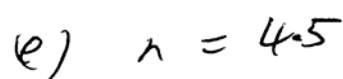
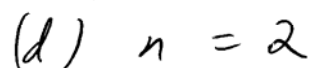
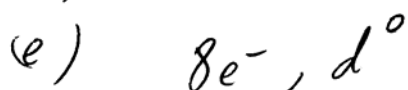
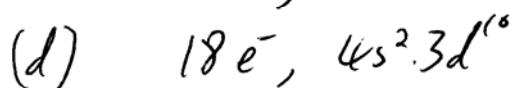
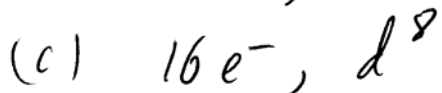
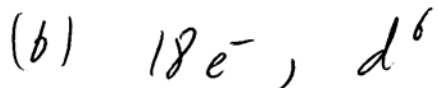
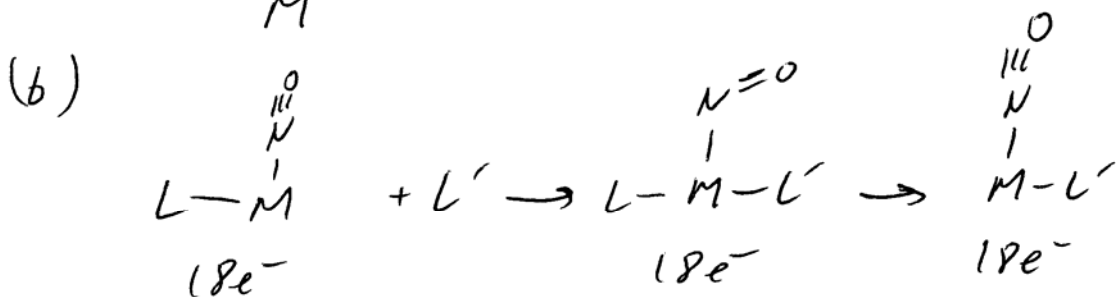


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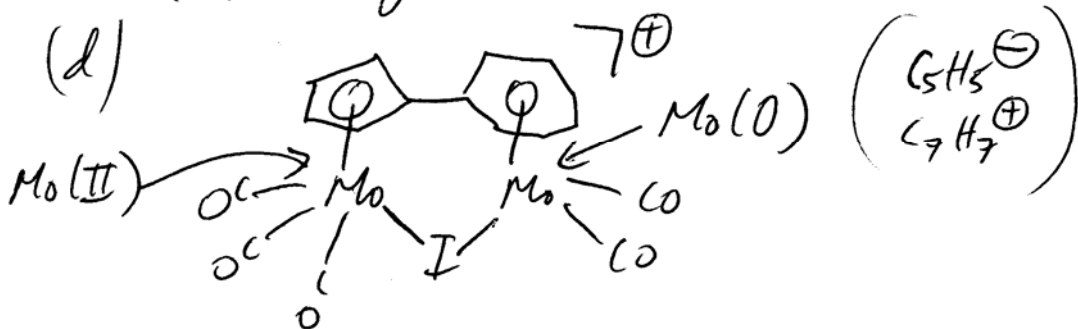


3. (a) aromatic stabilisation of the 6-membered ring:



Indenyl goes via the  $\eta^3$  intermediate shown in part (a).

- (c) (i) Steric crowding prevents dimerisation  
 (ii) late transition metal:  $d^8$ ,  $16e^-$ , square planar complex.  
 (iii) Early TM with a high OS.

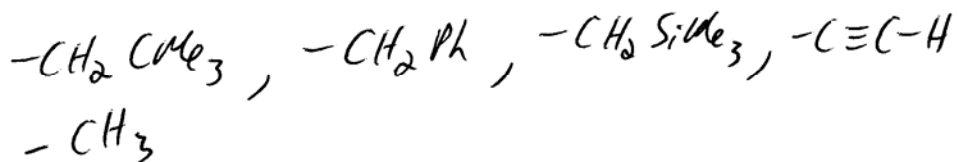


4. (a)

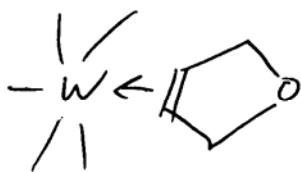


(b)  $CpV(CO)_2(RCCR)$  will show the longest C-C distance since it has a  $\pi$ -donor alkyne ligand ( $\sigma$  and  $\pi$ ).

(c) Require - a  $\beta$ -H atom  
 - coplanar M-C-C-H atoms  
 - vacant site in cis position  
 - electronically unsaturated metal ( $< 18e^-$ ).



5. (i)



I



II

(ii)

A = III

B = II

C = I

Cyclohexene is a very weak ligand and is rapidly replaced - however it is in large excess. II is a good  $\sigma$  donor and so gives the lowest  $\nu(\text{CO})$ .

(iii)

cyclohexane is a weak  $\sigma$  donor

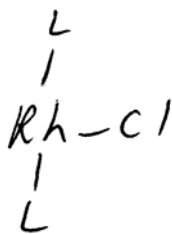
$\text{O}-\text{C}_6\text{H}_5$  is a strong  $\sigma$  donor

$\text{C}_6\text{H}_5-\text{C}$  is a  $\sigma$  donor/ $\pi$  acceptor

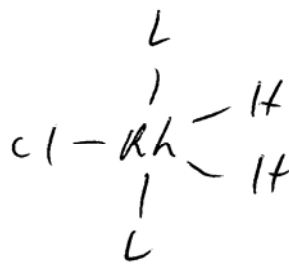
(iv) W has a low oxidation state and prefers  $\pi$ -acceptor ligands

(v) Sterically, it is easier for one O atom to get in and displace the cyclohexane than it is for 2 carbon atoms.

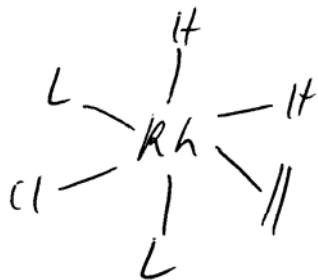
6. (a)



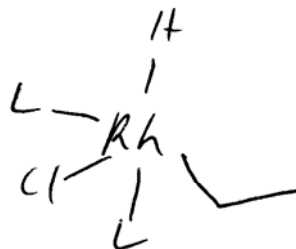
(A)



(B)

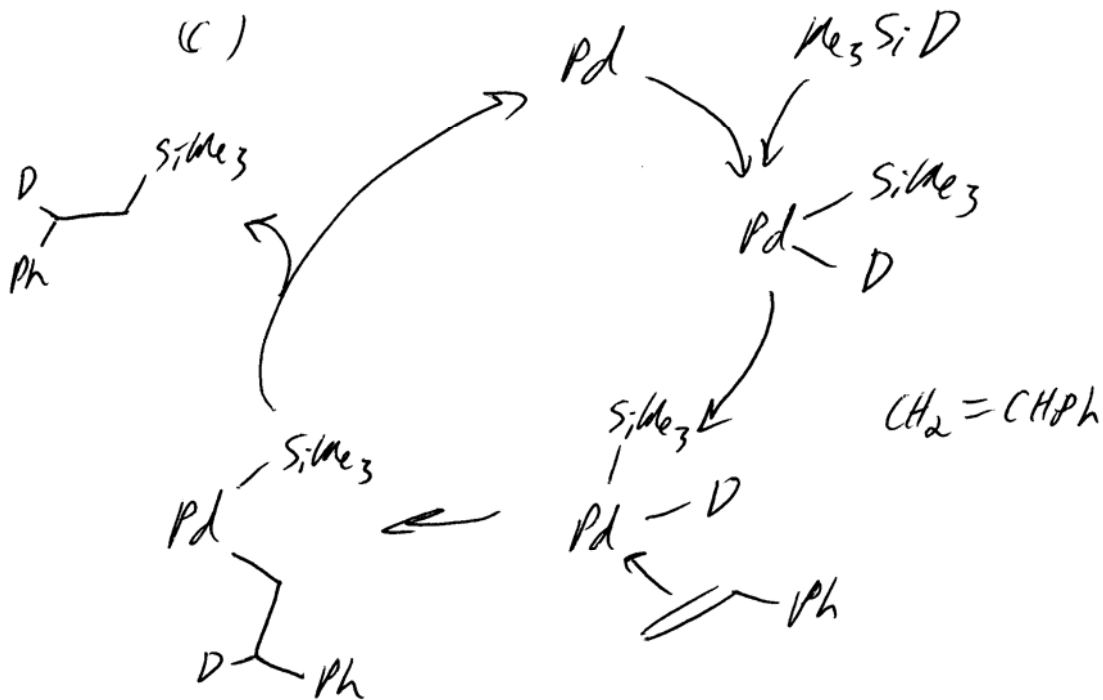


(C)



(D)

- (b)
- (1) oxidative addition
  - (2) ligand coordination
  - (3) reductive elimination



7. (a)  $\text{Mn}_2(\text{CO})_{10} + \text{Na}(\text{Hg})$   
 (b)  $\text{Mo}(\text{CO})_6 + \text{Cp}^-$   
 (c)  $\text{Co}_2(\text{CO})_8 + \text{Br}_2$   
 (d)  $\text{Cr}(\text{CO})_6 + \text{PhLi} \rightarrow \quad + \text{Me}_3\text{O}^+$   
 (e)  $\text{Fe}(\text{CO})_5 + \text{cod}$

