This feature focuses on a reagent chosen by a postgraduate, highlighting the uses and preparation of the reagent in current research.

**Introduction**

Ytterbium trifluoromethansulfonate [Yb(OTf)₃] has been widely used in organic syntheses in the last few years. Yb(OTf)₃ is a strong Lewis acid due to the hard character of Yb³⁺ ion and the presence of electron-deficient triflate in its coordination sphere. In contrast to traditional Lewis acids, such as AlCl₃, BF₃, TiCl₄, and SnCl₄, which are often used in stoichiometric amounts, only catalytic amounts of Yb(OTf)₃ are necessary. Moreover it can be easily recovered and reused without loss of activity. Interestingly, Yb(OTf)₃ remains catalytically active in the presence of many Lewis bases containing nitrogen, oxygen, phosphorus or sulfur atoms. The resulting water-compatibility of Yb(OTf)₃ is one of its well-known advantages, with respect to traditional Lewis acids that are very sensitive and easily decomposed or deactivated in the presence of small amounts of water. The most interesting point from a synthetic point of view is that Yb(OTf)₃-catalyzed reactions are clean, while Yb(OTf)₃ is regarded as environmentally friendly catalyst. Ytterbium triflate is prepared by heating ytterbium(III) oxide or chloride in an aqueous trifluoromethansulfonic acid solution (Scheme 1). Yb(OTf)₃ is prepared by heating ytterbium(III) oxide or chloride in an aqueous trifluoromethansulfonic acid solution (Scheme 1).

**Abstracts**

(A) **Friedel–Crafts Acylation:**

The acylation of 1-methylpyrrole has been reported recently. The reaction is carried out in [bpy][BF₄] with a catalytic amount of Yb(OTf)₃ (10 mol%) at room temperature. Good yields were obtained (80–93%), but the reaction fails without catalyst. Moreover, the catalyst can be recycled three times without loss of activity.

(B) **Crotonation:**

3-Acylacrylic acids possess a high potential in the synthesis of biologically or pharmaceutically active compounds, such as 4, 5, and 6. Recently, Gorobets and co-workers described a new facile protocol for the synthesis of aromatic and heteroaromatic 3-acylacrylic acids in good yields (54–78%). Aromatic ketones 7, glyoxylic acid monohydrate 8, and Yb(OTf)₃ (2.5 mol%) are reacted under microwave irradiation.
(C) Tosylation:
Most tosylations use triethylamine or pyridine as a base in the reaction of appropriate alcohols with the tosylating agents.17 In 2004, Schirrmacher and Comagic18 reported the low-yielding tosylation of 10 using TsCl and pyridine. Gratifyingly, when Yb(OTf)3 is used, the tosylation with Ts2O proceeded in excellent yield (85%). These conditions were also applied for several primary and secondary alcohols and provided the tosylates in good yields (75–89%).

(D) TEMPO-Mediated Oxidation:
Vatèle described a new method for the oxidation of alcohols with iodobenzylamine relying on the utilization of the TEMPO/PhIO system as oxidizing source.19 However, when 4-phenyl-butan-1-ol was treated with PhIO and TEMPO, only 5% of 4-phenyl-butan-1-ol was obtained. In the presence of Yb(OTf)3 (2 mol%), the expected aldehyde was obtained in good yield (83%). The triflate can also catalyze the oxidation of several primary or secondary alcohols into the corresponding aldehydes or ketones in good to excellent yields (76–94%).

(E) One-Pot Multicomponent Synthesis of Substituted Imidazoles:
Yb(OTf)3 has been used for the synthesis of substituted imidazoles through three-component condensation of benzil (E) 10 using Yb(OTf)3 as both a catalyst and oxidizing agent. This protocol also gave good yields (61–85%) when applied to primary alcohols and provided the tosylates in good yields (75–89%).

(F) Selective Anomeric Deacetylation:
Selective anomeric deacetylation is a key step in the oligosaccharide synthesis. Yb(OTf)3 can promote the selective anomeric deacetylation of compound 14 (an important synthon involved in the synthesis of heparin sulphate fragments).21 The reaction is carried out using catalytic amounts of Yb(OTf)3 (5 mol%) and gave compound 15 in good yield (75%). However, Nd(OTf)3 proved to be a superior catalyst. This protocol also gave good yields (61–85%) when applied to other sugar peracetics, such as α- or β-d-glucopyranose or β-d-xyllopyranose peracetates. One of the most striking features is that Yb3+ and Nd3+ catalyzed the transesterification of the anomeric acetate without catalyzing the methyl glycoside formation.

References