

# N-type pyrazine-based luminogens with aggregation-induced emission characteristics Ming Chen(11229038)<sup>a</sup>, Jing Zhi Sun<sup>a</sup>, Anjun Qin<sup>\*a</sup>, and Ben Zhong Tang <sup>\*a,b</sup>

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# INTRODUCTION

Among luminogens with aggregation-induced emission (AIE) characteristics, N-type molecules are rare in comparison with P-type and neutral species in despite of their indispensable roles in functions as electron transport and electron acceptor materials.<sup>1</sup> During our exploration for new AIE-active luminogens, we found that a pyrazine derivative features such property.<sup>2</sup> Furthermore, thanks to its electron-deficent property, this molecule could also serve as N-type material. In this work, N-type pyrazine-based triazoles with 1,4- and 1,5-regioregular isomers were prepared via Cu- and Ru-catalyzed azide-alkyne click reaction. These luminogens are AIE-active and could form charge-transfer complexes with triphenylamine (TPA).<sup>3</sup>

## **RESULTS AND DISCUSSION**

**Molecular Structures** 

### **Photo-physical Property**

### **Charge Transfer Complexes**







#### 1,5-isomer

**Scheme 1.** Synthetic routes to pyrazine-based 1,4-and 1,5-triazoles by click reaction.

### **Structure Characterization**



Wavelength (nm)

isity (au)

isomer (right).

Water fraction (%)

Wavelength (nm)

Water fraction (%)



С



**Figure 4.** (A) PL spectra of 1,4-isomer/TPA complex in THF and its aqueous mixtures. (B) Changes of PL intensity of 1,4-isomer/TPA (red) and 1,5-isomer/TPA (blue) complexes with various fractions of water content (C) Photographs of 1,4-isomer, 1,5-isomer and their complexes with TPA in solids. All isomer/TPA complexes here prepared with the molar ratio: 1:1





### CONCLUSIONS

(red) and 1,5- (blue) isomers.

N-type pyrazine-based 1,4- and 1,5-regioregular triazoles, synthesized via click reactions, exhibit a discriminative AIE features with pale blue and dark blue emission. These luminogens could form charge-transfer complexes with TPA and emit red light in the aggregate states. This work thus offers an attractive strategy to fine-tune the emission of AIE luminogens by chemical or physical approaches.

### ACKNOWLEDGMENTS

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