

# Cambridge International AS & A Level

CANDIDATE NAME					
CENTRE NUMBER			CANDIDATE NUMBER		

CHEMISTRY

Paper 4 A Level Structured Questions

May/June 2022

2 hours

9701/42

You must answer on the question paper.

No additional materials are needed.

#### **INSTRUCTIONS**

- Answer all questions.
- Use a black or dark blue pen. You may use an HB pencil for any diagrams or graphs.
- Write your name, centre number and candidate number in the boxes at the top of the page.
- Write your answer to each question in the space provided.
- Do **not** use an erasable pen or correction fluid.
- Do not write on any bar codes.
- You may use a calculator.
- You should show all your working and use appropriate units.

#### **INFORMATION**

- The total mark for this paper is 100.
- The number of marks for each question or part question is shown in brackets [].
- The Periodic Table is printed in the question paper.
- Important values, constants and standards are printed in the question paper.

## Answer **all** the questions in the spaces provided.

1	(a)	The	e solubility of the Group 2 hydroxides increases down the group.	
		Exp	plain this trend.	
			[	
	(b)	The	e solubility of Be(OH) <sub>2</sub> in water is $2.40 \times 10^{-6}  \text{g}  \text{dm}^{-3}$ at $298  \text{K}$ .	
		(i)	Write an expression for the solubility product, $K_{\rm sp}$ , of Be(OH) $_{\rm 2}$ and state its units.	
			$\mathcal{K}_{sp}$ =	
			units =	
				2]
		(ii)	Calculate the numerical value of $K_{sp}$ for Be(OH) <sub>2</sub> at 298 K.	
			$K_{sp} = \dots$	2]
		5		
	(C)		$(OH)_2$ is soluble in aqueous solutions containing an excess of hydroxide ions and forms the nplex ion $[Be(OH)_4]^{2-}$ . This complex ion has a similar shape to that of $[CuCl_4]^{2-}$ .	Ю
		(i)	Define the term complex ion.	
			[	11
		(ii)	Draw a three-dimensional diagram to show the structure of the complex ion [Be(OH) <sub>4</sub> ] <sup>2-</sup> .	
		(11)	Name the shape of the $[Be(OH)_4]^{2-}$ complex ion.	1
			shaper	 1]
			L L	ا ا

(d)	<b>(</b> i	i)	Explain why transition eler	ments can form co	mplex ions.		
							[11]
	(ii	i)	Complete Table 1.1 to she and overall polarities of the	ow the coordination	n number of each		
				Table 1.1			
			complex ion	shape	coordination number	polar or non-polar	
		cis	s-[Pt(H <sub>2</sub> NCH <sub>2</sub> CH <sub>2</sub> NH <sub>2</sub> )Cl <sub>2</sub> ]	square planar			
		[A	g(NH <sub>3</sub> ) <sub>2</sub> ] <sup>+</sup>			non-polar	
		[F	e(C <sub>2</sub> O <sub>4</sub> ) <sub>3</sub> ] <sup>3-</sup>		6		
							[2]
(e)	<b>(</b> i	i)	Define stability constant, <i>I</i>	ς <sub>stab</sub> .			
							[1]
	(ii	i)	Nickel can form complexes as shown.	with the ligands <i>en</i> ,	H <sub>2</sub> NCH <sub>2</sub> CH <sub>2</sub> NH <sub>2</sub> , a	and <i>tn</i> , H <sub>2</sub> NCH <sub>2</sub> CH	I <sub>2</sub> CH <sub>2</sub> NH <sub>2</sub> ,
			equilibrium 1 [Ni(H <sub>2</sub> O) <sub>6</sub> ]	<sup>2+</sup> + 3en ⇌ [Ni(	en) <sub>3</sub> ] <sup>2+</sup> + 6H <sub>2</sub> O	$K_{\text{stab}} = 6.76 \times 10$	17
			equilibrium 2 [Ni(H <sub>2</sub> O) <sub>6</sub>	$]^{2+} + 3tn \rightleftharpoons [Ni($	$(tn)_3]^{2+} + 6H_2O$	$K_{\text{stab}} = 1.86 \times 10$	12
			Construct an expression for State the units for $K_{\text{stab}}$ .	or the stability cons	stant, $K_{ ext{stab}}$ , for equ	uilibrium <b>1</b> .	
			$\mathcal{K}_{ ext{stab}} =$				
						units =	[2]
	(iii	i)	Describe what the $K_{\text{stab}}$ value and <b>2</b> . Use the $K_{\text{stab}}$ value stable.	lues indicate abou es to deduce whic	it the position of ech complex, [Ni(e	equilibrium for equing $n)_3$ ] <sup>2+</sup> or [Ni( $tn$ ) <sub>3</sub> ] <sup>2+</sup>	uilibrium <b>1</b> , is more
							[1]
							[Total: 16]

2	(a)	Explain why transition elements have variable oxidation states.	
			[1]
	(b)	Sketch the shape of a 3d <sub>xy</sub> orbital.	
		Z 	
		y x	
			[1]
	(c)	Explain why transition elements form coloured compounds.	

(d)	Aqueous solutions	s of copper(II)	) salts contain	$[Cu(H_2O)_6]^{2+}$ ions
(u)	Aqueous solutions	s or cobber(II)	) Sails Cortain	

Equilibrium 3 and equilibrium 4 show two reactions of these ions.

equilibrium 3 
$$[Cu(H_2O)_6]^{2+}(aq) + 2OH^-(aq) \rightleftharpoons Cu(OH)_2(s) + 6H_2O(l)$$
  
equilibrium 4  $[Cu(H_2O)_6]^{2+}(aq) + 4NH_3(aq) \rightleftharpoons [Cu(NH_3)_4(H_2O)_2]^{2+}(aq) + 4H_2O(l)$ 

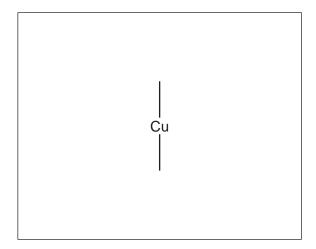
(i) State the colour of  $Cu(OH)_2(s)$  and  $[Cu(NH_3)_4(H_2O)_2]^{2+}(aq)$ .

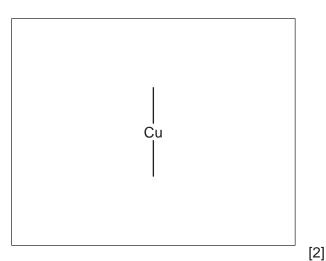
$colour or ca(Ori)_2(s) \dots$	(OH) <sub>2</sub> (s)
colour of $[Cu(NH_3)_4(H_2O)_2]^{2+}(aq)$	(NH <sub>3</sub> ) <sub>4</sub> (H <sub>2</sub> O) <sub>2</sub> ] <sup>2+</sup> (aq)[1]

(ii) Use Le Chatelier's principle to explain why a precipitate is formed when NaOH(aq) is added dropwise to  $[Cu(H_2O)_6]^{2+}(aq)$ .


(e) There are two possible stereoisomers with the formula  $[Cu(NH_3)_4(H_2O)_2]^{2+}$ .

Draw three-dimensional diagrams to show the two stereoisomers.





[Total: 9]

(a) (i)	Define first electron affinity.
	[2]
(ii)	The first electron affinity of an atom is usually an exothermic process, whereas the second electron affinity is an endothermic process.
	Suggest why.
	[1]
(iii)	Describe the general trend in first electron affinities for Cl, Br and I. Explain your answer.
	[2]

(b) Table 3.1 shows energy changes to be used in this question and in (c).

Table 3.1

energy change	value/kJ mol <sup>-1</sup>
standard enthalpy change of atomisation of zinc	+131
first ionisation energy of zinc	+906
second ionisation energy of zinc	+1733
standard enthalpy change of formation of ZnI <sub>2</sub> (s)	-208
lattice energy, $\Delta H_{\text{latt}}^{e}$ , of zinc iodide, $\text{ZnI}_{2}(s)$	-2605
first ionisation energy of iodine	+1008
second ionisation energy of iodine	+1846
I–I bond energy	+151
enthalpy change of sublimation of iodine, $I_2(s) \rightarrow I_2(g)$	+62

Calculate the first electron affinity for iodine. Use relevant data from Table 3.1 in your working
It may be helpful to draw a labelled energy cycle.
Show all working.

first alastron offinit	v for iodino -	k I mal-1	[2]
nirst electron animit	y for foatne =	kJ mol <sup>-1</sup>	131

(c) Predict how  $\Delta H_{\text{latt}}^{\bullet}$  of  $\text{CdI}_2(s)$  differs from  $\Delta H_{\text{latt}}^{\bullet}$  of  $\text{ZnI}_2(s)$ . Place a tick ( $\checkmark$ ) in the appropriate box in Table 3.2.

Table 3.2

$\Delta H_{\rm latt}^{\rm e}$ of ${\rm CdI_2}({\rm s})$ is less negative than $\Delta H_{\rm latt}^{\rm e}$ of ${\rm ZnI_2}({\rm s})$	$\Delta H_{\text{latt}}^{\bullet}$ of $\text{CdI}_2(s)$ is the same as $\Delta H_{\text{latt}}^{\bullet}$ of $\text{ZnI}_2(s)$	$\Delta H_{\text{latt}}^{\bullet}$ of $\text{CdI}_2(s)$ is more negative than $\Delta H_{\text{latt}}^{\bullet}$ of $\text{ZnI}_2(s)$

Explain your answe	er.	
		 [1]
		[Total: 9]

4 (a) Calcium carbonate decomposes on heating.

$$CaCO_3(s) \rightarrow CaO(s) + CO_2(g)$$

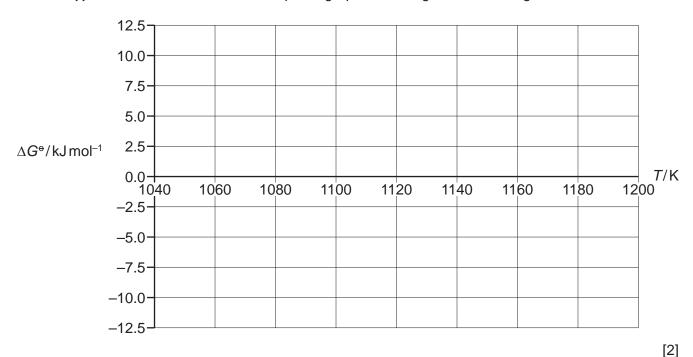
Table 4.1 shows the values of the Gibbs free energy change,  $\Delta G^{\circ}$ , for this reaction at various temperatures.

Table 4.1

T/K	$\Delta G^{\circ}/\text{kJ}\text{mol}^{-1}$
1050	9.9
1085	4.3
1120	-1.3
1148	-5.8
1176	-10.3

Assume the standard enthalpy change,  $\Delta H^{e}$ , and the standard entropy change,  $\Delta S^{e}$ , for this reaction remain constant over this temperature range.

(i) Use the data in Table 4.1 to plot a graph of  $\Delta G^{\circ}$  against T on the grid.



(ii) Calculate the gradient of your graph. Determine the  $\Delta S^e$  in  $J K^{-1} mol^{-1}$  for this reaction. Show all working.

$$\Delta S^{e} = .... J K^{-1} mol^{-1}$$
 [2]

(b)		oup 1 hydrogencarbonates, MHCO <sub>3</sub> , decompose on gentle heating to give the corresponding cal carbonate, carbon dioxide and water vapour.
	(i)	Write an ionic equation for the decomposition of the hydrogencarbonate ion.
		[1]
	(ii)	The thermal stability of Group 1 hydrogencarbonates increases down the group.
		Suggest an explanation for the trend in thermal stability of the Group 1 hydrogencarbonates.
		[2]
(c)	The	buffer system in seawater contains a mixture of HCO <sub>3</sub> <sup>-</sup> and H <sub>2</sub> CO <sub>3</sub> .
		equilibrium 5 $H_2CO_3 + H_2O \rightleftharpoons HCO_3^- + H_3O^+$
	(i)	Define a buffer solution.
		[2]
	(ii)	Construct <b>two</b> equations to show how equilibrium <b>5</b> acts as a buffer solution.
		roa
	/:::\	The IUCO -1/IU CO 1 ratio in a comple of economic at 44.4
(	(iii)	The [HCO <sub>3</sub> -]/[H <sub>2</sub> CO <sub>3</sub> ] ratio in a sample of seawater is 14.1.
		Calculate the pH of this sample. [p $K_a$ : H <sub>2</sub> CO <sub>3</sub> , 6.35]

[Total: 14]

**5 (a)** Complete Table 5.1 to predict the substance liberated at each electrode during electrolysis of the indicated electrolyte with inert electrodes.

Table 5.1

electrolyte	substance liberated at the anode	substance liberated at the cathode
PbBr <sub>2</sub> (I)		
concentrated NaCl(aq)		
Cu(NO <sub>3</sub> ) <sub>2</sub> (aq)		

[3]

**(b)** An electrolytic cell is set up to determine a value for the Avogadro constant, *L*. The electrolyte is dilute sulfuric acid and both electrodes are copper.

When a current of 0.600A is passed through the acid for 30.0 minutes, the anode decreases in mass by 0.350 g.

(i)	State the relationship between the Faraday constant, <i>F</i> , and the Avogadro constant, <i>L</i> .	
	[1	1]

(ii) Use the experimental information in (b) and data from the table on page 23 to calculate a value for the Avogadro constant, L.

Show all working.

Avogadro constant,  $L = \dots$  [4]

[Total: 8]

6 (a) The reagent and conditions required for the nitration of benzene, benzoic acid and phenol are shown in Table 6.1.

Table 6.1

compound	reagents and conditions for nitration
benzene	concentrated HNO <sub>3</sub> , 50 °C, concentrated H <sub>2</sub> SO <sub>4</sub> catalyst
benzoic acid	concentrated HNO <sub>3</sub> , 100 °C, concentrated H <sub>2</sub> SO <sub>4</sub> catalyst
phenol	dilute HNO <sub>3</sub> (aq), 20°C

Concentrated HNO<sub>3</sub> reacts with concentrated H<sub>2</sub>SO<sub>4</sub> to generate the electrophile NO<sub>2</sub><sup>+</sup>.

(i) Complete Fig. 6.1 to show the mechanism of the reaction between benzene and  $NO_2^+$ . Include all relevant curly arrows and charges.

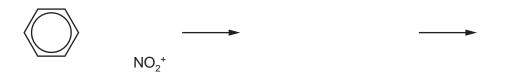


Fig. 6.1

[3]

(ii) Write an equation to show how H<sub>2</sub>SO<sub>4</sub> is regenerated.

(b) Draw the major products from the mononitration of benzoic acid and of phenol.

major product from benzoic acid

major product from phenol	

[2]

(c)		tration of benzene, benzoic acid and phere reference to the structures of the three	
	>	>	
	easiest		least easy
			[4]

(d) The azo compound Congo Red is used as an acid-base indicator and can be made by the route shown in Fig. 6.2.

In step 3 of this synthesis, compound  $\mathbf{Y}$  reacts with compound  $\mathbf{Z}$ . Compound  $\mathbf{Z}$  is made from compound  $\mathbf{X}$ . Assume that the  $-SO_3^-Na^+$  groups do not react.

Fig. 6.2

- (i) Suggest structures for compounds X, Y and Z and draw them in the boxes in Fig. 6.2. [3]
- (ii) Give the reagents and conditions for step 1 and step 2.

step 1	
step 2	
	[3]

[Total: 16]

			14		
(a	) S	ate the uses of T	MS and D <sub>2</sub> O in NMR spectroscopy	<i>t</i> .	
	Т	MS			
	D	<sub>2</sub> O			 [1]
(b	) T	ne three isomeric	: ketones with molecular formula $C_{\scriptscriptstyle{5}}$	<sub>5</sub> H <sub>10</sub> O are:	
	•	pentan-2-one pentan-3-one 3-methylbutar	one.		
	(i)		ole 7.1 to show the number of pein the carbon-13 NMR spectrum fo	eaks observed in the proton ( <sup>1</sup> H) Nor each compound listed.	NMR
			Table 7.1		
		ketone	number of peaks observed in the proton (1H) NMR spectrum	number of peaks observed in the carbon-13 NMR spectrum	
	p	entan-2-one			
	p	entan-3-one			
	3-m	nethylbutanone			
_					[2]
	(ii)	State <b>all</b> the k	etones with molecular formula C <sub>5</sub> H	<sub>10</sub> O that have:	
		a doublet in th	eir proton (¹H) NMR spectrum		
		a singlet in the	eir proton ( <sup>1</sup> H) NMR spectrum.		
					[2]

(c) Cortisone,  $C_{21}H_{28}O_5$ , is a naturally occurring chemical that contains chiral carbon atoms.

### cortisone

Fig. 7.1

(i)	Deduce the number of chiral carbon atoms in one molecule of cortisone.	
		[1]
(ii)	Cortisone is reacted with an excess of NaBH <sub>4</sub> .	
	State the molecular formula of the organic compound formed.	
		[1]
(iii)	Cortisone is an optically active molecule.	
	Explain what is meant by optically active.	
		[1]
	[Total	: 8]

[2]

8	(a)	Compare the relative acidities of ethanol, ethanoic acid, chloroethanoic acid and ph Explain your reasoning.	enol.
		most acidic   least a	
			[4]
	(b)	An excess of ethanedioic acid, HOOCCOOH(aq), is reacted with warm acidified KN	InO₄(aq).
		State the type of reaction undergone by ethanedioic acid. Describe what you would observe. Write an equation for this reaction.	
		Your equation can use [O] or [H] as necessary.	
		type of reaction	
		observations	
		equation	

(c) A section of a polyester is shown.

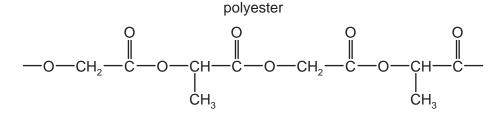


Fig. 8.1

Draw the structures of the two monomers that form this polyester.



**(d)** Serine can polymerise to form two different types of condensation polymer; a polyester and a polypeptide.

serine

Fig. 8.2

Draw the structure of the polypeptide showing  ${\bf two}$  repeat units. The peptide linkage should be shown displayed.

(e) Explain why condensation polymers normally biodegrade more readily than addition polymers.

[1]

[Total: 11]

[2]

**9** The structure of cyclohexylamine is shown in Fig. 9.1.

cyclohexylamine



Fig. 9.1

(a)	Compare the relative basicities of ammonia, cyclohexylamine and phenyl Explain your reasoning.	amine.
	most basic	least basic
		[3]

(b) Cyclohexylamine reacts with ethanoyl chloride to form the corresponding amide, L.

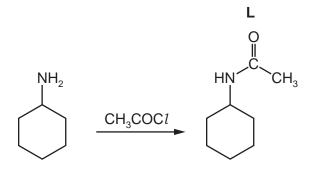


Fig. 9.2

(i) Name the mechanism for the reaction shown in Fig. 9.2.

.....[1]

(ii) Complete the mechanism of the reaction between cyclohexylamine and CH<sub>3</sub>COC*l*.

R-NH<sub>2</sub> is used to represent cyclohexylamine.

Include all relevant lone pairs of electrons, curly arrows, charges and partial charges.



 $R-NH_2$ 

[4]

(iii) The reaction between cyclohexylamine and an excess of  $CH_3COC_l$  forms compound M. Compound M has the molecular formula  $C_{10}H_{17}NO_2$ .

Suggest and draw the structure of M.

[1]

[Total: 9]

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## Important values, constants and standards

molar gas constant	$R = 8.31 \mathrm{J}\mathrm{K}^{-1}\mathrm{mol}^{-1}$
Faraday constant	$F = 9.65 \times 10^4 \mathrm{C}\mathrm{mol}^{-1}$
Avogadro constant	$L = 6.022 \times 10^{23} \mathrm{mol^{-1}}$
electronic charge	$e = -1.60 \times 10^{-19} \mathrm{C}$
molar volume of gas	$V_{\rm m} = 22.4 {\rm dm^3  mol^{-1}}$ at s.t.p. (101 kPa and 273 K) $V_{\rm m} = 24.0 {\rm dm^3  mol^{-1}}$ at room conditions
ionic product of water	$K_{\rm w} = 1.00 \times 10^{-14} \rm mol^2  dm^{-6}  (at  298  K  (25  {}^{\circ}C))$
specific heat capacity of water	$c = 4.18 \mathrm{kJ  kg^{-1}  K^{-1}}  (4.18 \mathrm{J  g^{-1}  K^{-1}})$

The Periodic Table of Elements

	~		ന	E ~		m	ç ⟨\		<u>_</u>	5 6		_	<u>5</u> 8		ď	E &		_	E	_	<u></u>	sson			
	18	2	Ĭ	heliu 4.0	10	ž	20.5	18	Ā	argon 39.9	36	Ž	krypt 83.8	54	×	xenc 131.	88	Ϋ́	radc	111	ŏ	ogane			
	17				6	Щ	fluorine 19.0	17	Cl	chlorine 35.5	32	ğ	bromine 79.9	53	П	iodine 126.9	82	At	astatine	117	<u>⊾</u>	tennessine -			
	16				80	0	oxygen 16.0	16	ഗ	sulfur 32.1	34	Se	selenium 79.0	52	<u>e</u>	tellurium 127.6	84	Ъо	polonium –	116	_	livermorium —			
	15				7	z	nitrogen 14.0	15	۵	phosphorus 31.0	33	As	arsenic 74.9	51	Sp	antimony 121.8	83	<u>.</u>	bismuth 209.0	115	Mc	moscovium			
	4				9	ပ	carbon 12.0	14	S	silicon 28.1	32	Ge	germanium 72.6	20	Sn	tin 118.7	82	Pp	lead 207.2	114	Εl	flerovium			
	13				5	Ф	boron 10.8	13	Ν	aluminium 27.0	31	Ga	gallium 69.7	49	In	indium 114.8	81	11	thallium 204.4	113	R	nihonium			
										12	30	Zu	zinc 65.4	48	g	cadmium 112.4	80	Ρ̈́	mercury 200.6	112	ပ်	copernicium			
										1	29	D O	copper 63.5	47	Ag	silver 107.9	62	Αu	gold 197.0	111	Rg	roentgenium -			
dn										10	28	z	nickel 58.7	46	Pd	palladium 106.4	78	Ŧ	platinum 195.1	110	Ds	darmstadtium -			
Group										0	27	ပိ	cobalt 58.9	45	R	rhodium 102.9	77	Ir	iridium 192.2	109	¥	meitnerium -			
		-	I	hydrogen 1.0						80	26	Ьe	iron 55.8	44	Ru	ruthenium 101.1	92	SO	osmium 190.2	108	¥	hassium			
					,					7	25	Mn	manganese 54.9	43	ည	technetium -	75	Re	rhenium 186.2	107	B	bohrium –			
									loc	SS			9	24	ပ်	chromium 52.0	42	Mo	molybdenum 95.9	74	≯	tungsten 183.8	106	Sg	seaborgium
						Key	atomic number	atomic symbo	name relative atomic mass			2	23	>	vanadium 50.9	41	g	niobium 92.9	73	Д	tantalum 180.9	105	9	dubnium	
								ato	rela			4	22	F	titanium 47.9	40	Zr	zirconium 91.2	72	Ξ	hafnium 178.5	104	쪼	rutherfordium -	
								_		က	21	Sc	scandium 45.0	39	>	yttrium 88.9	57-71	lanthanoids		89–103	actinoids				
	7				4	Be	beryllium 9.0	12	Mg	magnesium 24.3	20	Ca	calcium 40.1	38	လွ	strontium 87.6	56	Ba	barium 137.3	88	Ra	radium			
	_				8	:=	lithium 6.9	11	Na	sodium 23.0	19	×	potassium 39.1	37	Rb	rubidium 85.5	55	S	caesium 132.9	87	ь	francium —			

						_
71	3	lutetium 175.0	103	ت	lawrencium	I
		ytterbium 173.1			_	1
69	E	thulium 168.9	101	Md	mendelevium	_
89	ш	erbium 167.3	100	Fm	fermium	1
29	웃	holmium 164.9	66	Es	einsteinium	-
99	Ś	dysprosium 162.5	86	ŭ	californium	1
65	<u>م</u>	terbium 158.9	26	益	berkelium	_
49	р	gadolinium 157.3	96	CB	curium	1
63	П	europium 152.0	92	Am	americium	1
62	Sm	samarium 150.4	94	Pn	plutonium	_
61	Pm	promethium -	93	ď	neptunium	1
09	P	neodymium 144.4	92	$\supset$	uranium	238.0
29	Ā	praseodymium 140.9	91	Ра	protactinium	231.0
28	Ö	cerium 140.1	06	H	thorium	232.0
22	Ľ	lanthanum 138.9	89	Ac	actinium	-

lanthanoids actinoids

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