

Fe I oscillator strengths of lines of astrophysical interest

A. Bard, A. Kock^{***}, and M. Kock^{**}

Institut für Plasmaphysik, Universität Hannover, Callinstrasse 38, W-3000 Hannover 1, Federal Republic of Germany

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Abstract. Oscillator strengths of 114 Fe I lines in the wavelength range 260–1170 nm are presented which were obtained by means of emission measurements on a high-current hollow-cathode and by using a high-resolution Fourier-transform spectrometer, in conjunction with literature data of lifetime that are considered reliable. In order to comprise a large number of lines of astrophysical interest we have included some branching fractions from Bridges & Kornblith; May et al., and Kock et al. in this analysis. As a result the uncertainty of the f -values typically is between 7 and 12%, and it is 33% at the most. The data are compared with literature data, especially with those of the Oxford group.

Key words: atomic and molecular data – plasmas – spectroscopy – stars: abundances – Sun (the): abundances

1. Introduction

In recent papers Holweger et al. (1990) and Pauls et al. (1990) derived solar abundances from faint Fe II lines which, however, differ by more than 50% from each other. As discussed in detail by Holweger et al. possible reasons for this discrepancy may be inadequate assumptions made in the solar model and/or uncertain f -values.

For Fe I the data situation is fairly satisfactory. Since the publication by Garz & Kock (1969) the spread of the solar iron abundance $\log N(\text{Fe})$ obtained with Fe I f -values has been from 7.6 to 7.67 where the highest figure corresponds to the very precise f -values measured by the Oxford group (see, e.g. Blackwell et al. 1970, 1972, 1979a, b, 1980, 1982).

Since there is a continuing interest in f -values of faint solar Fe I lines we have resumed the investigation of Fe I spectra, using measurement and evaluation techniques comparable to those applied by Heise & Kock (1990) for the determination of Fe II f -values. In a preliminary abundance study Holweger (1990) obtained $\log N(\text{Fe}) = 7.50$ or even less by making use of our laboratory Fe I f -values, in accordance with the result based on

our Fe II data. We have taken this surprising outcome as an incentive to investigate still more Fe I lines. In this paper we present a large set of f -values and compare them with recent literature data. As a matter of fact, we also have been looking for systematic errors that might be hidden in the results given by other authors. However, whatever error source we conceived of, the magnitude of the effects were insufficient to explain the 50% discrepancy in the solar iron abundance.

2. Experimental procedure

The state of the art method for a reliable determination of f -values of lines of astrophysical interest is the combination of measurements of lifetimes and branching fractions (see, e.g. Huber & Sandeman 1988).

As to the lifetimes, modern laser techniques allow selective population of excited states, thus avoiding cascade effects. The influence of buffer gas atoms on the measurement can be accounted for. The accuracy of such lifetime measurements is in the percent range (see, e.g. Marek et al. 1979).

The measurement of branching fractions is a straight-forward technique that avoids virtually all systematic errors. Since only lines are involved originating from a common upper level, plasma parameters need not be known, and non-equilibrium light sources can be used. An absolute radiance calibration is not required. Only three error sources have to be considered:

i) If a line is not emitted from an optically thin layer selfabsorption will falsify the result. Spatially inhomogeneous plasma layers can also cause self-reversal in the lines, especially in transitions to the ground state and to metastable states.

ii) Strongly overlapping lines, or blends, may lead to erroneous results. Misidentification is still another problem.

iii) Faint to very faint lines may be recorded with a poor S/N ratio leading to large statistical uncertainties.

We have applied a high-current hollow-cathode (HC) as the light source a detailed description of which was given by Danzmann et al. (1988). The discharge has been burned in argon (1.6 hPa) at a current of 1.5 Å. The burning voltage was about 430 V. The iron atoms were sputtered from a metallic cylinder made of very pure iron (VP 99.95%).

Most of the measurements have been performed with the high-resolution Fourier-transform spectrometer (Brault 1985) at Kitt Peak, Arizona. By adding up several runs and by using band pass filters the S/N ratio in the spectra could be increased considerably. The line intensities were calibrated with a tungsten strip lamp (Optronics) and with a Deuterium lamp (Cathodeon).

Send offprint requests to: M. Kock

* Now with Max-Planck-Institut für Astronomie, Königstuhl 17, W-6900 Heidelberg, Germany

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Table 1. Absolute oscillator strength for Fe I

λ (nm)	Transition	E_i (cm^{-1})	A_{ki} (10^4s^{-1})	$\log(gf)$	Error (%)
$z^5D_1^\circ$ $E_k = 26479.39 \text{ cm}^{-1}$ $\tau_k = 83.5 \pm 5.0 \text{ ns}$ a)					
387.8570	a^5D_2	704.00	663	-1.348	10
390.6477	a^5D_1	888.13	105	-2.144	11
392.0256	a^5D_0	978.07	261	-1.744	10
540.5774	a^5F_2	7985.80	101.0	-1.877	8
545.5609	a^5F_1	8154.73	60.0	-2.098	8
1168.9997	a^5P_1	17927.41	6.0	-2.433	21
$z^5D_0^\circ$ $E_k = 26550.50 \text{ cm}^{-1}$ $\tau_k = 85.6 \pm 5.0 \text{ ns}$ a)					
389.5654	a^5D_1	888.13	986	-1.649	9
543.4522	a^5F_1	8154.73	176	-2.109	8
1159.3606	a^5P_1	17927.41	6.0	-2.930	33
$z^5F_4^\circ$ $E_k = 27166.84 \text{ cm}^{-1}$ $\tau_k = 64.9 \pm 4.0 \text{ ns}$ b)					
367.9911	a^5D_4	0.00	149	-1.564*	12
373.7129	a^5D_3	415.93	1380	-0.584*	12
493.9685	a^5F_5	6928.28	1.47	-3.315	9
505.1633	a^5F_4	7376.78	5.16	-2.750	9
514.2927	a^5F_3	7728.07	2.33	-3.080+	9
658.1211	a^3F_4	11976.26	0.0358	-4.679	10
$z^5F_3^\circ$ $E_k = 27394.70 \text{ cm}^{-1}$ $\tau_k = 65.1 \pm 4.0 \text{ ns}$ a)					
370.5564	a^5D_3	415.93	353	-1.294*	12
374.5559	a^5D_2	704.00	1170	-0.764*	12
499.4129	a^5F_4	7376.78	4.23	-2.956	9
508.3338	a^5F_3	7728.07	5.66	-2.814+	9
515.0838	a^5F_2	7985.80	3.57	-3.003	9
673.9523	a^3F_3	12560.95	0.0337	-4.794	15
$z^3P_2^\circ$ $E_k = 33946.97 \text{ cm}^{-1}$ $\tau_k = 37.8 \pm 2.0 \text{ ns}$ c)					
298.1442	a^5D_3	415.93	683	-1.342	9
300.7279	a^5D_2	704.00	286	-1.712	9
302.4030	a^5D_1	888.13	461	-1.500	9
381.2961	a^5F_3	7728.07	897	-1.010	10
385.0814	a^5F_2	7985.80	162	-1.745	9
387.6036	a^5F_1	8154.73	13.0	-2.835	10
616.3542	a^5P_2	17727.02	0.844	-3.619	8
624.0645	a^5P_1	17927.41	2.00	-3.233	7
642.1348	a^3P_2	18378.22	37.0	-1.945	7
694.5196	a^3P_1	19552.49	10.16	-2.435	7
899.9560	b^3P_2	22838.36	79.0	-1.321	7
908.8328	b^3P_1	22946.86	17.0	-1.986	7

Table 1 (continued)

λ (nm)	Transition	E_i (cm^{-1})	A_{ki} (10^4s^{-1})	$\log(gf)$	Error (%)
$z^3P_1^o$ $E_k = 34362.89 \text{ cm}^{-1}$ $\tau_k = 103.1 \pm 8.0 \text{ ns}$ c)					
297.0116	a^5D_2	704.00	274	-1.964	10
298.6454	a^5D_1	888.13	20.0	-3.088	11
299.4500	a^5D_0	978.07	159	-2.192	11
379.0092	a^5F_2	7985.80	292	-1.724	11
381.4522	a^5F_1	8154.73	73.0	-2.322	10
608.2714	a^5P_1	17927.41	1.53	-3.593	9
625.4262	a^3P_2	18378.22	20.0	-2.443	9
675.0150	a^3P_1	19552.49	12.0	-2.608	9
697.8863	a^3P_0	20037.86	15.0	-2.484	9
867.4761	b^3P_2	22838.36	47.0	-1.800	9
875.7208	b^3P_1	22946.86	25.0	-2.059	10
883.8446	b^3P_0	23051.79	25.0	-2.050	10
$z^3P_0^o$ $E_k = 34555.64 \text{ cm}^{-1}$ $\tau_k = 109.8 \pm 6.0 \text{ ns}$ c)					
296.9356	a^5D_1	888.13	423	-2.252	9
378.6672	a^5F_1	8154.73	322	-2.160	9
601.2204	a^5P_1	17927.41	1.69	-4.038	8
666.3428	a^3P_1	19552.49	53.0	-2.453	7
861.1804	b^3P_1	22946.86	107.0	-1.926	7
$z^5G_4^o$ $E_k = 35257.35 \text{ cm}^{-1}$ $\tau_k = 11.2 \pm 0.7 \text{ ns}$ a)					
283.5454	a^5D_4	0.00	60.0	-2.184	10
286.9305	a^5D_3	415.93	159	-1.752	10
358.5704	a^5F_4	7376.78	668	-0.936 ^{f)}	11
363.1462	a^5F_3	7728.07	4530	-0.094	13
429.4124	a^3F_4	11976.26	435	-0.966	11
440.4749	a^3F_3	12560.95	3000	-0.105	11
639.3604	a^3H_5	19621.04	67.0	-1.432	9
646.2729	a^3H_4	19788.28	7.62	-2.367	9
871.3198	b^3G_5	23783.65	3.33	-2.467	16
897.5413	b^3G_4	24118.85	5.38	-2.233	11
$y^3D_3^o$ $E_k = 38175.38 \text{ cm}^{-1}$ $\tau_k = 6.9 \pm 0.5 \text{ ns}$ d)					
261.8708	a^5D_4	0.00	475	-1.466	12
264.7555	a^5D_3	415.93	144	-1.976	12
266.7910	a^5D_2	704.00	32.0	-2.623	16
381.5839	a^3F_4	11976.26	11000	0.226*	13
390.2944	a^3F_3	12560.95	2410	-0.414*	13
396.6060	a^3F_2	12968.57	190	-1.504*	13
488.9001	a^5P_2	17727.02	16.0	-2.387	11
504.9820	a^3P_2	18378.22	169	-1.344+	10
570.1546	b^3F_4	20641.14	22.0	-2.131	10

Table 1 (continued)

λ (nm)	Transition	E_i (cm^{-1})	A_{ki} (10^4s^{-1})	$\log(gf)$	Error (%)
577.8457	b^3F_3	20874.52	1.04	-3.440	12
618.0206	a^3G_4	21999.17	6.32	-2.596	10
651.8370	b^3P_2	22838.36	7.78	-2.460	10
711.2172	b^3G_4	24118.85	1.89	-2.998	24
722.3663	c^3P_2	24335.80	11.0	-2.225	10
836.5655	a^3D_3	26225.03	12.0	-2.047	16
$y^3D_2^o$ $E_k = 38678.07 \text{ cm}^{-1}$ $\tau_k = 7.1 \pm 0.5 \text{ ns}$ d)					
261.2770	a^5D_3	415.93	138	-2.150	13
263.2591	a^5D_2	704.00	198	-1.987	12
264.5419	a^5D_1	888.13	59.0	-2.510	15
382.7830	a^3F_3	12560.95	10800	0.075*	13
388.8520	a^3F_2	12968.57	2570	-0.535*	13
477.1696	a^5P_2	17727.02	3.30	-3.249	20
492.4769	a^3P_2	18378.22	31.0	-2.256	11
522.7146	a^3P_1	19552.49	210	-1.367	23
561.5300	b^3F_3	20874.52	15.0	-2.453	10
608.5259	a^3G_3	22249.46	2.80	-3.110	12
631.1503	b^3P_2	22838.36	2.34	-3.156	10
635.5034	b^3P_1	22946.86	14.0	-2.365+	10
829.3490	a^3D_2	26623.73	13.0	-2.190	18
$y^3D_1^o$ $E_k = 38995.76 \text{ cm}^{-1}$ $\tau_k = 7.2 \pm 0.7 \text{ ns}$ e)					
261.0748	a^5D_2	704.00	148	-2.342	19
262.3364	a^5D_1	888.13	341	-1.976	16
262.9570	a^5D_0	978.07	200	-2.207	18
384.1050	a^3F_2	12968.57	13000	-0.065*	16
514.1735	a^3P_1	19552.49	91	-1.964	15
527.3377	a^3P_0	20037.86	87	-1.965+	13
556.7393	b^3F_2	21039.02	20.0	-2.564	13
622.9232	b^3P_1	22946.86	9.0	-2.805	20
627.0228	b^3P_0	23051.79	19.0	-2.464	13
794.1085	a^3D_1	26406.49	18.0	-2.286	17
e^5D_4 $E_k = 44677.01 \text{ cm}^{-1}$ $\tau_k = 15.6 \pm 0.9 \text{ ns}$ a)					
461.1185	$z^7F_4^o$	22996.69	7.0	-2.720	19
476.8398	$z^7P_4^o$	23711.47	11.0	-2.461	14
532.4181	$z^5D_4^o$	25900.00	2060	-0.103	7
539.3170	$z^5D_3^o$	26140.19	491	-0.715	7
561.5646	$z^5F_5^o$	26874.56	2640	0.050	7
570.9382	$z^5F_4^o$	27166.84	213	-1.028	7
578.4660	$z^5F_3^o$	27394.70	6.51	-2.532	7
640.0005	$z^5P_3^o$	29056.34	928	-0.290	7

Table 1 (continued)

λ (nm)	Transition	E_i (cm^{-1})	A_{ki} (10^4s^{-1})	$\log(gf)$	Error (%)
e^5D_3					
	$E_k = 45061.33 \text{ cm}^{-1}$		$\tau_k = 15.8 \pm 0.9 \text{ ns}$	a)	
478.7828	$z^7P_3^{\circ}$	24180.88	12.0	-2.530	26
521.7391	$z^5D_4^{\circ}$	25900.00	298	-1.070	7
528.3623	$z^5D_3^{\circ}$	26140.19	1262	-0.432	7
533.9931	$z^5D_2^{\circ}$	26339.71	753	-0.647	7
558.6759	$z^5F_4^{\circ}$	27166.84	2450	-0.096	7
565.8818	$z^5F_3^{\circ}$	27394.70	479	-0.793	7
571.2134	$z^5F_2^{\circ}$	27559.60	30.0	-1.990	7
624.6322	$z^5P_3^{\circ}$	29056.34	452	-0.733	7
641.1649	$z^5P_2^{\circ}$	29469.03	589	-0.595	7
e^5D_2					
	$E_k = 45333.88 \text{ cm}^{-1}$		$\tau_k = 15.7 \pm 0.9 \text{ ns}$	a)	
520.8596	$z^5D_3^{\circ}$	26140.19	623	-0.897	7
526.3308	$z^5D_2^{\circ}$	26339.71	636	-0.879	7
530.2302	$z^5D_1^{\circ}$	26479.39	904	-0.720	7
557.2844	$z^5F_3^{\circ}$	27394.70	2280	-0.275	7
562.4545	$z^5F_2^{\circ}$	27559.60	741	-0.755	7
565.8534	$z^5F_1^{\circ}$	27666.36	58.0	-1.856	8
614.1735	$z^5P_3^{\circ}$	29056.34	123.0	-1.459	7
630.1501	$z^5P_2^{\circ}$	29469.03	643	-0.718	7
640.8020	$z^5P_1^{\circ}$	29732.75	312	-1.018	7

a) Marek et al. (1979)

b) Averaged Lifetimes of Marek et al. (1979) and Kwiatkowski et al. (1983)

c) Averaged Lifetimes of Marek et al. (1979) and Figger et al. (1975)

d) Kwiatkowski et al. (1983). Because the authors didn't give any errors for their lifetime measurements, we estimated the errors to be 7% which is typical for laser fluorescence experiments of this type.

e) Andersen and Sørensen (1971)

f) Possibly blended in our spectra.

*) Lines for which ratios of f-values have been used which were taken from other authors (Bridges and Kornblith (1974), May et al. (1974), Kock et al. (1984)).

+) Reference lines used to link the data of the present study to those of the other authors.

The influence of selfabsorption in the lines was checked with two different methods:

i) By varying the discharge current the intensity ratios of lines with a common upper level have to be constant. This method has been applied to measurements in our laboratory because we were not able to resolve the line profiles with our grating monochromator.

ii) With the FTS the purely Doppler broadened lines of the HC are resolved so that their profiles can be used for a test: Lines affected by selfabsorption are broadened with respect to optically thin lines. We have fitted Gaussian profiles to the lines. If necessary the intensities have been corrected to optically thin layer by the following formula:

$$I_c = I\alpha/(1 - \exp(-\alpha)),$$

where α is the measured optical depth. Maximum correction of line intensity was no more than 10% for a few critical lines.

Because of the non-overlapping band filters some connecting measurements have been performed in our laboratory. The equipment is described elsewhere (Heise & Kock 1990; Bard 1989).

3. Data reduction

The Fe I spectra were analysed by means of the tables given by Moore (1952). We used the semiempirical calculations by Kurucz

& Peytremann (1975) for blend searching and for checking the contribution of faint lines not measured in an actual run.

For the highly resolved FTS spectra the area under a lines was determined by fitting a Gaussian function to the line profile. As mentioned above, we have corrected a few lines for selfabsorption. For the evaluation of the broadband scans the apparatus profile of the FTS, which is simply a sinc-function, has been taken into account.

We have measured a few connecting lines with our 2 m grating monochromator whose apparatus function with a width of 45 mÅ is too large to resolve the HC lines. Nevertheless, the profile could be approximated sufficiently well by a Gaussian one too, so that the same procedure has been applied.

A few line ratios (labelled in Table 1) have been taken from Bridges & Kornblith (1974), May et al. (1974) as well as from Kock et al. (1984) in order to increase the set of lines applicable for a solar abundance study. Such line ratios derived from literature data will not be affected by systematic errors other than those of types (i) through (iii) mentioned above. Indeed, there is no indication of a systematic error in the respective data.

As a result we present a set of more than 20 lines in Table 1 which can be applied for an abundance study. The uncertainty in the f -values is typically 7–33% depending on the line and on the way in which it has been linked.

4. Results and discussion

For the 114 Fe I lines listed in Table 1 we have used published lifetime data for the absolute calibration (data and references are given in the table).

We have compared our $\log(gf)$ values with literature data. Thus, mutual deviations of the data sets will be characterized by the mean difference Δ , which is a measure of the shift in the absolute scales. The average deviation in the relative scales will be given by an rms scatter σ .

The semi-empirical calculations of Kurucz & Peytremann (1975) yield smaller f -values ($\Delta = 0.24$ dex) compared to our data. The mutual scatter ($\sigma = 0.31$ dex) is very large.

Bridges & Kornblith (1974) have performed refined measurements on a cascaded arc. Their data set has been enlarged considerably by May et al. (1974) who performed only relative

measurements on a free-burning arc. A comparison with our $\log(gf)$ values is given in Fig. 1. Despite a larger scatter ($\sigma = 0.09$ dex) the data are consistent. We cannot find systematic errors.

Of special interest is a comparison with the NBS compilation (Fuhr et al. 1988) because a number of literature data are put together whereby the basic set is that of the Oxford group. Other sets have been renormalized to those precise f -values. The comparison is given in Fig. 2. The difference of the $\log(gf)$ values is plotted versus the energy of the lower states of the line transitions. Even though there is a large scatter ($\sigma = 0.12$ dex) the f -values are obviously dependent on the excitation energy of the lines. An explanation of such a systematic error is hardly to be expected from the NBS compilation; it should rather follow from a comparison with the basic set.

In Fig. 3 this comparison is made with the data of the Oxford group. Besides a small shift in the absolute scale ($\Delta = 0.034$ dex) and a remarkably small scatter ($\sigma = 0.06$ dex) no energy dependence can be seen. However, for sets like the lines from the level $z^3 P_2^0$ the same energy dependence is found as in Fig. 2. Kock et al. (1984) also described a systematic error of this amount and traced it back to a slight temperature bias in the experiments of the Oxford group. Yet we cannot find an error in the procedure by which the Oxford group has determined the axial temperature profile in their oven.

Diffusion, or rather the neglect of diffusion, may however give rise to a systematic error producing a dependence of the f -values on the excitation potential of the lines under consideration. It is clearly stated by Blackwell & Collins (1972): "In calculating the oscillator strength from the measured equivalent width it is necessary to integrate an expression of the form $n(T) \exp(-E/kT)$ along the tube." In other words: Not only the temperature distribution but also the local iron density $n(T)$ have to be known. Blackwell & Collins made use of the vapour pressure curve given by Nesmeyanov (1961) under the assumption that diffusion is not substantial. In our opinion, however, axial temperature gradients will cause diffusion of iron atoms to the cooler regions, thus increasing the iron density at the ends of the vapour column.

We took the temperature distribution and tube geometry from the papers of the Oxford group and applied a simple diffusion

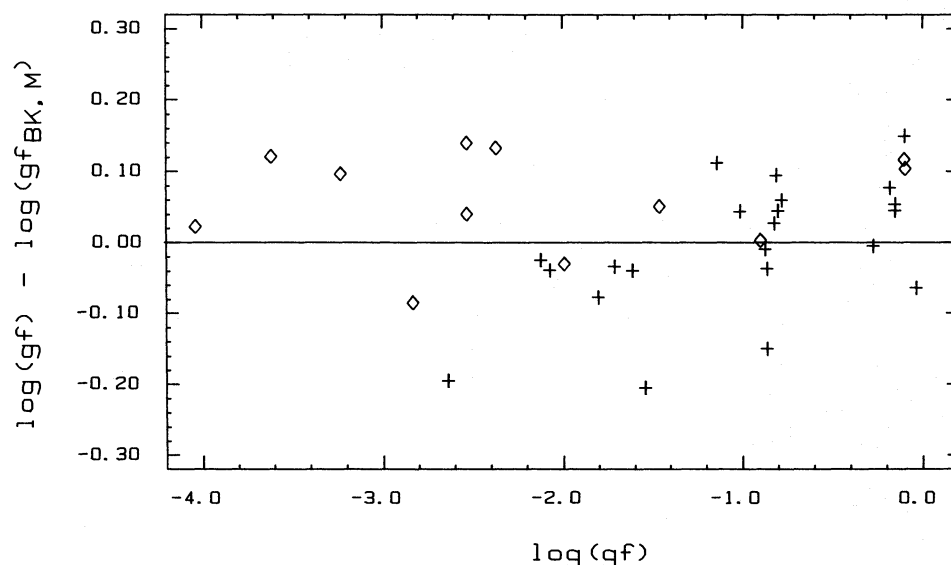


Fig. 1. Comparison of our $\log(gf)$ values with the data (+) of Bridges & Kornblith (1974) and the data (◇) of May et al. (1974) who have linked their f -values to the values of Bridges & Kornblith. For both sets the scatter is $\sigma = 0.02$ dex. The mutual deviations are characterized by $\Delta(+)=0.005$ dex and $\Delta(\diamond)=0.055$ dex

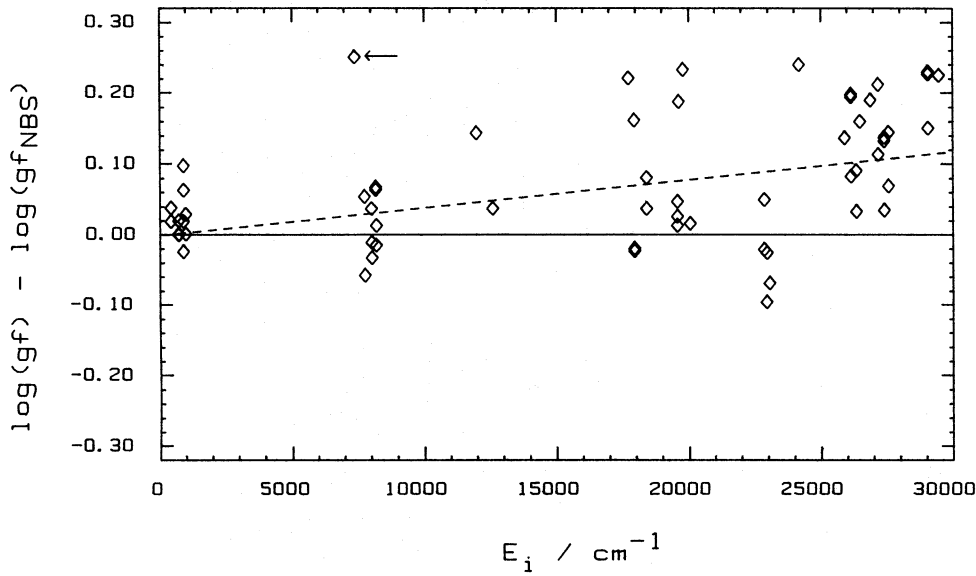


Fig. 2. Comparison of our $\log(gf)$ values with the data compiled by Fuhr et al. (1988). The difference is shown versus the energy of the lower states of the line transitions. The dashed line has been obtained by a linear regression. The point marked by an arrow refers to the line 358.57 nm which is possibly blended in our laboratory spectra (see also Fig. 3)

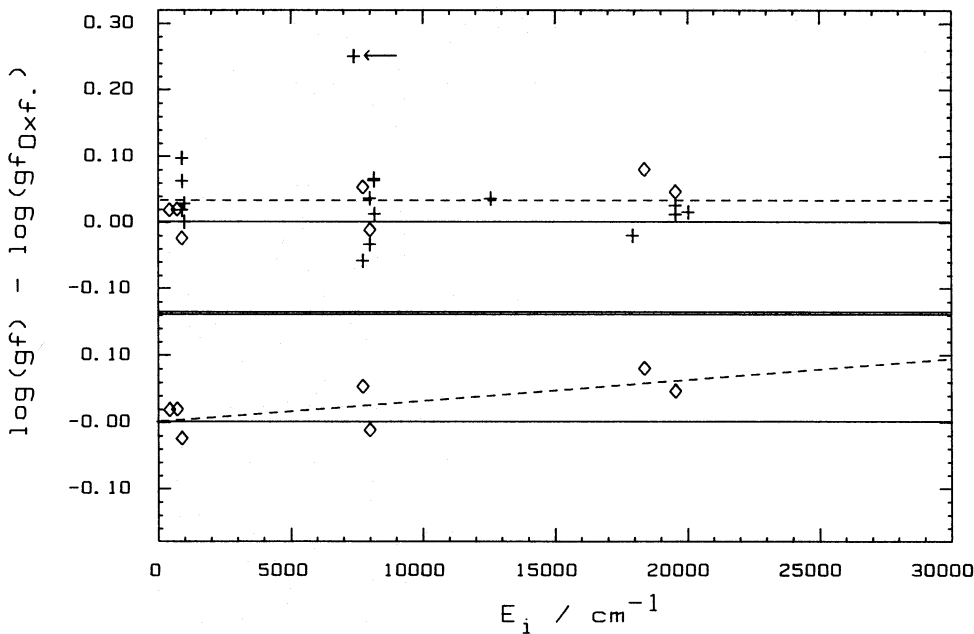


Fig. 3. Comparison of our $\log(gf)$ values with the data of the Oxford group (Blackwell et al. 1970, 1972, 1979a, b, 1980, 1982). For both frames the difference is shown versus the energy of the lower states of the line transitions. In the upper frame all lines in common are compared and the lines from the $z^2 P^0_2$ - level are marked thus: $\diamond$. The data point marked by an arrow belongs to the same line 358.57 nm as in Fig. 2. The dashed line gives the mean difference between both sets ($\Delta = 0.034$). In the lower frame a comparison is made only with lines having the $z^3 P^0_2$ - level in common. There is nearly the same energy dependence as in Fig. 2

model (Chen 1974). We then found a cosine-like decrease of the axial density instead of an exponential relationship used by the Oxford group. The resulting dependence of the f -values on the excitation potential of the lines readily matches the data in Figs. 2 and 3, as well as the result given by Kock et al. (1984). For the 2.7 eV lines, e.g., this would mean a 12% correction in the f -values of the Oxford group and hence a reduction in their solar iron abundance by 0.05 dex. That is still within our mutual scatter but clearly not sufficient to explain the 50% discrepancy.

It should be mentioned, that a deviation from LTE in the Oxford oven as well as a slightly different interpretation of the experimental vapour pressure data would also give contributions in the "right direction", namely larger f -values for lines with higher excitation potential. On the other hand, the f -values of our weak lines would be too large and hence the solar abundance too small, if we should have underestimated self-absorption in our strong lines.

5. Conclusion

By combining branching fraction measurements with reliable lifetime data we present a set of 114 Fe I f -values. More than 20 of the measured lines are well suited for a solar iron abundance study.

The agreement with the data of other authors is within the mutual limits of uncertainty. Although the agreement with the data of the Oxford group is by far the best we cannot rule out a possible small systematic error, depending on the excitation potential of the lines, which we trace back to axial diffusion of iron atoms in the Oxford oven. The large discrepancy of 50% in the solar iron abundance as published recently, however, can be explained in part by this possible error source. We conclude that at present a full solution for the discrepancy is still pending.

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