

# 1 Cascade emission in electron beam ion trap plasma of 2 $W^{25+}$ ion

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## 9 Abstract

10 Spectra of the  $W^{25+}$  ion are studied using the collisional-radiative model  
11 (CRM) with an ensuing cascade emission. It is determined that the cascade  
12 emission boosts intensities only of a few lines in the 10 – 30 nm range. The  
13 cascade emission is responsible for the disappearance of structure of lines at  
14 about 6 nm in the electron beam ion trap plasma. Emission band at 4.5 to  
15 5.3 nm is also affected by the cascade emission. The strongest lines in the  
16 CRM spectrum correspond to  $4d^9 4f^4 \rightarrow 4f^3$  transitions, while  $4f^2 5d \rightarrow 4f^3$   
17 transitions arise after the cascade emission is taken into account.

18 *Keywords:* Electron beam ion trap, collisional-radiative modelling, cascade  
19 emission, tungsten

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## 1. Introduction

Tungsten emission has been intensively studied over the last few decades due to its application in fusion devices [1]. The intense lines in tungsten spectra are observed at around 5 nm (mostly of N-shell ions) where a large number of transitions from several charge states contributes to the plasma emission. This region has attracted great attention due to its importance for the plasma power balance and possible diagnostic applications. The 10 – 30 nm region has been also investigated in the fusion and electron beam ion trap (EBIT) device plasma [2, 3, 4, 5, 6, 7]. In the fusion plasma, a complex structure of lines is observed in this region. Surprisingly, the EBIT spectra corresponding to the  $W^{15+}$  –  $W^{28+}$  ions feature only a few lines in the 13 – 18 nm wavelength range [5]. The emission from many tungsten ions contributes to the line of sight measurements in the fusion plasma, thus, such spectra contain many lines from the different ions. On the other hand, the EBIT devices provide an unique opportunity to study the emission from one or several neighboring ions. Therefore, analysis of their spectra is much easier compared with those from other plasma sources. The emission originating only from several ions can be the reason why the EBIT spectra are sparse of the lines in the 13 – 18 nm range. However, the corona modeling of the spectral line intensities provides complex structure of the lines for  $W^{25+}$  [8] in the aforementioned range. Since these calculations contradict the EBIT observations, it is necessary to check what kind of spectra corresponds to the collisional-radiative modeling (CRM). On the other hand, it is shown that the cascade emission boosts the intensities only for some lines of the  $W^{13+}$  ion in the EBIT plasma [9]. It has to be noted that the term “cascade emission” is used here instead of the radiative cascade in order to distinguish population of the levels from the higher-lying levels through the radiative cascade which is accounted in the corona model. The different population mechanisms appear on the scene in these two cases [9]. The cascade processes were mostly studied for the radiative and Auger decays when an inner-shell vacancy was created [10, 11, 12, 13].

Ions in the EBIT move in cycloidal orbits spending part of their time outside electron beam [14]. The cascade emission starts when the ions leave the electron beam and the interaction with the electrons ends. This effect is more pronounced for the ions in the low or intermediate ionization stages [15]. It was found that under the same conditions, the higher charge ions show less expansion in the radial direction. When the charge state of the ions

increases, the Coulomb's attraction force directed toward the electron beam also increases. It leads to the decrease of the time the ions spend outside the beam where the cascade emission depopulates the excited levels. The effective electron density is often introduced in order to reduce electron-impact collision rates [15, 16]. The time fraction which the ions spend inside and outside the electron beam depends on many parameters, such as ion temperature, electron beam energy, electron beam current, electric and magnetic fields. On the other hand, the range of the ion radius  $r_i$  ratio against the geometric electron radius  $r_e$  can be expressed through the effective charge  $Z_{eff}$  of the ion:  $1/(Z_{eff}/Z)^\alpha$  with  $1 < \alpha \leq 2$  [15]. For  $W^{25+}$ , one can estimate that the main paths of the ions span outside the electron beam:  $r_i/r_e \approx 3^\alpha$ .

The main aim of the current work is to study the emission spectra of the  $W^{25+}$  ion in the EBIT plasma by performing the CRM with ensuing cascade emission. The emission from  $W^{25+}$  has not deserved wide attention so far since calculations are complicated due to the open  $f$  shells. Systems with the open  $f$  shells is further of interest for the study of the complex multi-electron high- $Z$  ions. The present work focuses on the 2 – 30 nm region which accumulates the main emission from the  $W^{25+}$  ion [8]. As far as we know, influence of the cascade emission on the formation of lines in the EBIT plasma has not been studied for this ion before. Previous works concentrated on analysis of the spectral lines obtained from the CRM or the corona model [3, 8]. It was shown that the corona model is suitable for the low density EBIT plasma [17]. The CRM calculations included the  $4f^3$  and  $4f^25l$  ( $l = 0, 1, 2, 3$ ) configurations but omitted the important the  $4d^94f^4$  and  $4f^25g$  configurations [3]. Only the strongest lines were presented in their work. The current study has been extended to 19612 levels compared with 13937 levels used in the corona model calculations [8].

The rest of the paper is organized as follows. In the next section we present theoretical methods used to calculate atomic data and emission spectra. In Section 3, the determined emission spectra corresponding to the CRM and the cascade emission are discussed.

## 2. Theoretical methods

Energy levels, radiative transition probabilities, and electron-impact excitation rates for  $W^{25+}$  have been calculated using Flexible Atomic Code (FAC) [18] which implements the relativistic Dirac-Fock-Slater method. Previous study included 22 configurations [8] while the current work employs 43 con-

93 configurations:  $4f^3$ ,  $4f^25l$  ( $l = 0, 1, 2, 3, 4$ ),  $4f^26l'$ ,  $4f^27l'$  ( $l' = 0, 1, 2, 3, 4, 5$ ),  
 94  $4f^28l$ ,  $4d^94f^4$ ,  $4d^94f^35l''$  ( $l'' = 0, 1, 2$ ),  $4d^94f^25s^2$ ,  $4d^84f^5$ ,  $4f5s^2$ ,  $4f5s5l'''$   
 95 ( $l''' = 1, 2, 3, 4$ ),  $4f5p^2$ ,  $4f5p5d$ ,  $4f5s6l$ ,  $4p^54f^4$ ,  $4p^54f^35s$ . These configura-  
 96 tions produce 19612 levels. Configuration interaction has been taken into  
 97 account for all the considered configurations. The radiative transition prob-  
 98 abilities have been calculated for electric dipole, quadrupole, and octupole  
 99 and for magnetic dipole and quadrupole transitions.

100 Electron-impact excitation cross-sections are obtained within the dis-  
 101 torted wave approximation. Collision rates are calculated for 790 eV electron  
 102 beam energy which corresponds to the energy used in the spectra measure-  
 103 ments [5]. The Gaussian distribution function with a full width at a half-  
 104 maximum of 30 eV is used for the electron energy.

105 Populations of levels in the CRM have been obtained by solving the sys-  
 106 tem of coupled rate equations

$$\frac{dn_i(t)}{dt} = N_e \sum_k n_k(t) C_{ki} + \sum_{k>i} n_k(t) A_{ki}^r - N_e n_i(t) \sum_k C_{ik} - n_i(t) \sum_{j<i} A_{ij}^r \quad (1)$$

107 in the steady-state equilibrium approximation ( $\frac{dn_i}{dt} = 0$ ). Here  $n_i$  is the  
 108 population of the level  $i$ ,  $A_{ij}^r$  is the radiative transition probability from the  
 109 level  $i$  to the level  $j$ , and  $C_{ik}$  is the electron-impact excitation rate from the  
 110 level  $i$  to the level  $k$ ,  $N_e$  is the electron density ( $N_e = 1 \times 10^{12} \text{ cm}^{-3}$ ).

111 Total populations of the levels during the cascade emission can be found  
 112 by summation of the population in every step of the cascade:

$$n_i^{j+1} = \sum_{m>i} \frac{n_m^j A_{mi}^r}{\sum_{k<m} A_{mk}^r}, \quad (2)$$

113 where  $n_i^j$  corresponds to the population of the level  $i$  in  $j$  step of the cascade.  
 114 By the step of the cascade, we mean all possible radiative transitions from  
 115 every not zero-populated level to the other levels. Thus, transfer of the  
 116 population through the intermediate levels is not included in the single step.  
 117 Equation (2) means that radiative transition from the level  $m$  to the level  $i$   
 118 transfers only part  $A_{mi}^r / \sum_{k<m} A_{mk}^r$  of the population  $n_m^j$ . The same approach  
 119 was used analyzing Auger cascades [10, 12, 13]. Since the cascade emission  
 120 takes place after ions leave electron beam, the initial population of the levels  
 121 for the first step of the cascade is determined from the CRM.

122 Equation (2) determines the populations of the levels when all the higher-  
 123 lying levels are depopulated by the radiative decay. However, fraction of

124 depopulation strongly depends on the time which the ions spend outside the  
 125 electron beam. In this case, the population of the levels has to be determined  
 126 by solving the time-dependent rate equations which omits interaction with  
 127 the electrons:

$$\frac{dn_i(t)}{dt} = \sum_{k>i} n_k(t) A_{ki}^r - n_i(t) \sum_{j<i} A_{ij}^r. \quad (3)$$

128 The total population  $n_i(\Delta t)$  which leaves the level  $i$  during the time  
 129 interval  $\Delta t$  is found by integrating expression:

$$\frac{dn_i(t)}{dt} = n_i(t) \sum_{j<i} A_{ij}^r \quad (4)$$

130 which leads to

$$n_i(\Delta t) = \int_0^{\Delta t} \frac{dn_i(t)}{dt} dt = \int_0^{\Delta t} n_i(t) dt \sum_{j<i} A_{ij}^r. \quad (5)$$

131 Here  $\Delta t$  is the time the ions have spent outside the electron beam. Equation  
 132 (5) provides the populations for the time-integrated line intensities. The  
 133 total population obtained by summation of the population from every step  
 134 of the cascade in Eq. (2) corresponds to the integration taking  $\Delta t = \infty$  in  
 135 Eq. (5). Practically, however, convergence of the spectral line intensities  
 136 has to be obtained for the finite time values. The equation (2) is applied to  
 137 calculate the final spectra of the cascade emission. Thus, there is no need to  
 138 perform convergence check of the spectra obtained from Eq. (5).

### 139 3. Results

140 The CRM spectrum in the 2 – 30 nm range is presented in Fig. 1. The  
 141 strongest lines correspond to the  $4d^9 4f^4 \rightarrow 4f^3$  and  $4f^2 5d \rightarrow 4f^3$  transitions.  
 142 These lines form complex structure at about 5 nm. It has to be noted that  
 143 our CRM calculations succeeded to reproduce a smaller peak in the 5.5–6 nm  
 144 region. The similar peak but with larger intensity was obtained for  $W^{23+}$  in  
 145 the CRM spectra at various electron densities using Maxwellian distribution  
 146 for the electron velocities [7]. The spectra from fusion plasma contain this  
 147 additional structure of the lines [2]. The following lines mainly arise from  
 148 the  $4d^9 4f^4 \rightarrow 4f^3$  transitions in our calculations. However, this structure  
 149 is not seen in the EBIT plasma of tungsten ions [19] suggesting that some  
 150 other mechanisms are responsible for the line formation.

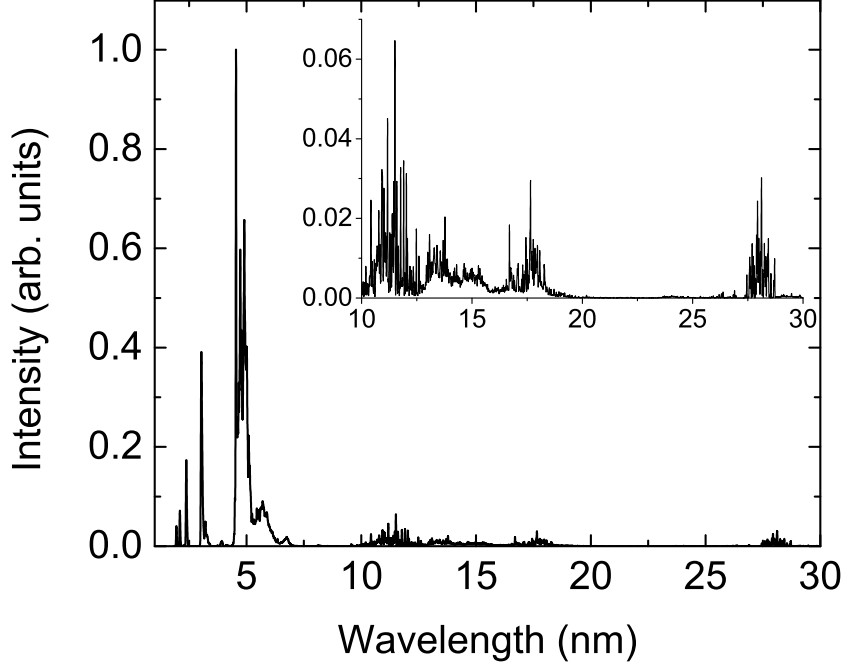


Figure 1: CRM spectrum of  $W^{25+}$ . The inset shows structure of lines in the range 10 – 30 nm.

151 The current CRM calculations (Fig. 1) and the previous results from  
 152 the corona model [9] present a complex structure of the emission lines in  
 153 the range 10 – 30 nm. In the current calculations, the number of configura-  
 154 tions has been increased to check the influence of higher-lying levels on the  
 155 formation of spectral lines. It has to be noted that the strongest lines in  
 156 the spectral range arise from the  $4f^25s \rightarrow 4f^3$  transitions which have wave-  
 157 lengths in the 10 – 12 nm region. The configuration  $4f^25s$  is the first excited  
 158 one which can decay to the ground configuration only through the electric  
 159 octupole transitions in a single-configuration approximation. Extended basis  
 160 of interacting configurations makes it possible for the electric dipole transi-  
 161 tions to occur. However, their transition probabilities are much lower than  
 162 those of other electric dipole transitions in the region. Other strong lines in  
 163 this region come from the  $4f^25f \rightarrow 4f^25d$  (12 – 16 nm),  $4f^25d \rightarrow 4f^25p$   
 164 (12 – 14, 16 – 18 nm), and  $4f^25p \rightarrow 4f^25s$  (16 – 19, 27 – 30 nm) transitions.  
 165 Unfortunately, the EBIT spectra exhibit just a few lines in the spectral  
 166 range from 13 to 18 nm [5]. As the theoretical spectra contain the com-  
 167 plex structure of lines compared with the observations, it was suggested that

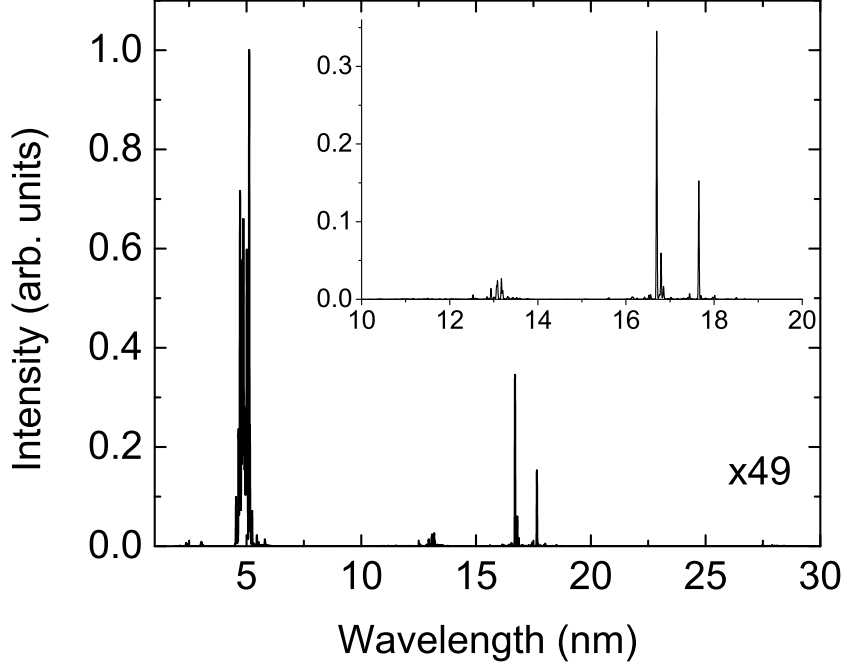


Figure 2: Cascade emission spectrum of  $W^{25+}$ . The inset shows structure of lines in the 10 – 20 nm range. The factor shows an increase of the line intensities compared to the CRM spectrum.

168 cascade emission process, which starts after ions leave the electron beam,  
 169 could be important in the formation of the spectral lines. It has been previ-  
 170 ously demonstrated that the cascade emission highlights only a few lines in  
 171 the spectrum [9]. However, such an effect has been determined for the low  
 172 ionization stage,  $W^{13+}$ . As was mentioned above, influence of the cascade  
 173 emission has to be larger for the lower ionization stages compared to the in-  
 174 termediate charge states, which, as far as we know, have never been studied  
 175 using the cascade emission process.

176 Figure 2 shows that the cascade emission highlights several lines in the  
 177 range 10 – 30 nm for the  $W^{25+}$  ion. In this case, the population of levels is  
 178 obtained using Eq. (2). In our view, the presented results demonstrate the  
 179 validity of our idea that the cascade emission is responsible for line formation  
 180 in the EBIT spectra. The strongest lines correspond to the  $4f^25d \rightarrow 4f^25p$   
 181 and  $4f^25p \rightarrow 4f^25s$  transitions in  $W^{25+}$  among the levels with high  $J$  values  
 182 (Table 1). Due to the the significantly smaller number of such levels and  
 183 selection rules for the electric dipole transitions, the cascade emission leads

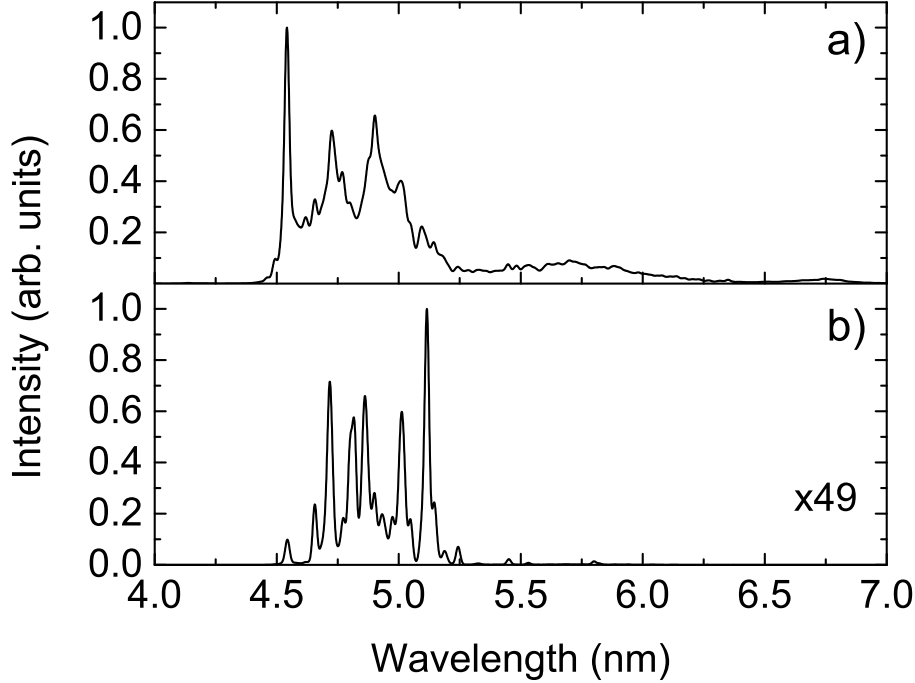


Figure 3: Theoretical spectra of the  $W^{25+}$  ion obtained a) from the CRM and b) from the CRM with ensuing emission cascade [Eq. (2)] in the 4 – 7 nm spectral range. The factor shows an increase of the line intensities compared to the CRM spectrum.

184 to the concentration of intensity.

185 The EBIT experiment for the  $W^{25+}$  ion also contains a few strong lines  
186 but with shorter wavelengths by about 2 nm compared with our calculations.  
187 The discrepancy between the theoretical and experimental wavelengths  
188 shows that important correlation effects are not taken into account for the  
189  $4f^25d \rightarrow 4f^25p$  and  $4f^25p \rightarrow 4f^25s$  transitions. The importance of the  
190 correlation effects for tungsten ions has been illustrated for magnetic dipole  
191 transitions [20, 21] using configuration interaction strength [22, 23] to build  
192 basis of the interacting configurations. However, these calculations are very  
193 cumbersome. Furthermore, for Er-like tungsten it was found that FAC can  
194 show discrepancy for wavelengths within 2 nm of the measured values when  
195 the correlation effects are not considered [24].

196 It has to be noted that influence of the  $4f^25s \rightarrow 4f^3$  transitions on the  
197 line formation is negligible in the cascade emission spectrum. The corona  
198 model has revealed that intensities of these lines strongly increase due to



199 contributions of the higher-lying levels through radiative cascade [8].

200 The relative intensities of the lines in the 4 – 7 nm region compared with  
201 the lines at the shorter wavelength side are strongly increased in the cas-  
202 cade emission spectrum (Fig. 2) compared with the CRM spectrum (Fig.  
203 1). These lines in the 2 – 4 nm range originate from the  $4f^25g \rightarrow 4f^3$ ,  
204  $4f^26g \rightarrow 4f^3$ ,  $4f^27g \rightarrow 4f^3$ , and  $4f^28g \rightarrow 4f^3$  transitions. The previous  
205 investigation showed that strong electron-impact excitations occur from the  
206 ground configuration to the  $4f^25g$  and  $4f^26g$  configurations [8]. Populations  
207 of these configurations are not affected by the cascade emission process be-  
208 cause the  $4f^25g$ ,  $4f^26g$ ,  $4f^27g$ , and  $4f^28g$  configurations are highly excited  
209 ones. The cascade emission is responsible for increase of the spectral line  
210 intensities at 5 nm.

211 Other interesting result of modeling is formation of lines in the 4.5 – 5.3  
212 nm region (Fig. 3). It seems that the structure of these lines is not so  
213 significantly affected by the cascade emission as in the 13 – 18 nm range  
214 because the emission lines overlap in the CRM and cascade emission spectra.  
215 However, it can be seen from Fig. 3 that the cascade emission spectrum is  
216 more structured than the CRM one and distribution of the line intensities  
217 is different. The CRM calculations show that the lines in the 4.5 – 5.3  
218 nm region correspond to the  $4d^94f^4 \rightarrow 4f^3$  and  $4f^25d \rightarrow 4f^3$  transitions.  
219 Nevertheless, other levels are involved in the line formation for the cascade  
220 emission spectrum compared to the CRM calculations. Figure 4 shows how  
221 line intensities in the region changes with time. It can be seen that many  
222 strong lines disappear from the spectrum while the other line intensities are  
223 significantly increased. The strongest lines in the CRM spectrum correspond  
224 to the  $4d^94f^4 \rightarrow 4f^3$  transitions (Table 2) while the  $4f^25d \rightarrow 4f^3$  transitions  
225 dominate in the cascade emission spectrum (Table 3). One can see that the  
226 distribution of the line intensities in the CRM calculations is more smooth  
227 compared with the cascade emission data. There are only a few strong lines  
228 in the cascade emission spectrum.

229 The spectral feature of lower intensity is visible at about 5.5 nm to 6 nm  
230 in the CRM spectrum but it is not seen in the cascade emission calculations.  
231 This additional lower intensity peak is presented in the fusion spectra [2]  
232 but it disappears from the EBIT spectra [19]. The obtained results illustrate  
233 importance of the cascade emission of ions outside the electron beam in the  
234 EBIT device. To the best of our knowledge, these differences in the fusion and  
235 EBIT spectra have not been explained before. It has to be noted that group  
236 of lines in 5.5 – 6.0 nm region seen in the CRM spectrum disappears from

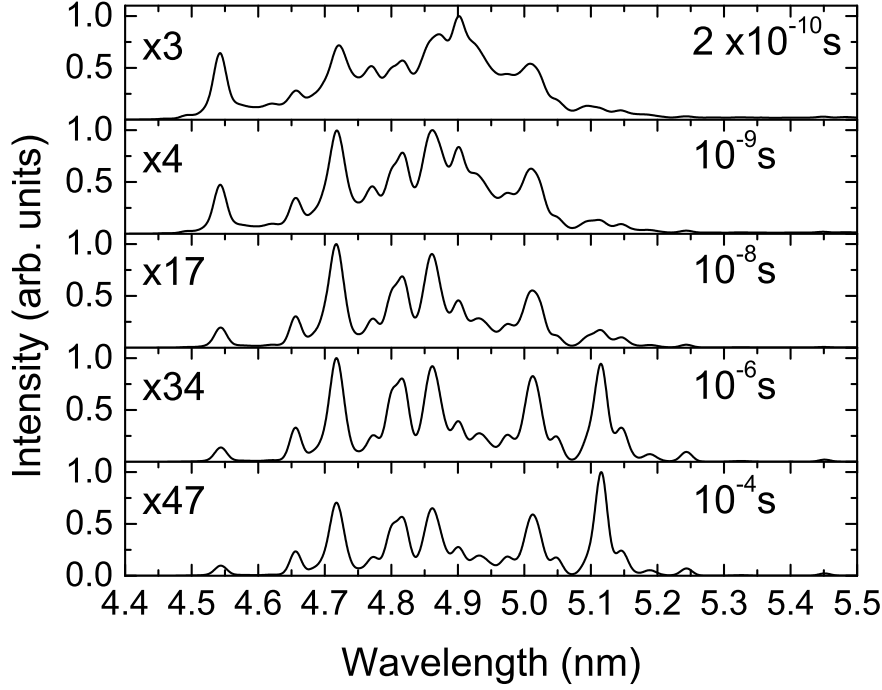


Figure 4: Time-integrated spectra of cascade emission in the 4.4 – 5.5 nm region. Times spent by ions outside the electron beam are shown. The factor shows an increase of the line intensities compared to the CRM spectrum.

the cascade emission spectrum after about  $10^{-8}$  s. The relative intensities of the lines decrease about two times after  $2 \cdot 10^{-10}$  s and four times after  $10^{-9}$  s compared to the intensity of the strongest line in the spectrum. Since this group of lines is not seen in the EBIT spectrum it implies that the ions spend outside the beam in average more than  $10^{-9}$  s.

In addition, we have estimated influence of charge exchange process on formation of spectral lines due to interaction with neutrals. The captured electron occupies a state with principal quantum number  $n_c \approx Z_{eff}^{3/4}$  ( $Z_{eff}$  is the effective charge of the ion) [25]. For the  $W^{26+}$  ion, one can derive  $n_c \approx 12$ . The angular momentum of the captured electron is defined by the ion charge and the relative collision velocity  $v$  (in atomic units):  $l = (5Z_{eff})^{0.5}v$  [25]; that leads to  $l = 0$  for the considered collision energy. Again, the cascade emission from the  $4f^212s$  configuration gives a large number of the lines in the 13 – 18 nm range. It indicates that the charge exchange process is not important for the formation of the spectral lines from the  $W^{25+}$  ion in the

EBIT plasma. The same result was obtained for the  $W^{13+}$  ion [9] and for the higher ionization stages of the tungsten ions [26, 27].

#### 4. Conclusions

The CRM with ensuing cascade emission have been studied for the  $W^{25+}$  ion. It is demonstrated that the cascade emission is responsible for formation of some lines in the EBIT spectra of the  $W^{25+}$  ion. These lines correspond to transitions among the levels with high  $J$  values.

The relative intensity of lines at 5 nm is strongly increased in the cascade emission spectrum compared to the lines at the shorter wavelength side which are not affected by the cascade emission. The cascade emission produces only a few strong lines in the region while the CRM calculations give more smooth distribution for the line intensities. The strongest lines in the CRM spectrum correspond to the  $4d^9 4f^4 \rightarrow 4f^3$  transitions while many lines from the  $4f^2 5d \rightarrow 4f^3$  transitions appear in the cascade emission calculations.

The CRM gives a spectrum with a complex structure of lines in the 13 – 18 nm region contradicting the observations as well as cascade emission spectrum. Calculations show that the lines belong to the  $4f^2 5d \rightarrow 4f^2 5p$  and  $4f^2 5p \rightarrow 4f^2 5s$  transitions.

The less intense line structure observed in fusion spectra at about 6 nm is reproduced by our CRM calculations. The missing structure of the lines in the EBIT measurements is explained by the cascade emission of ions outside the electron beam. The reason of the difference between the fusion and EBIT spectra for this wavelength region has never been determined before. Time-integrated study of the line intensities gives that the ions spend in average more than  $10^{-9}$  s outside the electron beam.

Finally, our results demonstrate that the cascade emission has to be taken into account for the ions in intermediate ionization stages when the spectra from the EBIT plasma are analyzed. The CRM alone does not provide a reasonable agreement with the measurements because it omits physical processes which occur after the ions leave the electron beam region.

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Table 1: The strongest lines of the cascade emission spectrum for  $W^{25+}$  in the 10 – 20 nm wavelength range. Wavelengths  $\lambda$ , relative intensities  $I$ , and indexes of initial  $i$  and final  $f$  levels are presented.  $J$  stands for the total angular momentum quantum number.

$\lambda$ (nm)	$I$	$i$	$f$	$J_i$	$J_f$	Initial level	Final level
16.698	100	1094	107	17/2	15/2	$4f_{7/2}^2$ (6) $5d_{5/2}^1$	$4f_{7/2}^2$ (6) $5p_{3/2}^1$
17.653	54	107	51	15/2	13/2	$4f_{7/2}^2$ (6) $5p_{3/2}^1$	$4f_{7/2}^2$ (6) $5s_{1/2}^1$
16.693	25	1152	127	17/2	15/2	$4f_{5/2}^1$ $4f_{7/2}^1$ (6) $5d_{5/2}^1$	$4f_{5/2}^1$ $4f_{7/2}^1$ (6) $5p_{3/2}^1$
16.796	21	1061	99	15/2	13/2	$4f_{5/2}^1$ $4f_{7/2}^1$ (5) $5d_{5/2}^1$	$4f_{5/2}^1$ $4f_{7/2}^1$ (5) $5p_{3/2}^1$
13.173	9	1029	76	15/2	13/2	$4f_{7/2}^2$ (6) $5d_{3/2}^1$ (3)	$4f_{7/2}^2$ (6) $5p_{1/2}^1$
13.085	8	1003	69	13/2	11/2	$4f_{5/2}^1$ $4f_{7/2}^1$ (5) $5d_{3/2}^1$	$4f_{5/2}^1$ $4f_{7/2}^1$ (5) $5p_{1/2}^1$
16.852	6	1087	109	15/2	13/2	$4f_{7/2}^2$ (6) $5d_{5/2}^1$	$4f_{7/2}^2$ (6) $5p_{3/2}^1$
13.062	5	1038	76	15/2	13/2	$4f_{7/2}^2$ (6) $5d_{3/2}^1$	$4f_{7/2}^2$ (6) $5p_{1/2}^1$
12.937	5	1109	85	15/2	13/2	$4f_{7/2}^2$ (6) $5d_{3/2}^1$	$4f_{5/2}^1$ $4f_{7/2}^1$ (6) $5p_{1/2}^1$

Table 2: The strongest lines of the CRM spectrum for  $W^{25+}$  in the 4 – 7 nm wavelength range. Wavelengths  $\lambda$ , relative intensities  $I$ , and indexes of initial  $i$  and final  $f$  levels are presented.  $J$  stands for the total angular momentum quantum number.

$\lambda$ (nm)	$I$	$i$	$f$	$J_i$	$J_f$	Initial level	Final level
4.546	100	1221	31	15/2	17/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (2) 3/2 4f_{7/2}^2 (6)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.656	99	1200	31	17/2	17/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 5/2 4f_{7/2}^2 (6)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.543	91	1186	6	13/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^1 (1) 4f_{7/2}^3 (15/2)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.544	91	1176	3	11/2	13/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 5/2 4f_{7/2}^2 (6)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.536	87	1204	13	11/2	13/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^3 (9/2) 3 4f_{7/2}^1$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
4.539	84	1220	27	11/2	13/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 5/2 4f_{7/2}^2 (4)$	$4f_{5/2}^1 4f_{7/2}^2 (4)$
4.543	82	1209	18	13/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^1 (1) 4f_{7/2}^3 (15/2)$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
4.544	79	1219	29	13/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^1 (1) 4f_{7/2}^3 (15/2)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.727	78	1137	13	13/2	13/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^3 (9/2) 4 4f_{7/2}^1$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
4.682	75	1188	29	15/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^1 (1) 4f_{7/2}^3 (15/2)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.701	72	1160	18	15/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^3 (9/2) 4 4f_{7/2}^1$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
4.721	68	1174	27	13/2	13/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 5/2 4f_{7/2}^2 (4)$	$4f_{5/2}^1 4f_{7/2}^2 (4)$
4.539	66	1167	1	9/2	11/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^3 (9/2) 3 4f_{7/2}^1$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
4.733	65	1121	6	15/2	15/2	$4f_{7/2}^2 (6) 5d_{3/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.738	59	1183	34	11/2	11/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^1 (1) 4f_{7/2}^3 (11/2)$	$4f_{7/2}^3 (11/2)$
4.727	55	1095	3	13/2	13/2	$4f_{5/2}^1 4f_{7/2}^1 4 5d_{3/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.572	51	1203	12	9/2	11/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 5/2 4f_{7/2}^2 (6)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.524	49	1223	34	9/2	11/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^1 (1) 4f_{7/2}^3 (11/2)$	$4f_{7/2}^3 (11/2)$
4.751	47	1054	1	11/2	11/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^3 (9/2) (4) 4f_{7/2}^1$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
5.116	44	989	31	19/2	17/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) (7/2) 4f_{7/2}^2 (6)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$



Table 3: The strongest lines of the cascade emission spectrum for  $W^{25+}$  in the 4 – 7 nm wavelength range. Wavelengths  $\lambda$ , relative intensities  $I$ , and indexes of initial  $i$  and final  $f$  levels are presented.  $J$  stands for the total angular momentum quantum number.

$\lambda$ (nm)	$I$	$i$	$f$	$J_i$	$J_f$	Initial level	Final level
5.116	100	989	31	19/2	17/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 7/2 4f_{7/2}^2 (6)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.716	62	1121	6	15/2	15/2	$4f_{7/2}^2 (6) 5d_{5/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.818	52	1061	6	15/2	15/2	$4f_{5/2}^1 4f_{7/2}^1 5 5d_{5/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.800	43	1152	31	19/2	17/2	$4f_{5/2}^1 4f_{7/2}^1 6 5d_{5/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.864	37	1029	6	15/2	15/2	$4f_{7/2}^2 (6) 5d_{3/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
5.007	29	1038	27	15/2	13/2	$4f_{7/2}^2 (6) 5d_{3/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (4)$
4.856	26	1087	18	15/2	15/2	$4f_{7/2}^2 (6) 5d_{5/2}^1$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
4.656	25	1200	31	17/2	17/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 5/2 4f_{7/2}^2 (6)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.727	23	1095	3	13/2	13/2	$4f_{5/2}^1 4f_{7/2}^1 4 5d_{5/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
5.147	23	899	6	17/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^3 (9/2) 5 4f_{7/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
5.014	22	956	6	17/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 7/2 4f_{7/2}^2 (6)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
5.024	21	1029	27	15/2	13/2	$4f_{7/2}^2 (6) 5d_{3/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (4)$
5.106	18	956	18	17/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^2 (4) 7/2 4f_{7/2}^2 (6)$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
5.048	17	1018	29	17/2	17/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^1 1 4f_{7/2}^3 (15/2)$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.901	16	1109	31	15/2	17/2	$4f_{7/2}^2 (6) 5d_{3/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
4.701	15	1160	18	15/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^3 (9/2) 4 4f_{7/2}^1$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
4.773	13	1087	6	15/2	15/2	$4f_{7/2}^2 (6) 5d_{5/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (6)$
5.094	10	949	13	15/2	13/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^1 (4) 7/2 4f_{7/2}^2 (4)$	$4f_{5/2}^2 (4) 4f_{7/2}^1$
4.975	8	1061	27	15/2	13/2	$4f_{5/2}^1 4f_{7/2}^1 5 5d_{5/2}^1$	$4f_{5/2}^1 4f_{7/2}^2 (4)$
5.244	8	899	18	17/2	15/2	$4d_{3/2}^3 (3/2) 4f_{5/2}^3 (9/2) 5 4f_{7/2}^1$	$4f_{5/2}^2 (4) 4f_{7/2}^1$