








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seconds!.27 kJ/mol) with a
stoichiometric reaction of

5.7 mol mol⁻¹ of glucose to g-C₃N₄. The g-C₃N₄/CD composite was also examined by X-ray diffraction (XRD), FTIR, UV-Vis absorption and N₂ sorption measurements. The XRD measurements showed a distinct low intensity diffraction peak at $2\theta = 26.43^\circ$, which was related to the (002) reflection of graphitic carbon nitride \[[@CR28]\]. FTIR spectrum of g-C₃N₄ and g-C₃N₄/CD composite are shown in Fig. [7](#Fig7) a. The broad peaks are observed at the range of

1350 cm^{-1} and 1660 cm^{-1} and the weak characteristic peaks are present in the range of 1040 cm^{-1} and 1480 cm^{-1} . The peaks were assigned to stretching vibration of C--H (2800 cm^{-1}), C--N (1600 cm^{-1}) and N--H (1000 cm^{-1}) vibrations \[\[@CR29 \]. The absorption bands in the range of 1000--1300 cm^{-1} are assigned to the stretching of N--H, and --NH and the band at 2800 cm^{-1} arises due to the stretching of --C--H bond in the graphitic structure of g-C~3~N~4~ \[\[@CR29 \],

[@CR30]\]. In the FTIR spectrum of the g-C₃N₄/CD composite, the band around 1650 cm⁻¹ corresponding to the C=N stretching vibration of g-C₃N₄ disappeared and two new peaks were observed at 1602 cm⁻¹ and 1522 cm⁻¹ which are assigned to the stretching vibration of -C=C- bending and vibration of C=C, respectively \[[@CR31]\]. All of the above results

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