

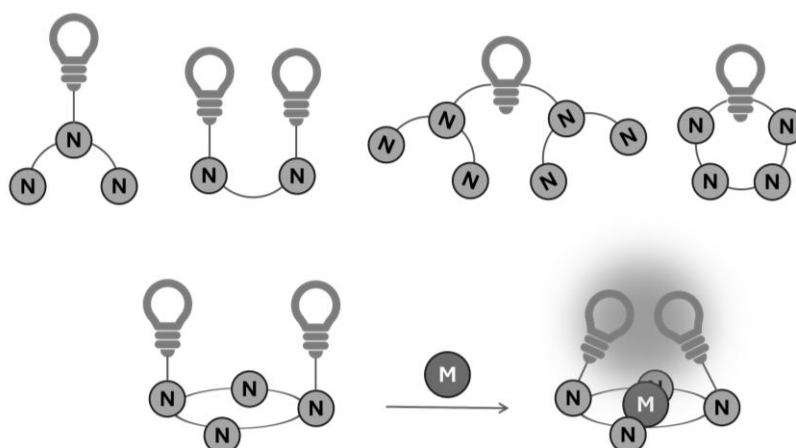
## Polyazaligands as tools for sensing

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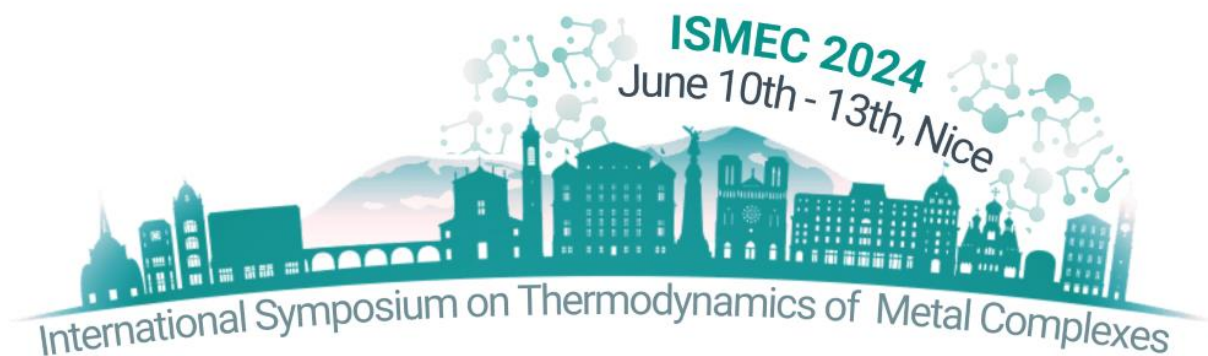
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The design of receptors able to selectively recognize and bind specific guests is a challenging and dateless target. Due to their easy functionalization and tunable topology, polyaza scaffolds are useful to this purpose, moreover they provide high solubility in aqueous solution.

Both open-chain and macrocyclic compounds could provide favourable properties. On one hand, open-chain polyamine ligands ensure flexibility, allowing for the accommodation of metal cations with different sizes and coordination geometries. Polynuclear complexes can form when the ligand features multiple aza-binding sites, and the distance between the metal centres can be tuned to have them cooperate or not in the formation of the active site [1]. Moreover, a preorganized polyaza-metal complex can become a metallo-receptor for additional guests [2–4]. On the other hand, macrocyclic polyamine ligands ensure high complex stability and selectivity towards the target metal cations, thanks to the preorganization provided by the stiffened system [5]. The insertion in the aza-ligands of chromophores or fluorophores can signal the occurred interaction with the guest [6,7]. The optical active group can be linked to the polyamine through a simple spacer or could be part of the macrocycle, forming a cyclophane [8–10] (Figure 1).



**Figure 1:** Scheme of possible polyaza-ligands and example of metal ion complexation



Our group has long been working on polyaza-ligands to be used as receptors for guests of different nature. In this contribution, some examples of open-chain, macrocycle or cyclophane polyamine ligands are reported, to show their peculiar behaviour towards selected metal ions.

#### References:

- [1] M. Formica, G. Ambrosi, V. Fusi, L. Giorgi, M. Arca, A. Garau, et al., *New J. Chem.* 2018, 42, 7869–7883.
- [2] E. Macedi, L. Giorgi, M. Formica, P. Rossi, D. Paderni, P. Paoli, V. Fusi, *Chempluschem* 2023, 88, e202200364.
- [3] P. Rossi, E. Macedi, M. Formica, L. Giorgi, P. Paoli, V. Fusi, *Chempluschem* 2020, 85, 1179–1189.
- [4] D. Paderni, L. Giorgi, E. Macedi, M. Formica, P. Paoli, P. Rossi, V. Fusi, *Dalt. Trans.* 2021, 50, 15433.
- [5] A. Garau, L. Lvova, E. Macedi, G. Ambrosi, M. C. Aragoni, et al., *New J. Chem.* 2020, 44, 20834–20852.
- [6] D. Paderni, D. Lopez, E. Macedi, G. Ambrosi, A. Ricci, et al., *Inorganica Chim. Acta* 2023, 549, 121400.
- [7] D. Paderni, E. Macedi, L. Lvova, G. Ambrosi, M. Formica, et al., *Chem. - A Eur. J.* 2022, 28, e202201062.
- [8] D. Paderni, G. Barone, L. Giorgi, M. Formica, E. Macedi, V. Fusi, *Dalt. Trans.* 2023, 3716–3724.
- [9] D. Paderni, L. Giorgi, M. Voccia, M. Formica, L. Caporaso, E. Macedi, V. Fusi, *Chemosensors* 2022, 10, 188.
- [10] G. E. Giacomazzo, D. Paderni, L. Giorgi, M. Formica, L. Mari, R. Montis, et al., *Molecules* 2023, 28, 2031.