

HIGH-SENSITIVITY INTRACAVITY LASER SPECTRUM ANALYZER OPERATING IN THE WAVELENGTH REGION 770–820 nm

A.N. Kolerov

*All-Union Scientific-Research Institute of Physico-Technical
and Radio-Technical Measurements, Moscow*

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The sensitivity of an intracavity laser spectrometer was increased by two orders of magnitude in the near-IR ($\Delta\lambda \sim 770\text{--}820$ nm). Using this spectrometer, 132 new absorption lines were recorded in this spectral interval, which is normally considered to be an atmospheric transmission window.

An intracavity laser (ICL) spectrum analyzer, based on a Cr^{3+} : GSGG laser,¹ sensitive to absorption coefficients κ as low as $\sim 10^{-6} \text{ cm}^{-1}$, has made it possible to record absorption lines in the range of $\Delta\lambda \sim 770\text{--}820$ nm. This spectral range is one of the "transmission windows" of the atmosphere, and instrumentation with high sensitivity is needed for a detailed analysis of the fine structure of these lines. Earlier,² using a spectrum analyzer based on a Ti^{3+} : Al_2O_3 laser, it became possible to increase the sensitivity of ICL spectroscopy in this range down to $\kappa \sim 10^{-8} \text{ cm}^{-1}$. However, this laser has a number of special features which complicate its use in these measurements (a short lifetime of excitation $\tau \sim 4 \mu\text{s}$, the necessity of using a "spectral transformer"² with flashlamp pumping, etc.). In this paper, therefore, the

possibility is considered of achieving a higher sensitivity of the ICL spectrum analyzer based on a quasi-continuous Cr^{3+} : GSGG laser.

The optical block diagram of the experimental setup is shown in Fig. 1. The basis of the laser crystal was GSGG: Cr^{3+} 6 mm in diameter and 65 mm in length, placed in a quartz silvered reflector cooled liquid refrigerant with the additives R6G and OX-17 organic dyes. The crystal was excited one or two xenon lamps with maximum energy of electrical pumping $W \leq 2$ kJ. To obtain a resonator configuration close to confocal, spherical mirrors were used (or flat in combination with ordinary ones, and also weakly absorbing lenses). The spectral reflection coefficients of the mirrors were 100% in the range $\Delta\lambda \sim 700\text{--}900$ nm.

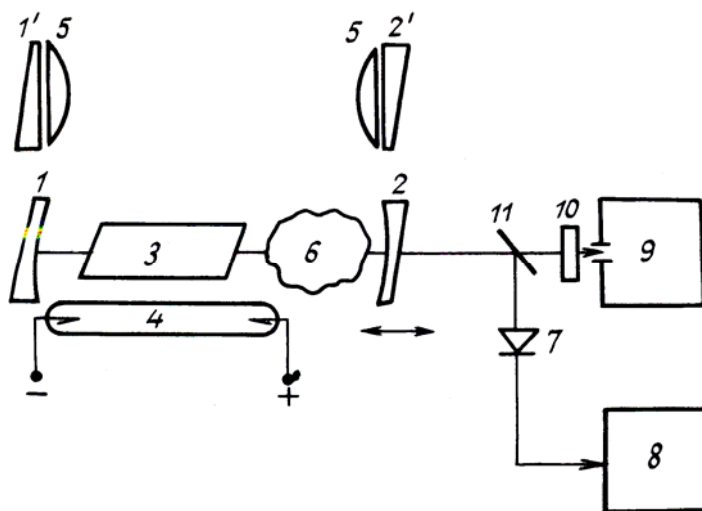


FIG. 1. Optical block diagram of the setup: 1, 2 – resonator mirrors; 3 – active medium; 4 – xenon lamp; 5 – lenses; 6 – investigated object; 7 – photodetector; 8 – recording oscilloscope; 9 – spectrograph; 10 – light filters; 11 – lightsplitter.

The selection of the wavelength interval of operation of the spectrum analyzer was accomplished by substituting mirrors with the corresponding spectral

reflection coefficients. The kinetics of pulse generation was recorded by a photodetector connected to a recording oscilloscope. This made it possible to ex-

perimentally record the value of the duration of quasicontinuous generation, which finally determines the sensitivity of the ICL spectrum analyzer. The laser emission spectrum was recorded on I-810 film behind the spectrograph ($R \sim 6 \cdot 10^5$, $D \sim 1$ nm/cm). To eliminate possible parasitic flare spots, S3S-20 and KS-19 filters, which cut off radiation below 720 nm, were mounted at the input of the spectrograph.

The main results obtained in these experiments were the following. In the free generation mode of the laser, the duration of an individual spike did not exceed 1–3 μ s with the total pulse duration on the order of 150–200 μ s. The maximum possible sensitivity of the spectrum analyzer in this case, allowing for 20% resolution of the absorption line, did not exceed $\sim 3 \cdot 10^{-6}$ cm $^{-1}$ (Ref. 1). Change of the base length ($L \sim 0.5$ –1 m) of the cavity and the pumping rate, as in Refs. 2 and 3, led to a transformation from the free generation mode to quasicontinuous generation. This, in its turn, caused an increase in the pulse

duration of quasicontinuous generation and an increase in the sensitivity of the ICL spectrum analyzer.

As the cavity configuration was changed bringing it closer to a confocal one, the depth of the amplitude pulsations of the generation pulse changed and it became possible to record weaker and weaker atmospheric absorption lines in the emission spectrum (see Fig. 2). For an amplitude pulsation depth (Δ) of ~ 30 –40%, it was possible to record absorption lines whose sensitivity of detection was lower than the values $\kappa \sim 5 \cdot 10^{-8}$ cm $^{-1}$ listed in the atlases.^{4,5} For $\Delta \sim 15$ –20%, a number of absorption lines, lacking in the atlases,^{4,5} were recorded, which testifies that the ICL spectrum analyzer is capable of a sensitivity of $\kappa \sim 10^{-8}$ cm $^{-1}$. The absorption lines corresponding to the spectra in Fig. 2 are listed below (identification of known lines is given in Refs. 4 and 5). 132 new absorption lines were recorded within a rather small range ($\Delta\lambda \sim 8$ nm). Thus, this spectrum analyzer may find application in analytical spectroscopy.

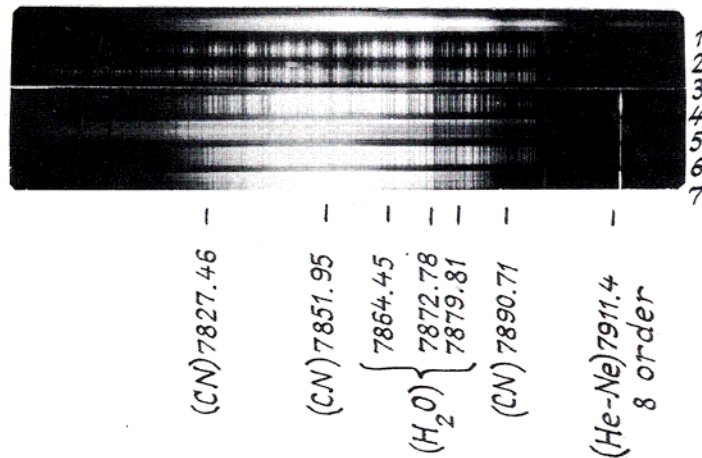


FIG. 2. Fragments of absorption spectra with 1) $\Delta \sim 100\%$ ($\kappa \sim 10^{-6}$ cm $^{-1}$), 2) $\Delta \sim 10\%$ ($\kappa \sim 10^{-8}$ cm $^{-1}$), 3) $\Delta \sim 20\%$ ($\kappa \sim 3 \cdot 10^{-8}$ cm $^{-1}$), 4) $\Delta \sim 40\%$ ($\kappa \sim 8 \cdot 10^{-8}$ cm $^{-1}$), 5) $\Delta \sim 60\%$ ($\kappa \sim 10^{-7}$ cm $^{-1}$), 6) $\Delta \sim 80\%$ ($\kappa \sim 5 \cdot 10^{-7}$ cm $^{-1}$), 7) $\Delta \sim 90\%$ ($\kappa \sim 8 \cdot 10^{-7}$ cm $^{-1}$).

789.637	789.369	CN	789.101	H ₂ O	788.712	H ₂ O	788.346		
789.602	H ₂ O	789.348	H ₂ O	789.071	CN	788.682	H ₂ O	788.309	CN
789.581	789.311	788.997	788.623	H ₂ O	788.290	CN			
789.555	H ₂ O	789.284	CN	788.941	H ₂ O	788.574	788.233	CN	
789.513	CN	789.249	788.907	788.528	CN	788.210	CN		
789.486	H ₂ O	789.212	CN	788.882	H ₂ O	788.505	H ₂ O	788.930	H ₂ O
789.451	789.187	H ₂ O	788.847	788.446	CN	788.163			
789.412	CN	789.161	788.815	788.422	788.089				
789.407	CN	789.111	H ₂ O	788.779	CN	788.361	CN	788.073	H ₂ O
788.040	CN	786.908	785.807	784.661	783.178				
787.987	H ₂ O	786.861	CN	785.750	784.627	H ₂ O	783.150		
787.935	786.818	785.703	784.591	783.098					
787.895	CN	786.803	785.663	784.551	783.077	CN			
787.871	786.759	CN	785.647	784.514	783.046				
787.829	786.733	CN	785.596	784.484	783.009				

787.811	786.702	785.559	784.444	782.971
787.780	786.667 H ₂ O	785.506	784.357	782.925
787.734	786.608 H ₂ O	785.467	784.260	782.893
787.709 CN	786.567	785.437 CN?	784.209	782.847
787.671 H ₂ O	786.515	785.401 H ₂ O	784.174	782.746 CN
787.653 H ₂ O	786.494	785.379	784.152	782.702
787.608 H ₂ O	786.469	785.346	784.113	782.635 CN
787.589 CN	786.445 H ₂ O	785.313	784.053	782.606
787.531 H ₂ O	786.428	785.255	784.015	782.583
787.990 CN	786.375	785.229	783.962 H ₂ O	782.561
787.450	786.375	785.195 CN	783.895	782.521
787.433	786.321 H ₂ O	785.171	783.880	782.453
787.405	786.309	785.145	783.837	782.418 CN
787.395 CN	786.270	785.129	783.811	782.387
787.337 CN	786.236	785.087	783.777	782.321
787.301	786.206	785.049 CN	783.740	782.298
787.278 H ₂ O	786.188 CN?	785.029 CN	783.709	782.177 CN
787.265 CN	786.127	785.995 CN	783.670 CN	782.125
787.201	786.104	785.938	783.643	782.074
787.168	786.076 H ₂ O	785.903 CN	783.578 CN?	782.041
787.141	786.015	784.875 H ₂ O	783.562 CN	781.965
787.114	785.983	784.847	783.460 CN	781.950
787.056 H ₂ O	785.947 CN	784.838	783.414	781.923
787.009	785.915	784.797	783.374	781.902
786.991 H ₂ O	785.882 CN	784.764 CN?	783.351	781.857
786.962	785.849	784.735	783.314	781.817
786.950	785.826	784.691	783.251	781.724 CN

Note: wavelength in nm.

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