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## Answers

# Practice exam questions

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Check your answers to the questions in this issue.

## Laundry: washing clean and smelling fresh (pp. 2–6)

- 1 In each case you only need to provide one suitable test, either spectroscopic or chemical. Both forms of answers are given here.
  - a To distinguish between samples of nonanal and nonanoic acid, add a solution of sodium hydrogen carbonate (or sodium carbonate). You will see effervescence and evolution of  $\text{CO}_2$  with the nonanoic acid, but no reaction with the aldehyde (nonanal).  
  
**Alternative:** adding a magnesium ribbon and a little water produces effervescence and evolution of  $\text{H}_2$  with the acid, but no reaction with the aldehyde.  
  
 A positive test for the aldehyde would be to use ammoniacal silver nitrate solution, which on warming would give a silver mirror with nonanal, but would not react with the acid.  
  
**Spectroscopic method:** Both compounds would give a sharp peak in the  $\text{C}=\text{O}$  stretching region (around  $1700\text{ cm}^{-1}$ ) in the infrared (IR) spectrum. Only nonanoic acid would also give a very broad, strong peak in the range  $2500\text{--}3500\text{ cm}^{-1}$  in the IR spectrum, due to the  $\text{O}\text{--}\text{H}$  bond.
  - b The best way of distinguishing between heptanal and heptan-2-one using a chemical reaction is to separately warm them with ammoniacal silver nitrate solution. This would give a silver mirror with heptanal (an aldehyde), but there would be no reaction with heptan-2-one (a ketone).  
  
 Heptanal and heptan-2-one are isomers, so have the same molecular peak (114) in a mass spectrum. The infrared spectra are similar. The  $^1\text{H}$  NMR spectrum is the best instrumental means to distinguish them. In heptanal, all carbon atoms have neighbouring carbons bearing hydrogens, so all signals are multiplets due to spin-spin splitting. In heptan-2-one, the terminal  $\text{CH}_3$  next to  $\text{C}=\text{O}$  has no hydrogens on the neighbouring carbon, so its signal is a singlet.
  - c 3-Methylbutanal and pentanal are isomers. They are very hard to distinguish by chemical reactions, since both are aldehydes. A traditional way would be to separately add methanolic 2,4-dinitrophenylhydrazine solution to each sample, filter off the precipitated 2,4-dinitrophenylhydrazone derivative, recrystallise and dry the crystals,

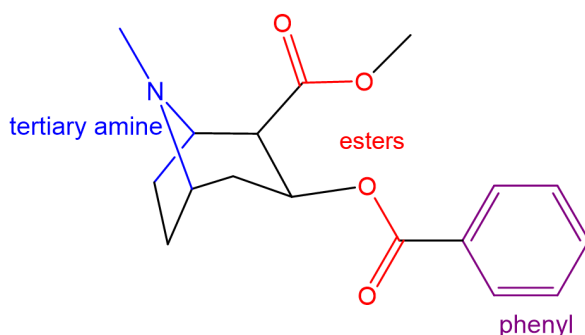
then determine their melting points. The melting point for the derivative of pentanal is 107°C and the value for the derivative of 3-methylbutanal is 123°C.

The best way to distinguish between 3-methylbutanal and pentanal is to use  $^{13}\text{C}$  NMR spectroscopy. The two terminal methyl groups on the alkyl chain in 3-methylbutanal are equivalent, so that there are just four different carbon environments in the 3-methylbutanal molecule. This means that there are four lines in its (proton decoupled)  $^{13}\text{C}$  NMR spectrum. All five carbon environments are distinct in pentanal, so there are five lines in its (proton decoupled)  $^{13}\text{C}$  NMR spectrum.  $^1\text{H}$  NMR spectra would similarly show the different number of environments.

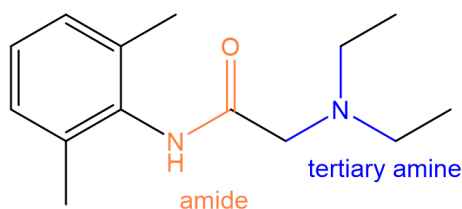
## Forensic science: testing for cocaine (pp. 22–24)

1

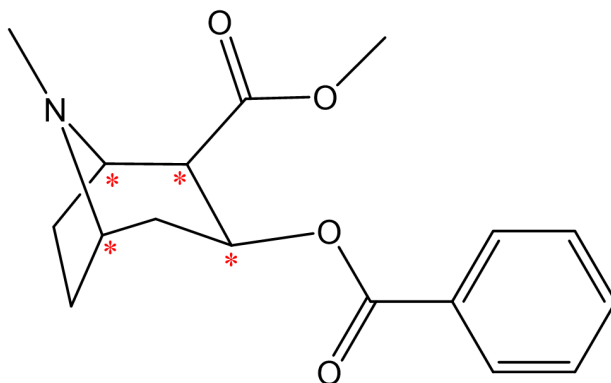
**a** Cocaine:



**b** Lidocaine:



2

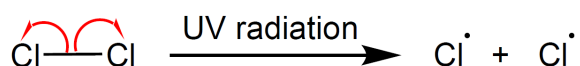


3 The  $m/z$  of a molecular ion corresponds to the molecular mass of the compound.

- a The  $m/z$  of a molecular ion of cocaine is 303.
- b The  $m/z$  of a molecular ion of lidocaine is 234.

## Curly arrows (pp. 28–29)

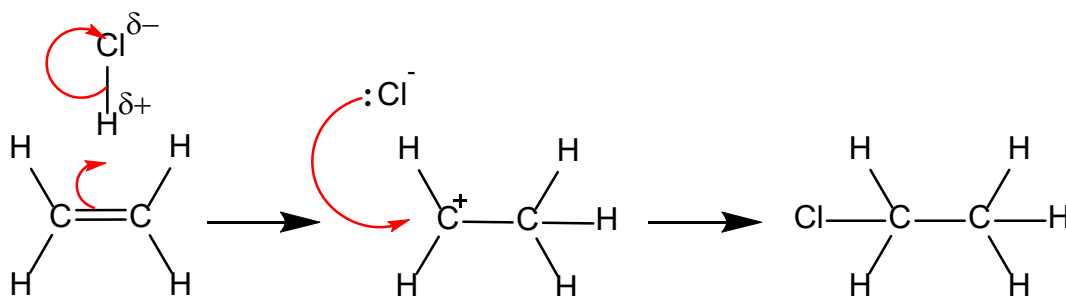
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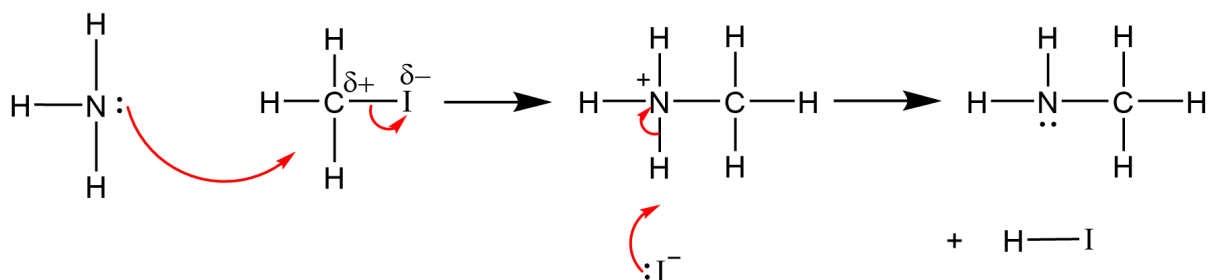
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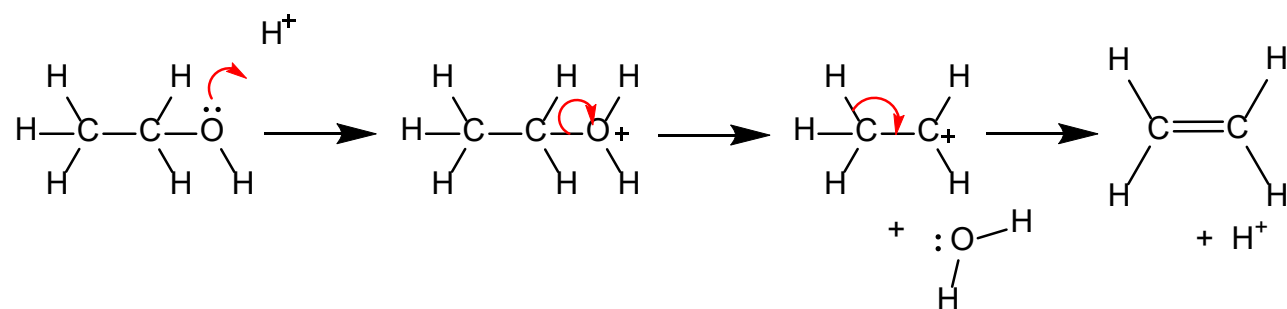
3



4



5



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