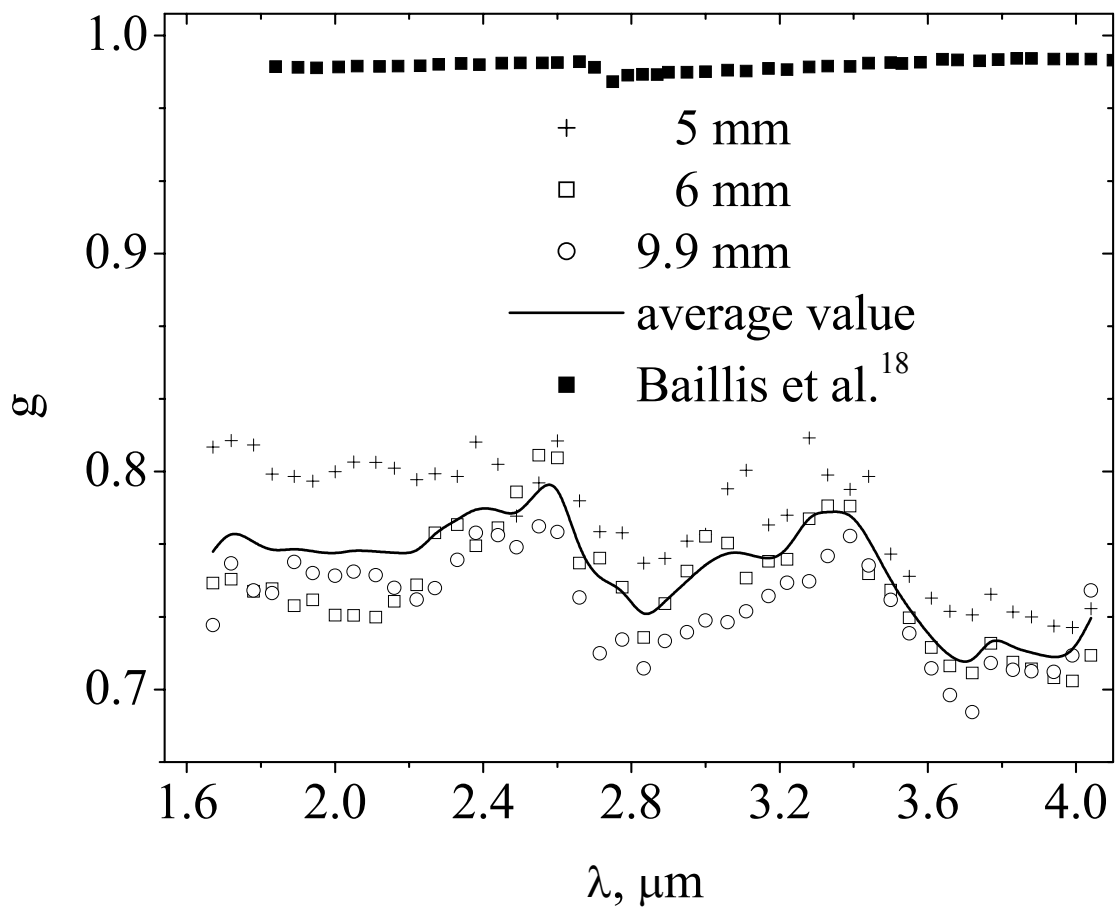


Fig. 9f, Randrianalisoa et al.

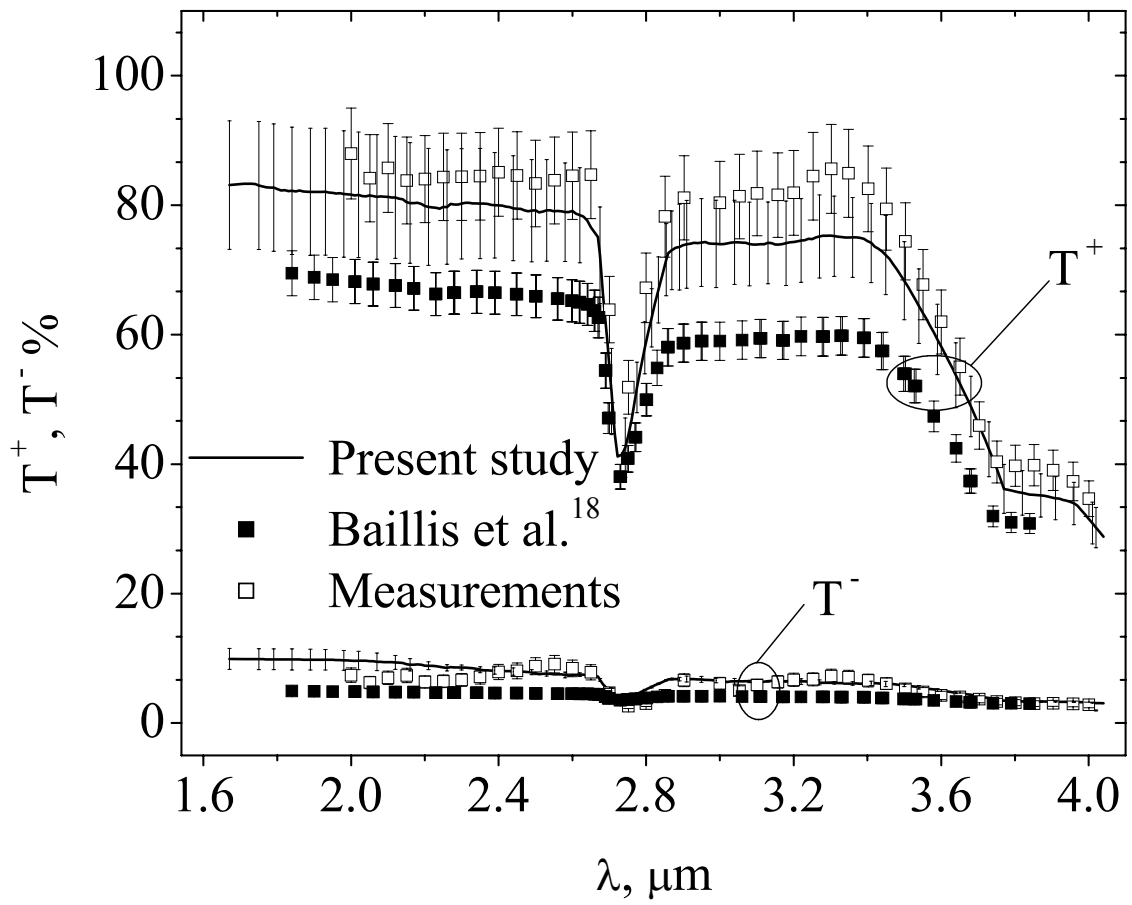


Fig. 10a, Randrianalisoa et al.

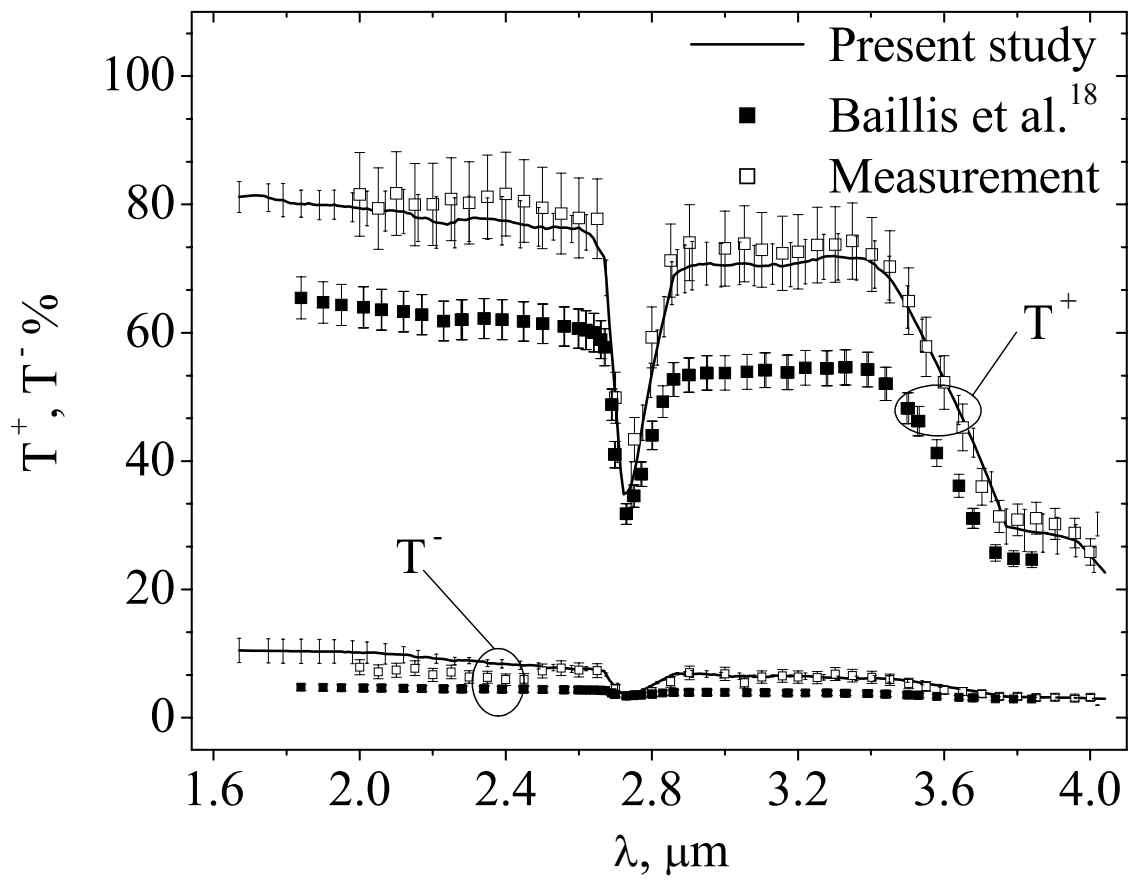


Fig. 10b, Randrianalisoa et al.

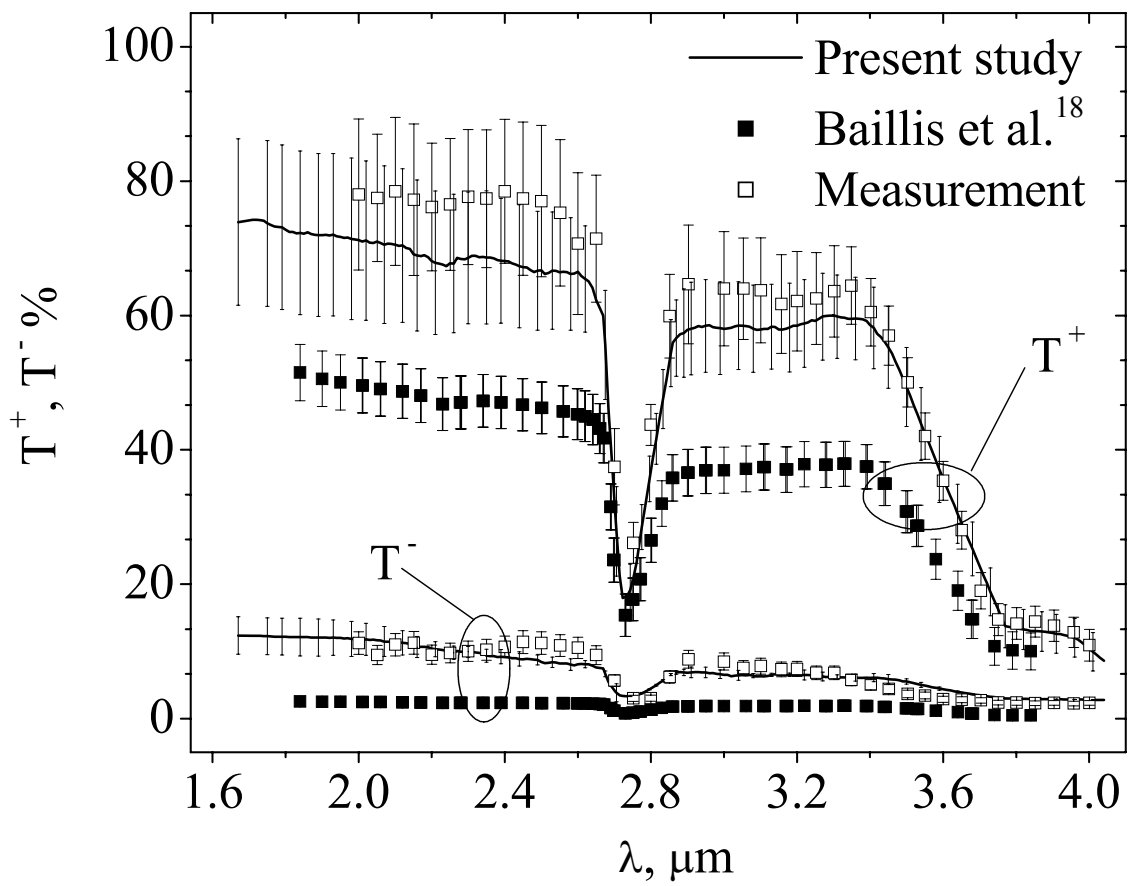


Fig. 10c, Randrianalisoa et al.