

EMISSION IN ABSORPTION LINES: RESULTS OF THE SL9 L NUCLEUS IMPACT WITH JUPITER

MIREL BÎRLAN^{1,2}

¹ *Observatoire de Paris-Meudon, DESPA
F-92195 Meudon Cedex, France
E-mail: Mirel.Birlan@obspm.fr*

² *Astronomical Institute of the Romanian Academy
Str. Cuștilor de Argint, RO-75212 Bucharest 28, Romania*

Abstract. High-resolution spectra of the impact sites and impact of the comet Shoemaker-Levy 9 with Jupiter have been performed at the Pic-du-Midi Observatory. The excitation of several chemical elements (Fe, Ca, Ba, Na, Mn, Mg, etc.) has been identified during the analysis of the L nucleus impact spectra obtained in visible and near-IR. The article presents the atomic lines and the time evolution of nine of them.

Key words: spectroscopy – comets – atomic lines.

1. INTRODUCTION

One of the major astronomical events of 1994 was the impact of the comet Shoemaker-Levy 9 with Jupiter. The astronomical community observed this event within the framework of the coordinated program; several ground-based and space instruments have been involved. SL9/Jupiter impact was a unique event (until now); before the show “live” of the impact SL9 comes inside Jupiter’s Roche limit which broken the nucleus in 22 fragments.

The first impact for each nucleus occurred on an unfavourable geometry, on the hidden part of Jupiter, not far from the limb. However, relevant data concerning the impacts were collected as soon the impact effects become visible.

2. OBSERVATIONS

The paper presents the spectroscopy in the wavelength range of visible and near-IR (5460-8750 Å) performed for the L impact site. The observations were performed with the 2-meter Bernard Lyot telescope from the Pic du Midi Observatory. The spectra were recorded by a 1024×1024 Thomson CCD chip, and

the spectral resolution was 36 000. The fiber of the spectrograph has 50 microns, which corresponds to a field of view of 2.2". This field of view is small enough to obtain high quality spectra only from the impact site (the apparent diameter of Jupiter is about 38"). The guidance software of the telescope allows both automatic/manual tracking during the exposure.

3. DATA REDUCTION

The pre-treatment of the observed data was performed using MUSICOS software. MUSICOS makes the calibration pixel-wavelength for the intensities spectra. Each spectrum was split in several wavelength intervals (named "orders") which little overlap between the adjacent orders. For the analysis in the absorption line we choose as target on Jupiter the L impact site for July 19/20 at 22:30:55 UT (referred to as S167), and July 20/21 (referred to as S213), 1994. As reference, the Jupiter spectrum was taken on July 20/21 (referred to as R216). The spectrum S167 emission lines have already been analysed in several papers (Roos-Serote et al. 1995a,b; Barucci et al. 1995).

The MIDAS software procedures for spectroscopy were employed. The goal of this work was to check the atomic absorption line depth one day after the impact moment and to see the excitation of different atoms of Jupiter-SL9 L impact plume. To reach this goal, all the orders of both spectra of impact site were compared with the reference spectrum R216. Then the results S213/R216 and S167/R216 in each order were compared.

In this treatment, the major problem of the differential rotation of the atmosphere of Jupiter occurred. From an order to another, in different spectra, the same atomic line presents a slight shift in wavelength, following the expression:

$$\frac{\Delta\lambda}{\lambda} = \frac{v_{diff}}{c},$$

where v_{diff} denotes the differential speed of Jupiter and c stands for the speed of light.

Thus, an automatic procedure of shift cannot be taken into account. For a good preliminary result in some orders the spectra were rebined. Then, shifting the lines in such a way made the subtraction or division between spectra, so that the minima of the lines are at the same wavelength.

The cosmic ray signatures represented another problem that occurred during the treatment. This was skipped manually, each time when the spectrum of Jupiter had no lines in this region, and one given atomic line had an abnormal profile.

At least, we cannot omit the presence of the terrestrial lines, even after a major part of them were eliminated by an automatic procedure. Their presence could alter our qualitative analysis and they were carefully analysed and skipped.

4. RESULTS

The comparison of the L impact site spectra with the unperturbed Jupiter's atmosphere spectrum was made in order to minimize any ambiguity. Then the shifted spectra S167 and S213 were compared in each order. To obtain a good signature for each excited atomic line, the ratio S167/S213 has been analysed. As presented in Fig. 1, only the signatures of atomic lines with amplitude larger than three times the noise amplitude were considered (three sigma relevance).

The analysis reveals orders on which the spectral lines were not perturbed by the impact of the comet (as seen on Fig. 2). At the opposite, there are spectral intervals where almost all of the atomic lines were perturbed.

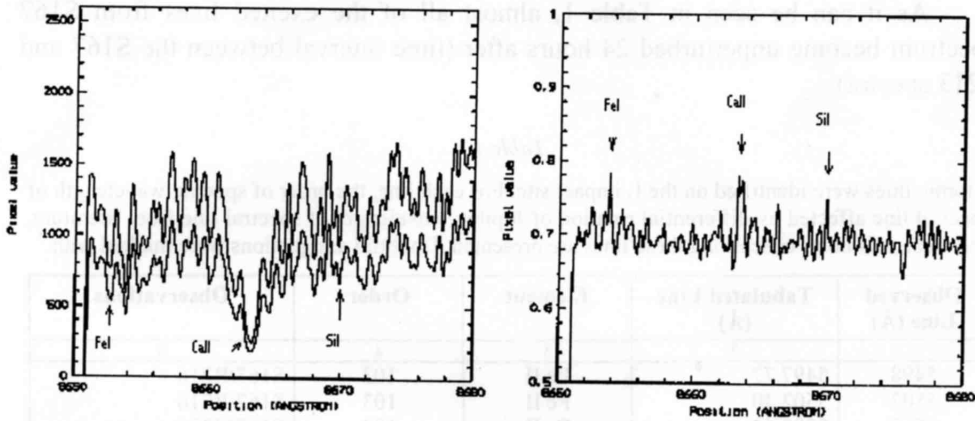


Fig. 1 – Left: Spectra of the 65-th order of R216, and S213 (upper and lower spectrum, respectively). On the center of this order the Paschen $n = 13$ absorption line of Ca II ion. Right: Signatures of Ca II, Fe I, and Si I after the S213/R216 computation.

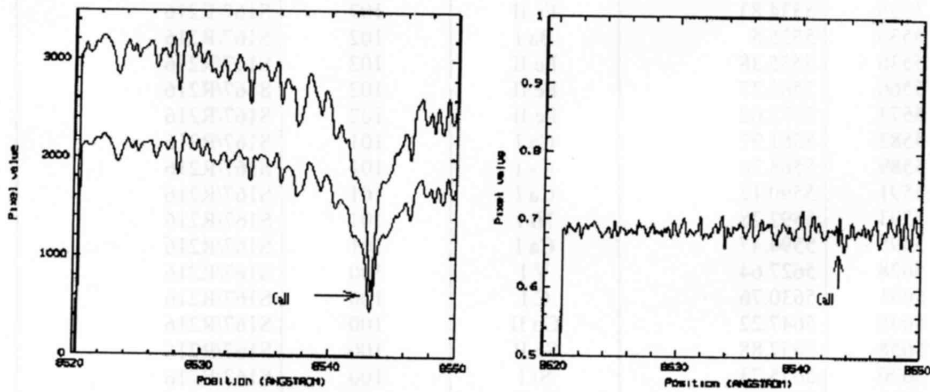


Fig. 2 – “Quiet” order. Spectra of the 66-th order of R216, and S213 (upper and lower spectrum, respectively) containing Paschen line $n = 15$ of Ca II ion (left plot). The result of the S213/R216 computation (right plot).

For the S213, almost of the orders are “quiet”. At the opposite, the S167 spectrum presents high differences for several atomic lines (Fig. 3). Thus, the Fe I, II, and III lines are excited on almost all of orders as well as the representative lines of Ca I, and II ions.

Table 1 lists the excited lines. In a first time the line identification was made taking as origin the profile of spectral lines from *The High Resolution Spectral Atlas of the Solar Irradiance* by Beckers, Bridges and Gilliam (1976), and *The Solar Spectrum from λ 6000 to λ 13495* by Babcock and Moore (1947). Then our identification was refined using the articles of Morton (1991, 2000), Morton and Noreau (1994), and the VizieR database of atomic lines (<http://vizier.u-strasbg.fr>), Reader and Corliss, and Hirata catalogues.

As it can be seen in Table 1, almost all of the excited lines from S167 spectrum become unperturbed 24 hours after (time interval between the S167 and S213 spectra).

Table 1

Atomic lines were identified on the L impact site. For each line, the order of spectra, wavelength of spectral line affected by differential rotation of Jupiter, wavelength of spectral line from literature, element, and notes concerning these lines are presented. Doubtful assignments were marked with *

| Observed Line (Å) | Tabulated Line (Å) | Element | Order | Observations |
|-------------------|--------------------|---------|-------|--------------|
| 1 | 2 | 3 | 4 | 5 |
| 5498 | 5497.77 | Fe II | 103 | S167/R216 |
| 5502 | 5502.30 | Fe II | 103 | S167/R216 |
| 5507 | 5506.44 | Fe II | 103 | S167/R216 |
| 5511 | 5512.98 | Ca I | 103 | S167/R216 |
| 5511 | 5510.61 | Cr I | 103 | S167/R216 |
| 5527 | 5526.8 | Sc II | 102 | S167/R216 |
| 5533 | 5535.05 | Mo I | 102 | S167/R216 |
| 5535 | 5534.83 | Fe II | 102 | S167/R216 |
| 5536 | 5535.5 | Ba I | 102 | S167/R216 |
| 5536 | 5535.38 | Fe II | 102 | S167/R216 |
| 5566 | 5565.37 | Fe II | 102 | S167/R216 |
| 5573 | 5572.62 | Fe II | 102 | S167/R216 |
| 5582 | 5581.97 | Ca I | 101 | S167/R216 |
| 5589 | 5588.76 | Ca I | 101 | S167/R216 |
| 5591 | 5590.12 | Ca I | 101 | S167/R216 |
| 5593 | 5592.28 | Ni I | 101 | S167/R216 |
| 5595 | 5594.47 | Ca I | 101 | S167/R216 |
| 5628 | 5627.64 | V I | 100 | S167/R216 |
| 5631 | 5630.76 | C I | 100 | S167/R216 |
| 5648 | 5647.22 | Co II | 100 | S167/R216 |
| 5658 | 5657.88 | Sc II | 100 | S167/R216 |
| 5676 | 5675.73 | Si I | 100 | S167/R216 |
| 5710 | 5708.93 | Fe II | 99 | S167/R216 |
| 5737 | 5736.75 | Ca I | 99 | S167/R216 |
| 5788 | 5787.9 | Cr I* | 98 | S167/R216 |

Table 1 (continued)

| Observed Line (Å) | Tabulated Line (Å) | Element | Order | Observations |
|----------------------|-----------------------|-----------------|-------|----------------------|
| 1 | 2 | 3 | 4 | 5 |
| 5853 | 5853.45 | Fe II | 97 | S167/R216 |
| 5858 | 5857.45 | Ca I | 97 | S167/R216 |
| 5863 | 5862.89 | Fe I | 97 | S167/R216 |
| 5890 | 5889.95 | Na I | 96 | S167/R216 |
| 5896 | 5895.92 | Na I | 96 | S167/R216 |
| 5915 | 5914.97 | Fe II | 96 | S167/R216, S213/R216 |
| 5984 | 5983.86 | Fe II | 95 | S167/R216 |
| 6014 | 6013.5 | Mn I | 94 | S167/R216 |
| 6017 | 6016.6 | Mn I | 94 | S167/R216 |
| 6066 | 6065.83 | Fe II | 93 | S167/R216 |
| 6123 | 6122.22 | Ca I | 92 | S167/R216 |
| 6142 | 6141.72 | Ba II | 92 | S167/R216 |
| 6163 | 6162.17 | Ca I | 92 | S167/R216 |
| 6176 | 6176.05 | N II | 92 | S167/R216 |
| 6210 | 6209.73 | Fe I | 91 | S167/R216 |
| 6226 | 6225.92 | Cr II* | 91 | S167/R216 |
| 6243 | 6242.87; 6242.9 | Ca I*; Mn I* | 91 | S167/R216 |
| 6245 | 6244.47 | Si I | 91 | S167/R216 |
| 6319 | 6318.66 | Fe II | 90 | S167/R216 |
| 6359 | 6358.76 | Fe II | 89 | S167/R216 |
| 6439 | 6439.07 | Ca I | 88 | S167/R216 |
| 6451 | 6449.81; 6450.24 | Ca I+Co I | 88 | S167/R216 |
| 6472 | 6471.66 | Ca I | 87 | S167/R216, S213/R216 |
| 6494 | 6493.78 | Ca I | 87 | S167/R216 |
| 6496 | 6495.78 | Fe I | 87 | S167/R216, S213/R216 |
| 6498 | 6498.75 | Ba I | 87 | S167/R216, S213/R216 |
| 6501 | 6499.65 | Ca I | 87 | S167/R216, S213/R216 |
| 6563 | 6562.85 | H I | 86 | S167/R216 |
| 6574 | 6572.78 | Ca I | 86 | S167/R216 |
| 6679 | 6678.9 | Fe II | 85 | S167/R216 |
| 6708 | 6707.91; 6707.76 | Li I | 84 | S167/R216 (double) |
| 6978 | 6978.48 | Cr I | 81 | S167/R216 |
| 7289 | 7288.88 | Fe II | 78 | S167/R216 |
| 7290 | 7290.26 | Si I | 78 | S167/R216 |
| 7326 | 7326.15 | Ca I | 77 | S167/R216, S213/R216 |
| 7327 | 7325.51 | Mn I | 77 | S167/R216, S213/R216 |
| 7853 | 7852.86 | C I* | 72 | S167/R216 |
| 7858 | 7858.09 | Fe III | 72 | S167/R216 |
| 7938 | 7938.06 | Fe II | 71 | S167/R216 |
| 8095 | 8094.93 | Fe I | 70 | S167/R216 |
| 8187 | 8186.97; 8186.99 | F III*, Mn II* | 69 | S167/R216 |
| 8195 | 8194.70 | Fe I | 69 | S167/R216 |
| 8405 | 8404.77; 8404.84 | Mn III*, Fe II* | 66 | S167/R216 |
| 8415 | 8414.89; 8414.95 | Fe II*, F I* | 66 | S167/R216, S213/R216 |
| 8664 | 8662.14 | Ca II | 65 | S167/R216, S213/R216 |
| 8683 | 8683.4 | N I | 65 | S167/R216, S213/R216 |
| 8710 | 8710.03 | Fe I | 65 | S167/R216, S213/R216 |
| 8737 | 8736.02; 8736.48 | Mg I, Mn I | 65 | S167/R216 |
| 8790 | 8789.34 | Fe I | 65 | S167/R216 |

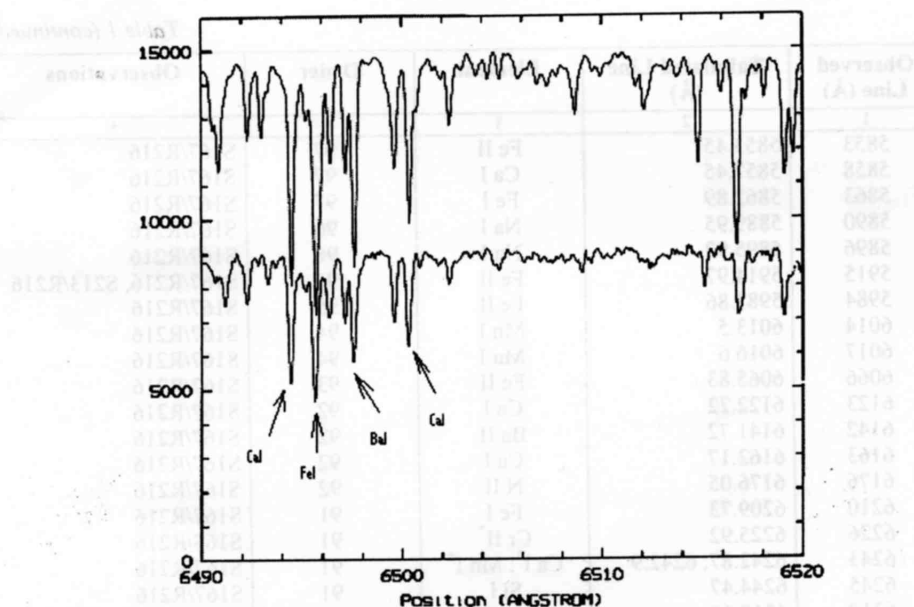


Fig. 3 – Order 87 on S167 (top), and R216 (bottom) spectra. We can distinguish the perturbed lines of Fe I, Ca I, and Ba I lines (marked by arrows).

However, some excited lines are present in both spectra (S167 and S213). In order to have their temporal evolution, casually the orders of a third spectrum were analysed. This spectrum (S166) was taken on July 19/20, 1994, just at the moment that L impact site and plume appeared on the Jupiter visible hemisphere. Table 2 presents such an evolution in the case of some important lines. The percentage values given by the last three columns of Table 2 are computed only from the geometrical consideration, taking into account the depth of the line in the R216 spectrum.

Table 2

The evolution of some spectral lines in three analysed spectra. The spectral line contains the differential rotation shift imposed by Jupiter's rotation.

| Element | Wavelength (Å) | S166/R216 | S167/R216 | S213/R216 |
|------------|----------------|-----------|-----------|-----------|
| Ca I | 6472 | 26% | 25% | 7% |
| Ca I | 6494 | 13% | 20% | – |
| Fe I | 6496 | 26% | 29% | 13% |
| Ba I | 6498 | 16% | 20% | 1% |
| Li I | 6708 | 3% | 24% | – |
| Ca II | 8664 | 84% | 70% | 10% |
| N I | 8683 | 15% | 33% | 13% |
| Fe I | 8790 | 44% | 35% | – |
| Mg I, Mn I | 8737 | 16% | 25% | – |

We have paid a special attention to the hydrogen quadrupolar momentum lines. We have searched for the S(0) – S(3) H signatures (6270.24 Å, 7959.77 Å, 8150.67 Å, and 8272.67 Å, respectively), but our search has not provided positive results.

5. DISCUSSION

The greatest part of the absorption atomic lines of the solar spectrum (of our spectral interval) remained unperturbed after the reflection on Jupiter. The Jovian atmosphere does not present metallic compounds on the analysed spectral interval, the profile of Jupiter's spectrum reproduces the solar spectrum. The perturbed spectra come from the L impact site. Therefore, we can formulate the conclusion that the most significant part of this excitation was released by the cometary material.

Various mechanisms could be responsible of the presence of the atom excitation. The perturbation of atomic lines could be explained as the effect of such a mechanism (or several such mechanisms). As long as the goal of this article is to present the qualitative results of spectra analysis, these mechanisms will be only remembered. However, the author intimate conviction is that different excited lines could be explained only by individual theories.

The main known mechanisms are: the resonant fluorescence, the thermal collision (if we consider the temperature on the impact site and plume greater than 1000 K), and the electronic recombination. The resonant fluorescence mechanism is unanimously accepted as the responsible of the presence of metallic lines on the cometary's spectra as well as for the presence of molecular bands. The thermal collisions could contribute to atomic lines only if the collision rate of atoms is high enough to produce transitions between the corresponding energy levels, and the electronic recombination could be efficient for the excitation of atoms in metastable states.

6. CONCLUSIONS

The analysis of the L impact spectra reveals several excited atomic lines. Alkaline lines (Li, Na, Ba, Ca, Mg) as well as line of transition metals (Fe, Cr, Ni, Co, Mn) are listed in Table 1. Nitrogen (N) and maybe fluorine (F) and carbon (C) lines are presented and rest to be confirmed by further analysis and astronomical observations. The major part of the atomic lines disappear 24 hours after the impact instant, which shows the efficiency of the de-excitation mechanism and energy dissipation on Jupiter's atmosphere.

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