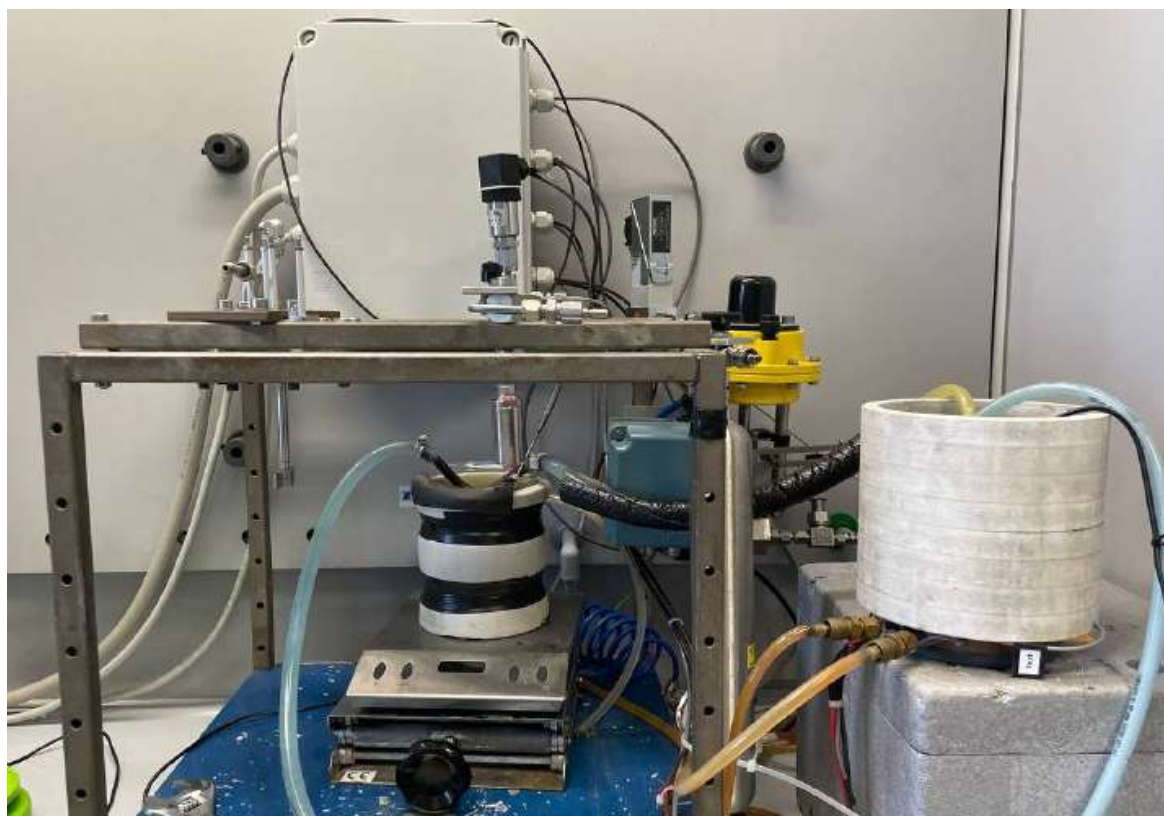
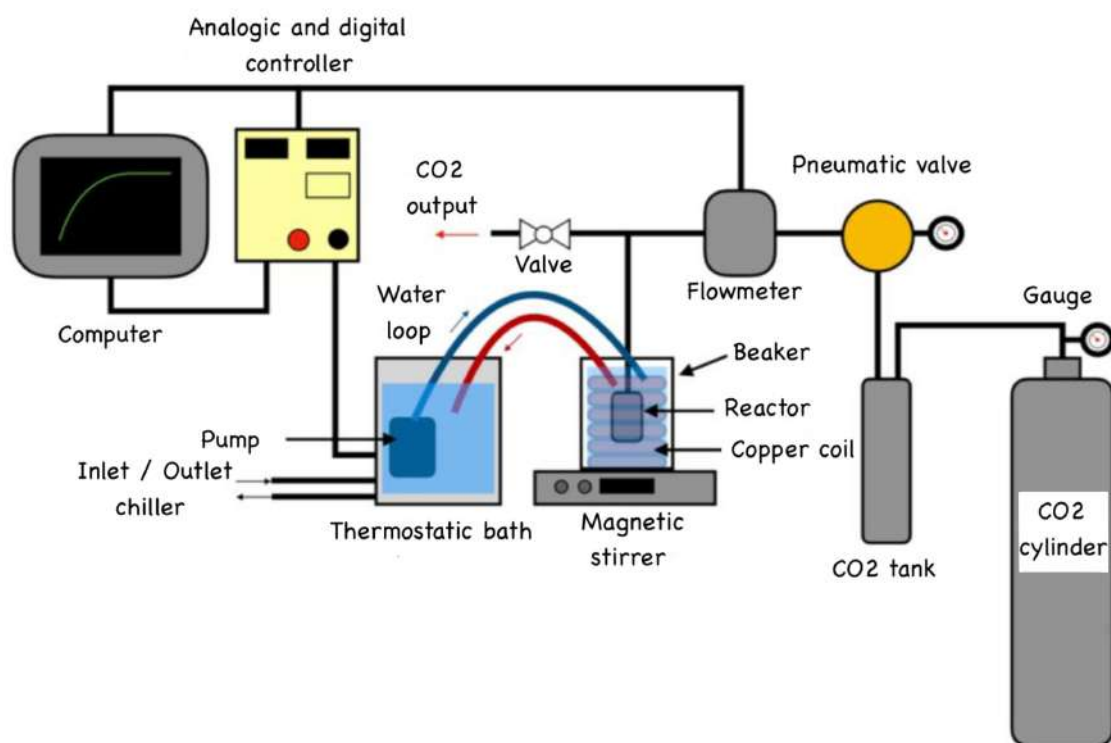


EXPERIMENTAL SECTION



This device allows us to determine when a solution is capable of capturing CO₂ by measuring the amount of gas absorbed using the flowmeter. The principle of this instrument is based on maintaining constant pressure and temperature inside the reactor which contains the analyzed solution; the pressure must be kept constant in the reactor because when the solution absorbs the CO₂ it would tend to decrease.

To supply the CO₂ to the system, a cylinder containing the gas is used to load the smallest tank of the instrument, this tank is connected to a pneumatic valve which, thanks to the flowmeter, determines the quantity of CO₂ entering the reactor. The reactor has a maximum capacity of 10 mL and is immersed in a beaker containing water and a copper coil which maintains the desired temperature.

This water loop forms a thermostatic bath having a pump and is placed on an elevator, the beaker is also placed above a magnetic stirrer. The flowmeter and the thermostatic bath are regulated by an analogical and digital controller which allows us to establish the temperature and pressure at which the analyzes will be carried out.

Solutions were prepared by mixing debole base, ethanol and water at different ratios. The first mixture was prepared with a ratio of debole base to water 1:2 and the concentration of debole base is 8M. Solution 1 is prepared by solubilizing 5.44g of debole base in 34g of H₂O and adding ethanol until a total volume of 10 mL is reached. Solution 1 was added into the stainless steel reactor and the stir bar was added into the mixture. The reactor is connected to the "CCUS machine" and the CO₂ tank is filled. Before the analysis started the system was purged to remove any air in the sample. The analysis started and we waited for the next day.

Analyzing the graph, we realized that it did not reflect the trend we expected, because the quantity of CO₂ released continued to increase and the solution could not absorb so much CO₂. Thinking about the incident we realized that it was a leak in the CO₂ outlet valve that we had to replace in order to repeat the experiment. To try again to make the analysis we had reused the same solution of the first attempt given that it had already released the absorbed CO₂, therefore the solution had been put under vacuum and then in the reactor. The carbon dioxide stream has been started to begin the analysis. The trial with solution 1 was repeated again.

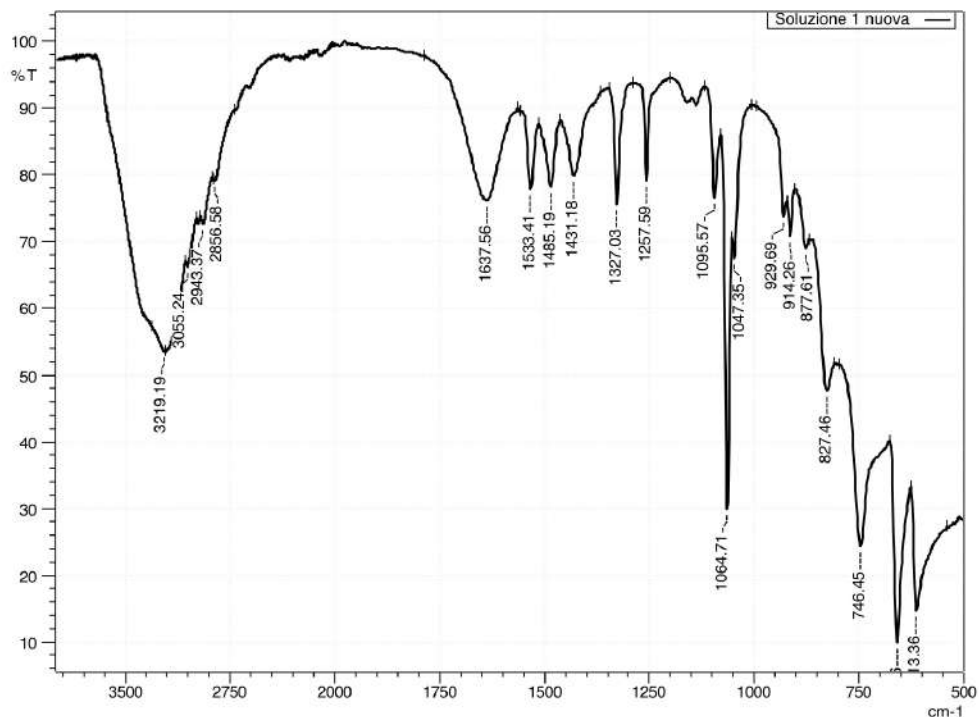
Other solutions with the ratio of debole base to H₂O 1:3 were prepared. To prepare solution 2, 4.08g of debole base and 3.24g of water were weighed out with 6M debole base concentration and ethanol was added until 10mL. While we were waiting for the results of the analysis of solution 2 we had passed the results of analysis of solution 1 to Excel.

We have to obtain the mg of carbon dioxide that had been absorbed by the sample excluding the mass of CO₂ that was needed to keep the pressure constant. The mg of CO₂ absorbed by the sample had been calculated by adding an inert material to the reactor (of the same volume as the sample inserted) and bringing the system to 20bar. The CO₂ that had been used to bring the system to 20bar has a mass of 306mg, the value obtained had been subtracted from the analysis values.

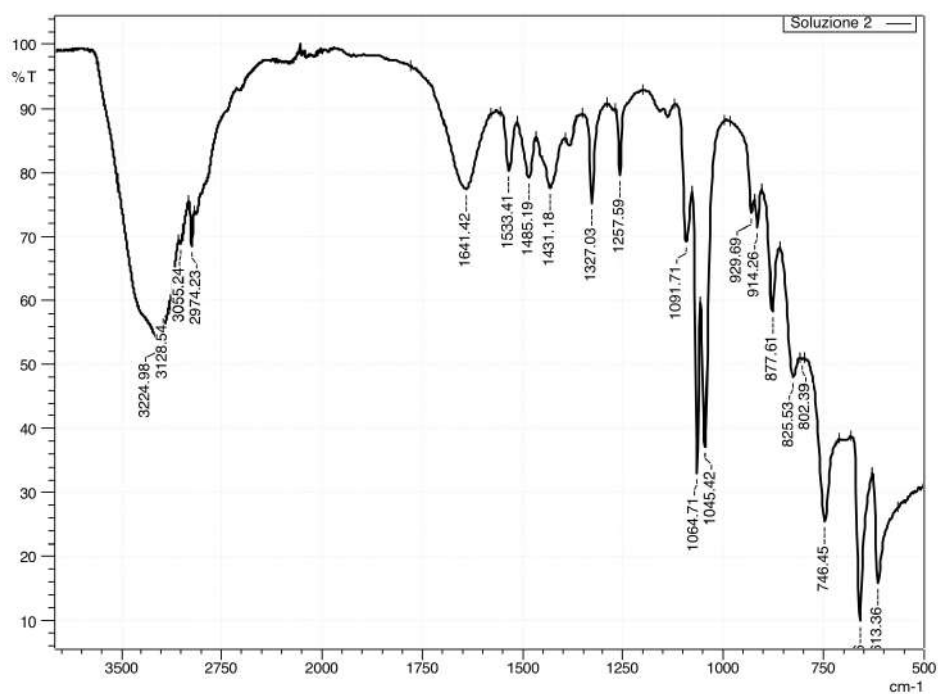
We thought it was usefull to prepare a solution similar to solution 2 but with an 8M debole base concentration so all analyzes were repeated with solution 3.

To conclude the experiment, the analyzes with Raman and infrared spectroscopy were carried out on all the prepared solutions and the spectra obtained are reported in "RESULTS AND DISCUSSION".

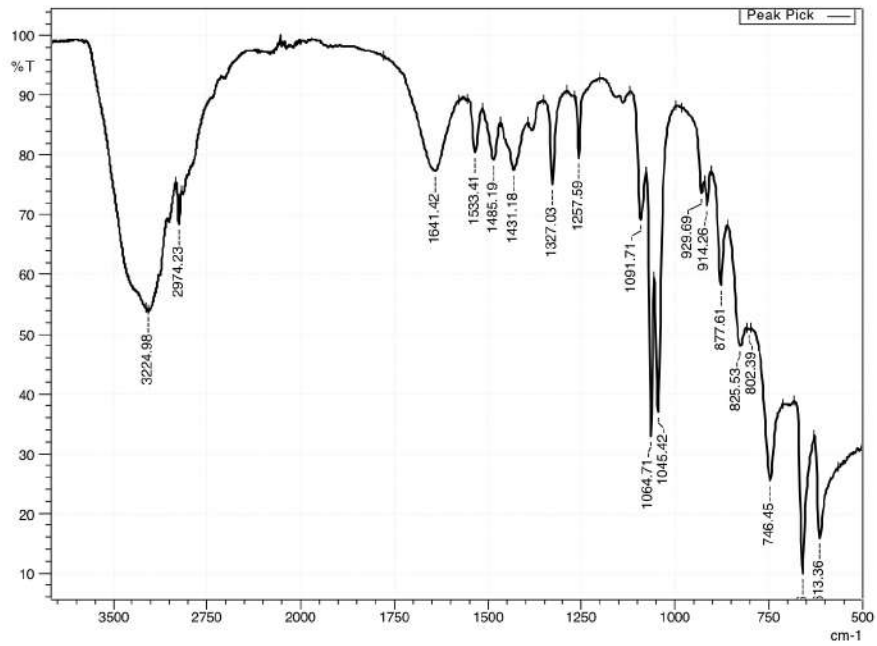
RESULTS AND DISCUSSION



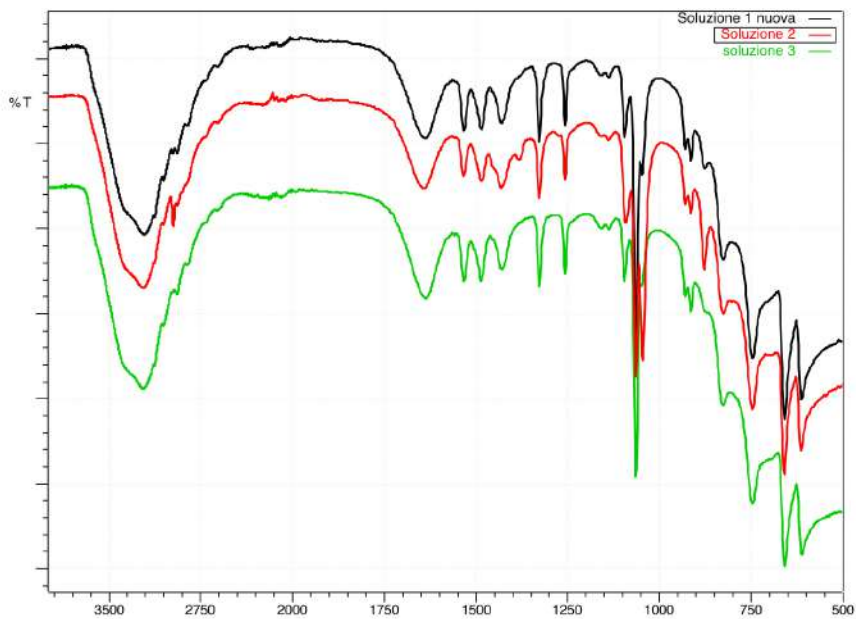
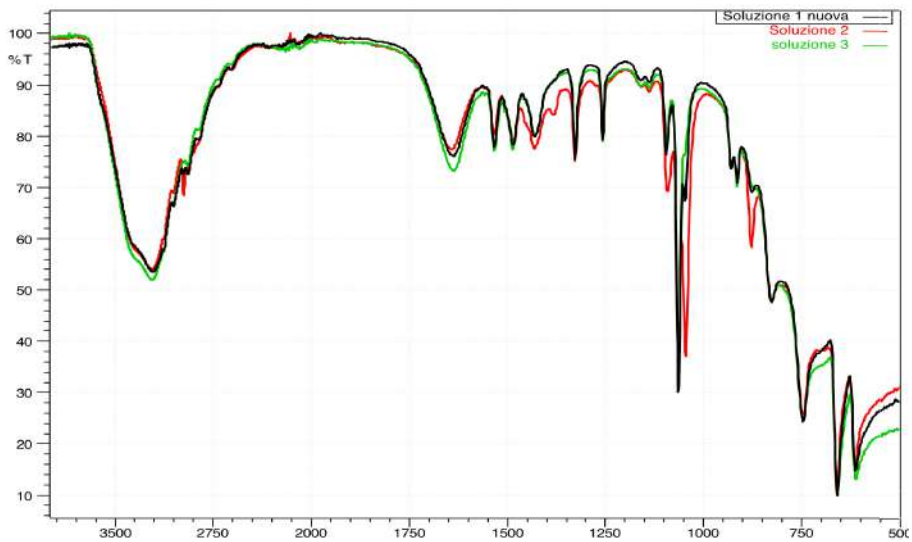
↑ IR spectrum solution 1



↑ IR spectrum solution 2



↑ IR spectrum solution 3



STRETCHING O-H (REFERRED TO ETHANOL)= 3200 cm^{-1}

STRECHING C=C (REFERRED TO AROMATHIC RING OF DEBOLE BASE)=~1500 cm^{-1}

STRETCHING C-N (REFERRED TO DEBOLE BASE)= 1060 cm^{-1}

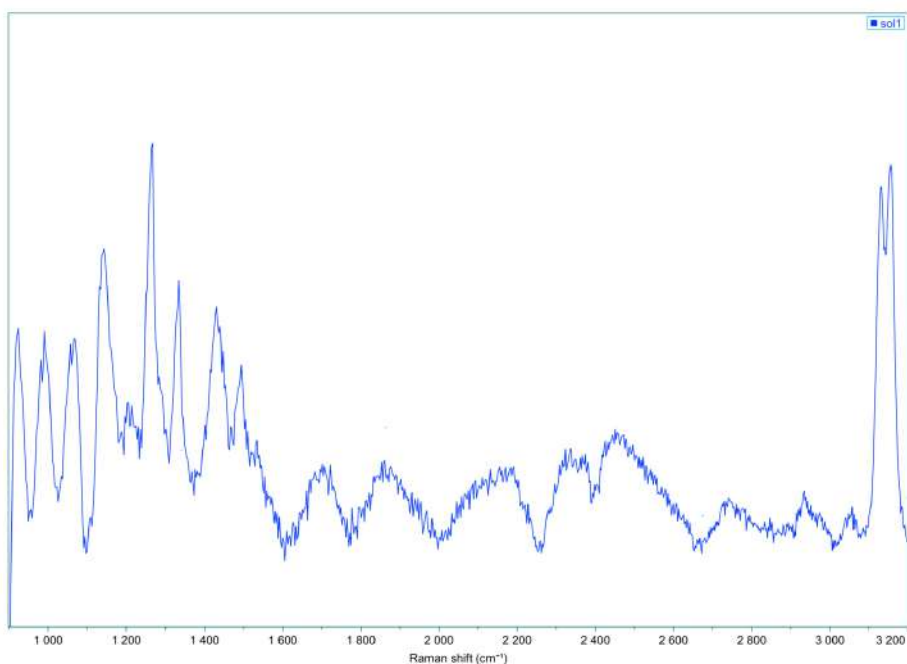
BENDING C-H (REFERRED TO DEBOLE BASE)= 1450 cm^{-1}

BENDING CH₂ (REFERRED TO ETHANOL)= 1390 cm^{-1}

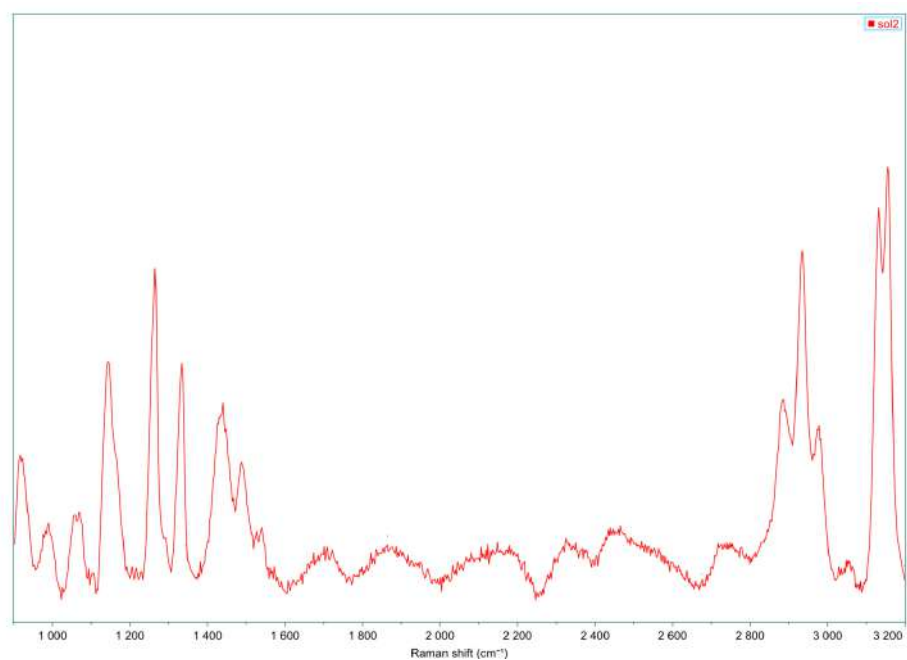
STRETCHING C-O (REFERRED TO ETHANOL)=1100-1040 cm^{-1}

BENDING N-H (REFERRED TO ETHANOL)=1635 cm^{-1}

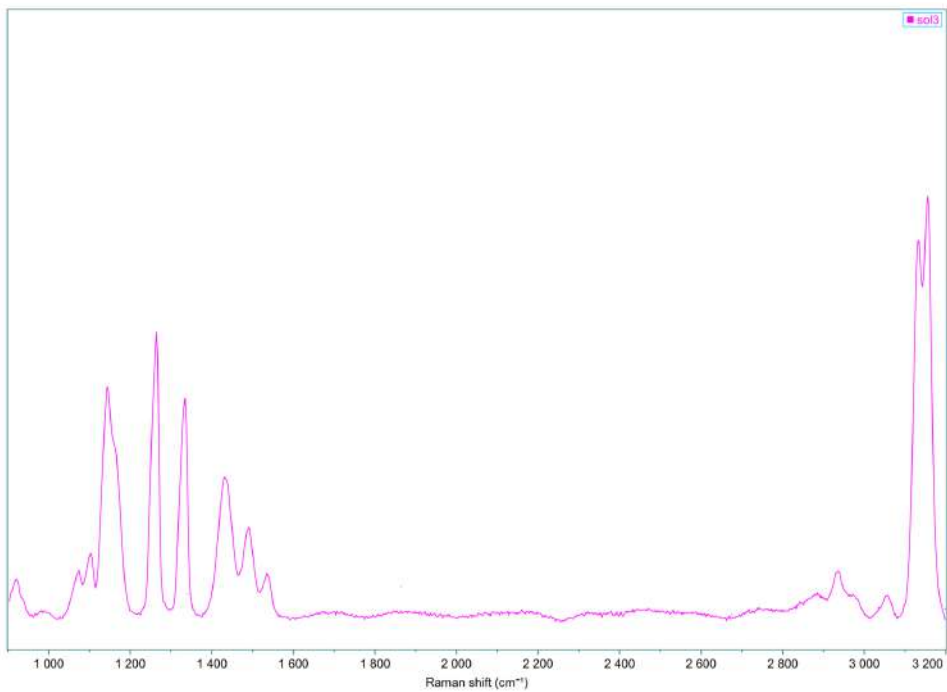
Stretching of the N-H bond is superimposed by stretching of the O-H bond.



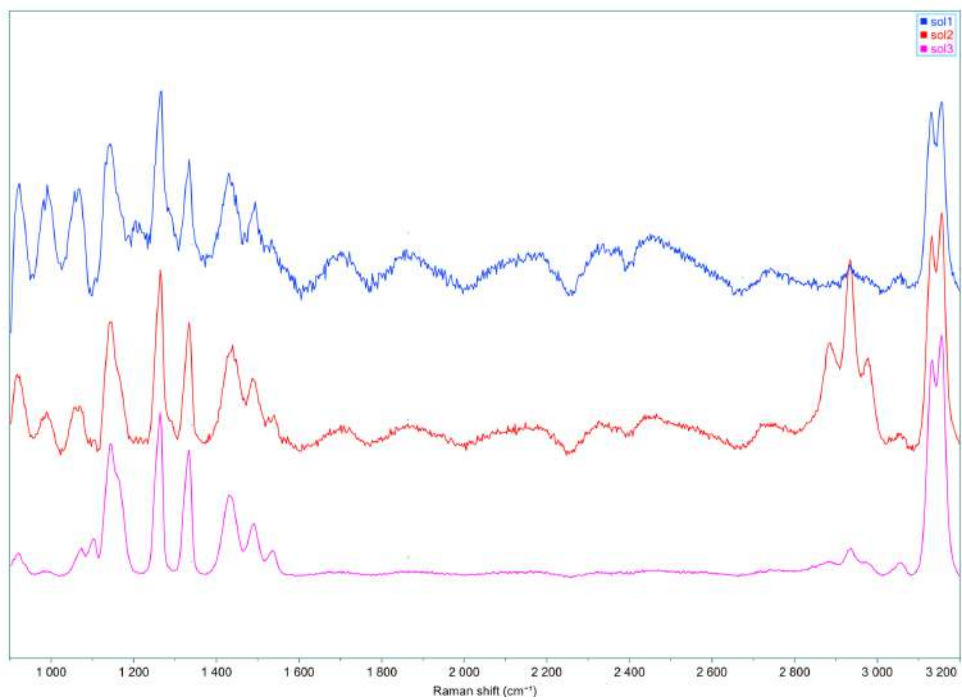
↑ Raman spectrum solution 1



↑ Raman spectrum solution 2



↑ Raman spectrum solution 3



~ 980= ring deformation;

~ 1100= C-O stretching;

~ 1250-1350= C-C stretching;

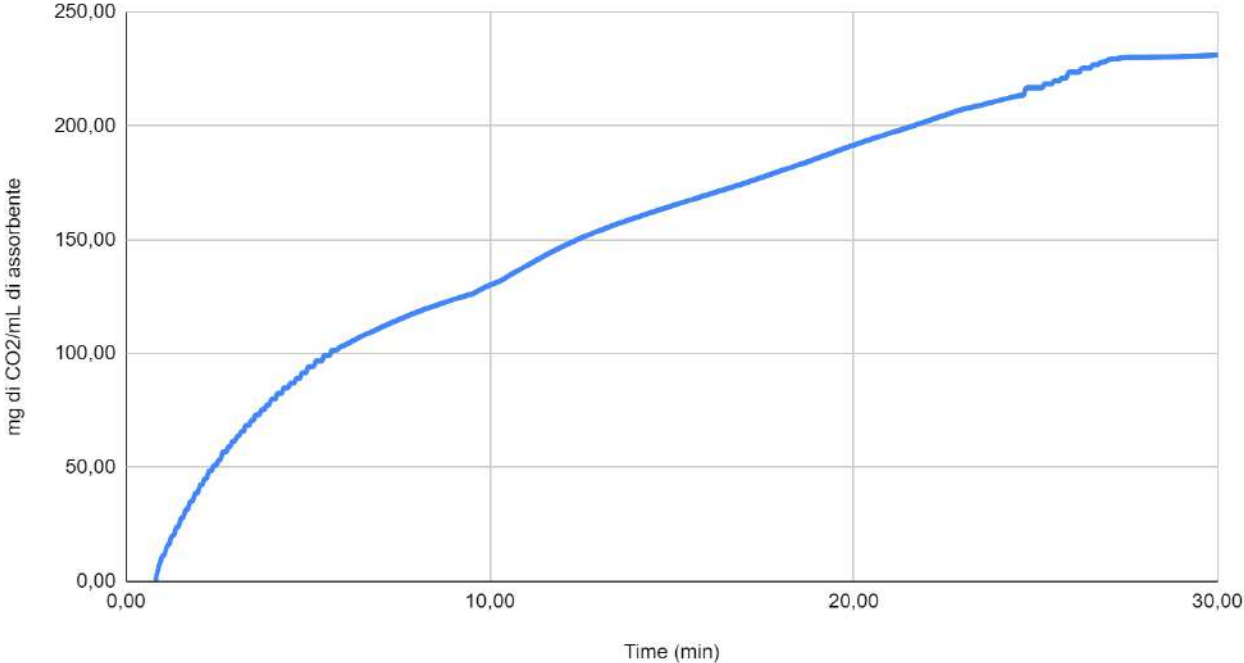
~ 1400-1600= C-N stretching + C-H wagging + N-H wagging;

~ 1600-2600= fluorescence bands that fade with increasing concentration (interference);

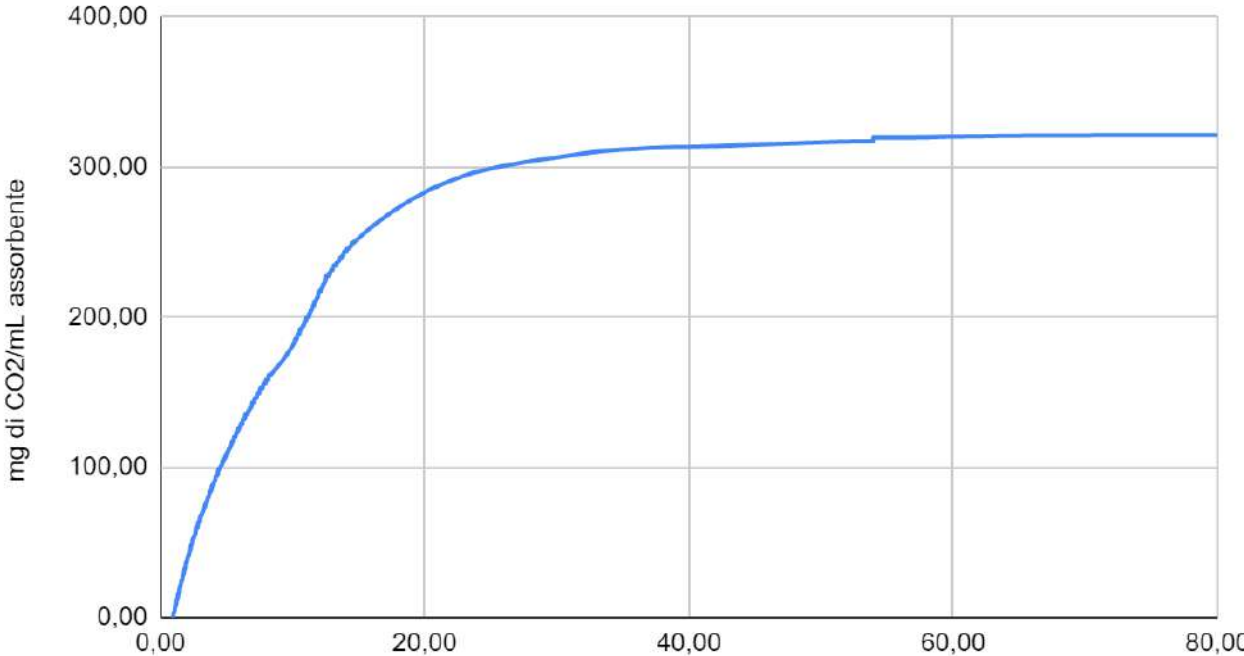
~ 2900- 3100= the peaks are seen only in the second solution because there is a higher concentration of ethanol and they represent the C-H stretches;

~ 3200=N-H stretching;

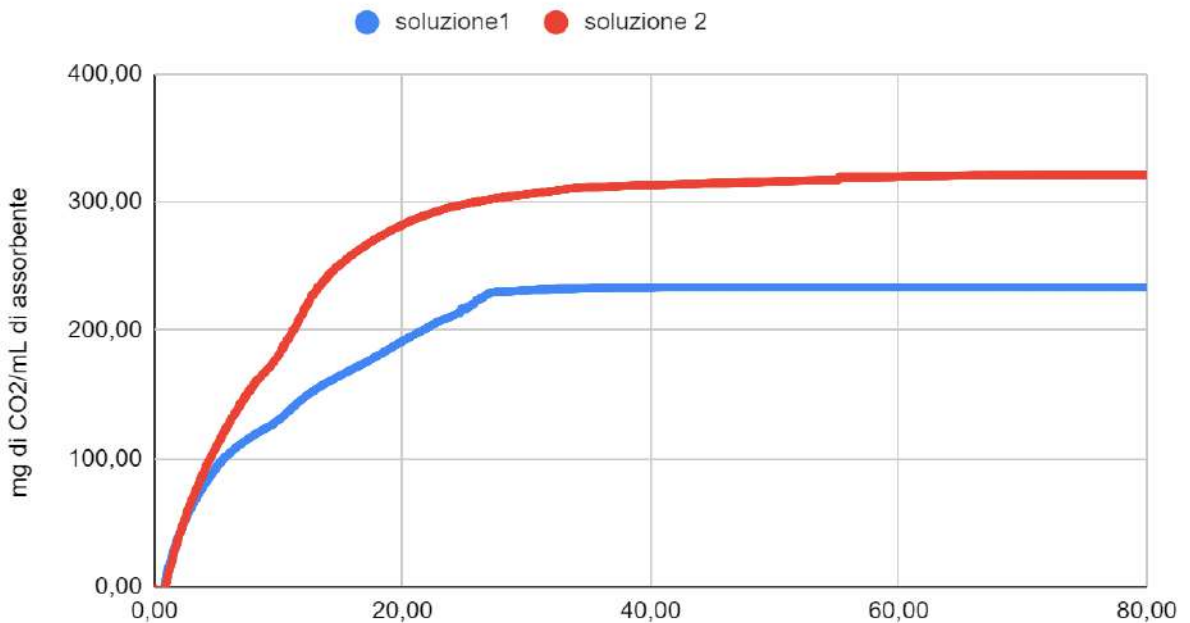
mg di CO2/mL di assorbente soluzione 1



mg di CO2/mL assorbente soluzione 3



mg di CO₂/mL di assorbente



↑ The trend in the graph shows the absorption of CO₂ as a function of time and the mg of CO₂ absorbed by the solutions.

CONCLUSIONS

The method that we had studied to capture CO₂ was especially effective with the 8M solution with the ratio of debole base to water 1:3. The solutions had been shown to be effective for both the capture and release of carbon dioxide due to the choice of a weak base which makes the reactions reversible. Since CO₂ is weakly acidic compared to other waste gases and has a closer affinity to debole base, this method is therefore selective towards CO₂. The studied method has advantages when applied to industries because if they produce a mixture of waste gases only CO₂ will be captured. This method of capturing CO₂ is environmentally sustainable as the solution after it releases the CO₂ will be able to absorb for several cycles.

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