

Fig. 5.

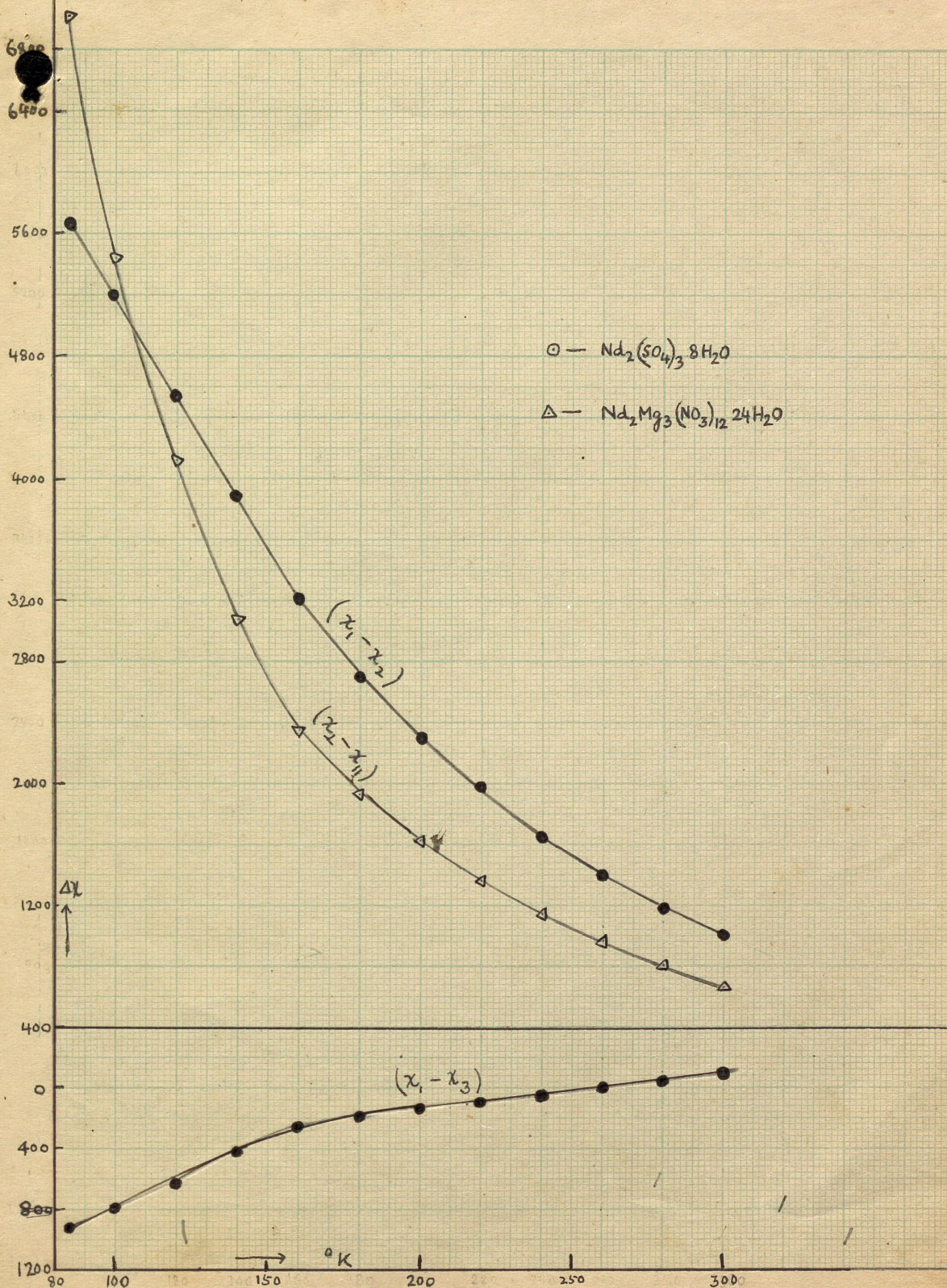


fig 6

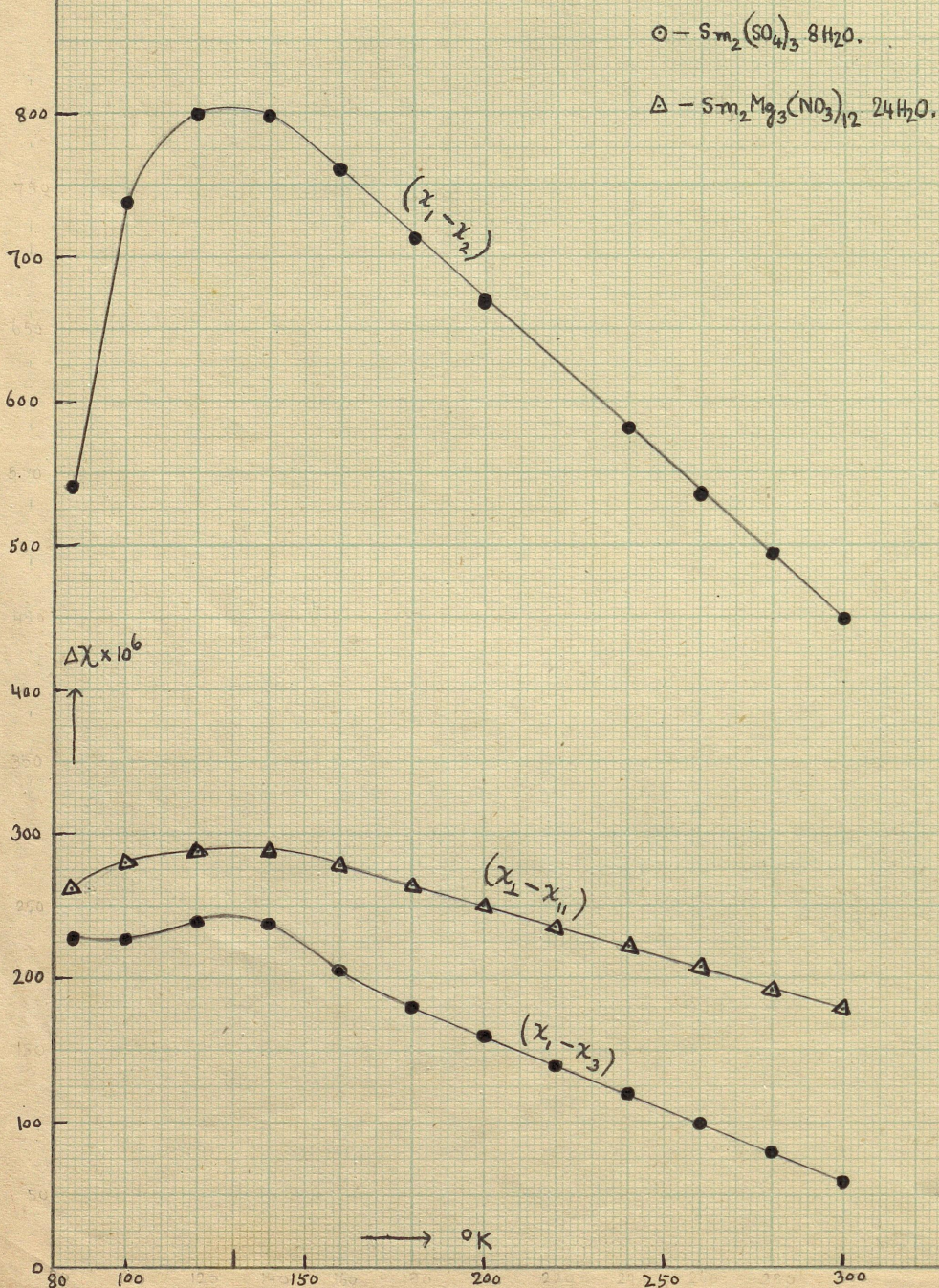


fig 7

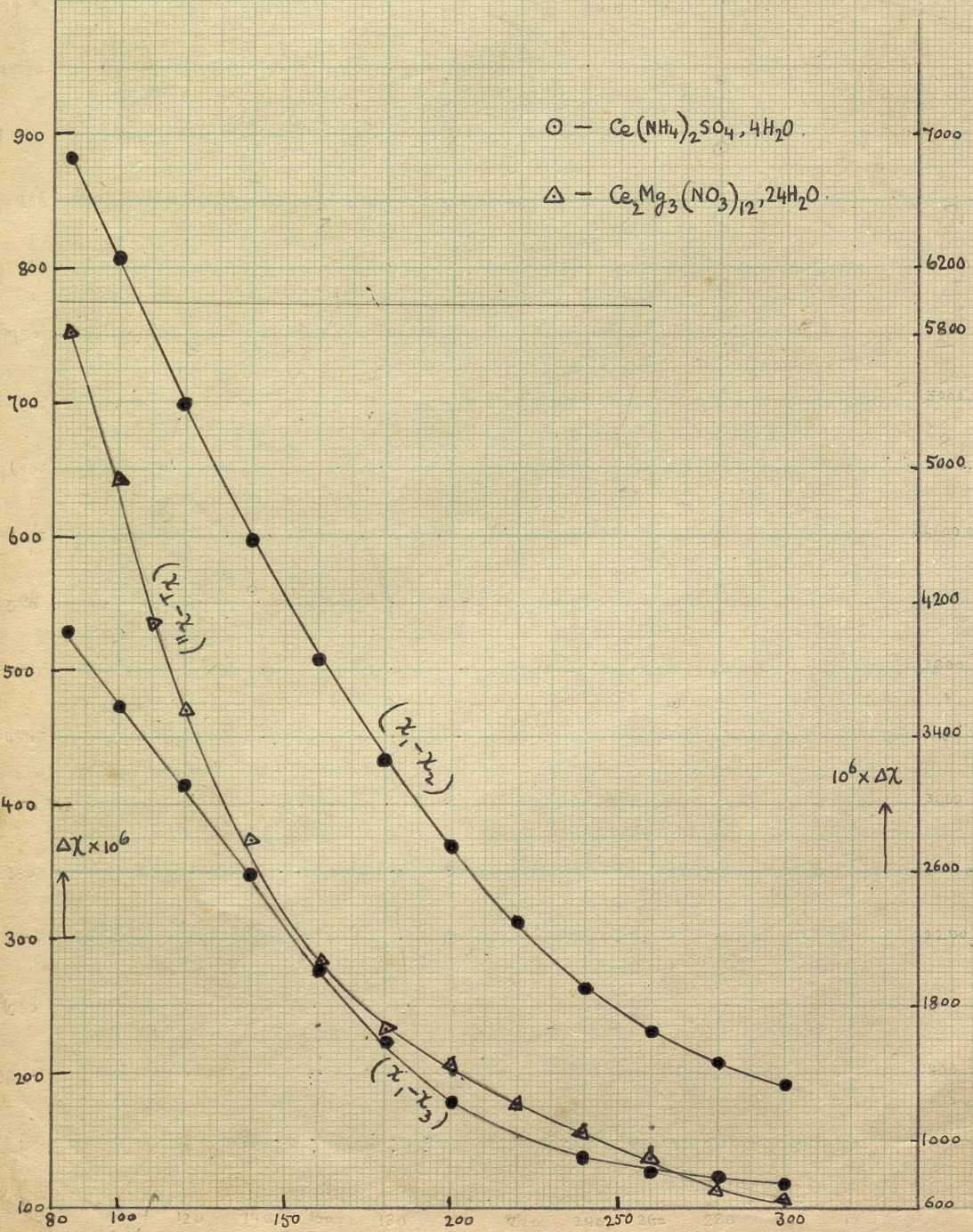


fig. 8

Temperature Variation
of
the square of the
effective moments.

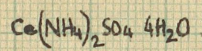
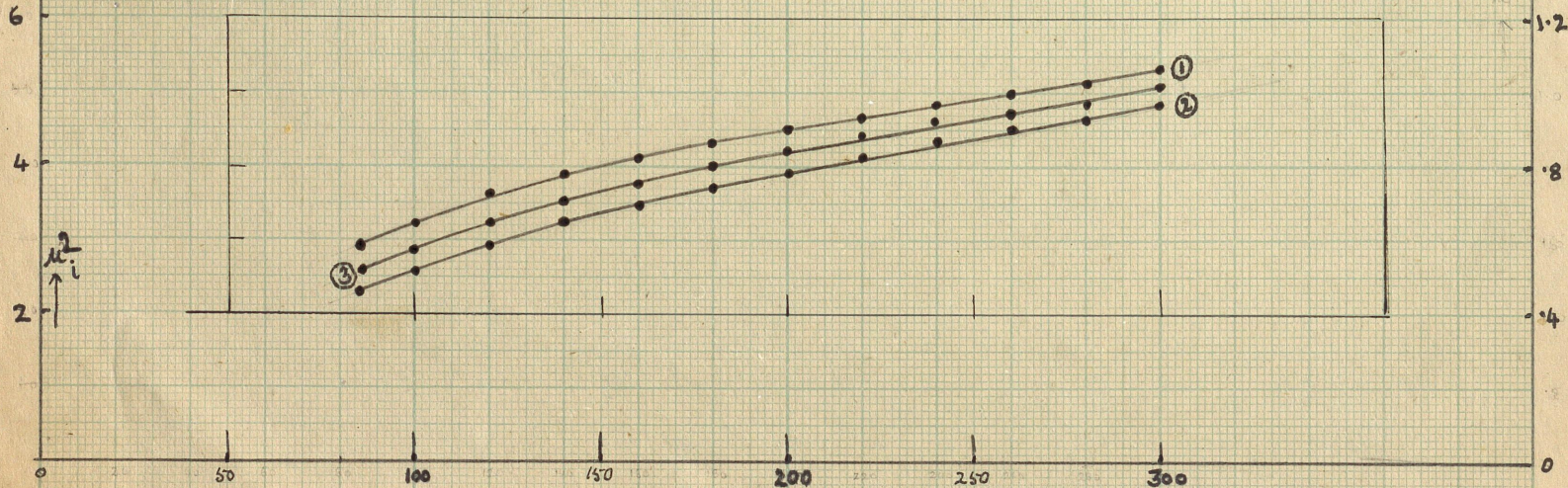


Fig (9)



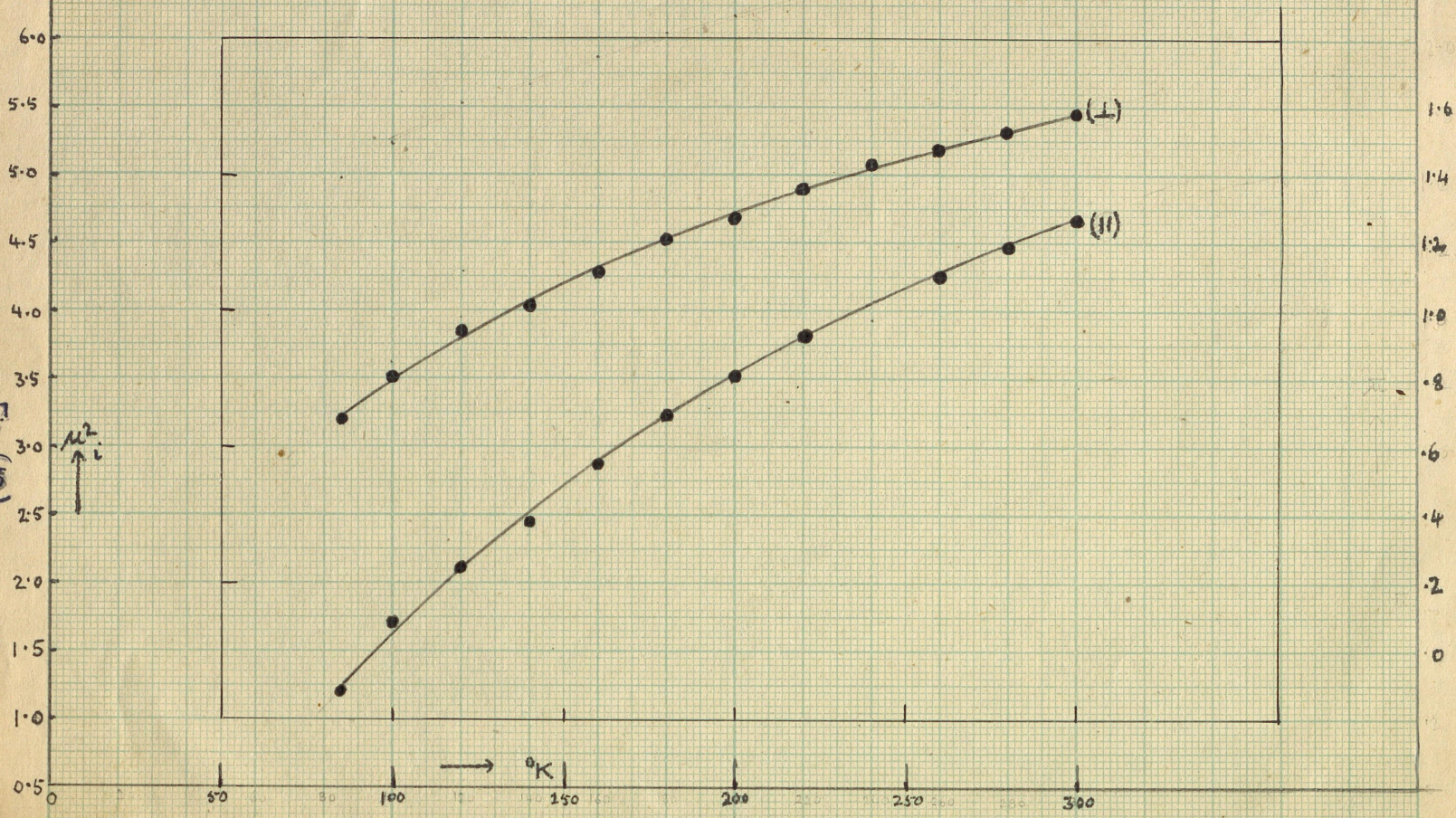
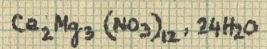
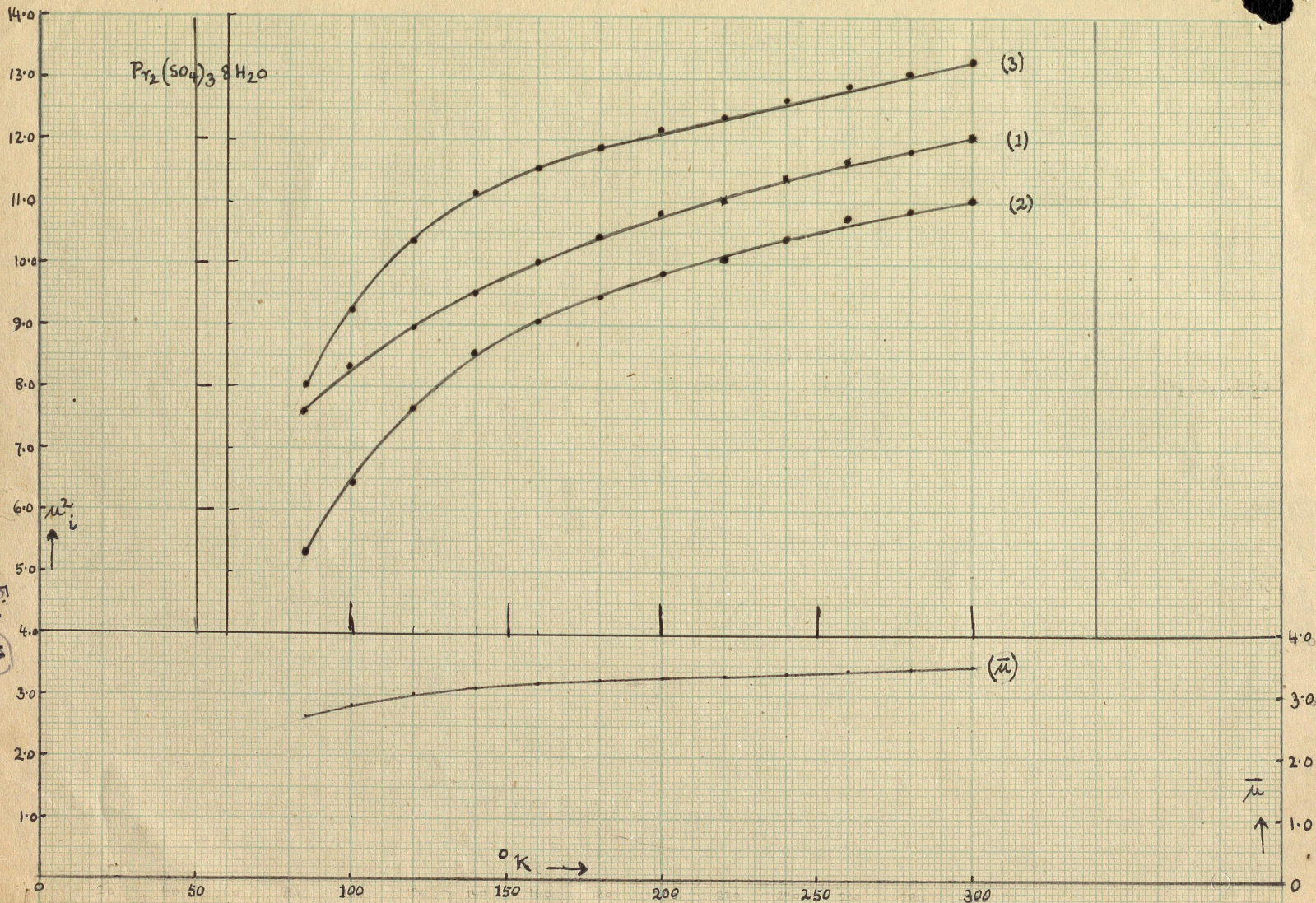
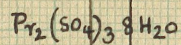


Fig (10)



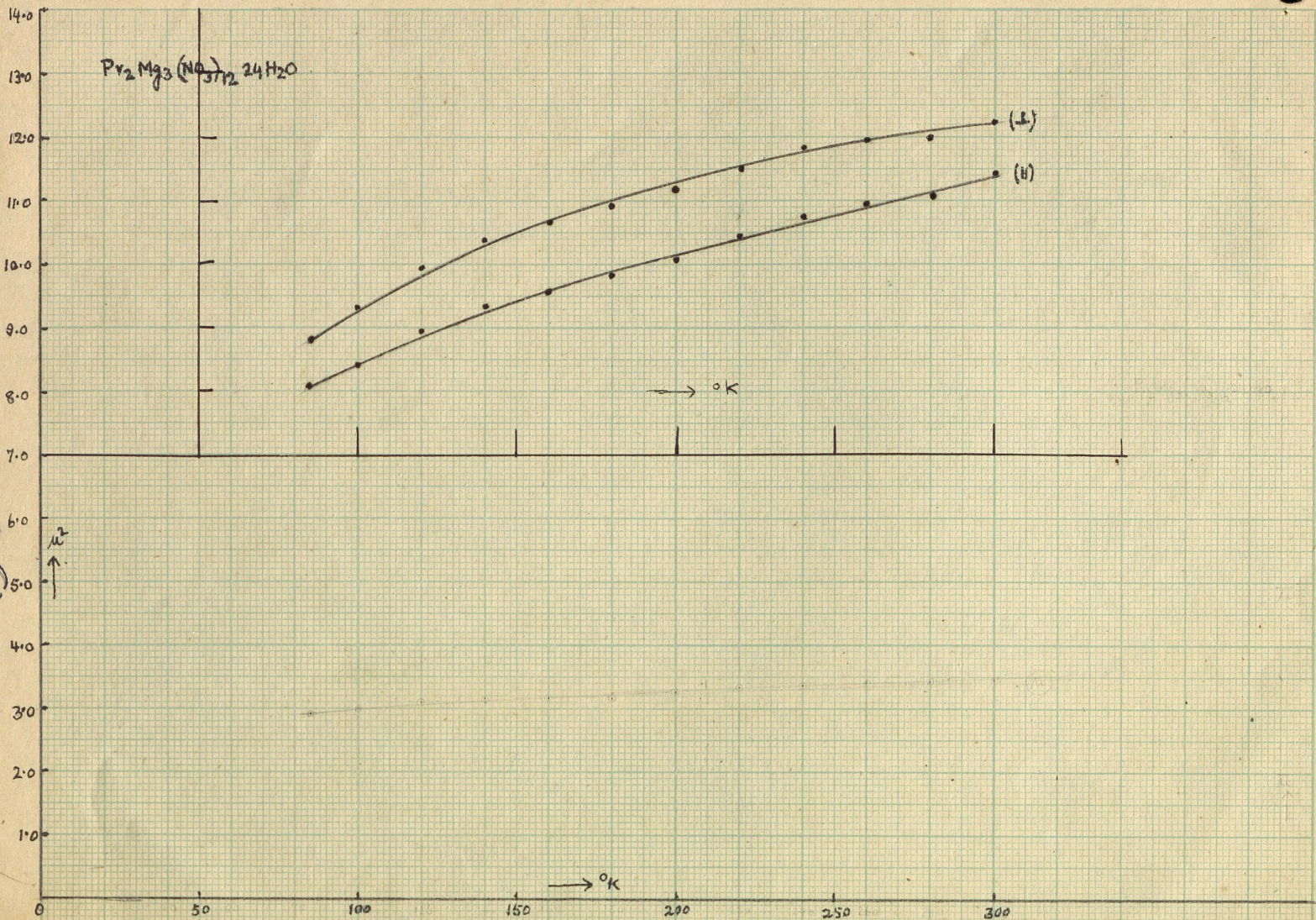
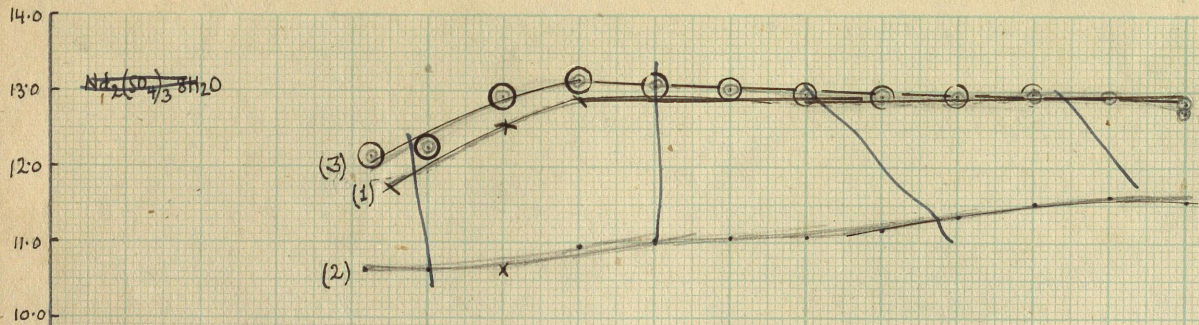
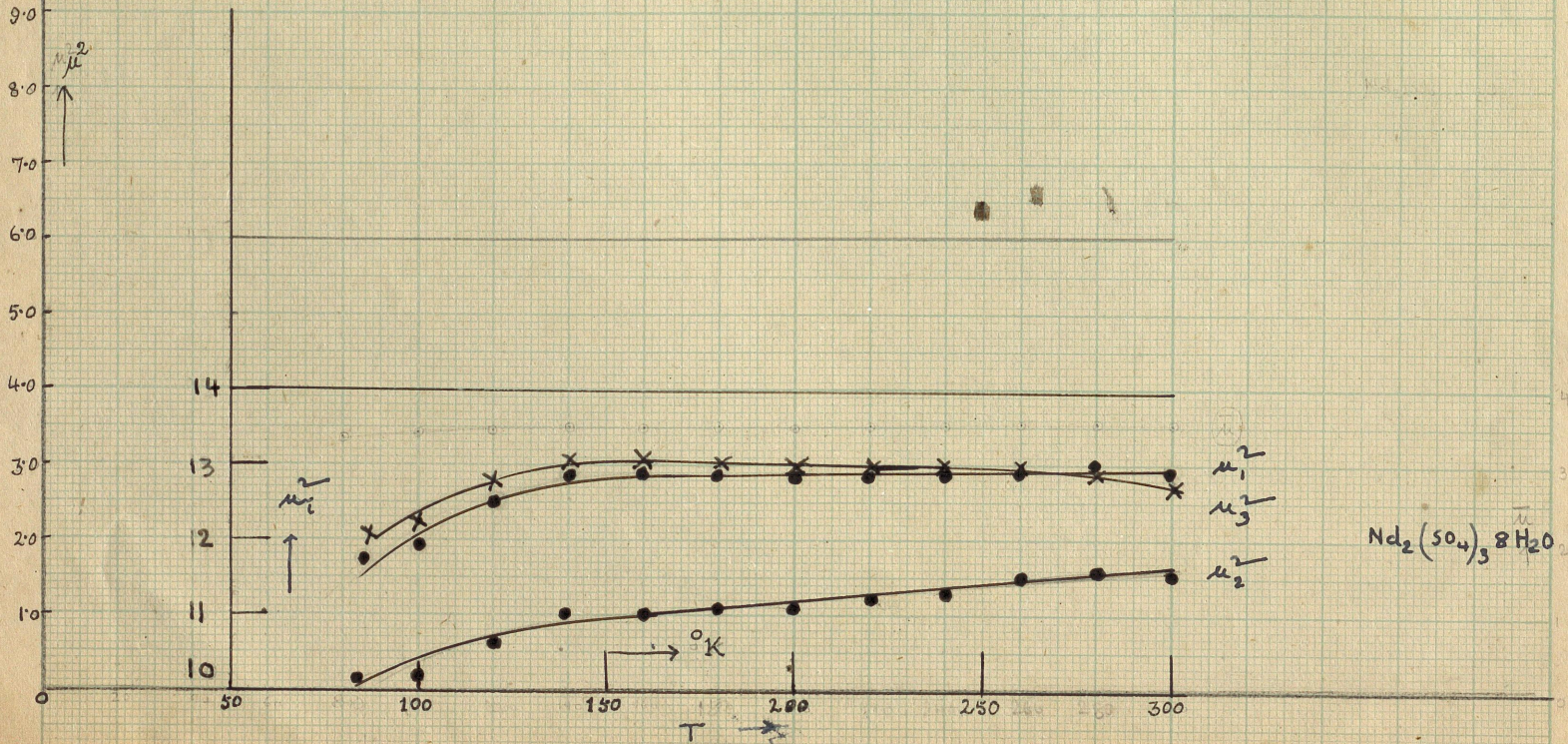


Fig (12)



(Fig 13)



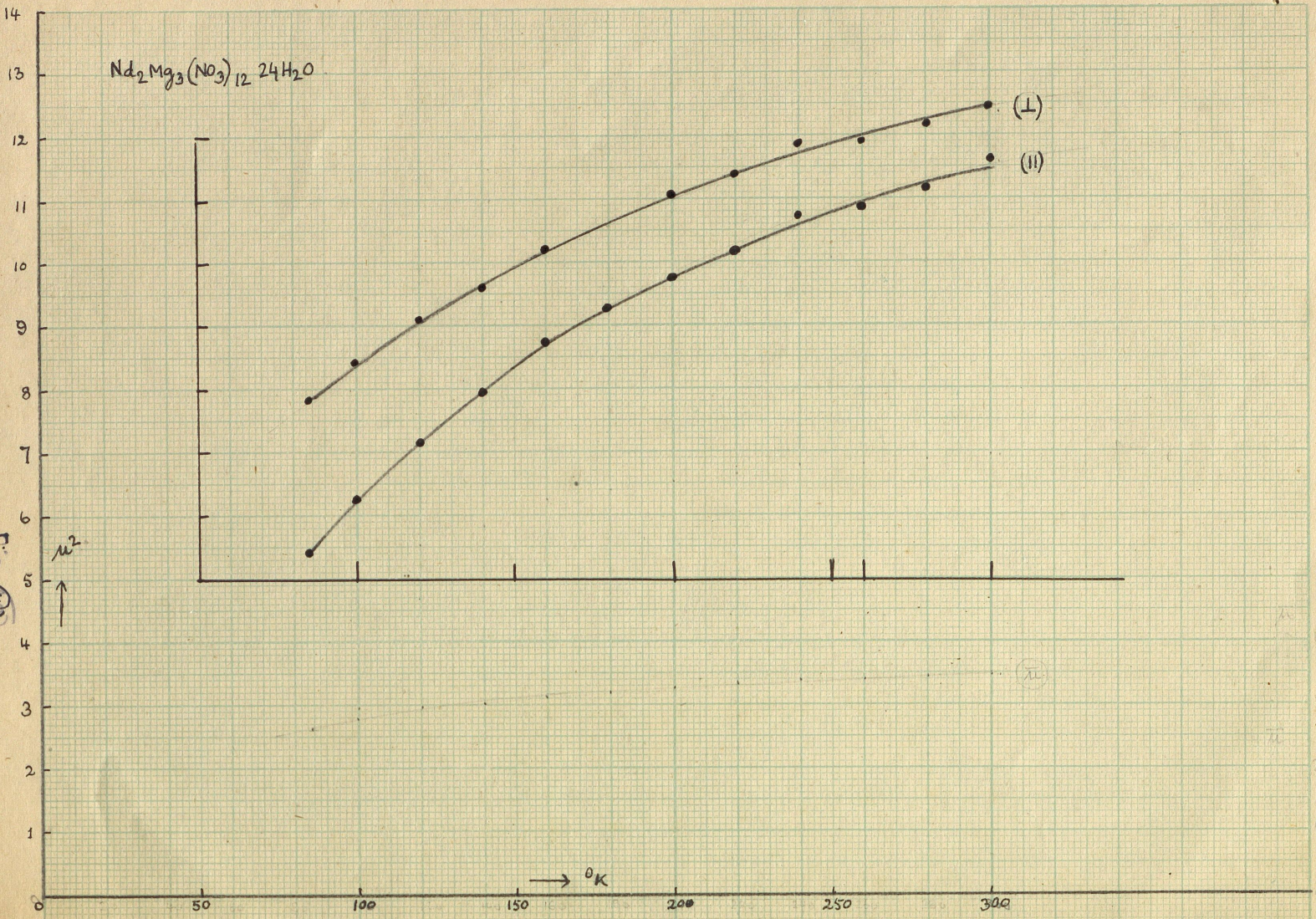
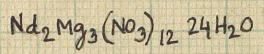


Fig. (14)

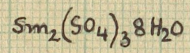
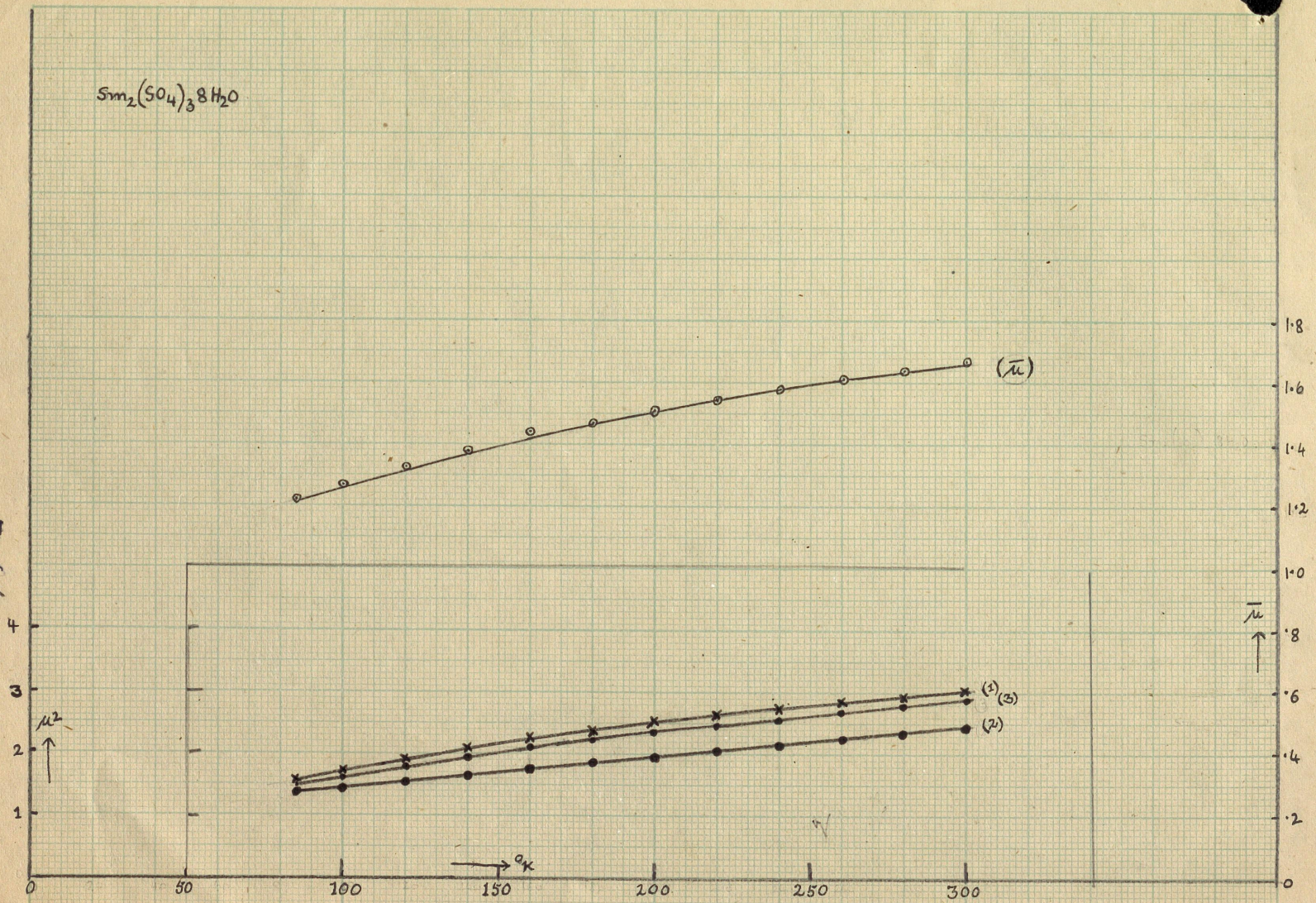


Fig (15)



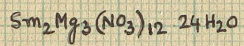


Fig 160

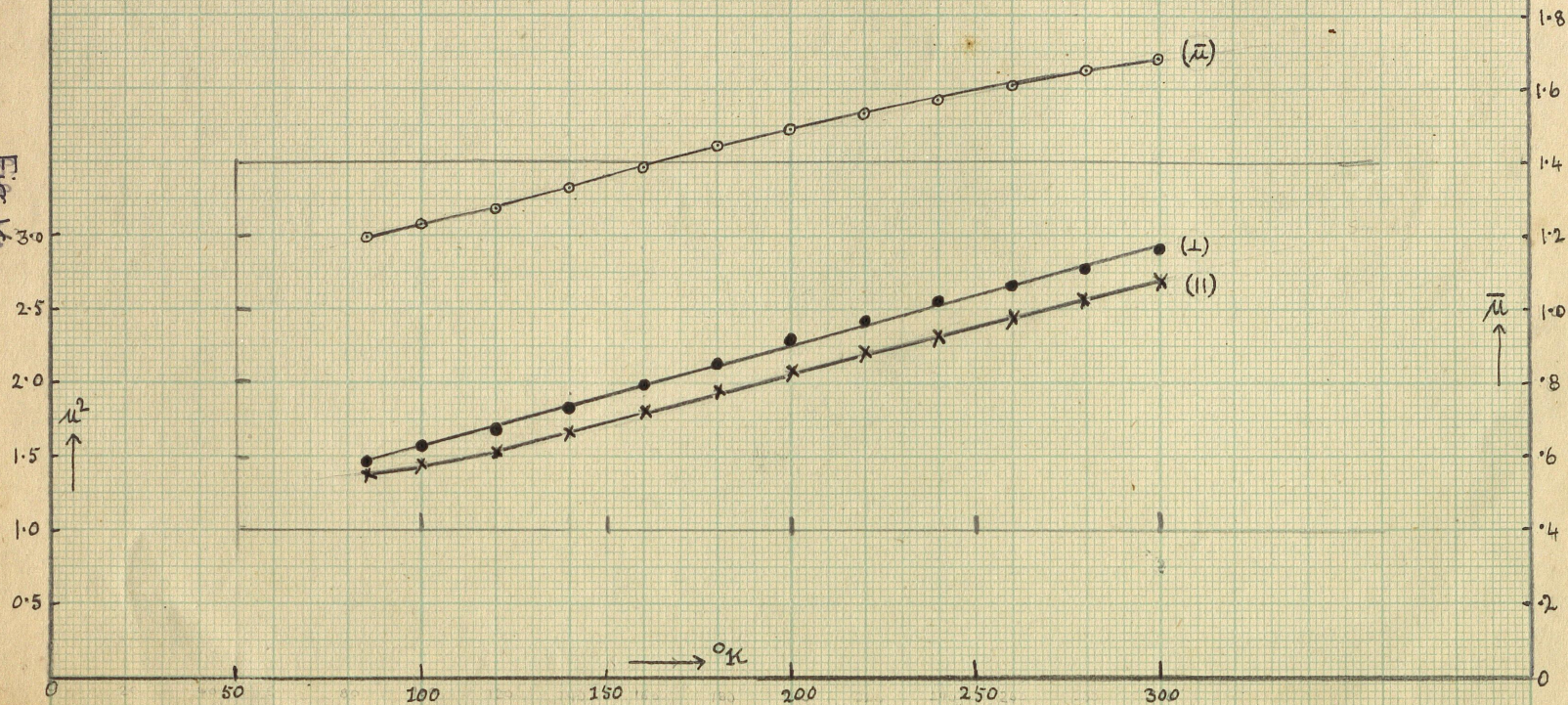
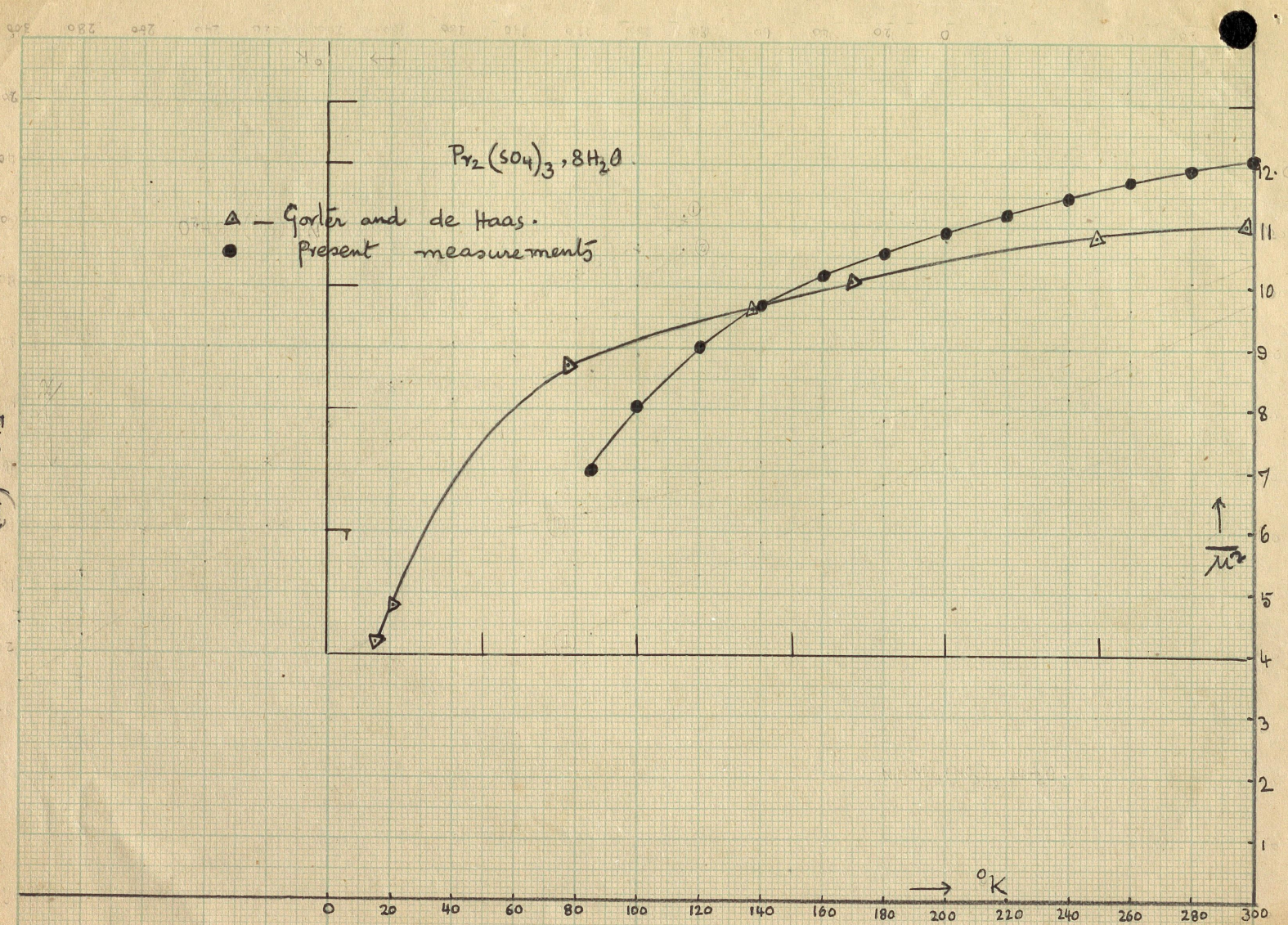


Fig. - (17)



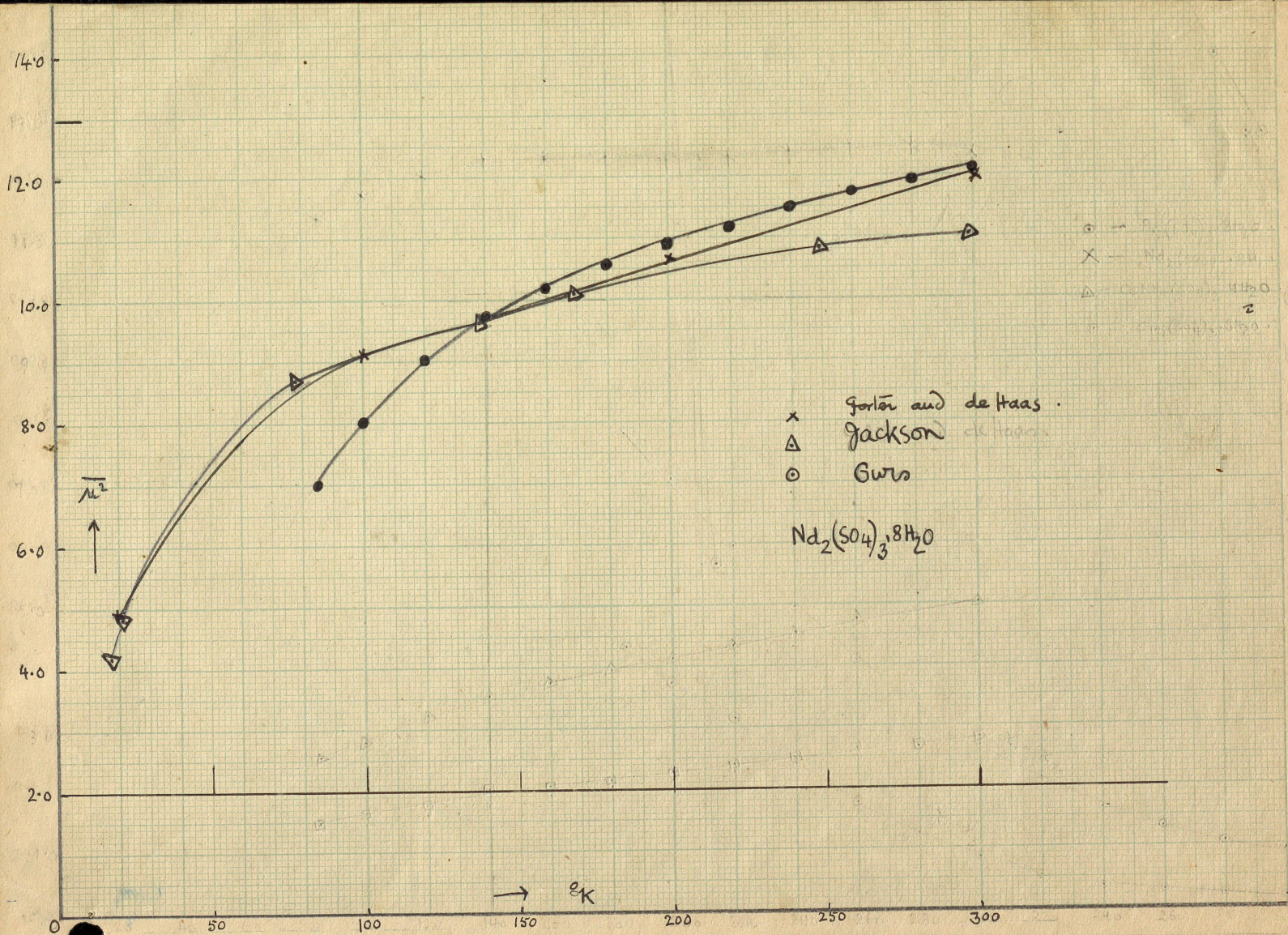
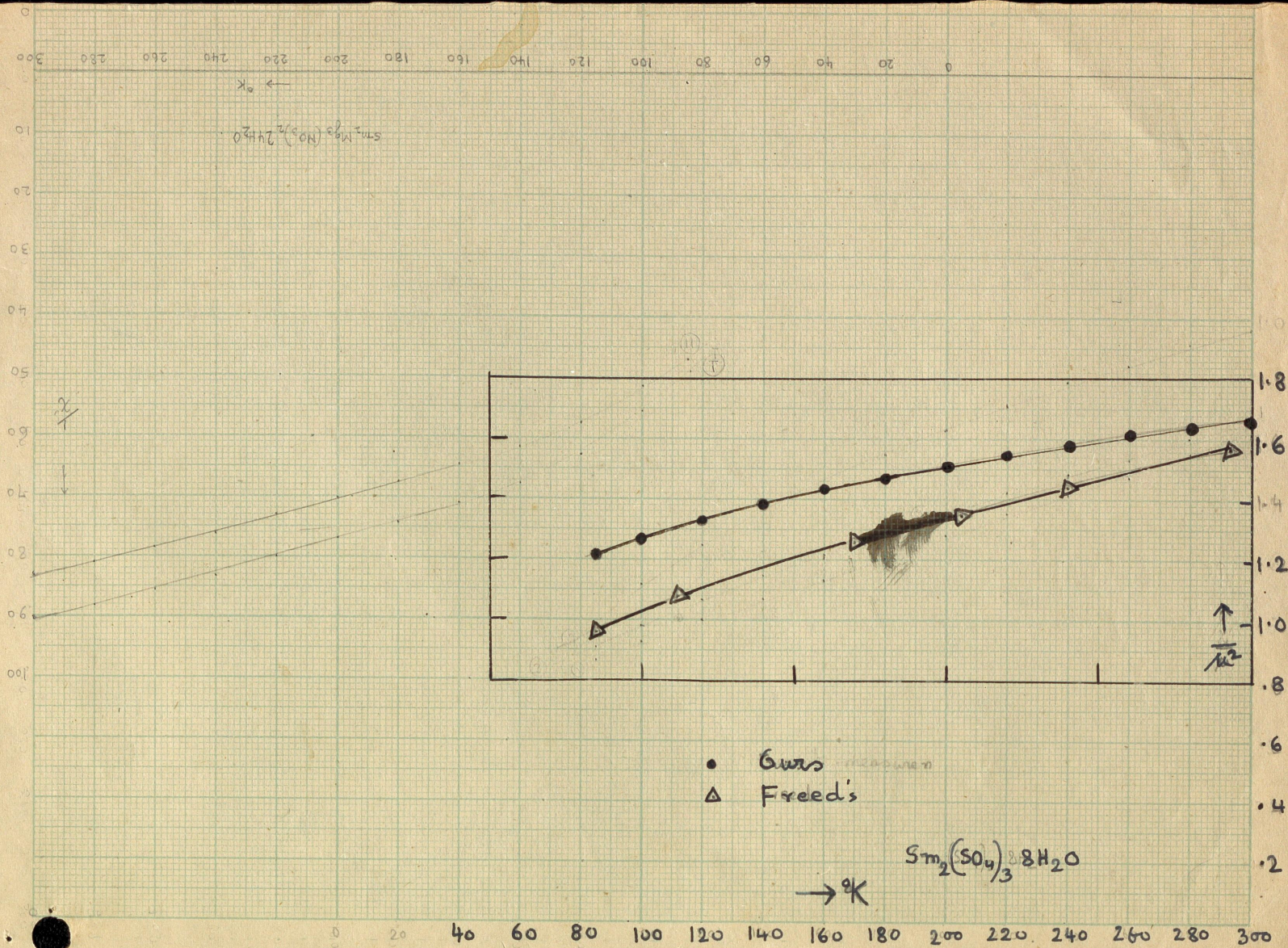


Fig. 18



• Guss
 Δ Fried's

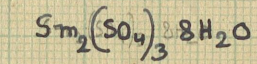


Fig (19)

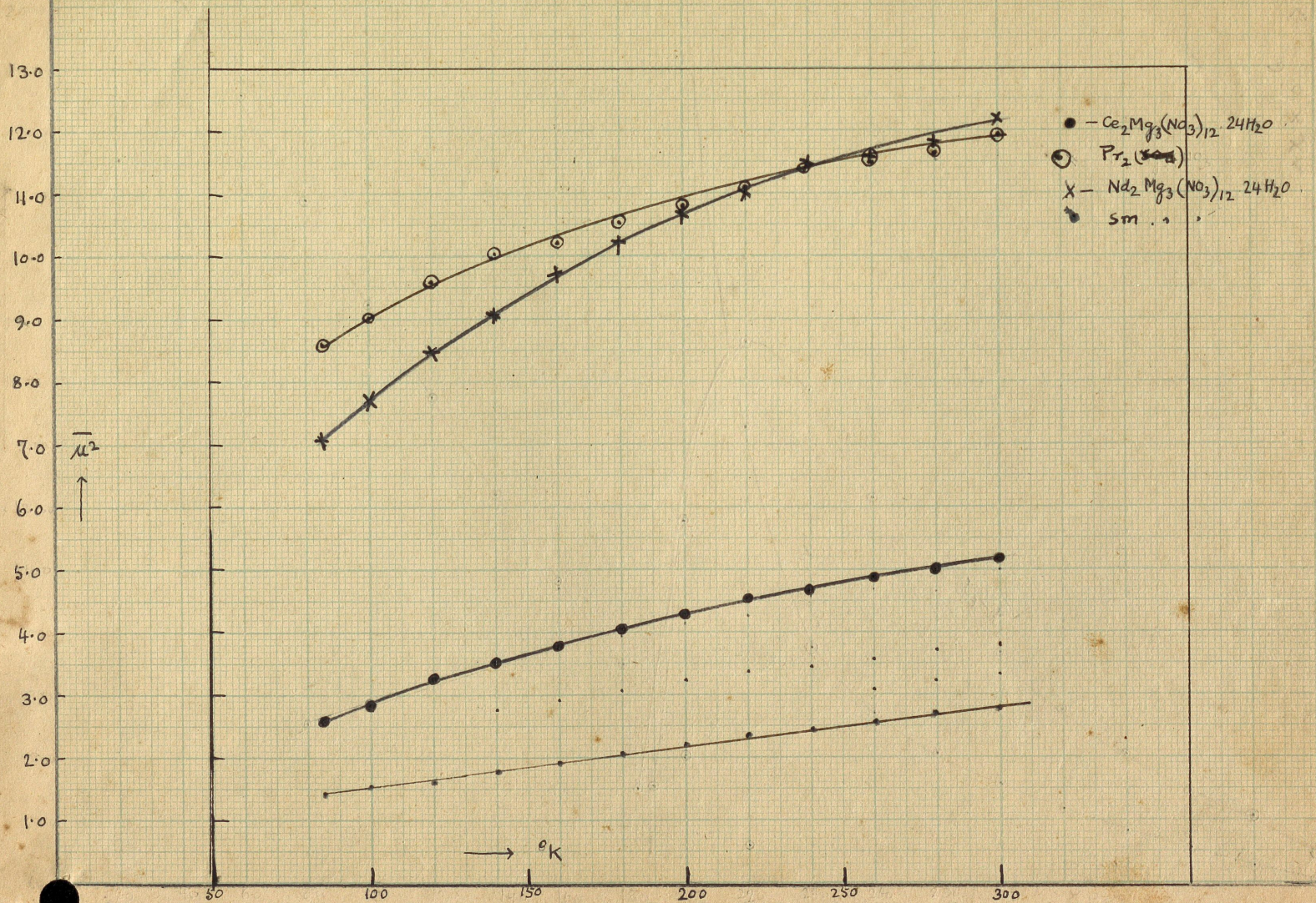


Fig 20

The observed paramagnetism of many of the salts of the Fe and the Rare Ear. groups shows that the elementary magnets in these ~~solids~~ crystals, namely the metallic ions, are capable of turning round + taking up preferred orientations in the magc. field. If the elementary magc. were completely free, i.e. free from each other's influence + influence of their environments they ^{per} ~~follow~~ would follow the Curie law. But in a crystal, unlike a gas, this will not be the case: Result, ^{consequence} deviation from Curie law.

Early days - powdered crystals - deviation attributed to \neq internal field of the Weiss type magc. field $\propto T$. ^{which is connected to} Exch. int. betw magnetic particles

$$\chi = \frac{C}{T - \theta} \quad \text{of the same type as in Fe. above } (\oplus)$$

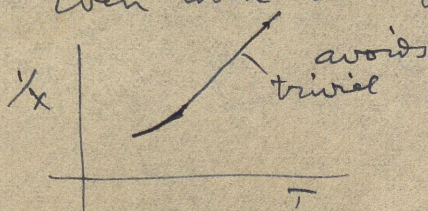
We now know that the observed deviations in crystals can not be attributed to such internal fields.

High dilution \Rightarrow 1) θ ^{appreciable} large even for magnetically dilute ones, quite indep of dilution, 1 atom in 30.

2) θ both + + -ve

3) ~~Even when $\theta = 0$ μ not constant~~

3) Even when θ large + +ve



$$\chi = \frac{C_1}{T} + \frac{C_2}{T^2} + \dots - \frac{C_1}{T - \theta}$$

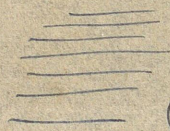
2

Large θ associated with large ΔX & large L

We now know that the local fields which ~~also~~ influence the mag. ions are wholly electronic due to the ~~A~~ - val charge atoms surrounding the ion. The theory of paramag. behav. under these fields has been worked out by Kramer, Bethe and in great detail by Van Vleck. The theory predicts ~~some of them quite uncannily~~ I may hasten to add the extensive exp. data on single crystal support

the theory:

Main features of the theory



$(2J+1)$ m states

Curie law

Actually part of the degeneracy removed.

- 1) Deviation from Curie law: Complicated
- 2) Local fields asymmetric anisotropy:
- 3) The important part wh. L plays.

Major divisions.

- 1) Feeble: Rare earths $(2J+1)$ m. Each with $(2J+1)$ degeneracy. Each J a good quantum no.
 spin.
 Each with $(2J+1)$ degeneracy
- 2) Moderate strong Russell-Saunders. L, S comp. = broken $\Sigma L \rightarrow L$ $\Sigma S \rightarrow S$ remains. From Salt
 $C \neq 1$
 $T = \theta_0$
- 3) L, S also broken. \neq complex salts. D.
 Feni

1. location of the axes
2. ΔX 's describe method
3. χ

Cryostat.

$$\chi_i = \frac{c}{T} \left(1 + \frac{\Delta_i}{T} \right) \quad \frac{\Delta X}{\chi} \sim \frac{\Delta_i}{T} \quad \sim 1^\circ \text{ or } \text{cm}^{-1}$$

$$\chi = \frac{c}{T} \quad \text{Mn salts - Fe}^{+++}$$

Gd $\frac{\Delta X}{\chi}$ was 10% extremely sensitive to impurities

Trombe pre from Sm + Tb

Europrum < 0.1%

only impurities diamagnetic yttrium salt 1.4%

$$\chi_1 - \chi_2 = 36$$

$$\chi_3 - \chi_1 = 16$$

$$\frac{\Delta X}{\chi} \sim 0.1\%$$

$$\Delta_i = 0.3^\circ \text{K}$$

$$0.2 \text{ cm}^{-1}$$

overall splitting.

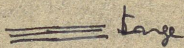
$$\Delta X \propto \frac{1}{T^2}$$

(Mn Gd.)

F state



D $\frac{+ve}{-}$



Octahedral.

Cr⁺⁺⁺

- 1) Large deviation for Curie law
- 2) large " spin value
- 3) anisotropy

$\text{NiSO}_4 \cdot 6\text{H}_2\text{O}$ - 330 cm^{-1}

Specti. - 335

Fe - 340

Ni Am. - { 315
348

370

360

K

Cryst. field constants same for all of Ni salts

Fluoborate

Be F_4 345

360.

Absolute value. Cr^{+++} +87 cm^{-1}

Field const.:



not only cubic but also rhombic

Variation: of ψ small.

Liquid state (solns)

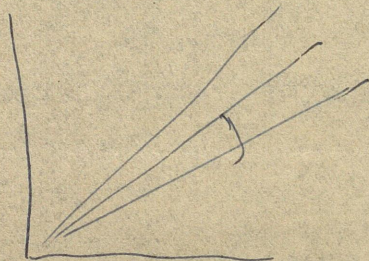
Jahn: Teller

Co_2CoCl_4
from mag. - 220 cm^{-1}
meas. - 180 cm^{-1}

Co_3CoCl_5

much closer to spin only value Curie law.

Cu
D state
Fe



2 different Curie constants

Co six coordinated
=

→ Fe.