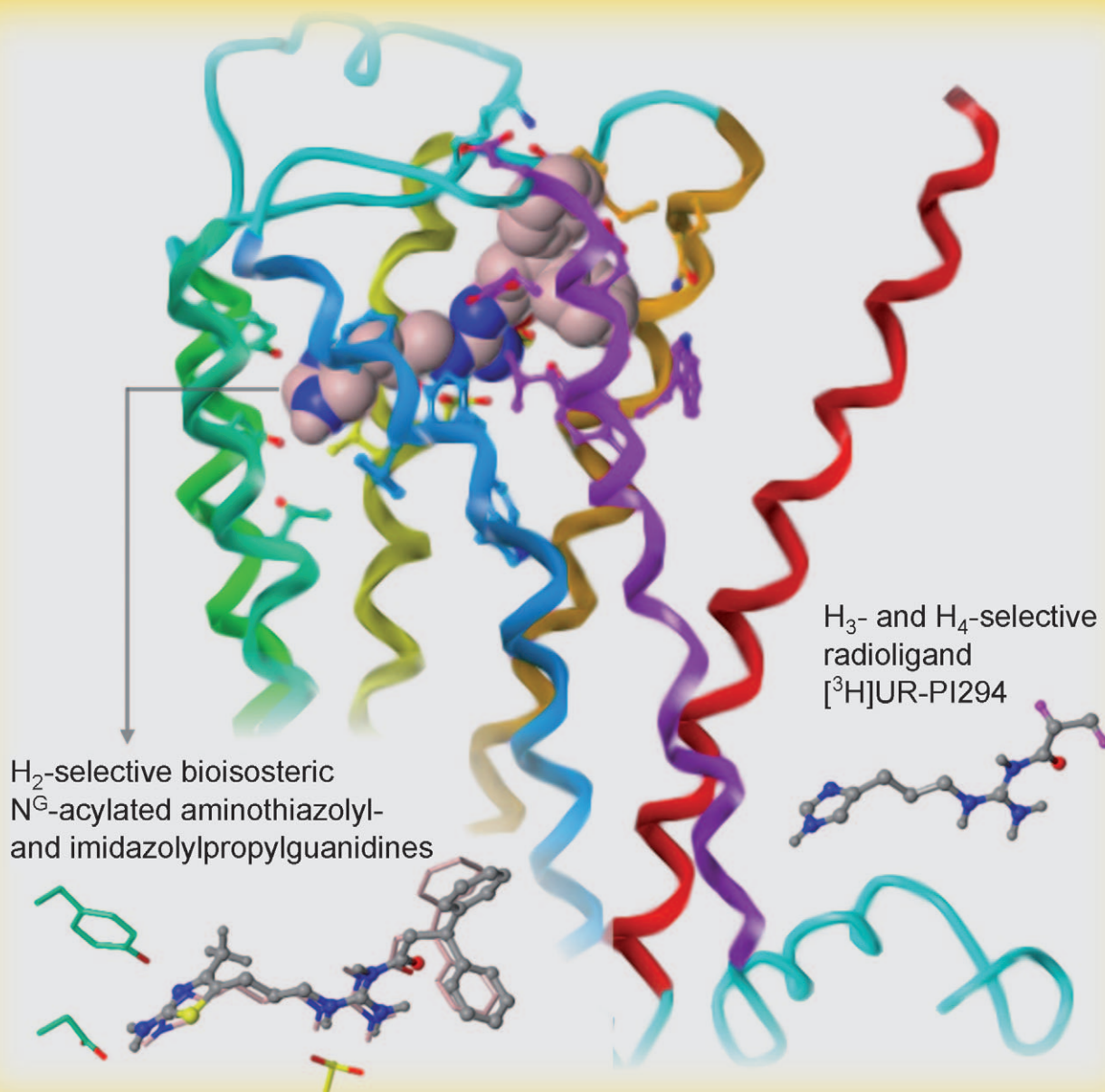


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The inside cover picture shows the human histamine H₂ receptor and the H₂ and H₃/H₄ receptor-selective ligands identified. Bioisosteric replacement of the imidazole ring of the N^G-acyl-imidazolylpropylguanidine with 2-aminothiazole yielded potent and selective H₂ receptor agonists. In contrast, modification of the acyl group gave a high-affinity H₃/H₄-selective tritiated radioligand. For more details, see the Full Papers by A. Buschauer et al. on p. 225 ff and p. 232 ff.

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Inside Cover

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The inside cover picture shows the human histamine H_2 receptor and the H_2 and H_3/H_4 receptor-selective ligands identified. Bioisosteric replacement of the imidazole ring of the N^G -acyl-imidazolylpropylguanidine with 2-aminothiazole yielded potent and selective H_2 receptor agonists. In contrast, modification of the acyl group gave a high-affinity H_3/H_4 -selective tritiated radioligand. For more details, see the Full Papers by A. Buschauer et al. on p. 225 ff and p. 232 ff.

