A FULL-SPECTRUM k -DISTRIBUTION LOOK-UP TABLE FOR RADIATIVE TRANSFER IN NONHOMOGENEOUS GASEOUS MEDIA

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Abstract

A full-spectrum k -distribution (FSK) look-up table has been constructed for gas mixtures within a certain range of thermodynamic states for three species, i.e., CO₂, H₂O and CO. The k -distribution of a mixture is assembled directly from the summation of the linear absorption coefficients of three species. The systematic approach to generate the table, including the generation of the pressure-based absorption coefficient and the generation of the k -distribution, is discussed. To efficiently obtain accurate k -values for arbitrary thermodynamic states from tabulated values, a 6-D linear interpolation method is employed. A large number of radiative heat transfer calculations have been carried out to test the accuracy of the FSK look-up table. Results show that, using the FSK look-up table can provide excellent accuracy compared to the exact results. Without the time-consuming process of assembling k -distribution from individual species plus mixing, using the FSK look-up table can save considerable computational cost. To evaluate the accuracy as well as the efficiency of the FSK look-up table, radiative heat transfer via a scaled Sandia D Flame is calculated to compare the CPU execution time using the FSK method based on the narrow-band database, correlations, and the look-up table. Results show that the FSK look-up table can provide a computationally cheap alternative without much sacrifice in accuracy.

Keywords

Radiative heat transfer, full-spectrum k -distribution, look-up table, accuracy, efficiency

Nomenclature

a	nongray stretching function in FSK method
$\bar{a}$	weight function in SLW method
C_{abs}	molar absorption cross section (cm^2)
f	full-spectrum k -distribution
F_s	absorption distribution function
g	cumulative full-spectrum k -distribution or quadrature point
I	radiative intensity ($\text{W}/(\text{m}^2 \cdot \text{sr})$)
k	k -value (cm^{-1})
L	slab width (m)
N_{pt}	number of nominal k -values
p	total pressure (bar)
R_u	universal gas constant ($\text{J}/(\text{mol} \cdot \text{K})$)
T	temperature (K)
w	quadrature weight
x	mole fraction

Greek Symbols

η	wavenumber (cm^{-1})
κ_η	spectral absorption coefficient (cm^{-1})
σ_s	scattering coefficient (cm^{-1})
$\underline{\phi}$	vector of local thermodynamic state variables
Φ	scattering phase function
Ω	solid angle (sr)

Subscripts

b	blackbody
max	maximum
min	minimum
p	pressure-based

Superscripts

0	reference state

Abbreviations

ADF	absorption distribution function
FSK	full-spectrum k -distribution
FSCK	full-spectrum correlated k -distribution
HITEMP	high-temperature spectroscopic absorption database
LBL	line-by-line
RTE	radiative transfer equation
SLW	spectral-line-based weighted-sum-of-gray-gases

1. Introduction

Radiative properties of molecular gases are of great importance for the study of combustion of hydrocarbon fuels and atmospheric radiation [1]. The absorption coefficient of gases, in particular for H_2O and CO_2 , varies dramatically across most of the spectrum, which is difficult to handle in practical applications. Therefore, accurate and efficient models for gaseous radiative properties are highly desirable. The most accurate radiative approach are line-by-line (LBL) calculations, which rely on very detailed knowledge of every single spectral line [2], but require vast amounts of computer resources and computational time [3], and remain unpractical for industrial applications. Therefore, the LBL method is used only for benchmark solutions to validate approximate methods.

An inspection of the spectral distribution of gaseous absorption coefficients reveals that the oscillatory absorption coefficients have the same value at many different wavenumbers. Recognition of this fact has led to the reordering of the absorption coefficients across the whole spectrum into a monotonic absorption coefficient distribution against normalized artificial wavenumber, which is much more efficiently integrated across the spectrum [1]. In order to treat radiation in nongray media accurately as well as efficiently, several models have been proposed based on the idea of the reordering concept, including the spectral-line-based weighted-sum-of-gray-gases (SLW) method [4], the absorption distribution function (ADF) method [5] and the full-spectrum k -distribution (FSK) method [6]. The first two, developed before the FSK, are approximate schemes, in which the absorption coefficients are reduced to a few discrete values (chosen by user), and integration over the spectrum is achieved by adding contributions from the “gray gases”.

The FSK method, on the other hand, is an exact method for homogeneous media using a continuous k -distribution over the entire spectrum. For nonhomogeneous media with variations in pressures, temperatures and species concentrations, a correlated absorption coefficient is assumed, i.e., the absorption coefficient from a thermodynamic state can essentially be correlated to that of another, resulting in the full-spectrum correlated k -distribution (FSCK) method [1]. With spectral integration performed by using high-accuracy Gauss quadrature, the FSK method can reduce the number of required radiative transfer equation (RTE) evaluations from over one million to around ten without losing spectral line information. With this obvious advantage, the FSK has become perhaps the most promising method for radiative heat transfer calculations in nongray media and has been employed in many applications. Consalvi and Liu [7] used several different spectral models to calculate radiation in axisymmetric pool fires and the results showed that the FSCK method achieves the best compromise in terms of accuracy and computational efficiency for a 34 kW pool fire. Clements et al. [8] investigated the FSK method under oxyfuel conditions and proved that the method can achieve good agreement for the test cases. Cai et al. [9] assembled full-spectrum k -distributions from a narrow-band database and developed high fidelity radiative heat transfer models to simulate high-pressure laminar hydrogen–air diffusion flames and they also applied the FSK method to the determination of radiation in fluidized-bed coal combustion [10].

FSK results tend to be very accurate, usually being within a few percent of “exact” LBL calculations. However, the drawback of the method is that FSKs are very cumbersome to calculate. For each state, say, with three radiatively participating gases (CO_2 , H_2O and CO) the method requires the assembly of mixture k -distributions or, for a CFD domain with several hundred thousand cells, the same amount of mixture FSKs. To overcome this drawback, Wang and Modest [11] described a narrow-band database for individual species, from which FSKs can be assembled efficiently by applying an appropriate mixing model. Cai and Modest [12] regenerated the narrow-band database using the newest HITEMP 2010 [13]. However, the mixing process for gas mixtures is computationally expensive, and creation of a single FSK for a two-gas mixture still requires between 0.0549 s to 0.469 s, depending on the

mixing model [14]. Clearly, if one million FSKs are needed per time step/iteration this becomes computationally prohibitive. This has prompted the development of simple correlation formulas for individual species at atmospheric pressure [15-20], from which k -distributions can be efficiently obtained; however, their accuracy are rarely satisfactory and sometimes may lead to serious errors [14]. Recently, Pearson et al. [21,22] tabulated ADF data (similar to FSK tabulations, except that they use absorption cross-sections instead of absorption coefficients) for CO₂, H₂O and CO over the full spectrum as a tool used in the SLW method, providing an alternative to solve the radiative property of nongray gas with excellent accuracy. However, this tabulation does not include gas mixtures and is relatively coarse, making it difficult to use for mixtures.

Therefore, to improve the calculation efficiency while retaining accuracy for radiative heat transfer, an FSK look-up table for gas mixtures with k -distributions assembled directly from the summation of the linear absorption coefficients has been constructed. Examples show that the FSK look-up table can be a computationally cheap alternative without much sacrifice in accuracy.

2. Theoretical background

2.1. Full-Spectrum k -Distributions

A full-spectrum k -distribution accounts for the variations of the Planck function and is defined as [1]

$$f_{\underline{\phi},T}(k;\underline{\phi},T) = \frac{1}{I_b(T)} \int_0^\infty I_{b\eta}(T) \delta(k - \kappa_\eta(\underline{\phi})) d\eta \quad (1)$$

where κ_η is the absorption coefficient calculated from a spectroscopic database, $\delta(\cdot)$ is the Dirac-delta function, $\underline{\phi}$ is a vector of local thermodynamic state variables including pressure, temperature and species concentration, $f_{\underline{\phi},T}(k;\underline{\phi},T)$ is a Planck-function-weighted k -distribution with absorption coefficient evaluated at the local state $\underline{\phi}$ and a Planck function temperature T , $I_b(T)$ and $I_{b\eta}(T)$ are the Planck function and the spectral Planck function, respectively, at temperature T , and η is the wavenumber.

The cumulative full-spectrum k -distribution is defined as

$$g_{\underline{\phi},T}(k;\underline{\phi},T) = \int_0^k f_{\underline{\phi},T}(k';\underline{\phi},T) dk' \quad (2)$$

Thus, $g_{\underline{\phi},T}$ represents the fraction of the spectrum whose absorption coefficient lies below the value of k and therefore, $0 \leq g_{\underline{\phi},T} \leq 1$. Inverting Eq. (2), a smooth, monotonically increasing function $k(g)$ can be obtained, with minimum and maximum values identical to those of κ_η .

For a nonhomogeneous mixture, the spectral variable is reordered according to the absorption coefficient at a reference state $\underline{\phi}^0$. Furthermore, to reduce the impact of uncorrelation, Cai and Modest [12] developed a new implementation to determine the correlated k -values. Applying this new implementation, the RTE can be reordered at the reference state $\underline{\phi}^0$ as

$$\frac{dI_g}{ds} = k_{\underline{\phi},T}^*(g_{\underline{\phi}^0,T^0}) \left[a(g_{\underline{\phi}^0,T^0}; T, T^0) I_b(T) - I_g \right] - \sigma_s(\underline{\phi}_s) \left(I_g - \frac{1}{4\pi} \int_{4\pi} I_g(s') \Phi(\underline{\phi}, s, s') d\Omega' \right) \quad (3)$$

where

$$I_g = \int_0^\infty I_\eta \delta(k - \kappa_\eta(\underline{\phi}^0)) d\eta / f_{\underline{\phi}^0,T^0}(k) \quad (4)$$

$$a(g_{\underline{\phi}^0,T^0}; T, T^0) = \frac{f_{\underline{\phi}^0,T}(k)}{f_{\underline{\phi}^0,T^0}(k)} = \frac{dg_{\underline{\phi}^0,T}(k)}{dg_{\underline{\phi}^0,T^0}(k)} \quad (5)$$

while σ_s is the scattering coefficient and Φ is the scattering phase function.

This is known as the full-spectrum correlated- k (FSCK) method. Usually, the total intensity is evaluated through an N -point numerical quadrature (with quadrature points g_i and weights w_i)

$$I = \int_0^1 I_g dg_{\underline{\phi}^0,T^0} = \sum_{i=1}^N w_i I_{g_i} \quad (6)$$

and for each quadrature point g_i , Eq. (3) can be rewritten as

$$\frac{dI_{g_i}}{ds} = k^*(g_i) \left[a(g_i) I_b(T) - I_{g_i} \right] - \sigma_s(\underline{\phi}_s) \left(I_{g_i} - \frac{1}{4\pi} \int_{4\pi} I_{g_i}(s') \Phi(\underline{\phi}, s, s') d\Omega' \right) \quad (7)$$

where $k^*(g_i)$ is the k -value corresponding to the quadrature point g_i for a gas mixture and $a(g_i)$ is obtained from Eq. (5).

2.2. Spectral-line-based Weighted-sum-of-gray-gases

If Eq. (3) is integrated using a crude trapezoidal scheme, i.e., the variable absorption coefficient $k_{\underline{\phi},T}^*(g_{\underline{\phi}^0,T^0})$ is replaced by a single, constant value $\tilde{k}_i(T, \phi)$ for the i th finite range of F_s spanning across $F_{s,i-1} < F_s \leq F_{s,i}$, integration of Eq. (3) over the i th F_s -range leads to

$$\frac{dI_i}{ds} = \tilde{k}_i(T, \phi) \left[\bar{a}_i(T, T^0) I_b(T) - I_i \right] - \sigma_s(\underline{\phi}_s) \left(I_i - \frac{1}{4\pi} \int_{4\pi} I_i(s') \Phi(\underline{\phi}, s, s') d\Omega' \right), \quad i = 1, \dots, N \quad (8)$$

This is known as the spectral-line-based weighted-sum-of-gray-gases (SLW) method developed by Dension and Webb [4]. Generally, the values of $\tilde{k}_i(T, \phi)$ and $\bar{a}_i(T, T^0)$ are obtained from the absorption distribution function (ADF) [20-23]. Thus, the weight function $\bar{a}_i(T, T^0)$ is evaluated from [24]

$$\bar{a}_i(T, T^0) = F_s(C_{\text{abs},i}, T, T^0) - F_s(C_{\text{abs},i-1}, T, T^0) \quad (9)$$

where F_s is the ADF and defined as

$$F_s(C_{\text{abs}}, T, T^0) = \frac{1}{I_b(T)} \sum_i \int_{\Delta\eta \in (C_{\text{abs},\eta}(T^0) < C_{\text{abs}})} I_{b\eta}(T) d\eta \quad (10)$$

The molar absorption cross section $C_{\text{abs},\eta}$ in Eq. (10) is related to the absorption coefficient by

$$C_{\text{abs},\eta} = \frac{R_u T^0}{p x_s} \kappa_\eta \quad (11)$$

where R_u is the universal gas constant, p is the total pressure and x_s is mole fraction of species s .

As Modest and Singh [19] pointed out that the F_s in the SLW method and the g in the FSK method can be expressed using the same equation, and the SLW method is just the crudest possible implementation of the FSK method. Therefore, the values of $\tilde{k}_i(T, \phi)$ and $\bar{a}_i(T, T^0)$ can also be obtained from the FSK look-up table developed in this paper, just replacing $\tilde{k}_i(T, \phi)$ and $\bar{a}_i(T, T^0)$ by $k^*(g_i)$ and $a(g_i)$, respectively. From this perspective, except for the order of quadrature, the SLW method is equivalent to the FSK method.

The purpose of this paper is to precalculate these $k^*(g_i)$ and $a(g_i)$ values for different gas mixtures and to tabulate them into an FSK look-up table for the convenience of accurate and rapid radiative calculations by using either the FSK method or the SLW method.

3. Construction of FSK look-up table

3.1. Generation of the Pressure-based Absorption Coefficient

In the present FSK look-up table, three species are taken into account, i.e., CO_2 , H_2O and CO . To avoid the mixing process during FSK generation, the mixture k -distribution is generated directly from the mixture linear absorption coefficients for different thermodynamic states including all three gases, which can be expressed as

$$\kappa_\eta = (x_{\text{CO}_2} \kappa_{p\eta\_ \text{CO}_2} + x_{\text{H}_2\text{O}} \kappa_{p\eta\_ \text{H}_2\text{O}} + x_{\text{CO}} \kappa_{p\eta\_ \text{CO}}) \cdot p \quad (12)$$

where x_{CO_2} , $x_{\text{H}_2\text{O}}$ and x_{CO} are the mole fractions of CO_2 , H_2O and CO , respectively, p is the total pressure and the subscript ‘ p ’ represents pressure-based absorption coefficient.

Since the linear absorption coefficient is converted from the pressure-based absorption coefficient as shown in Eq. (12), the first step in the construction of the FSK look-up table is the development of the detailed absorption spectrum as a function of relevant thermodynamic properties for the gases of interest. The detailed approach to generation of pressure-based absorption coefficients for individual species has been previously given [25], and outlined here for completeness. The HITEMP 2010 database was employed to generate the spectral pressure-based absorption coefficients, using the Voigt line profile [1] as it has been shown to have higher accuracy than the Lorentz profile for cases of sub-atmospheric pressure or high temperature where both collision and Doppler broadening are significant.

3.2. Generation of the k -Distribution

In order to cover most conditions found in industrial practices, a large range of pressures ranging from 0.1 bar to 80 bar, temperatures ranging from 300 K to 3000 K and a full mole fraction range (from 0 to 1 for CO₂ and H₂O, and from 0 to 0.5 for CO, which is not expected to occur at higher fractions in combustion system) is considered here. Because of both mixing and self-broadening, the k -distributions may show nonlinearities as shown in Fig. 1, in which the lines of exact k -values corresponding to different g -values may deviate from the respective linear dash-dot lines. To decrease these nonlinear effects, a large number of small mole fractions is included in the FSK look-up table. The nonlinear effects due to mixing mainly result from the overlap of absorption bands among three species with the strongest nonlinearity appearing at low mole fractions (Fig. 1a), leading to the need of including several low mole fraction data points in the database for the generation of mixture k -distributions. In a gas mixture with CO₂, CO, H₂O and N₂, the nonlinear effects due to self-broadening are very strong for H₂O (Fig. 1b) because of the large molecule size difference between H₂O and N₂, plus nonlinear effects due to mixing, resulting in more mole fraction points needed for H₂O to generate the k -distribution; for CO₂, the size of the molecule is a little larger than N₂, leading to a smaller effect of self-broadening compared to H₂O, therefore, fewer mole fraction points are needed for CO₂ to reduce the interpolation error resulting from nonlinear effects; for CO, the molecule size is about the same size as N₂ and the effect of self-broadening can be ignored, thus only five mole fraction points are needed for CO to decrease the nonlinear effect due to mixing. Table 1 summarizes the pressures, gas temperatures, reference temperatures and mole fractions stored in the present FSK look-up table.

Fig. 1. Exact k -values corresponding to 8th, 12th and 16th g -quadrature values with increasing mole fraction using a 16-point Gauss-Chebyshev quadrature scheme [dash-dot lines represent linear interpolation lines connecting the first and last k -values; ($\times 100$), ($\times 20$), ($\times 50$) and ($\times 10$) imply amplification factors].

Table 1
Precalculated thermodynamic states of FSK look-up table

Parameters	Range	Values	Number of points
Species	CO ₂ , H ₂ O and CO		3
Pressure (total)	0.1~0.5 bar	Every 0.1 bar	34
	0.7 bar	0.7 bar	
	1.0~14.0 bar	Every 1.0 bar	
	15.0~80 bar	Every 5 bar	
Gas temperature	300~3000 K	Every 100 K	28
Reference temperature	300~3000 K	Every 100 K	28
Mole fraction of CO ₂	0.0~0.05	Every 0.01	10
	0.25~1.0	Every 0.25	
Mole fraction of H ₂ O	0.0~0.05	Every 0.01	13
	0.1~0.2	Every 0.05	
	0.25~1.0	Every 0.25	
Mole fraction of CO	0.0~0.5	[0.0, 0.01, 0.05, 0.1, 0.25, 0.5]	6

When employing Eqs. (1) and (2) to calculate the cumulative k -distributions, a set of nominal k -values between the local maximum and minimum linear absorption coefficients must be chosen for each thermodynamic state. In this work, a power distribution is used and may be expressed as [11]

$$k_i = \left[\kappa_{\min}^{\beta} + \frac{i-1}{N_{pt}-1} \cdot (\kappa_{\max}^{\beta} - \kappa_{\min}^{\beta}) \right]^{1/\beta} \quad (0 < \beta < 1, 1 \leq i \leq N_{pt}) \quad (13)$$

With Eq. (13), a skewed distribution that places more points at smaller k -values is generated. After considerable numerical experimentation, a value of $\beta = 0.1$ was found to be most satisfactory [11]. To avoid errors when the two adjacent linear absorption coefficients occur in the same k -value interval, a large number of points $N_{pt} = 5000$ were employed. Therefore, the k -distributions obtained from LBL data can be regarded as essentially error-free.

For every thermodynamic state, the cumulative k -distribution g with 5000 k -values have extremely high accuracy; however, this leads to large storage, which is not acceptable for practical calculations. Since the cumulative k -distribution g always varies between 0 and 1, fixing the g -values is a logical choice to compact the present table. Since n -node Gauss quadrature has the highest order of accuracy, a set of the abscissas of the Gauss-Chebyshev quadrature (I) described in [11] were chosen for the fixed g -values.

To guarantee the accuracy for any optical conditions, 32 Gauss-Chebyshev quadrature points were chosen for the fixed g -values for all full-spectrum k -distributions. Thus, only the corresponding k -values and a -values were tabulated in the FSK look-up table. However, it still makes the total size of the FSK look-up table close to 5 GB. For practical applications, loading such a huge database into memory is both time-consuming and wasteful because generally only a portion of data is used for certain calculations. To avoid this drawback and also based on the fact that all records in the FSK look-up table are of the same length (there are 32 records at each thermodynamic states for both k -values and a -values), the direct data access method [11] is employed so that only the data needed by the calculations are loaded into the memory, saving both the time for reading the whole table and the

memory when carrying out certain calculations. This memory management approach that guarantees the lowest memory cost without affecting the runtime efficiency during certain calculations is known as dynamic loading.

To meet the demands of different users, k -values and a -values are stored separated by both pressure and mole fraction of three species requiring 100 KB for each file. Therefore, one can assemble a small FSK look-up table for special cases; for example, a table with a single constant pressure and a mole fraction range from 0 to 0.25 can be assembled from 700 (7 CO₂ mole fractions $\times$ 10 H₂O mole fractions $\times$ 5 CO mole fractions for both k -values and a -values) individual files, leading to a table of only 68.4 MB; furthermore, for RTE evaluations, usually 8~16 quadrature points are sufficient, making the size of the table even smaller, for example, a table with a single constant pressure with 16 quadrature points is only 34.2 MB in size.

3.3. Interpolation Method

For an arbitrary thermodynamic state, a full-spectrum k -distribution is specified by pressure, gas temperature, reference temperature and three concentrations. Therefore, a six-dimensional interpolation method is required to obtain the full-spectrum k -distribution from the look-up table. To achieve relatively small computational cost, the simplest and fastest 6-D linear interpolation method with a $2 \times 2 \times 2 \times 2 \times 2 \times 2$ domain is employed to obtain both k -values and a -values. Since both gas and reference temperature ranges are equally spaced, the domain values of these two parameters can be directly obtained; while the domain values of pressure and three species need to be searched due to the unequally spaced ranges, requiring some extra computational time.

Since the effect of self-broadening is remarkably larger at both high pressure and low temperature, especially for H₂O, it is appropriate to compare the negative radiative heat source $\nabla \cdot q$ calculated by two mixture k -distributions obtained from different sources to test the accuracy of 6-D linear interpolation. As shown in Fig. 2, the k -distributions shown as a red dash-dot line are obtained from 6-D linear interpolation among the tabulated values in the FSK look-up table; and the k -distributions shown as a black solid line are calculated by an exact FSK, for which the k -distributions are directly calculated from the LBL absorption coefficient database [25]. It can be seen that two values of $\nabla \cdot q$ are in excellent agreement at both high pressure and high temperature, and the maximum error stays around 2%. Therefore, the 6-D linear interpolation method was deemed good enough to obtain the k -values from the FSK look-up table.

Fig. 2. Comparison of $\nabla \cdot q$ calculated by k -distributions obtained by an exact evaluation and 6-D linear interpolation from tabulated data in the FSK look-up table; 16-point quadrature.

4. Results and discussions

4.1. Sample calculations

4.1.1. Accuracy investigation

To test the accuracy of the FSK look-up table, a large number of sample calculations were carried out for one-dimensional slabs of a homogeneous gas mixture bounded by cold black walls, with $\nabla \cdot q$ predicted by the FSK method using the look-up table. Since the FSK method is exact for homogeneous media, the reference solution is provided by the exact FSK. Thus, any error generated by using the FSK look-up table is due to inaccuracies from interpolation of k -distributions from the FSK look-up table. For these cases, the analytical solution of the RTE exists [1] and is used for comparison. Mixtures of three species (CO_2 , H_2O , CO) in different thermodynamic states are shown in Table 2. All FSK calculations employ a 16-point Gauss-Chebyshev scheme.

Table 2
Investigated gas mixture states and corresponding fixed parameters

Investigated gas mixture states		Fixed parameters						
		p (bar)	T (K)	x_{CO_2}	$x_{\text{H}_2\text{O}}$	x_{CO}	x_{N_2}	L (m)
p (bar)	0.85, 2.5, 17.5, 37.5, 72.5	—	1650	0.150	0.625	0.030	0.195	0.1
T (K)	450, 1050, 1650, 2250, 2850	17.5	—	0.375	0.375	0.075	0.175	0.1
L (m)	0.001, 0.01, 0.1, 0.5, 1	17.5	1650	0.625	0.125	0.175	0.075	—

Figs. 3 through 5 show the representative curves of $\nabla \cdot q$ across the slab varying with pressures, temperatures and slab widths, respectively. For better readability, only FSK exact results and errors made by database-assembled k -distribution calculations from the FSK look-up table have been plotted. Fig. 3 shows the effects of total pressure on overall accuracy, keeping all other parameters fixed. For

low pressures such as 0.85 bar, $\nabla \cdot q$ is essentially constant across the slab, since the gas is so optically thin that all emission escapes from the medium without self-absorption. For high pressures, the gas becomes optically thick and, therefore, energy emitted near the centre cannot escape, leading to the strong gradients toward the boundaries. The same phenomenon can be also seen in Fig. 4, in which the temperature T is increased, and in Fig. 5, in which the slab width L is increased. For all cases, the maximum error across the slab stays around or below 2%, implying that the FSK look-up is sufficiently accurate for radiative calculations.

Fig. 3. Negative radiative heat source for a 15.0% CO₂-62.5% H₂O-3% CO-19.5% N₂ mixture at varying pressures; 16-point quadrature.

Fig. 4. Negative radiative heat source for a 37.5% CO₂-37.5% H₂O-7.5% CO-17.5% N₂ mixture at varying temperatures; 16-point quadrature.

Fig. 5. Negative radiative heat source for a 62.5% CO₂-12.5% H₂O-17.5% CO-7.5% N₂ mixture at varying slab thicknesses; 16-point quadrature.

4.1.2. Efficiency investigation

To test the efficiency of the FSK look-up table, CPU times of generating k -distributions for arbitrary gas mixtures at arbitrary thermodynamic states are compared using FSK generation based on exact FSK, assembly from the narrow-band database [12], from correlations [18,19] and from the look-up table developed in this paper, respectively. Since the correlations can be used only at 1 bar, the pressure for all arbitrary k -distributions is fixed to 1 bar. The temperature range for arbitrary thermodynamic states is 300 K ~ 3,000 K. Three gases (CO₂, H₂O and CO) with mole fractions ranging from 0 to 0.25 are included in the generation of arbitrary k -distributions. However, correlations are only available for CO₂ and H₂O; therefore, the k -distributions generated using correlations include only two gases.

Table 3

CPU time comparisons of generating 10,000 arbitrary k -distributions based on different databases

Database	Mixing model	CPU (s)
Exact FSK	—	31216
Narrow-band	Multiplication	1390
	MRmixing	5905
Correlations	Multiplication	0.41
	MRmixing	8.97
Look-up table	—	0.26

Table 3 shows CPU time comparisons for generating 10,000 arbitrary k -distributions based on four different databases. Two mixing models are included in the comparisons for the k -distributions assembly from both the narrow-band database and correlations, i.e., the multiplication model [26] and the Modest and Riazzi mixing (MRmixing) model [27]. Since the exact FSK is calculated from the LBL absorption coefficient database, the required CPU time for generation using the exact FSK, while large, is considerably lower than that when calculated directly from HITEMP 2010. Using the

narrow-band database saves considerable CPU time because k -distributions have been precalculated for individual species. However, CPU time of generation using the narrow-band database is still large due to the mixing process. As listed in Table 3, even employing the more efficient but less accurate multiplication model, using the narrow-band database still requires 1390 s to generate 10,000 arbitrary k -distributions. CPU time for generation using correlations is much smaller than that using the narrow-band database; it also depends on the mixing model: CPU time employing the MRmixing model is much larger than that for the multiplication model. Without both assembly of k -distributions and mixing, CPU time using the look-up table is remarkably cheap, which is even faster than that using correlations with the more rapid multiplication model.

Comparisons of k -distributions calculated from different databases in Table 3 at one thermodynamic state are shown in Fig. 6 (left). The k -distributions calculated from both narrow-band database with MRmixing model and look-up table overlap very well with the exact results, which are directly calculated by exact FSK. By contrast, those k -distributions calculated from the narrow-band database with the multiplication model and from correlations with the two different mixing models show some discrepancies compared to the exact results. Employing those k -distributions, the negative radiative heat source and errors compared to exact FSK results across the 1D slab are shown in Fig. 6 (right). It is clear that with accurate k -distributions, the results using the narrow-band database with the MRmixing model and the look-up table are much better than those using the correlations and/or the multiplication mixing model, showing less than 2% error across the slab compared to exact FSK results. In addition, comparing the CPU times shown in Table 3 makes it clear that the look-up table is a more appropriate way for radiative calculations than using the narrow-band database with the MRmixing model.

Fig. 6. k -distributions for a gas mixture (left) and corresponding negative radiative heat source and errors compared to exact FSK for a 1D slab (right) using different databases; 16-point quadrature.

4.2. Flame calculation

In order to evaluate accuracy and efficiency of the FSK look-up table, radiation from a realistic flame is considered. The flame is derived from Sandia Flame D [28] by artificially quadrupling the jet diameter, resulting in strong radiation effects that are found in practical combustion configurations. The flame is simulated on a two-dimensional axisymmetric mesh, and only the quasi-steady time-

averaged temperature and species mass fraction fields shown in Fig. 7 are used here for radiation calculation comparisons. Hence, no turbulence-radiation interaction is considered, and $\nabla \cdot q$ is determined using the FSK method based on three different databases (narrow-band, correlations and look-up table) without feedback to the flame. The RTE is solved by the P_1 method. The total pressure is 1 bar and three species (CO_2 , H_2O and CO) are included in the calculations except for the calculations using the correlations, which contains only CO_2 and H_2O . All FSK calculations employ a 16-point Gauss-Chebyshev quadrature scheme and are compared with LBL calculations at three downstream locations.

Fig. 7. Time-averaged spatial profile of temperature, CO_2 mass fraction, H_2O mass fraction and CO mass fraction of enlarged Sandia Flame D.

The results using the narrow-band database employed the MRmixing model have been presented by Cai and Modest [12]. As shown in Fig. 8, at all three locations, using the narrow-band database with the MRmixing model provides excellent performance compared to LBL results. The discrepancies between the results from the narrow-band database with the MRmixing model and LBL results mainly come from the quadrature errors, interpolation errors and mixing errors. When employing the quicker multiplication model, the accuracy using the narrow-band database dramatically decreases due to the increase of mixing errors. Calculations using the correlations with either multiplication model or MRmixing model show the worst performance among the three databases. Since radiation of CO is not very dominant, the large discrepancies between the correlation results and LBL results are mainly due to the inaccurate k -distributions obtained from the correlations. Without mixing, only quadrature errors and interpolation errors are present when using the look-up table. Since the quadrature errors are very tiny when employing 16-point quadrature and the interpolation errors are around 2 % as shown in Section 4.1, calculations using the look-up table are very close to the LBL results, showing almost the best performance among three databases. At the locations where radiation is dominated by the

Fig. 8. Comparisons of FSK calculations using different databases (right column) along with the species mass fraction and the temperature (left column) at three locations, $z = 0.5$ m (top), $z = 1.0$ m (middle), and $z = 1.4$ m (bottom); Multi: multiplication model, MR: MRmixing model.

emission, e.g. $z = 0.5$ m and $z = 1.0$ m, calculations using the look-up table even overlap with LBL results. This illustrates that in practical applications using the look-up table can achieve a LBL accuracy in predicting of radiative heat source.

CPU times of the LBL calculations and the FSK calculations using the three databases are compared in Table 4. Because of the one million evaluations of the RTE, using the LBL database requires a large amount of CPU time. Using the narrow-band database can save much CPU time compared to LBL calculations. However, even employing the more rapid multiplication model, the CPU time using the narrow-band database is still about a factor of 970 higher than that using the look-up table due to the assembly of mixture k -distributions from individual species. Thus, for a single calculation without feedback, it is quite time-consuming to use the narrow-band database in practical applications. When employing dynamic loading, considerable time for reading the whole look-up table into memory is saved, making the CPU time using the look-up table even shorter than that using correlations with the multiplication model. Therefore, together with the accuracy discussions above, using the look-up table appears to be a better alternative for radiative calculations, and is computationally cheaper than using correlations but with high accuracy, which is even better than using the narrow-band database.

Table 4
Comparison of CPU times for calculations of the Sandia D flame case using different databases

Database	Mixing model	CPU (s)
LBL	–	87331.98
Narrow-band	Multiplication	1513.57
	MRmixing	8422.88
Correlations	Multiplication	1.76
	MRmixing	13.34
Look-up table	–	1.56

5. Conclusions

To avoid the time-consuming process of assembling mixture full spectrum k -distributions from spectroscopic databases plus mixing, an FSK look-up table has been constructed, from which k -distributions can be obtained by interpolation. The following conclusions are drawn from this work:

1. The FSK look-up table presented in this paper includes three species (CO_2 , H_2O and CO) and has a large range of pressures ranging from 0.1 bar to 80 bar and temperatures ranging from 300 K to 3000 K. For both CO_2 and H_2O , the considered mole fraction range is between 0 and 1; while for CO , it is between 0 and 0.5.
2. A large number of calculations were carried out to investigate the effects of total pressure, temperature and slab width on the accuracy of the FSK look-up table. It was demonstrated that the full-spectrum k -distributions of the gas mixture at arbitrary thermodynamic states can be obtained accurately from the FSK look-up table by 6-D linear interpolation scheme presented in the paper.
3. The major advantage of the FSK look-up table is efficiency, requiring CPU time that is even cheaper than that using correlations with the simple (and least accurate) rapid multiplication model but achieving almost the same accuracy as using the LBL database. Considering both accuracy and efficiency, the FSK look-up table appears to be a very effective tool for practical applications. The

look-up database is available from the corresponding author's website upon request at <http://eng.ucmerced.edu/people/mmodest>.

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