Eliminação de água a partir de 1-butilciclo-hexanol

Análogo ao procedimento abaixo:

Experiment 5.12 2-METHYLBUT-2-ENE (in admixture with 2-METHYLBUT-1-ENE)

$$Me_2C(OH)\cdot CH_2\cdot Me \xrightarrow{H_3PO_4} Me_2C=CH\cdot Me + CH_2=C(Me)\cdot CH_2\cdot Me$$

Place 25.0 g (31 ml, 0.28 mol) of 2-methylbutan-2-ol and 10 ml of 85 per cent orthophosphoric acid in a 100-ml, round-bottomed flask and swirl to mix thoroughly. Fit the flask with a 20-cm fractionating column filled with glass helices, a Claisen still-head and a condenser leading to a 50-ml receiving flask cooled in a beaker of iced water (Fig. 2.104). Add a few pieces of porous porcelain and heat the reaction mixture gently with a Bunsen burner. Collect the alkene fraction which distils in the range 35-38 °C during a period of 30 minutes. Dry the distillate with 1-2 g of magnesium sulphate. Wash and dry the distillation apparatus, decant the dried distillate into a 50-ml flask and redistil in the reassembled apparatus. Collect the fraction boiling at 37–38 °C; the yield is 12.5 g (64%). Record the infrared spectrum of the product using a fixed path-length cell (0.025 mm). The stretching bands of the terminal (1645 cm⁻¹) and non-terminal (1670 cm⁻¹) carbon-carbon double bonds can both be observed; bands at 890 and 805 cm⁻¹ (=C-H deformation) also establish the presence of both terminal and non-terminal olefinic systems. Analyse the product by g.l.c. on a Silicone oil column at 30 °C; 2methylbut-1-ene appears first, closely followed by 2-methylbut-2-ene; the areas under the peaks are in the ratio of 1:4.

Utilizar todo o 1-butilciclo-hexanol obtido no experimento anterior e calcular as quantidades correspondentes.