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Citation	The 27th International Astronomical Union (IAU) General Assembly, Rio de Janeiro, Brazil, 3-4 August 2009. In IAU Proceedings, 2009, v. 5, p. 508-509
Issued Date	2009
URL	http://hdl.handle.net/10722/130528
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SpS1-Infrared and submillimetre-wave spectroscopy as probes of stellar evolution

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1. Introduction

For over a hundred years, optical spectroscopy has been the main tool to study stellar structure and evolution. Photospheric spectra of the electronic transitions of atoms and ions are used to determine the temperature and elemental abundance. Beyond atomic and ionic lines, only the electronic transitions of a few simple molecules (C_2 , CN, H_2O , TiO, CH, etc.) appear in the optical photospheric spectra. With the recent development of infrared and submm spectroscopy, a wide range of molecules have been observed, specially in cool atmospheres of red giants and brown dwarfs. We also realize that beyond the photosphere, a stellar system consists of chromosphere, corona, and stellar wind. Both young and evolved stars possess extensive circumstellar regions and the atoms, molecules, and solid particles in this environment radiate a wide range of lines and bands observable at infrared and submm wavelengths.

2. Atoms and ions

Many of the fine structure lines of common neutral atoms and ions lie in the infrared part of the spectrum. The ground state of neutral carbon (C I) has two fine-structure lines ${}^{3}P_{1} - {}^{3}P_{1}$ (370 μ m) and ${}^{3}P_{1} - {}^{3}P_{0}$ (609 μ m). Neutral oxygen (O I) has fine-structure lines ${}^{3}P_{1} - {}^{3}P_{2}$ (63 μ m) and ${}^{3}P_{0} - {}^{3}P_{1}$ (145 μ m). Other common fine-structure lines of ions include the ${}^{2}P_{3/2} - {}^{2}P_{1/2}$ line of C II at 158 μ m, the ${}^{3}P_{2} - {}^{3}P_{1}$ (122 μ m) and ${}^{3}P_{1} - {}^{3}P_{0}$ (205 μ m) lines of N II, and the ${}^{3}P_{2} - {}^{3}P_{1}$ (52 μ m) and ${}^{3}P_{1} - {}^{3}P_{0}$ (88 μ m) of [O III]. Atoms with an odd number of nucleons can have hyperfine lines observable in the infrared. Examples are the 158 μ m ${}^{2}P_{3/2} - {}^{2}P_{1/2}$ F = 2-1 line of [¹³C II], the 9.0 μ m ${}^{3}P_{1} - {}^{3}P_{2}$ line of [Na IV] and the 3.66 μ m line of [Al VI].

3. Molecules

Since the development of mm/submm receivers, the rotational transitions of over 60 molecules have been discovered in the circumstellar envelopes of evolved stars. These include inorganics (e.g., CO, SiO, SiS, NH₃, AlCl, etc.), organics (CH₄, H₂CO, CH₃CN, etc.), radicals (CN, C₂H, C₃, HCO⁺, etc.), chains (e.g., HCN, HC₃N, HC₅N, etc.), and rings (C₃H₂). The rotational transitions of many of the lighter molecules lie in the submillimeter range. Some examples are the $N = 2 \rightarrow 1 \ ^2\Pi_{3/2} \rightarrow ^2\Pi_{1/2}$ at 149.390 μ m ($- \rightarrow +$) and 149.091 μ m ($+ \rightarrow -$) of CH, the ground-state ortho rotational transition $1_{10} - 1_{01}$ (537.7 μ m) of H₂O, the (N, J) = 3, 3 \rightarrow 1, 2 (615.7 μ m) of O₂, the $1_{11} - 0_{00}$ (127.65 μ m) line of methylene (CH₂).

Infrared spectroscopy from space has opened up the possibility of detecting the stretching and bending modes of molecules. For example, the fundamental vibrational mode of HCN consists of the stretching modes ν_1 (100 \leftrightarrow 000) at 4.8 μ m and ν_3 (001 \leftrightarrow 000) at 3 μ m) and bending mode ν_2 (010 \leftrightarrow 000) at 14 μ m. The 13.7 μ m ν_5 *cis*-bend of acetylene (C₂H₂) is commonly observed in absorption in extreme carbon stars. For CO₂, in addition to the 01¹0 \leftrightarrow 00⁰0 fundamental ν_2 bending mode at 14.97 μ m, hot bends (030 \leftrightarrow 020 at 15.40 μ m, 020 \leftrightarrow 010 at 16.18 μ m) and combination bands (100 \leftrightarrow 010 at 13.87 μ m and 110 \leftrightarrow 020 at 13.48 μ m) have been observed. C₄H₂ at 15.9 μ m, ν_{11} of C₆H₂ at 16.1 μ m), ν_4 of benzene (C₆H₆) at 14.84 μ m, and cyanopolyynes (ν_5 of HC₃N at 15.1 μ m, ν_7 of HC₅N at 15.6 μ m) have been detected in absorption in the proto-planetary nebula AFGL 618. Pure carbon chains (C_n) have no permanent electricdipole moment and therefore have no permitted rotational transitions, but their stretching and bending modes can be observed in the infrared. For example, the stretching modes of C₃ (ν_3 at 4.90 μ m) and C₅ (ν_5 at 4.62 μ m) have been detected in absorption in carbon star spectra. The ν_2 (0,1¹,0) \leftrightarrow (0,0⁰,0) bending mode of C₃ near 158 μ m can be observed with the *Herschel Space Observatory*. J = 2 - 0 at 28.2188 μ m) and ortho H₂ (S(1), J = 3 - 1 at 17.0348 μ m) have been observed by the Spitzer Space Telescope. Since HD is no longer homonuclear and has a dipole moment, the $v = 0, J = 1 \rightarrow 0, J = 2 \rightarrow 1$, and $J = 3 \rightarrow 2$ rotational transitions at 112.072, 56.2, and 37.7 μ m can be used to provide accurate determination of HD and to infer the deuterium abundance.

4. Solids

The 9.7 μ m Si–O stretching and the 18 μ m O–Si–O bending modes of amorphous silicates are the most commonly observed solid-state features in stars, with over 4000 such spectra recorded by the *IRAS Low Resolution Spectrometer*. The O–H stretching mode (3 μ m) and bending mode (6 μ m) of amorphous water ice is also widely seen. Other common forms of ice features include the 4.67 μ m C–O stretch of CO, the 15.3 μ m O–C–O bend of CO₂, the 7.7 μ m C–H deformation mode of CH₄. A family of infrared emission features at 3.3, 6.2, 7.7, 8.6, and 11.3 μ m have been identified as due to stretching and bending modes of aromatic compounds. Features at 3.4 and 6.9 μ m can be attributed to the C–H stretching and bending modes of aliphatic units. Broad emission plateaus at 8 and 12 μ m are likely to be due to the in-plane and out-of-plane bending modes of a mixture of aliphatic groups attached to aromatic rings. Amorphous carbonaceous solids with mixed sp^2/sp^3 structures are now known to be rapidly synthesized in the circumstellar environment of evolved stars.

5. Conclusions

With new observing facilities such as *Herschel*, *SOFIA*, and *ALMA*, we will have powerful tools in the infrared and submm wavelengths to probe the physical and chemical structures of stellar environments.

Acknowledgements

This work is supported by a grant from the Research Grants Council of the Hong Kong Special Administrative Region, China (Project No. HKU 7020/08P).

References

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