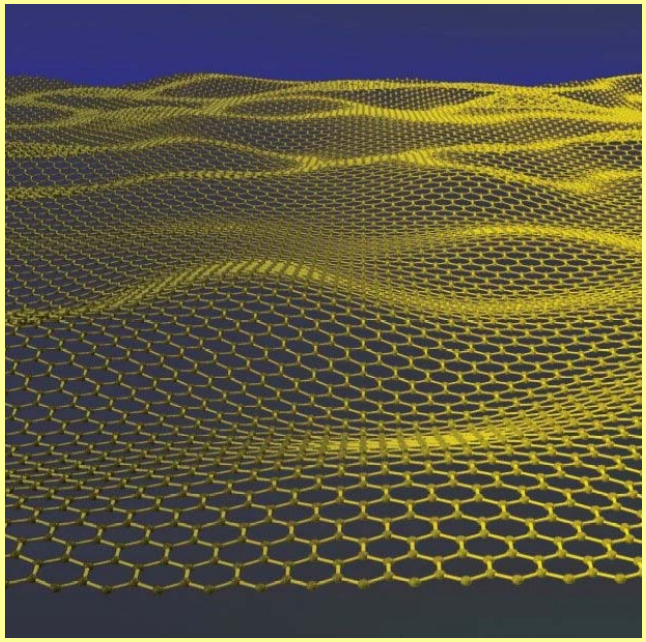
Layered Compounds

2D = Two-dimensional layers



Graphite and Graphene

Clay Minerals, Mica

Layered Double Hydroxides (LDHs)

Layered Zirconium Phosphates and Phosphonates

Layered Metal Oxides

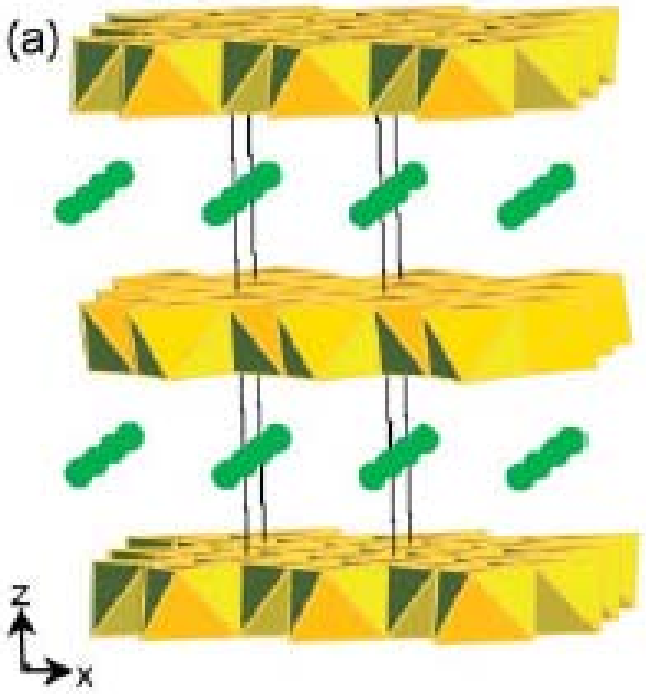
Layered Metal Chalcogenides - MS_2, MPS_3 (M = Ti, V, Mo, W, Mn, Fe, Co, Ni, Zn)

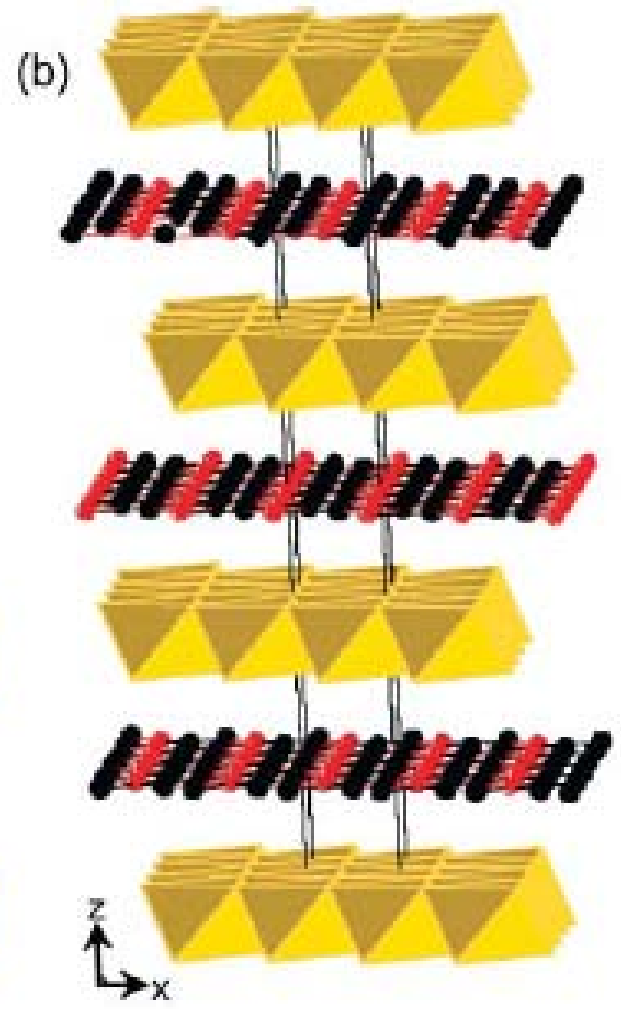
Alkali Silicates and Crystalline Silicic Acids

Layered Compounds

Intralayer bonding - strong (covalent, ionic)

Interlayer bonding - weak (H-bonding, vdWaals)

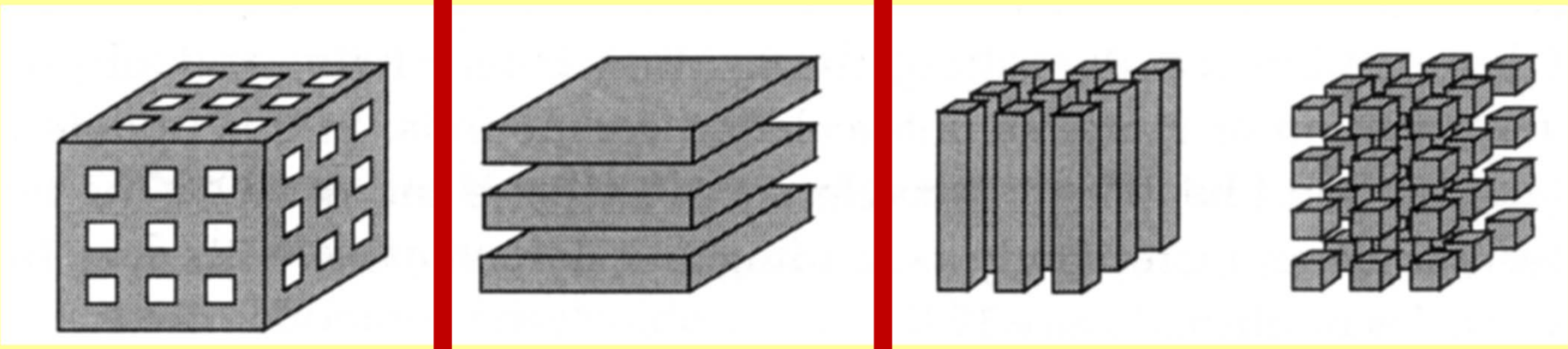




Host-Guest Structures

Host dimensionality

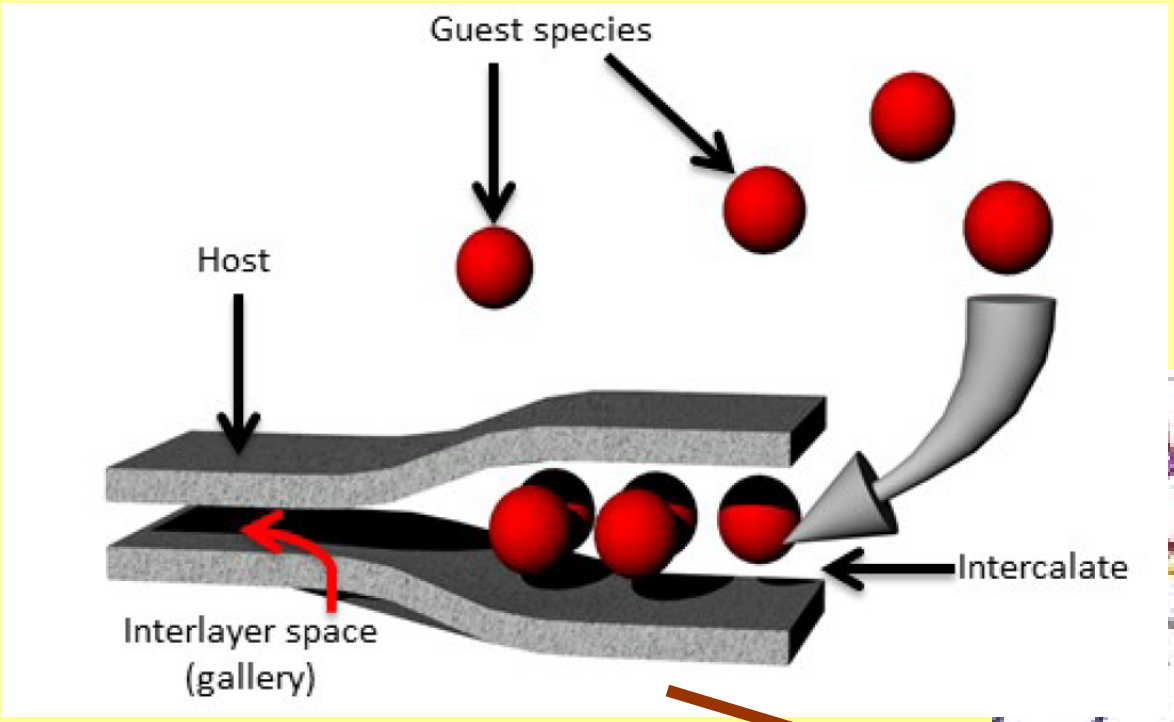
3D | 2D | 1D | 0D

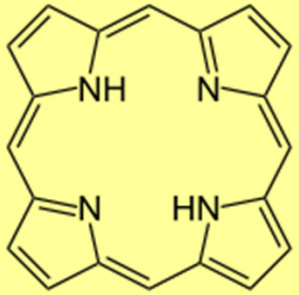


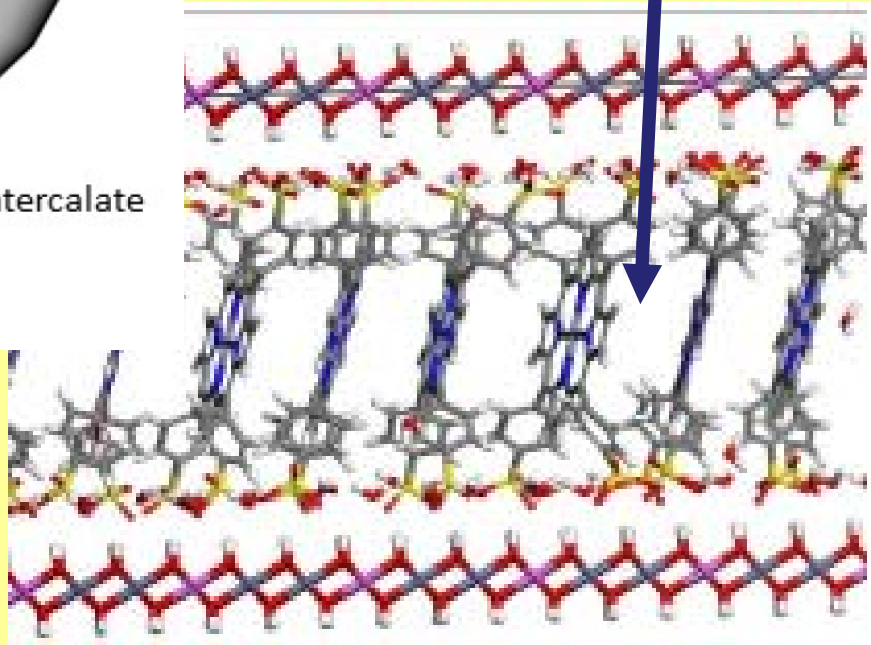
TOPOTACTIC SOLID-STATE REACTIONS = modifying existing solid state structures while maintaining the integrity of the overall structure

Intercalation

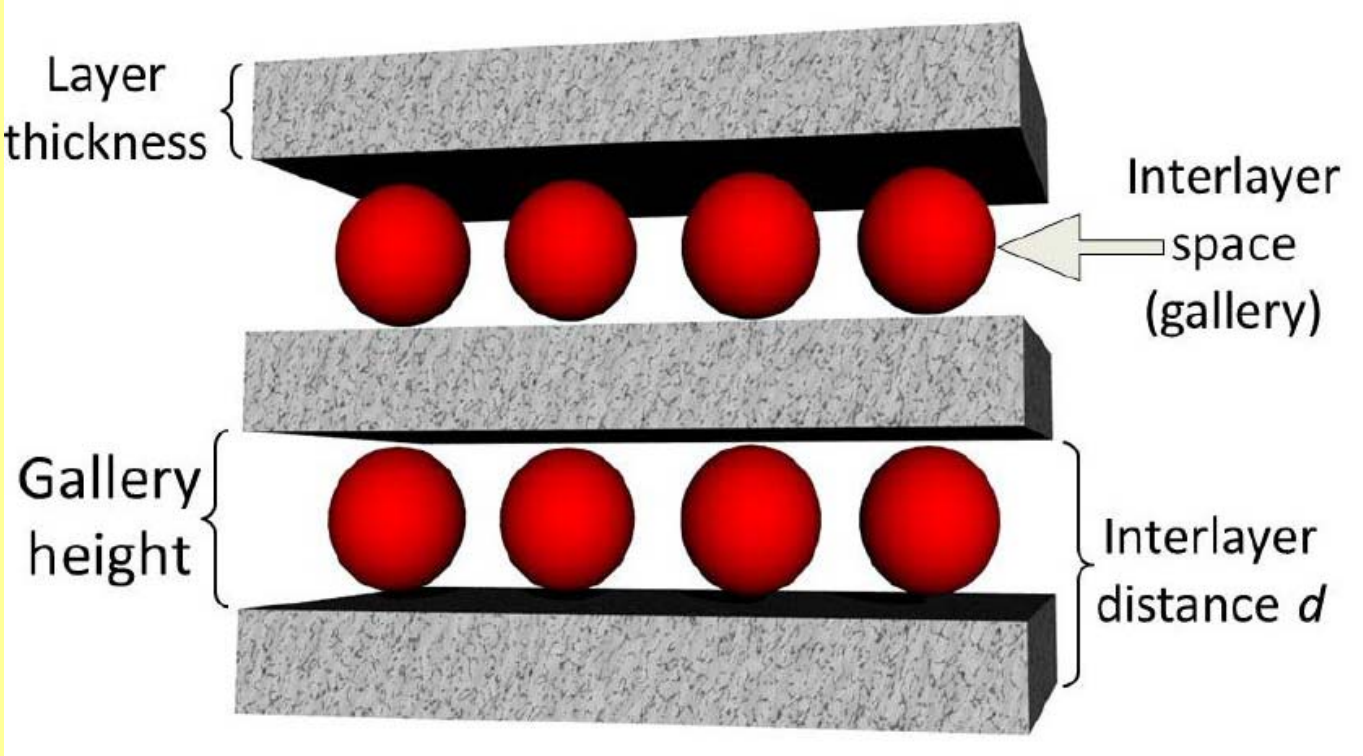
Intercalation **Insertion of molecules between layers**



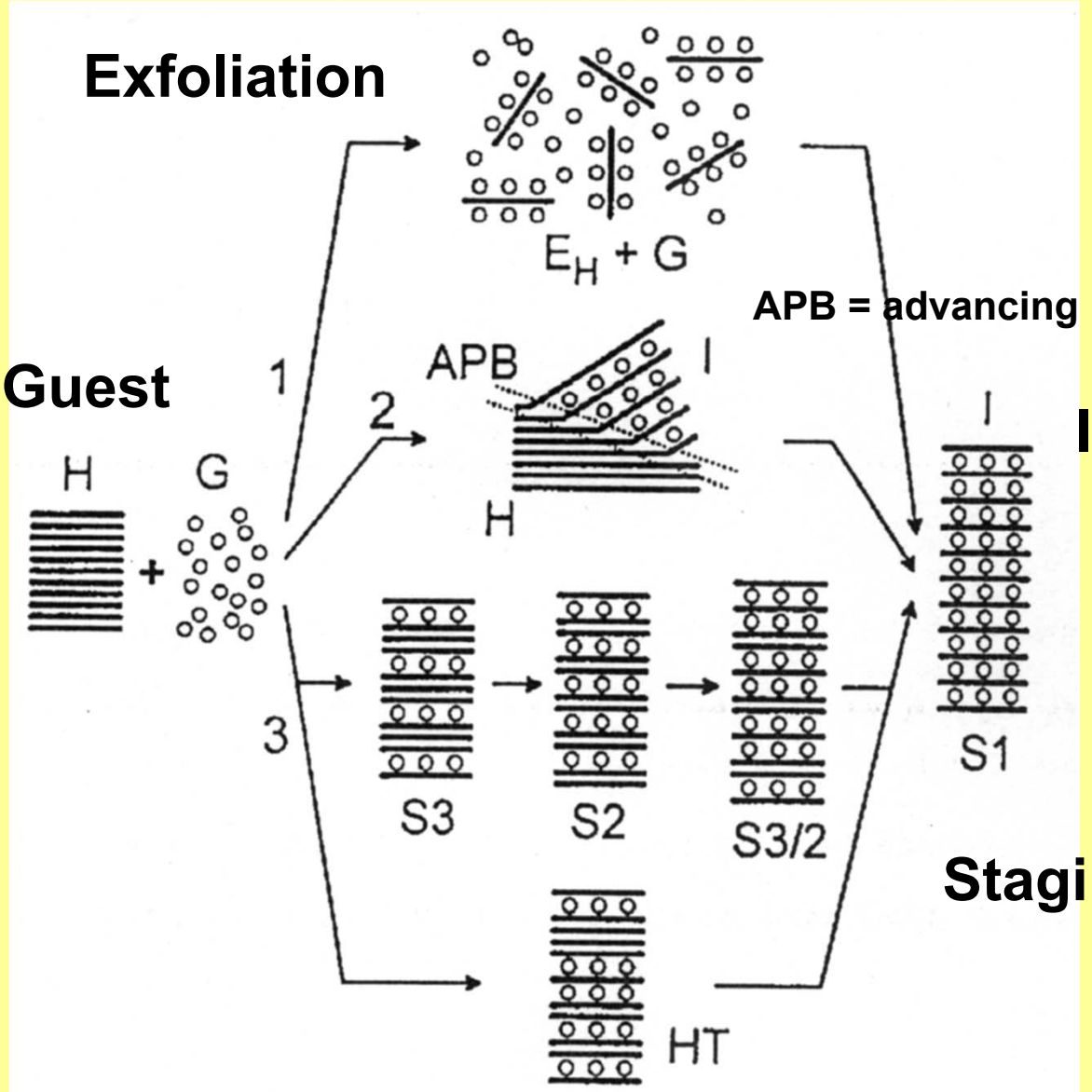




Intercalation

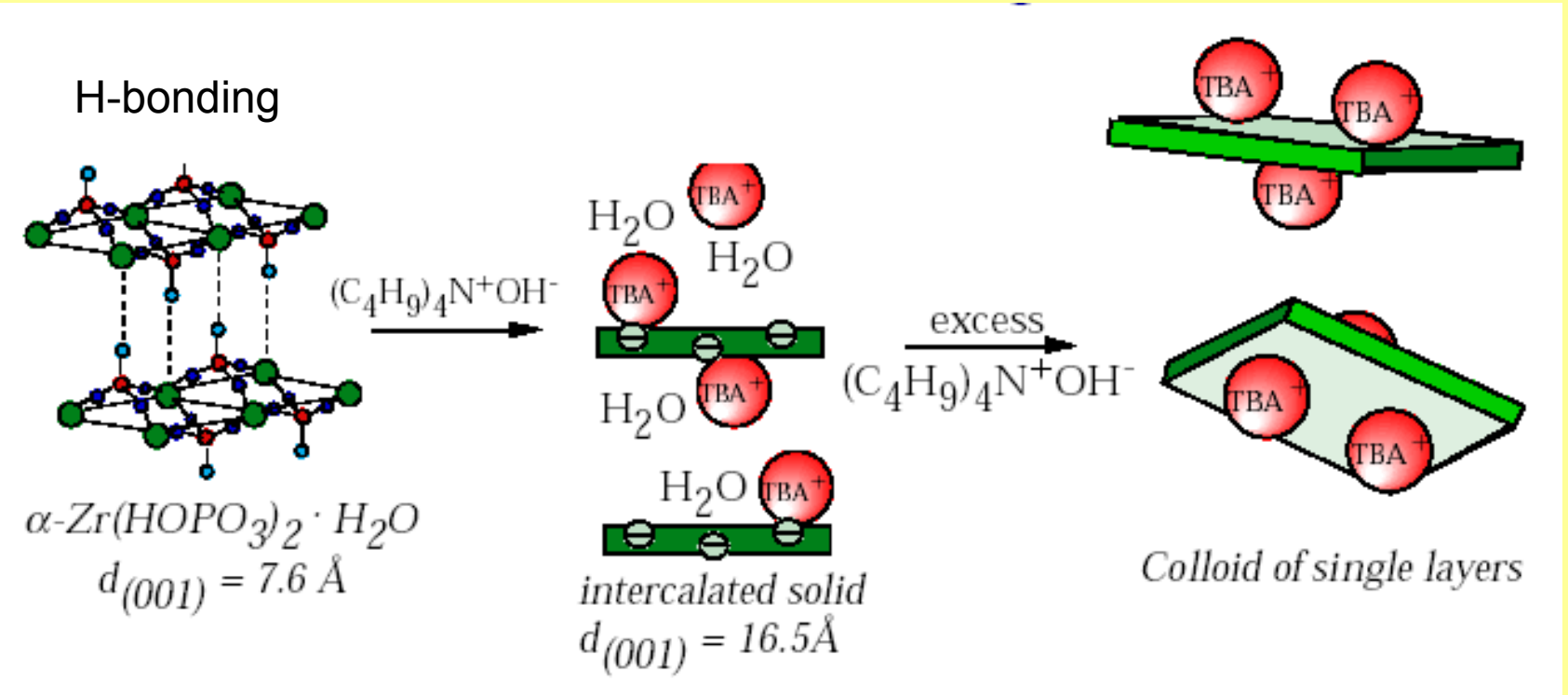


Intercalation



Exfoliation

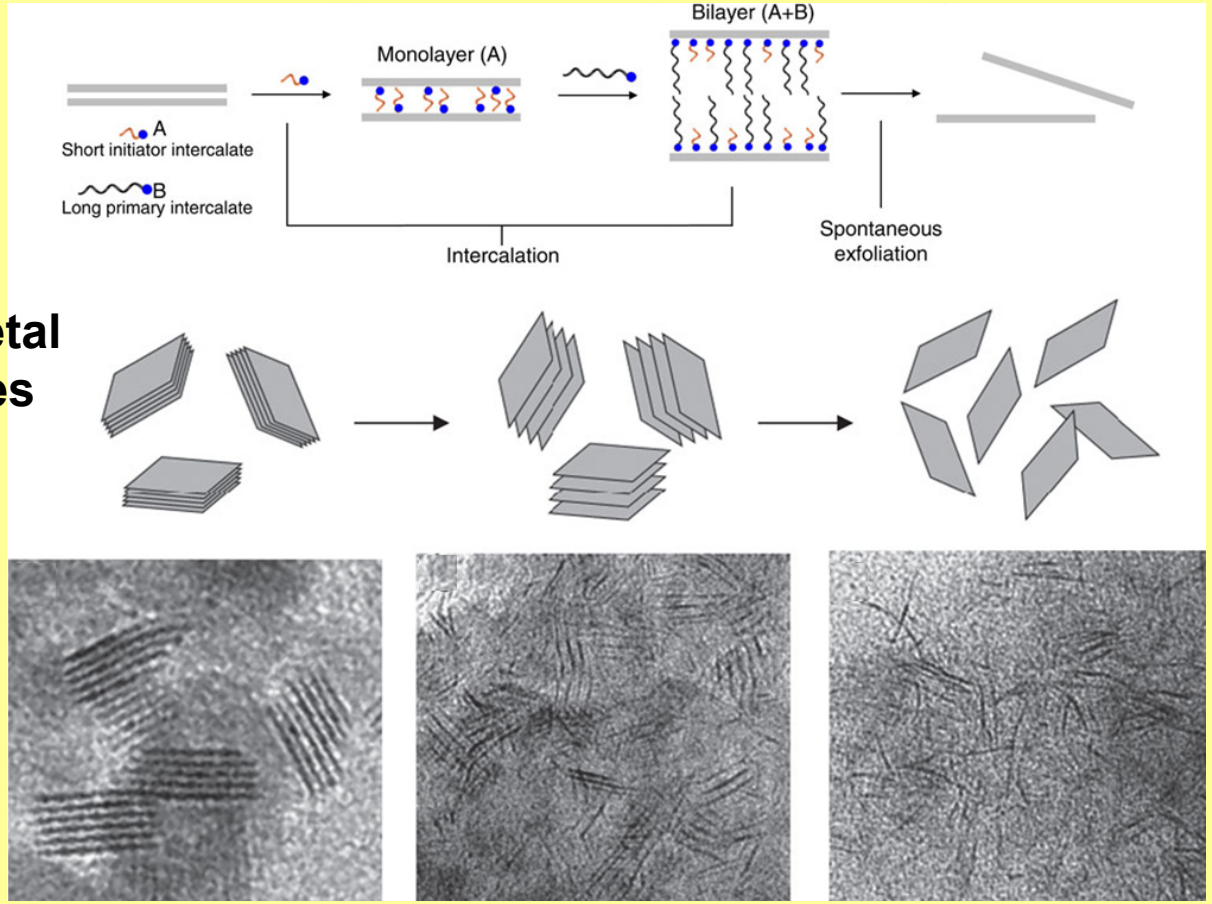
Decrease attractive forces between layers
Separate layers



Tandem Molecular Intercalation

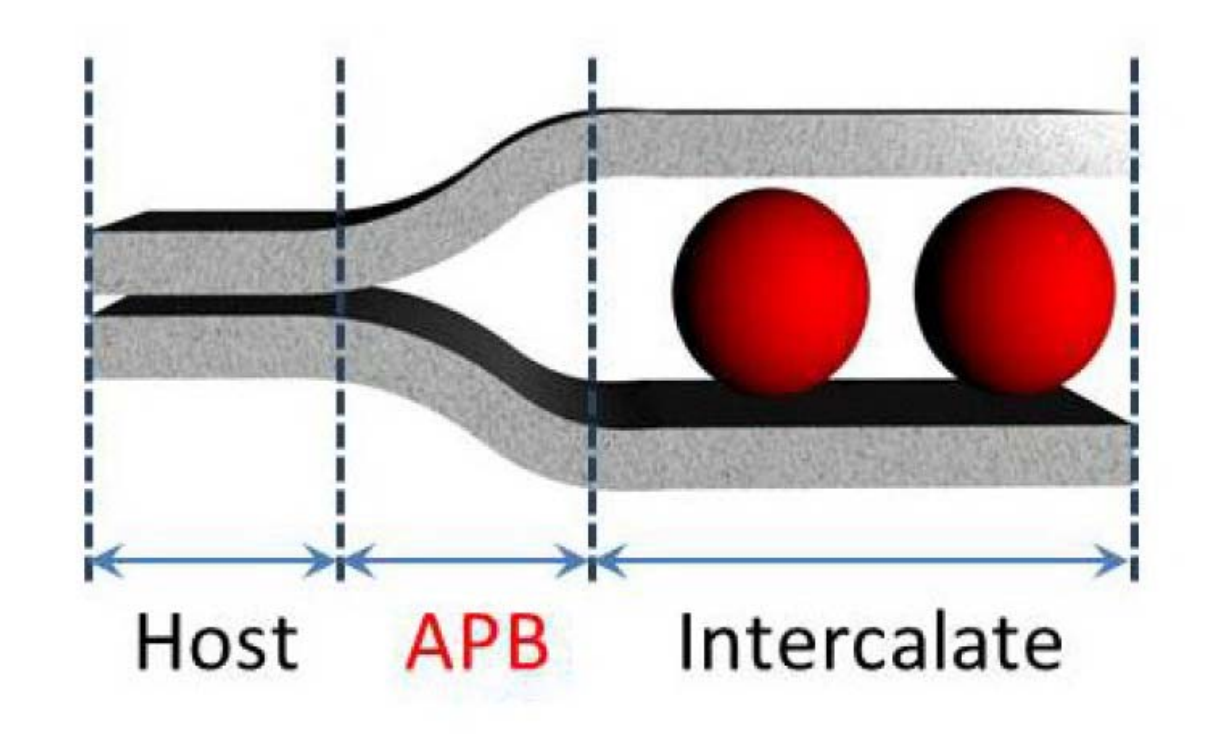
Lewis base short initiator first intercalates to open up the interlayer gap, and the long molecules then bring the gap to full width and overcome the interlayer attractive force resulting in spontaneous exfoliation

Transition-Metal Chalcogenides



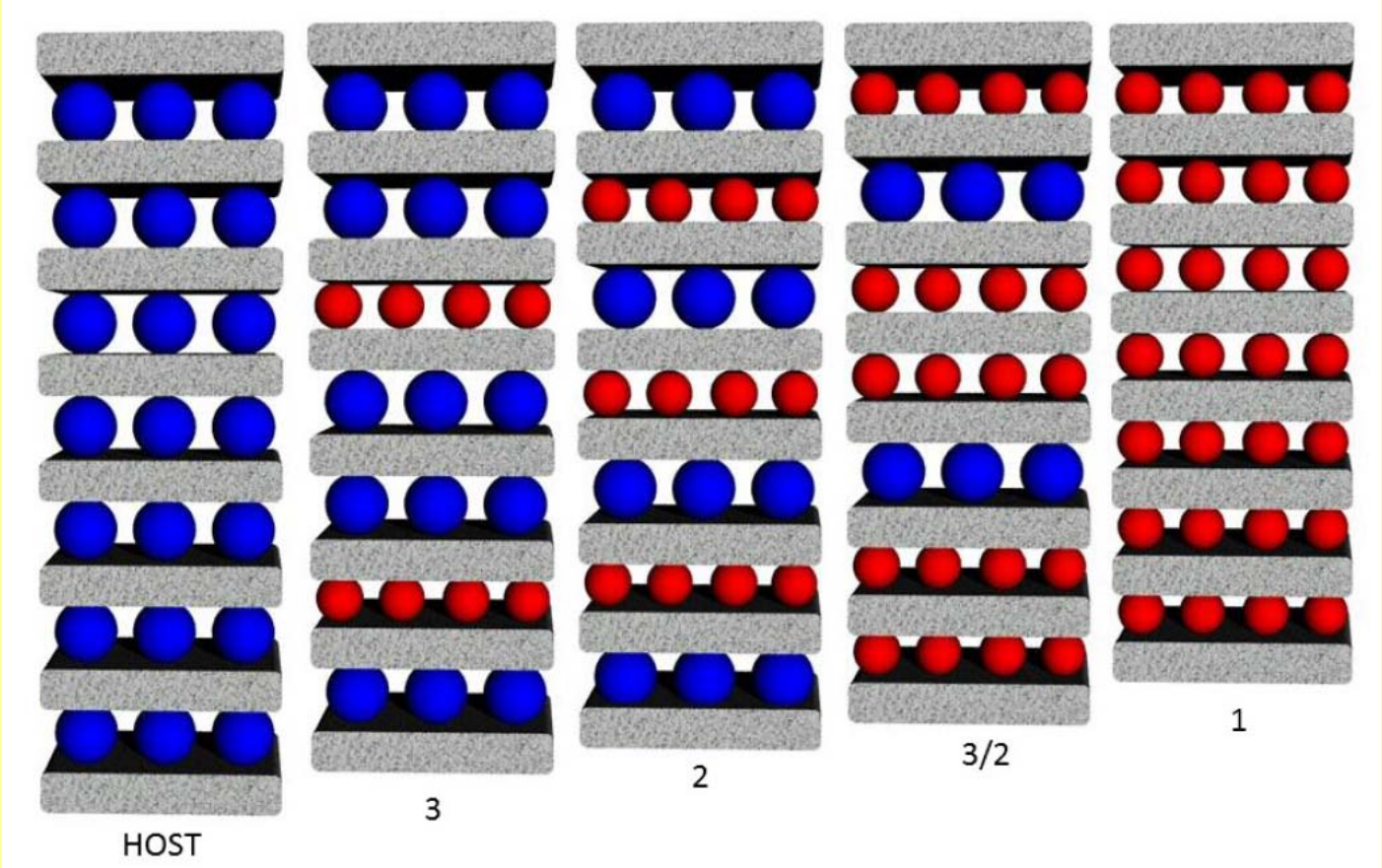
Nat. Commun. 2015,6, 5763, Jan. 9, 2015

APB = Advancing Phase Boundary

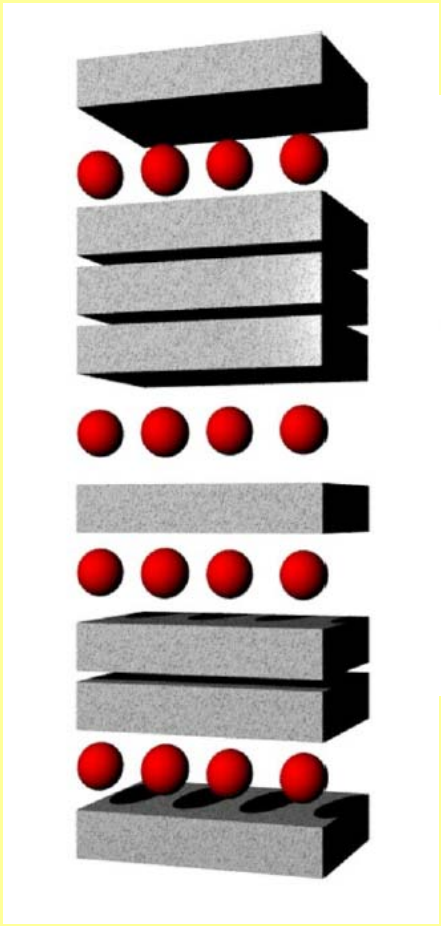


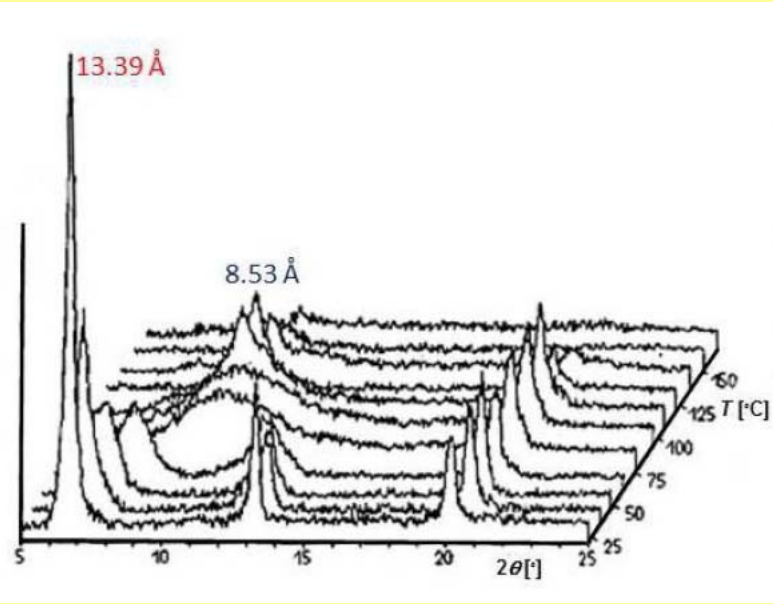
APB = advancing phase boundary

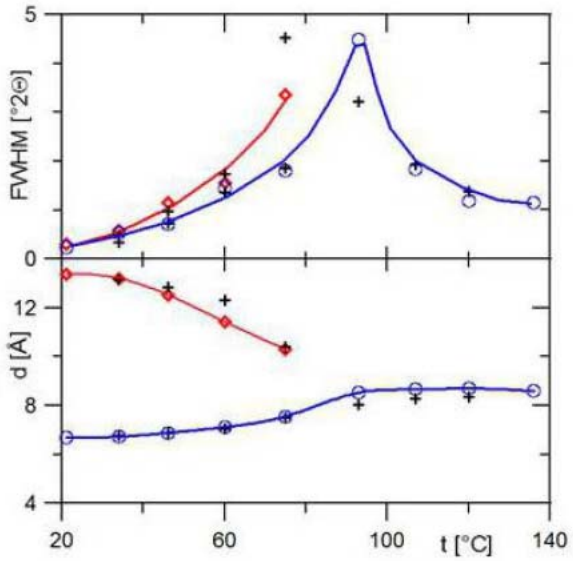
Staging



Hendricks-Teller Effect

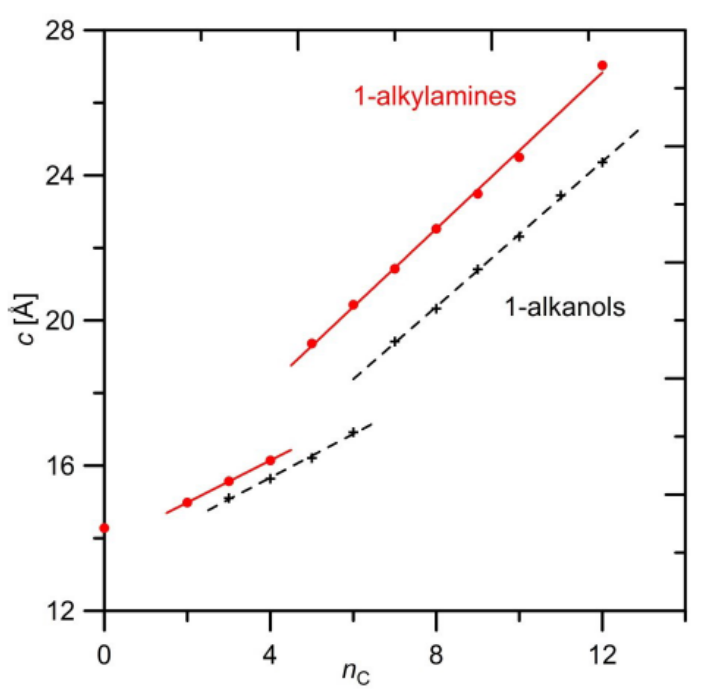


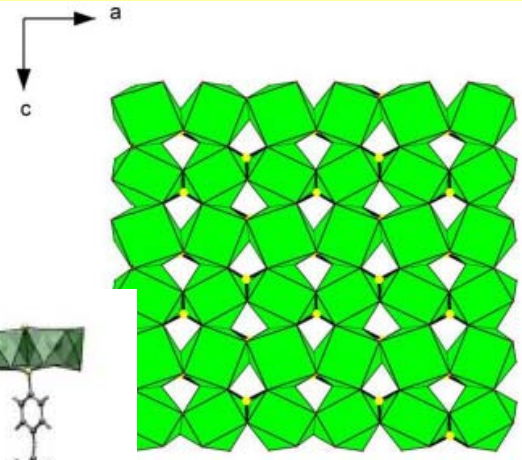


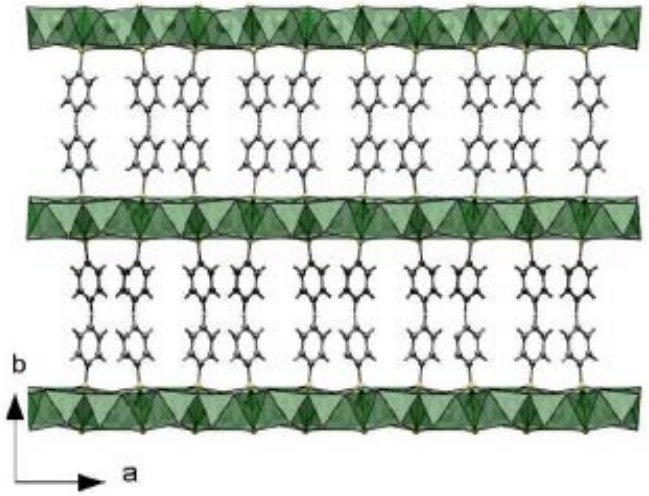


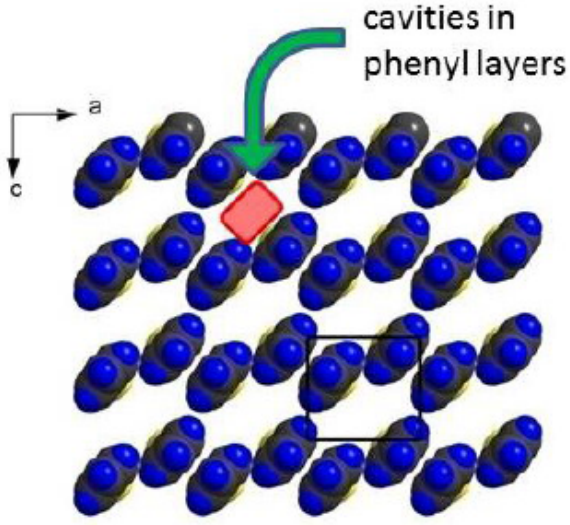
HT = galleries are filled randomly

Intercalation





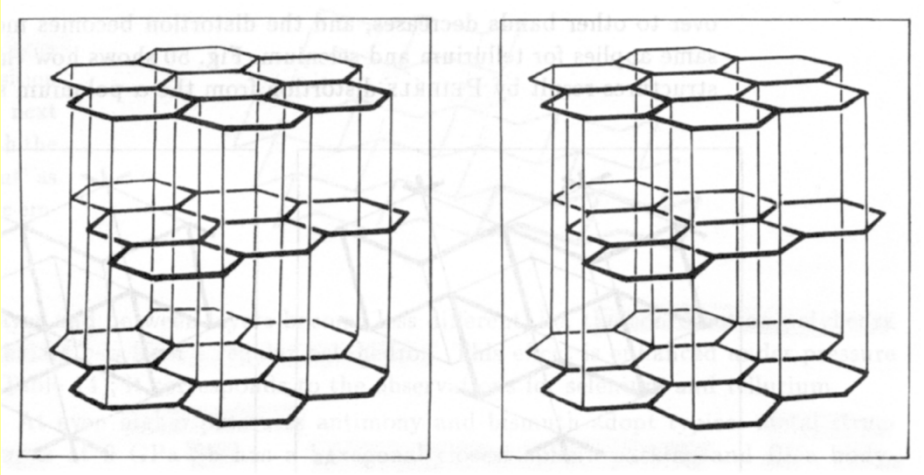


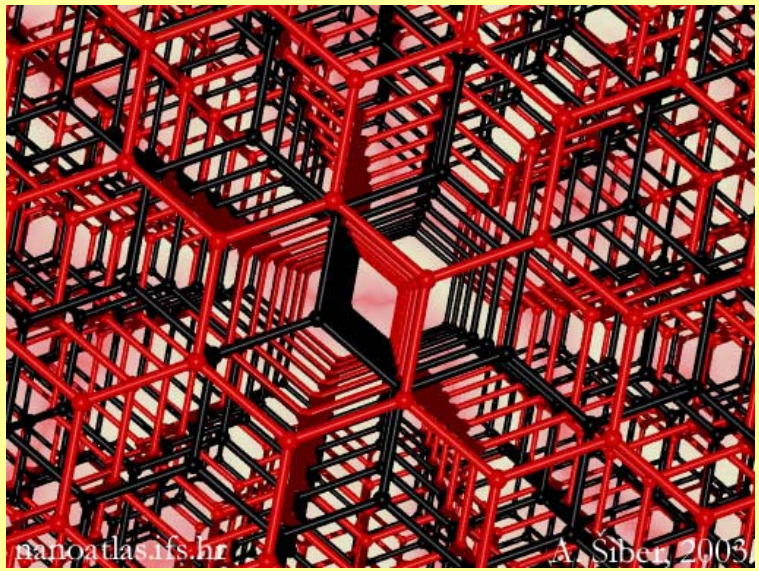


Dependence of the basal spacing of the intercalates of the alkylamines (circles) and alkanols (crosses) on the number of carbon atoms n_C in $SrC_6H_5PO_3 \cdot 2H_2O$

Graphite

ABABAB





Graphite sp^2 sigma-bonding in-plane p-p-bonding out of plane
Hexagonal graphite = two-layer ABAB stacking sequence

SALCAOs of the p-p-type create the valence and conduction bands of graphite, very small band gap, metallic conductivity properties in-plane, 10^4 times that of out-of plane conductivity

Graphite

GRAPHITE INTERCALATION

G (s) + K (melt or vapour) $\rightarrow$ C_8K (bronze)

C_8K (vacuum, heat) $\rightarrow$ $C_{24}K \rightarrow C_{36}K \rightarrow C_{48}K \rightarrow C_{60}K$

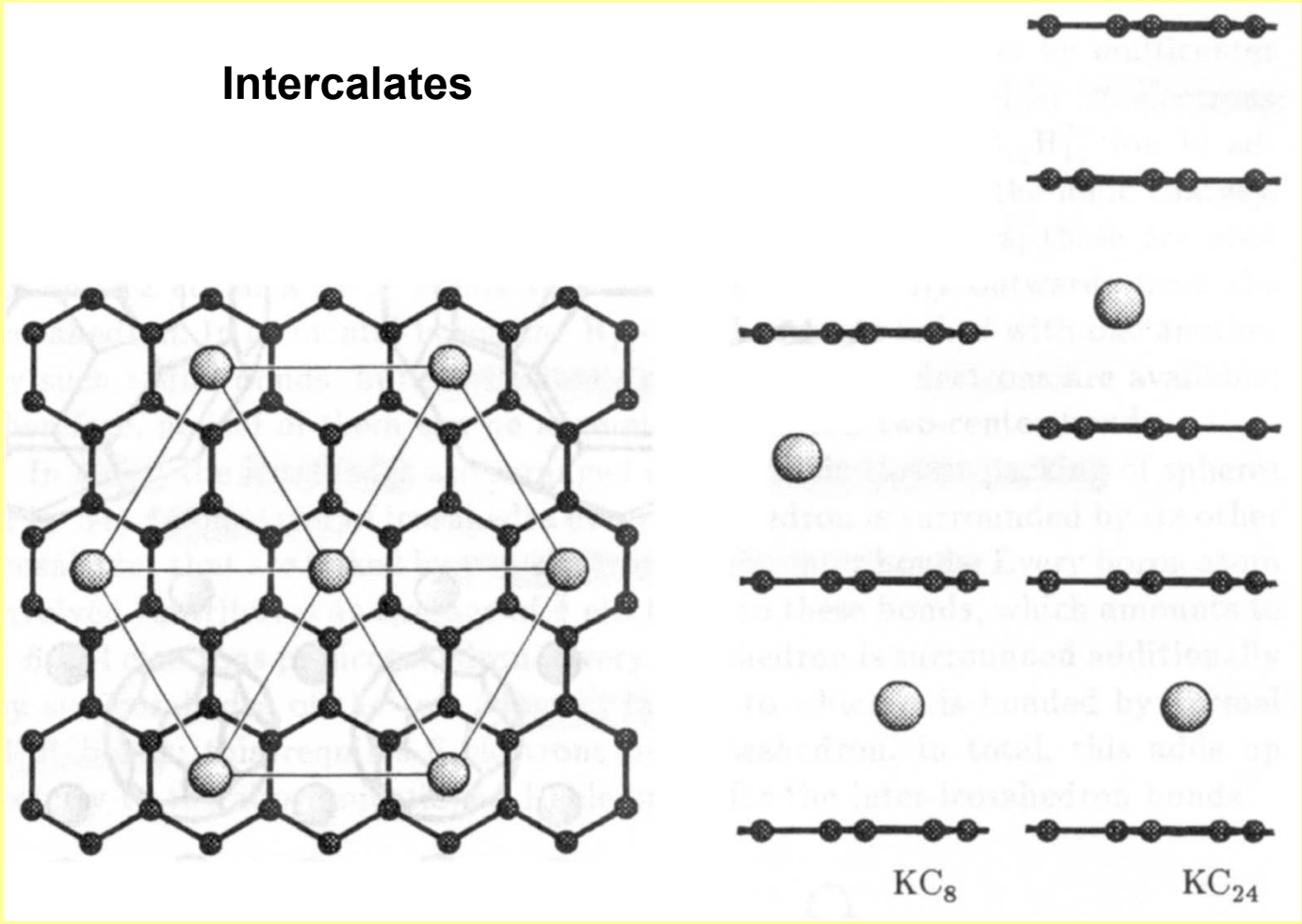
C_8K potassium graphite ordered structure

Ordered K guests between the sheets, K to G charge transfer

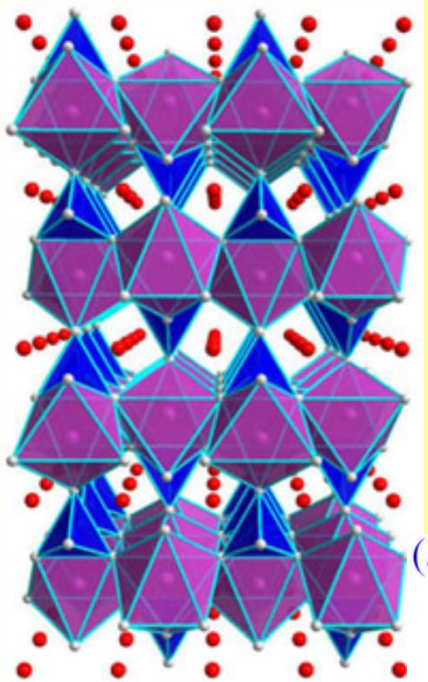
AAAA stacking sequence
reduction of graphite sheets, electrons enter CB
K nesting between parallel eclipsed hexagonal planar carbon six-rings

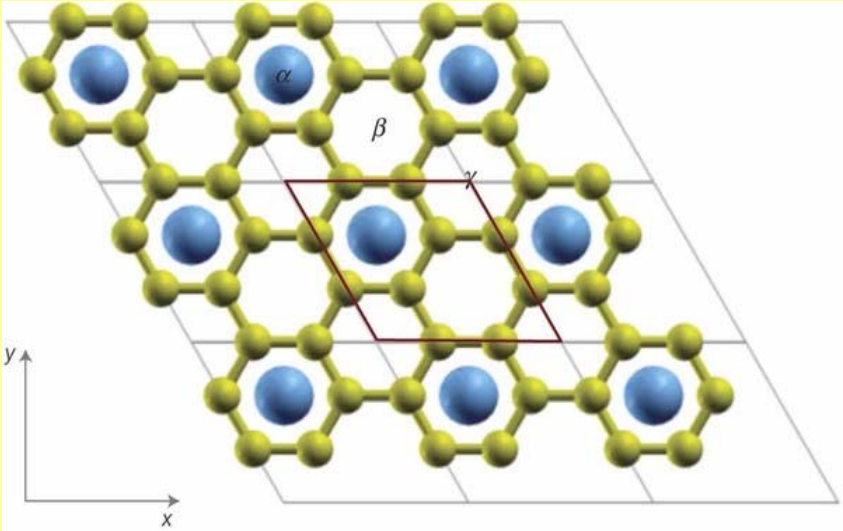
Graphite

Intercalates

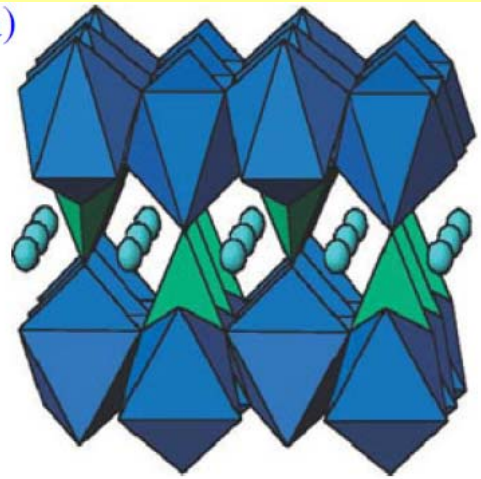


Intercalation in Li-ion Cells



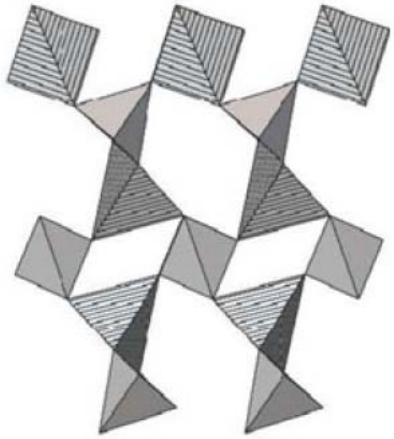


(a)



Charge

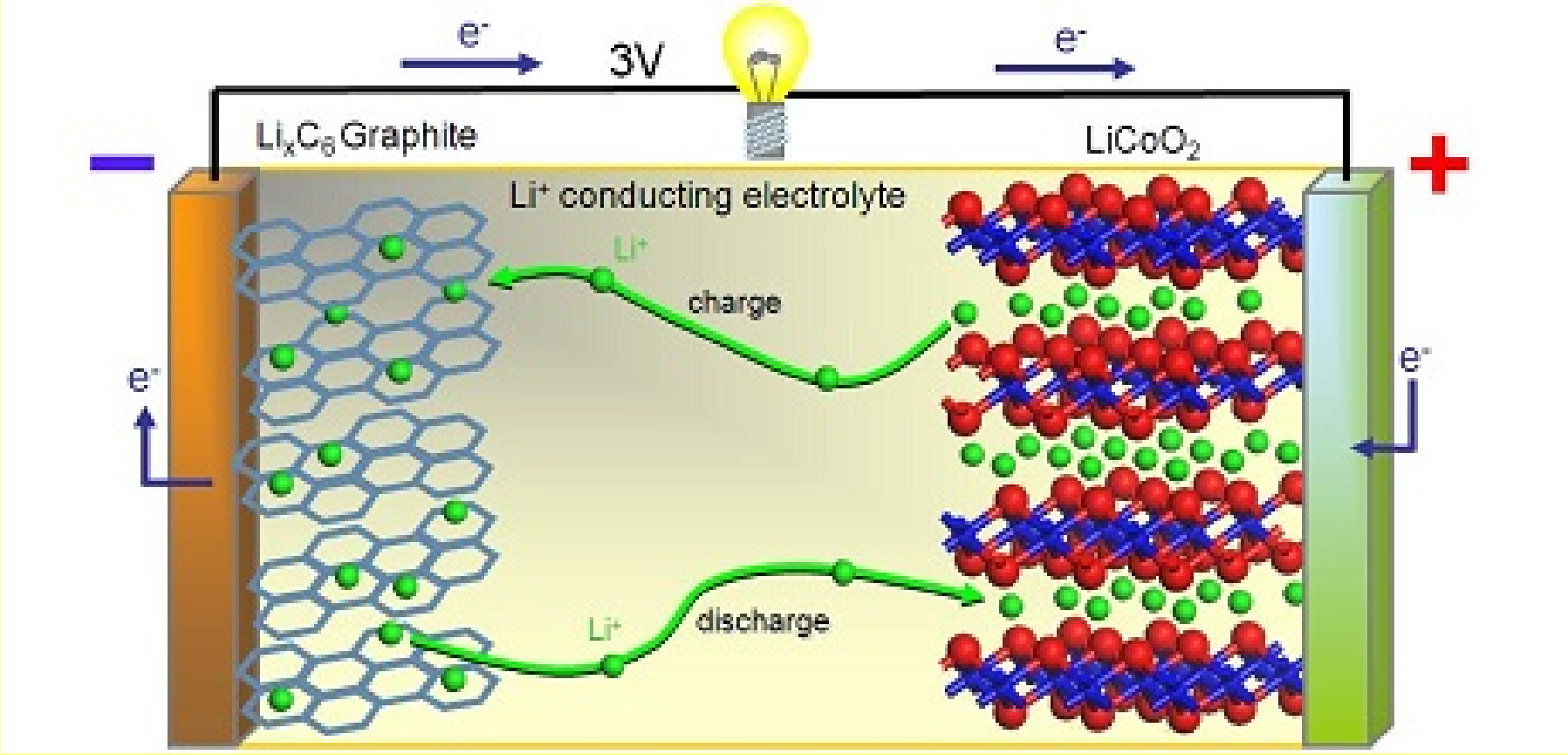
Discharge



$LiFePO_4$

$FePO_4$

Intercalation in Li-ion Cells

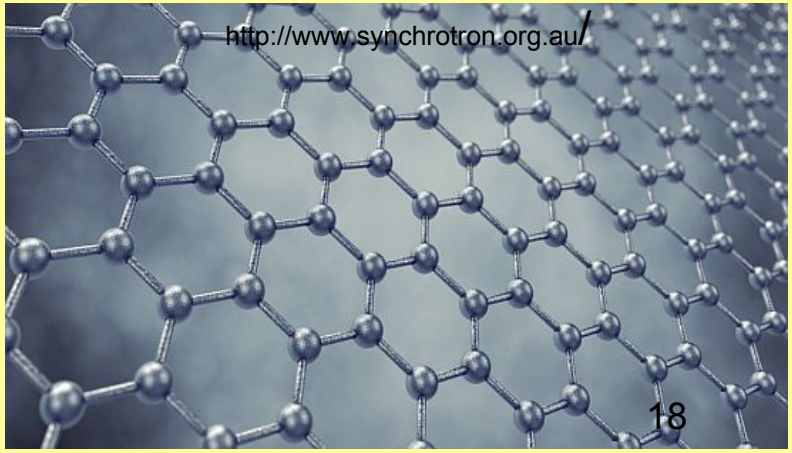


Graphene

- **Discovery – 2004**
- **Exotic properties:**
  - **Firm structure**
  - **Inert material**
  - **Hydrofobic character**
  - **Electric and thermal conductivity**
  - **High mobility of electrons**
  - **Specific surface area (theoretically): 2630 m^2g^{-1}**

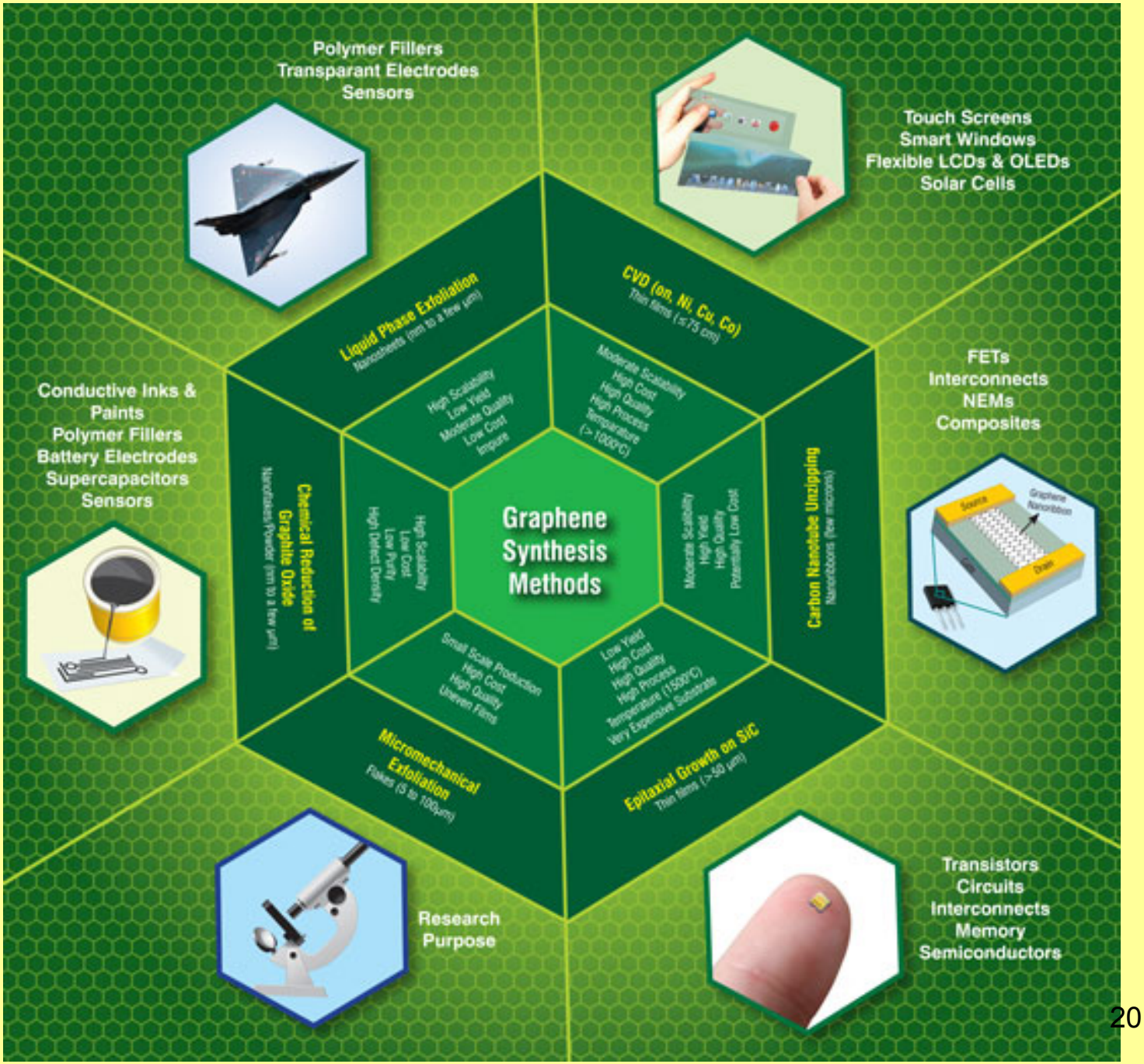
K. Novoselov

A. Geim



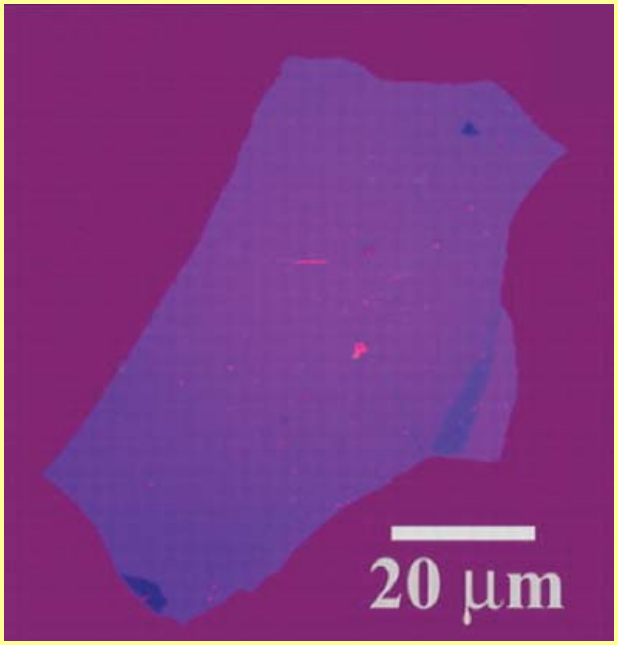
Sythesis of graphene

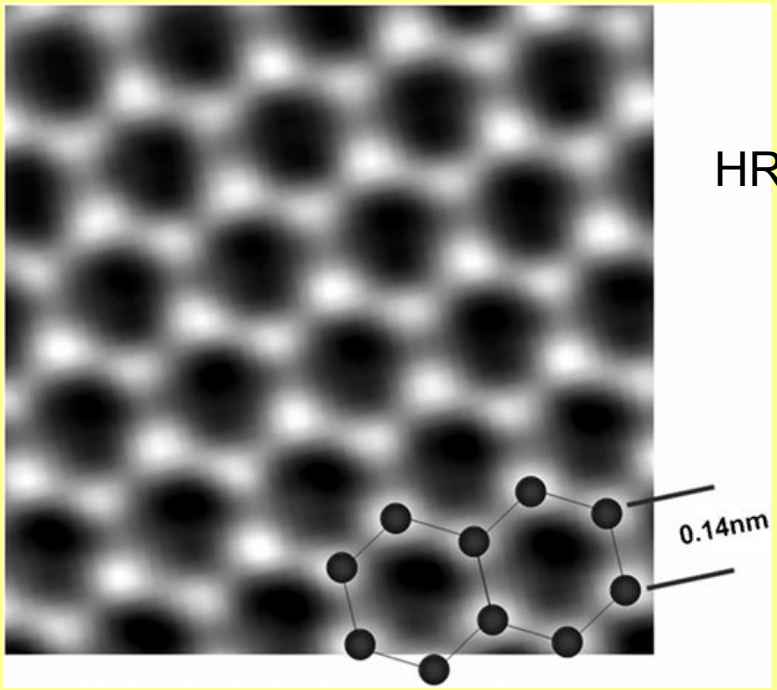
- **Top down**
  - **Mechanical exfoliation**
  - **Chemical exfoliation**
- **Bottom up**
  - **CVD, epitaxial growth, …**
- **Defects**
- **Application: diodes, sensors, solar cell, energy storage, composites, …**



http://www.nanowerk.com/what_is_graphene.php

Graphene





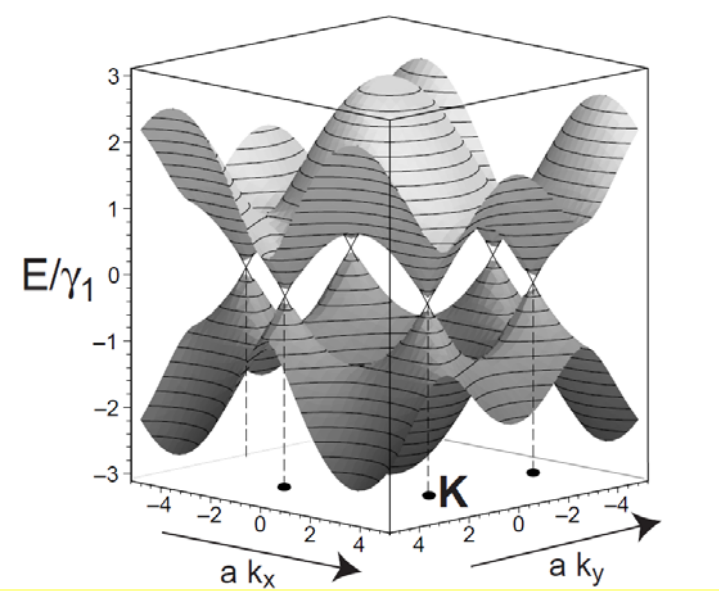
HRTEM

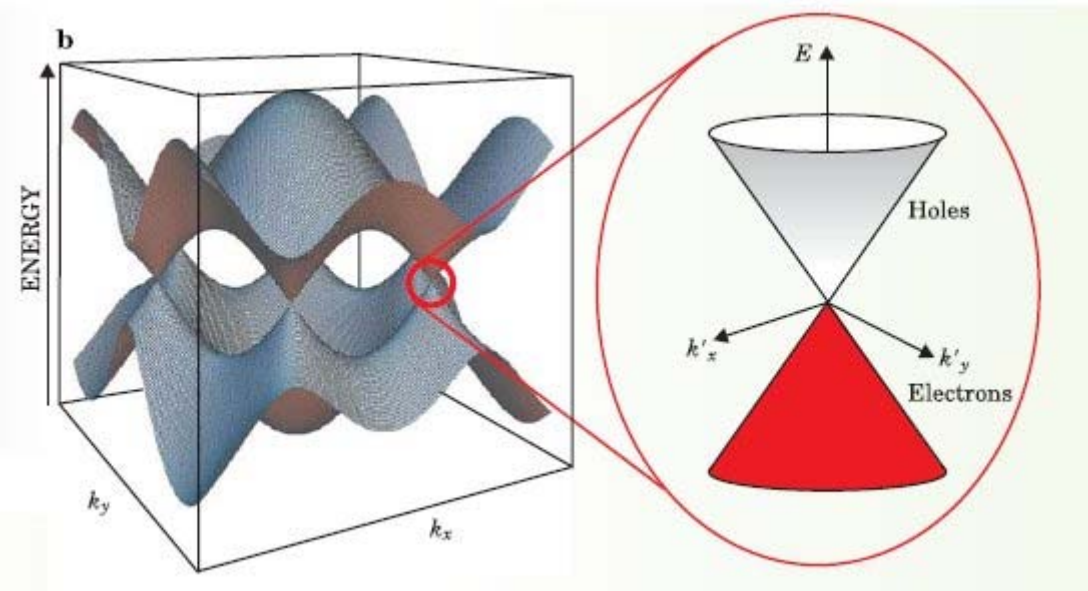
High electric conductivity (metallic)

Optically transparent – 1 layer absorbs 2.3% of photons

High mechanical strength

Graphene



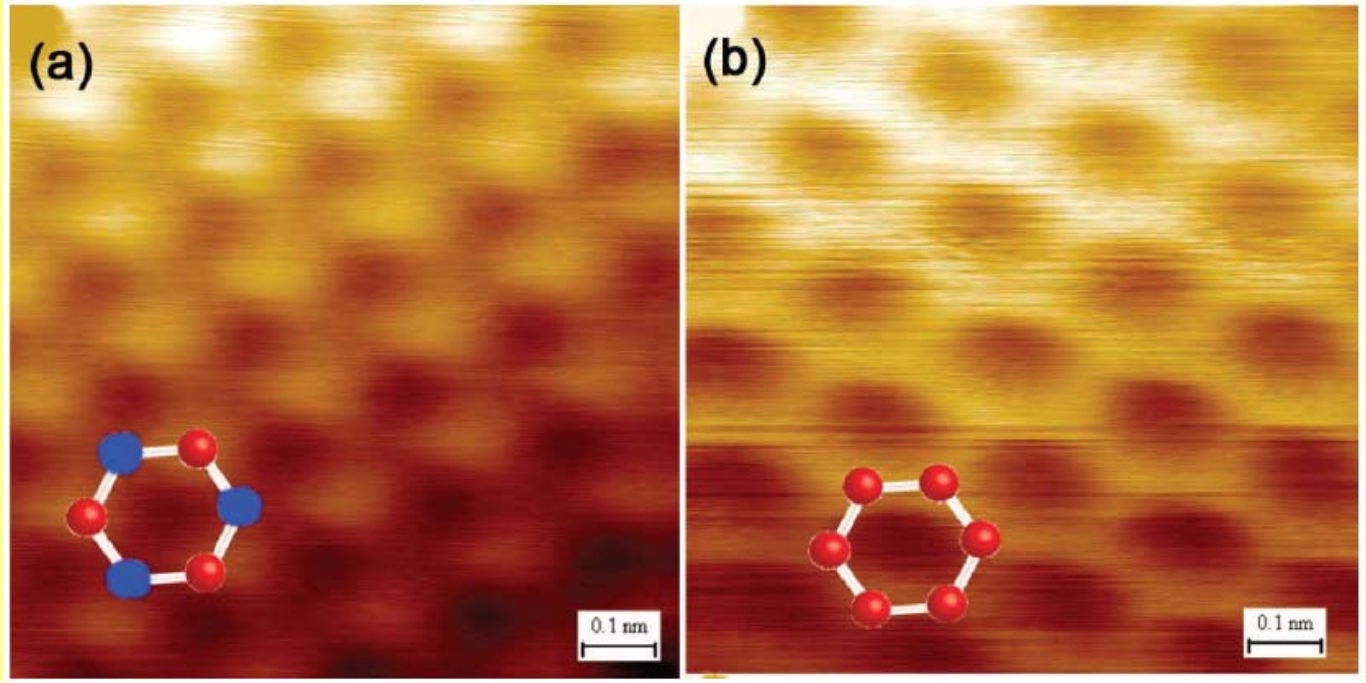


LCAO-band structure of graphene

Graphene

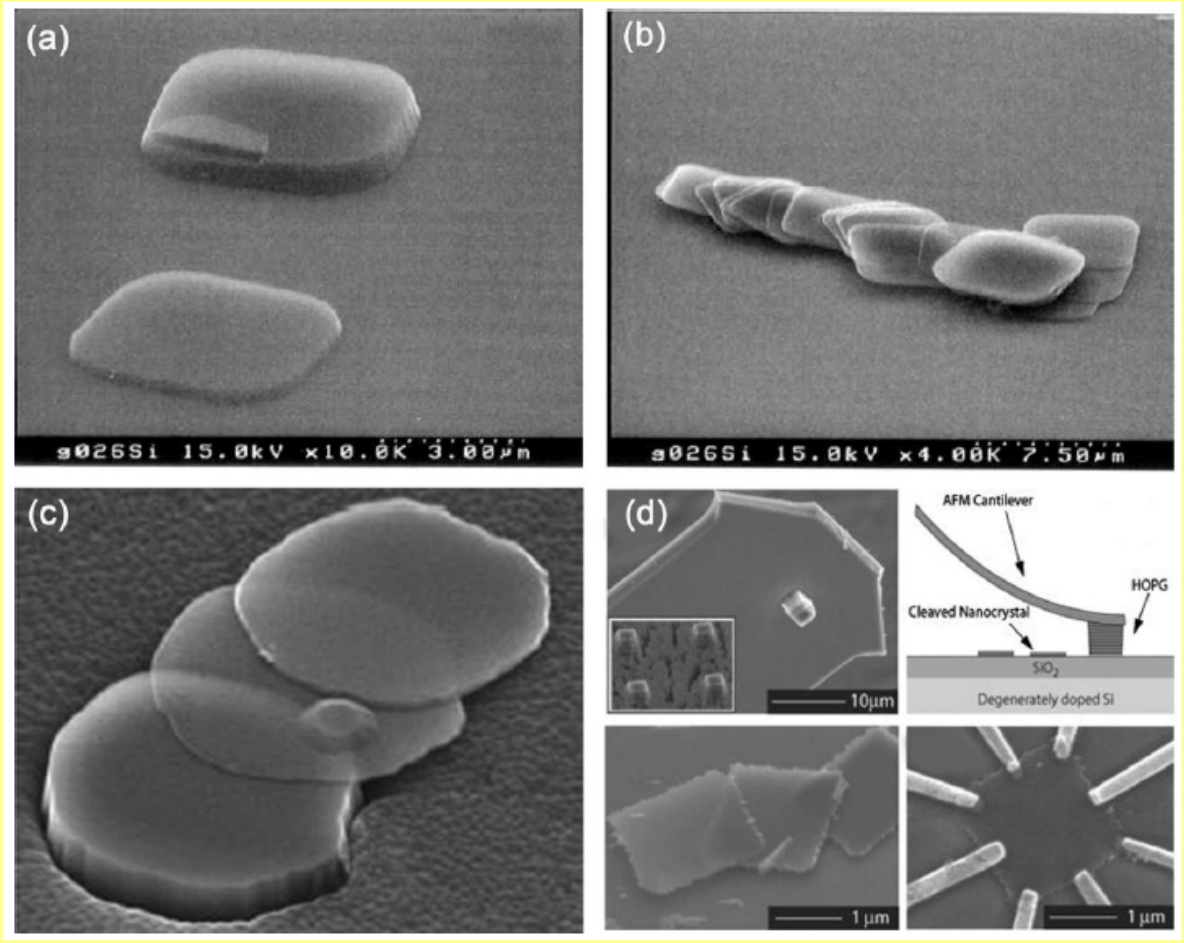
Preparation:

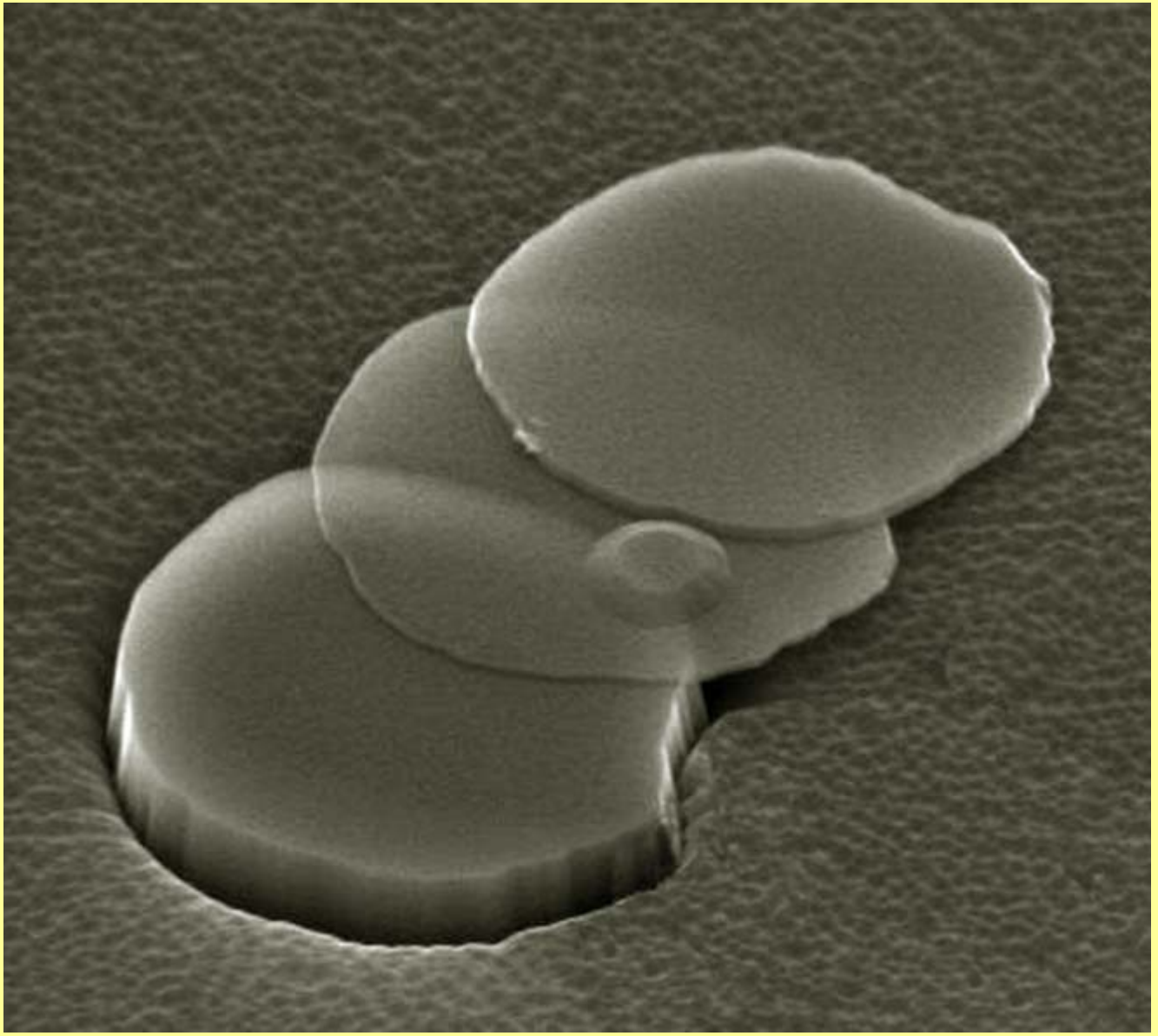
- **Scotch tape – layer peeling, flaking**
- **SiC pyrolysis – epitaxial graphene layer on a SiC crystal**
- **Exfoliation of graphite (chemical, sonochemical)**
- **CVD from CH_4, CH_2CH_2, or CH_3CH_3 on Ni (111), Cu, Pt surfaces**



Scotch tape – Layer peeling

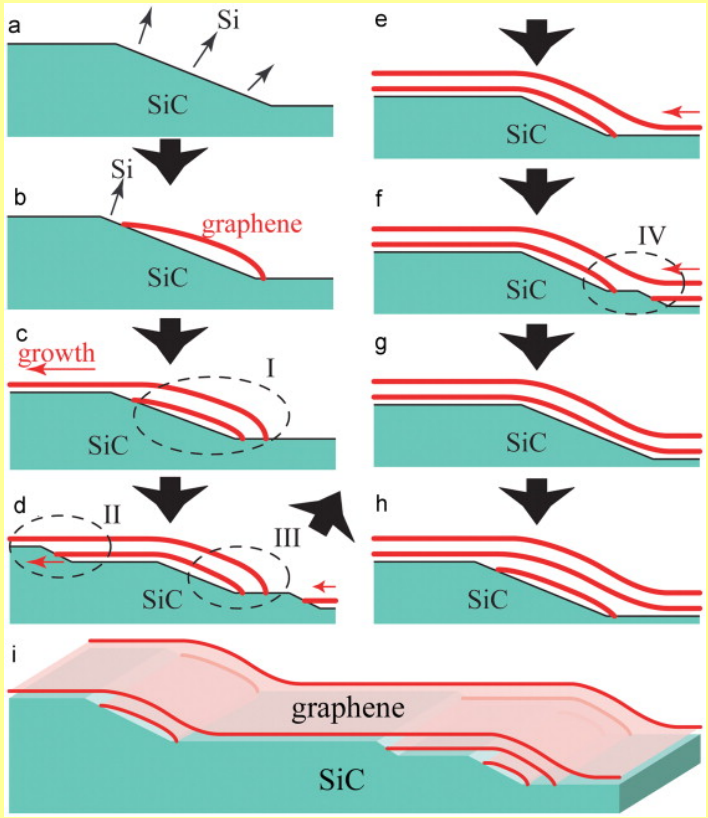
Mechanical exfoliation

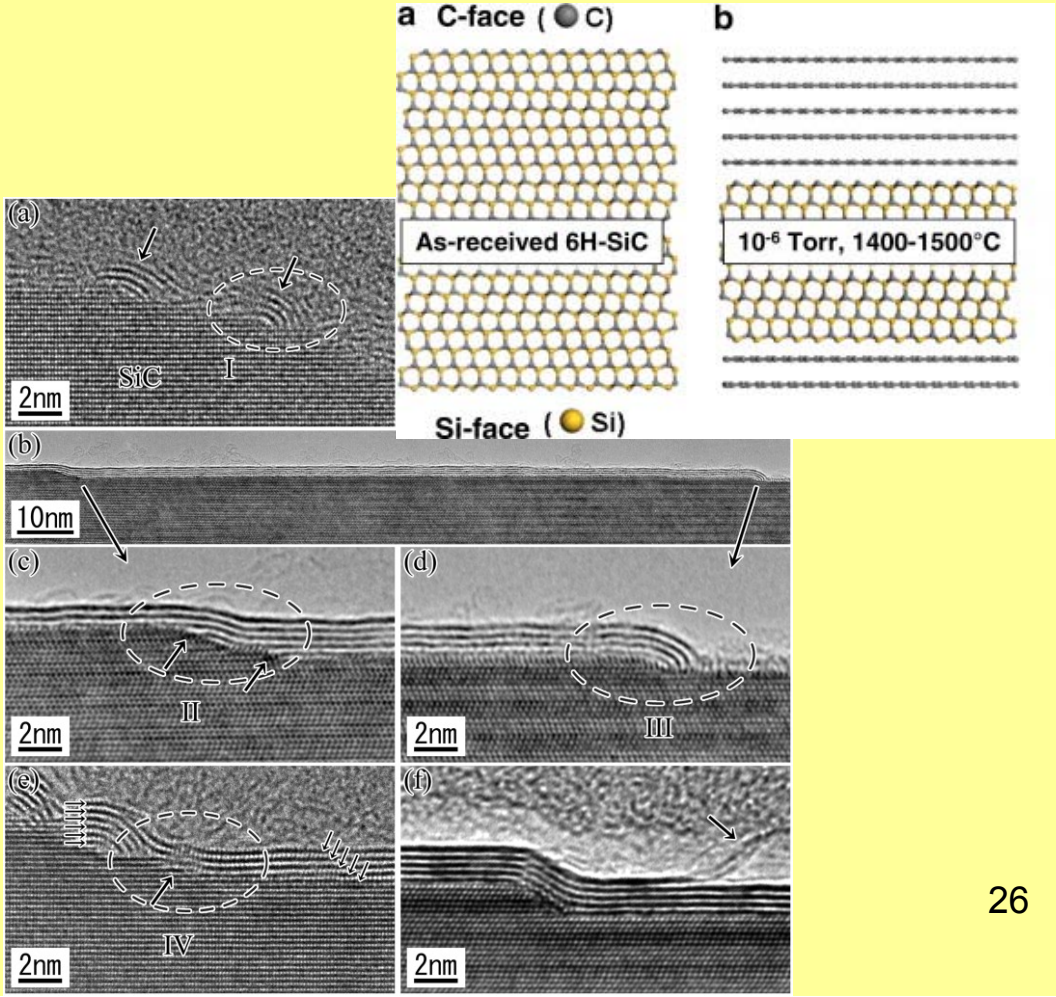


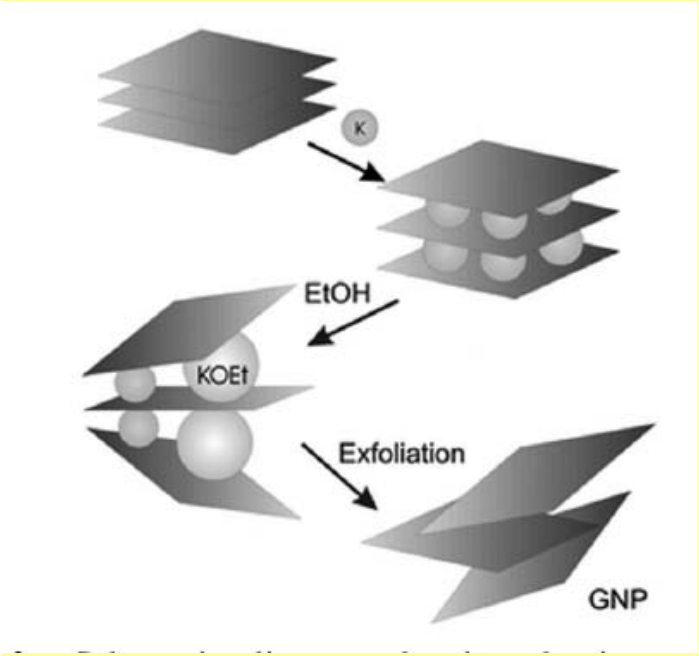


SiC pyrolysis

- Annealing of the SiC crystal in a vacuum furnace (UHV 10^{-10} Torr)
- Sublimation of Si from the surface at 1250 - 1450 °C
- The formation of graphene layers by the remaining carbon atoms



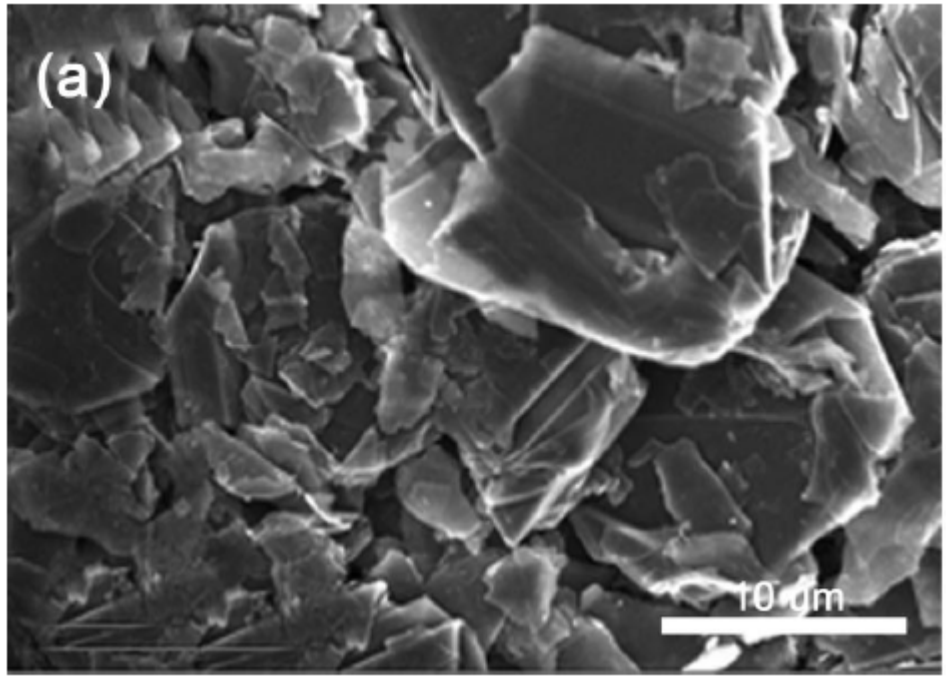


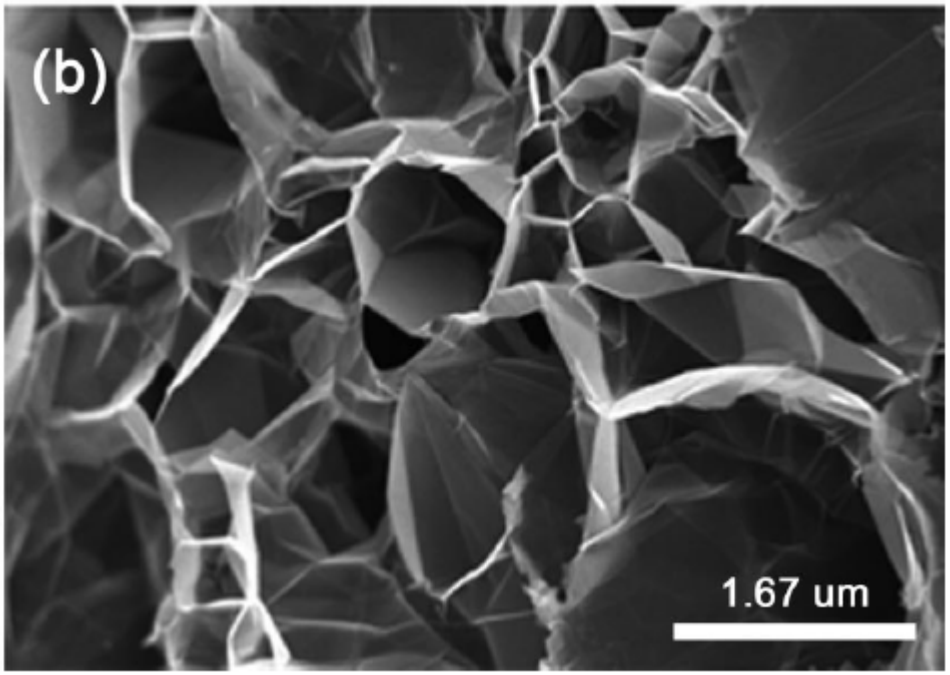


Exfoliation

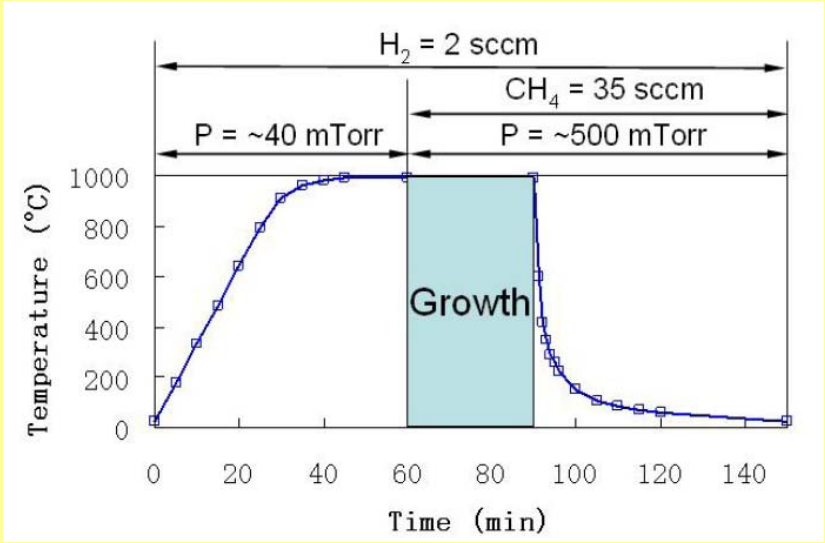
Chemical exfoliation (surfactant)

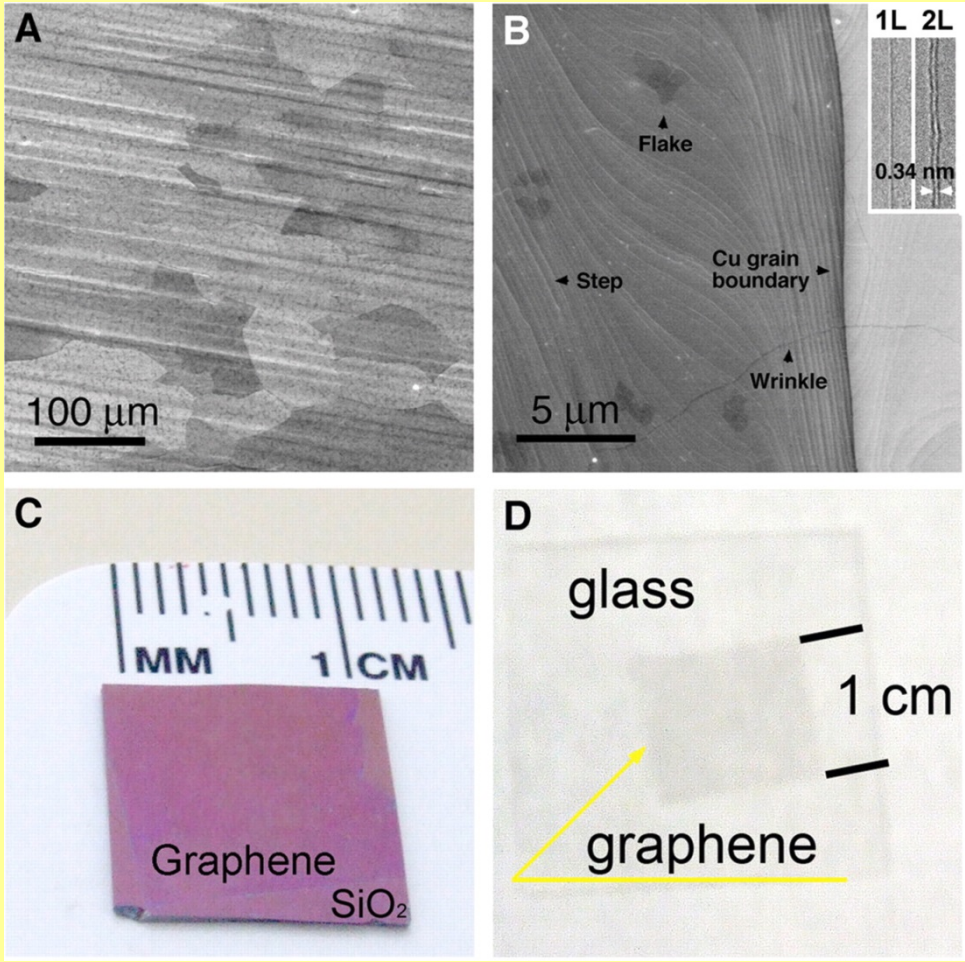
Sonochemical exfoliation





CVD from CH_4 / H_2 on Metal Surfaces





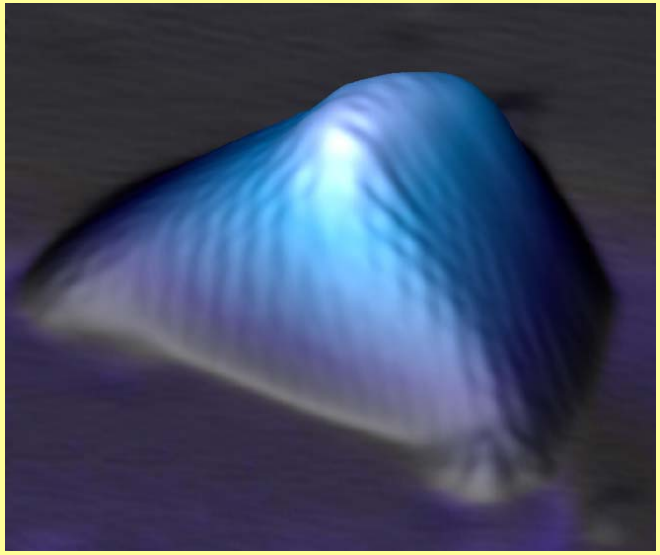
(A) SEM - graphene on a copper foil
(B) High-resolution SEM - Cu grain boundary and steps, two- and three-layer graphene flakes, and graphene wrinkles. Inset (B) TEM images of folded graphene edges. 1L, one layer; 2L, two layers.

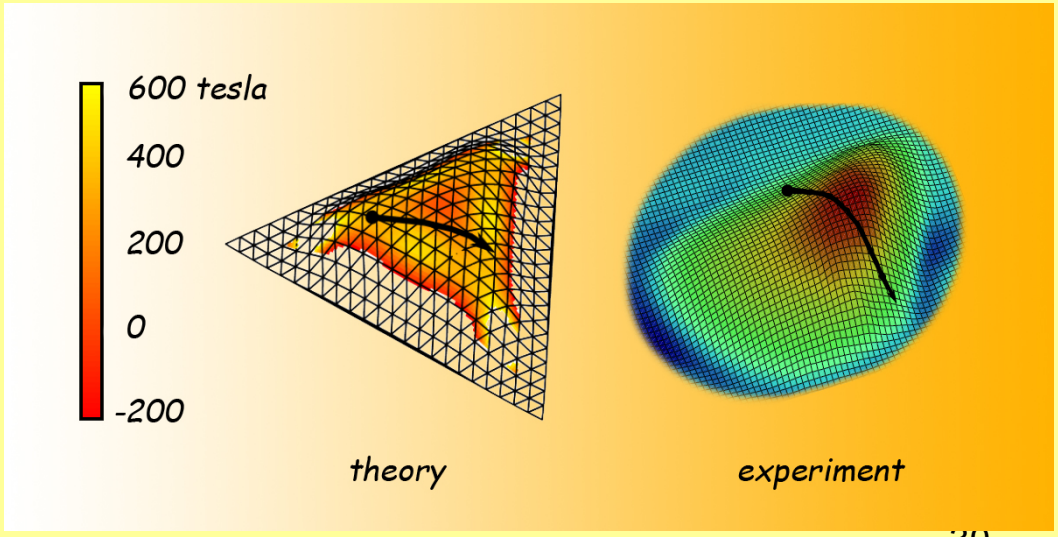
Graphene transferred onto
(C) a SiO_2/Si substrate
(D) a glass plate

Graphene on SiO_2

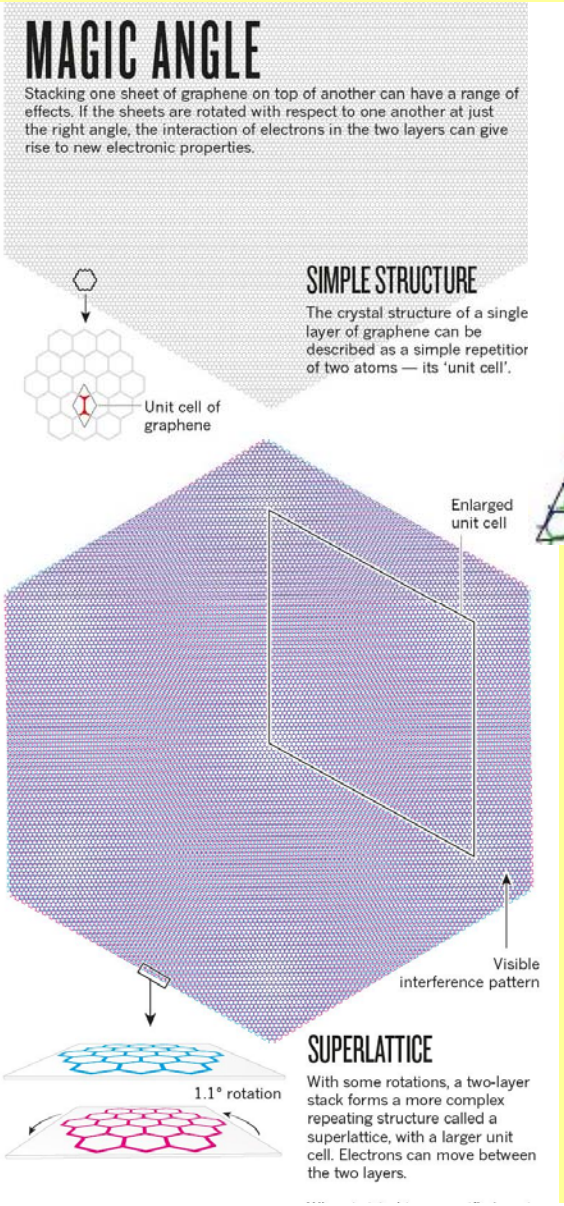
Pseudo-magnetism

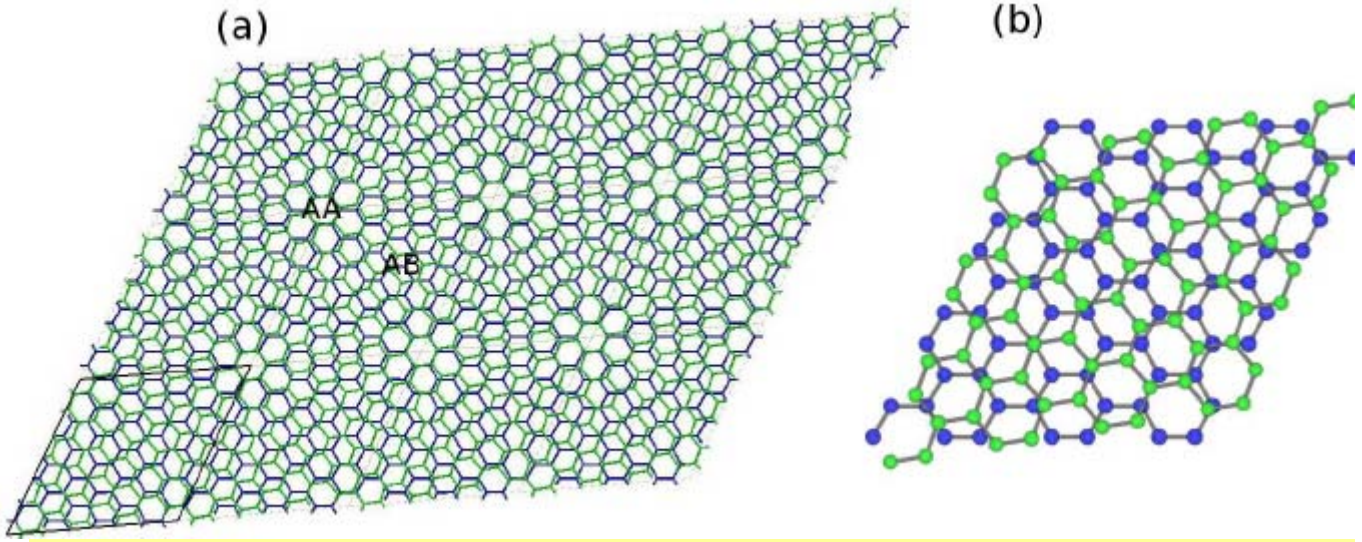
Graphene on platinum grown from ethylene at high temperatures.
Cooled to low temperature to measure STM to a few degrees above absolute zero.
Both the graphene and the platinum contracted – but Pt shrank more, excess graphene pushed up into bubbles, size 4-10 nm x 2-3 nm
The stress causes electrons to behave as if they were subject to huge magnetic fields around 300 T
(record high in a lab, max 85 T for a few ms)

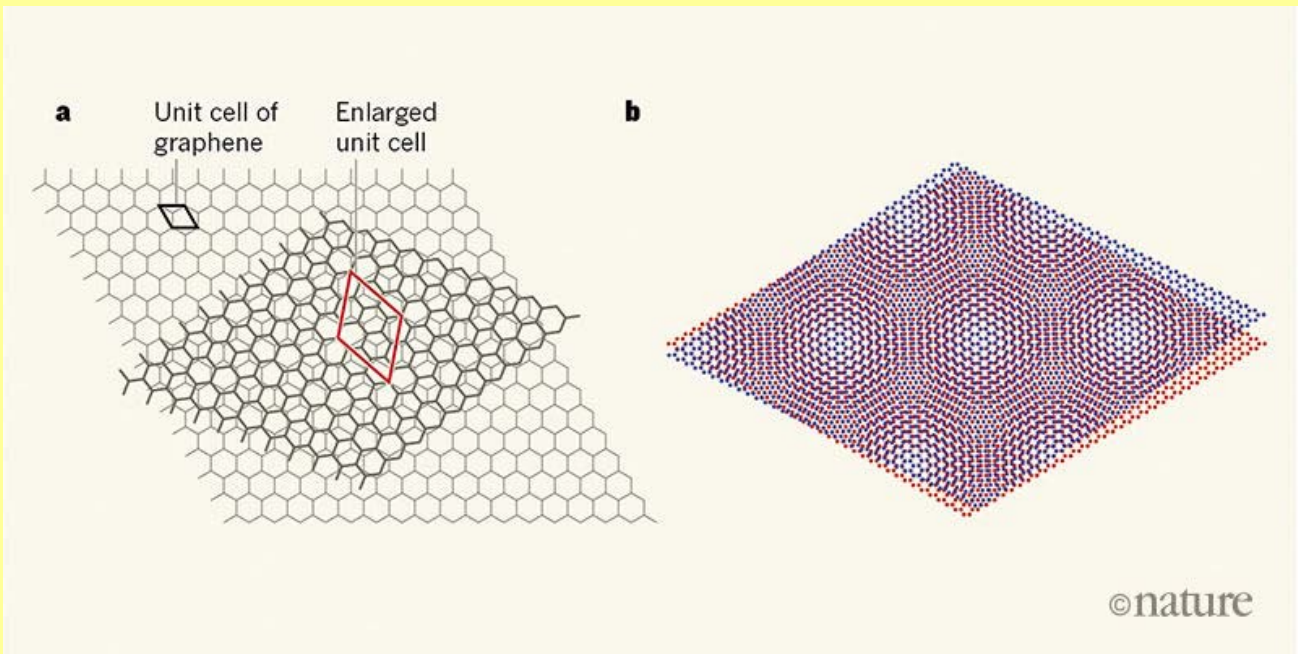




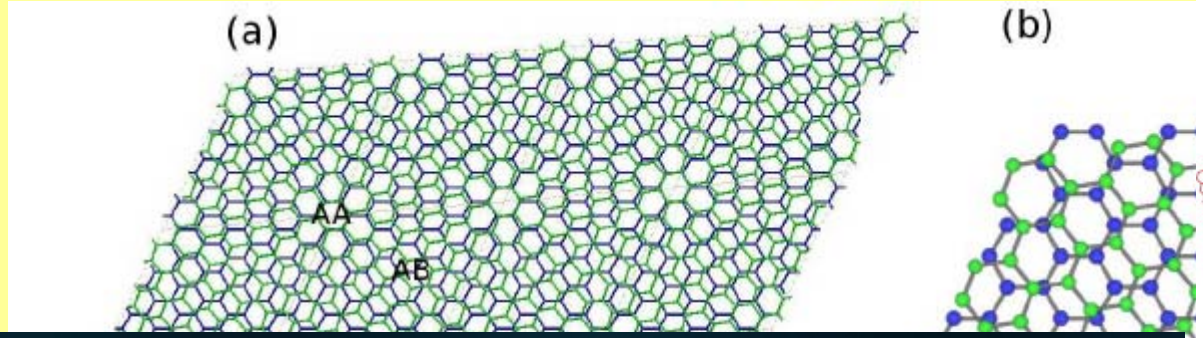
Twisted Bilayer Graphene

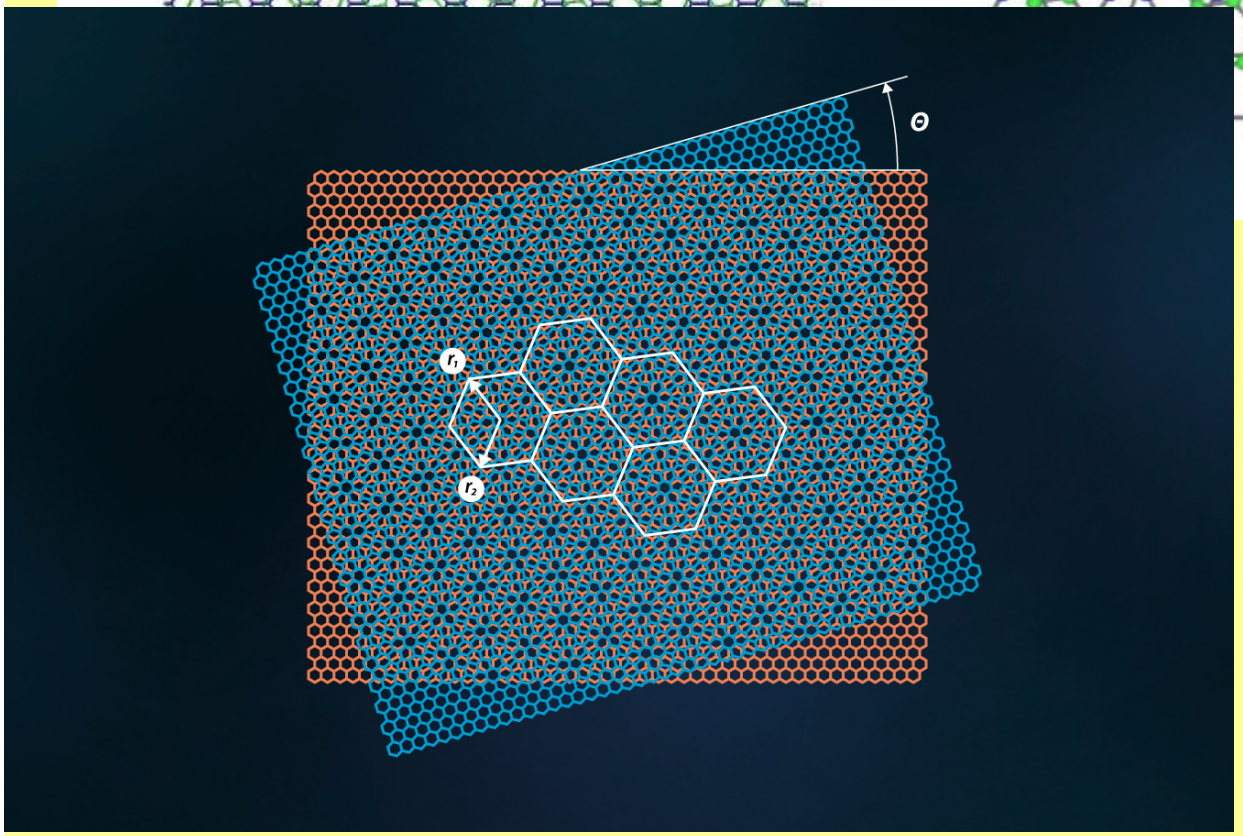


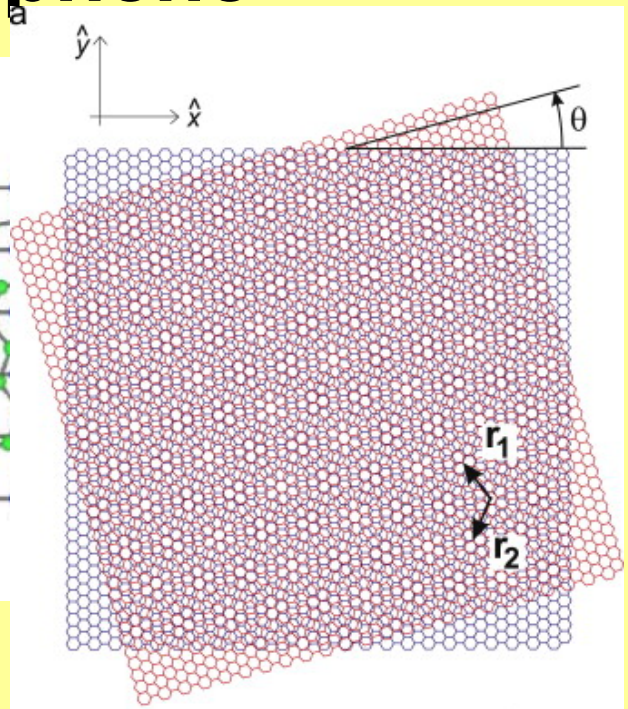


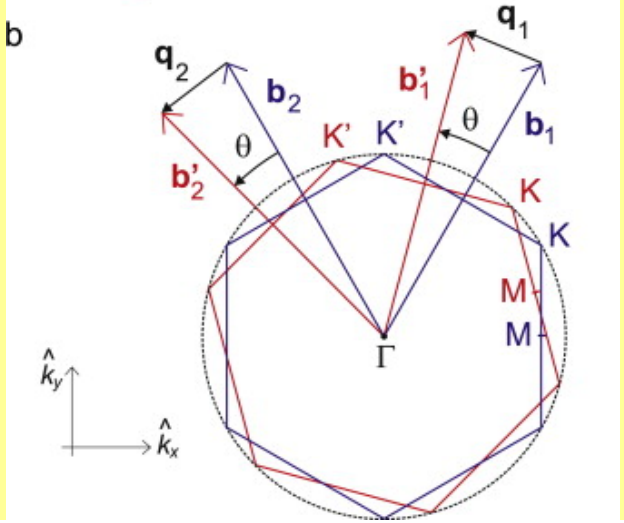


Twisted Bilayer Graphene









Graphene Family

Graphene

h-BN

BCN

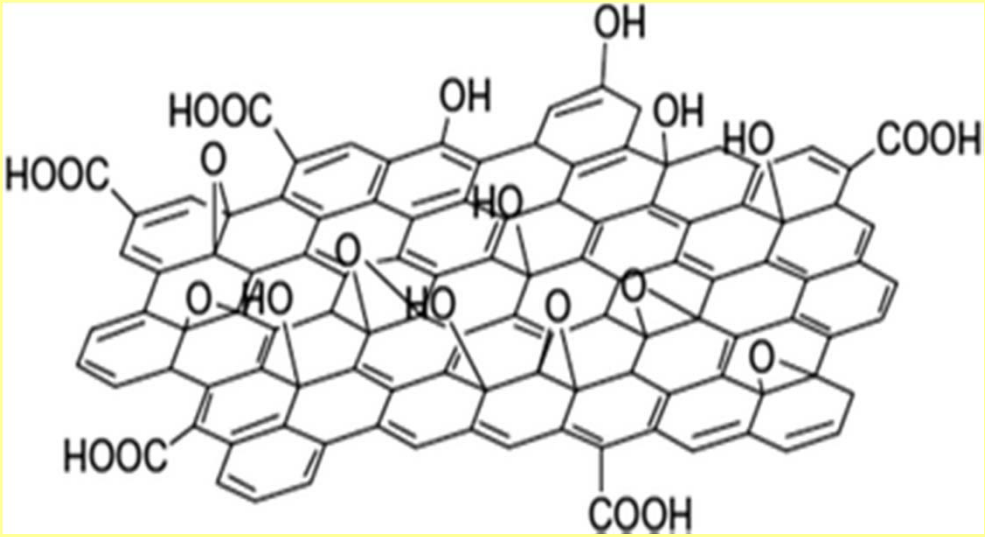
Fluorographene

Graphene oxide

C_3N_4

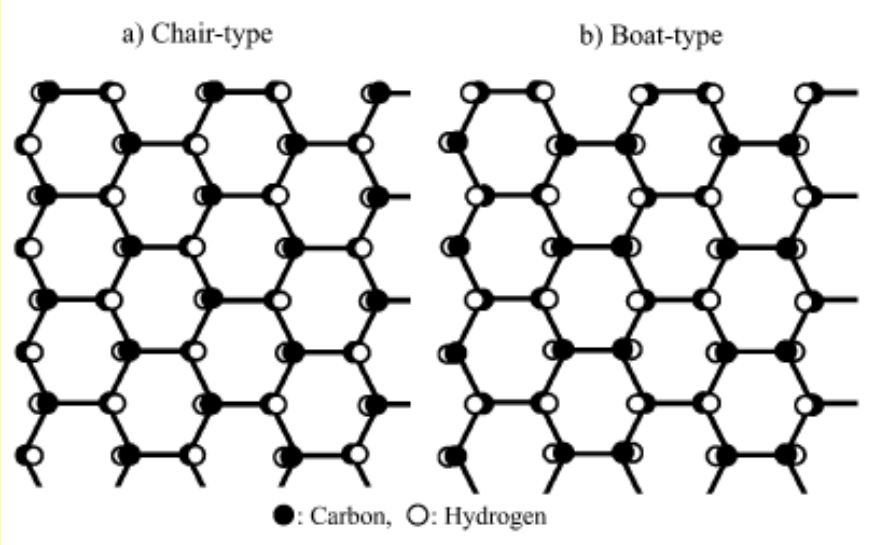
Graphene Oxide

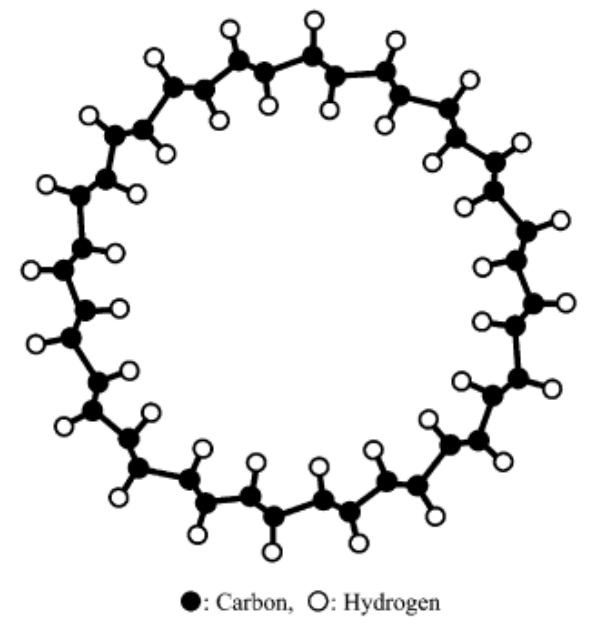
- More reactive than graphene
- Presence of oxygen groups: -OH, -COOH, =O, -O- hydrophilic character
- Electric insulator
- Specific SA (theoretically): 1700-1800 m^2g^{-1}
- Hummers method



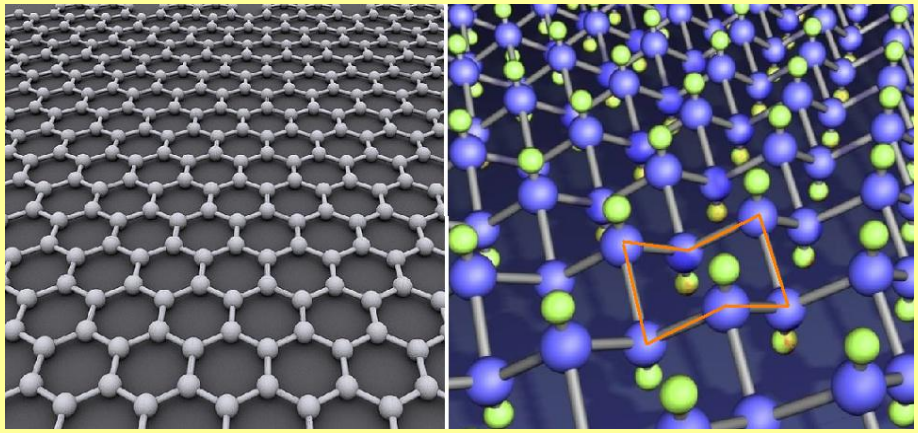
Graphane – Hydrogenated Graphene

- 2009 (graphene + cold hydrogen plasma)
- Two conformations: chair x boat
- Calculated binding energy = most stable compound with stoichiometric formula CH
- Chair type graphane insulating nanotubes



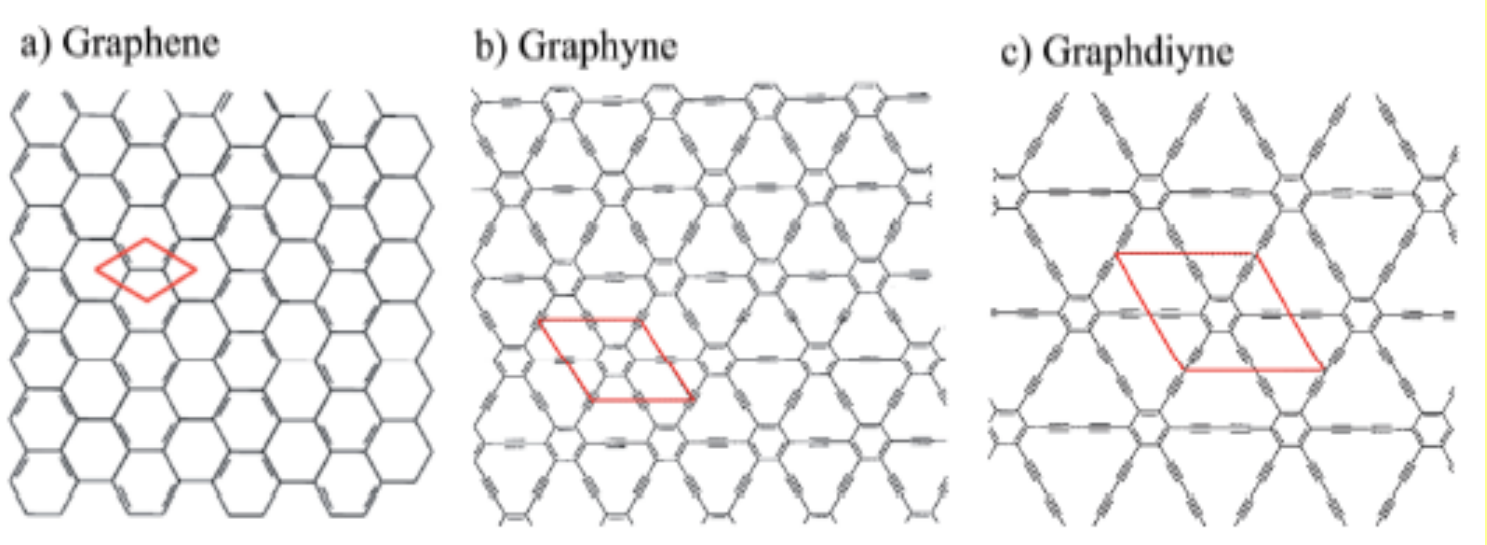


Fluorographene



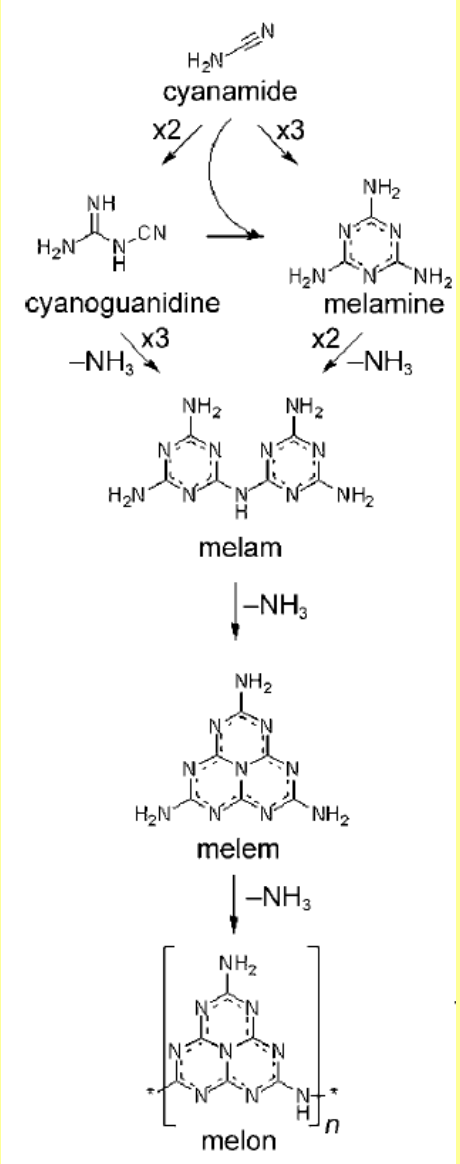
- **Monolayer of graphite fluoride**
- **Chair type x boat type-strong repulsion**
- **Sythesis:**
  - **Graphene + XeF_2/CF_4 (room temperature)**
  - **Mechanical or chemical exfoliation of graphite fluoride**
  - **By heating graphene in XeF_2 gas at 250 °C**
- **Graphene + XeF_2 at 70 °C – high-quality insulator, stable up to 400 °C (resemblence with teflon)**

Graphyn, Graphydiyn

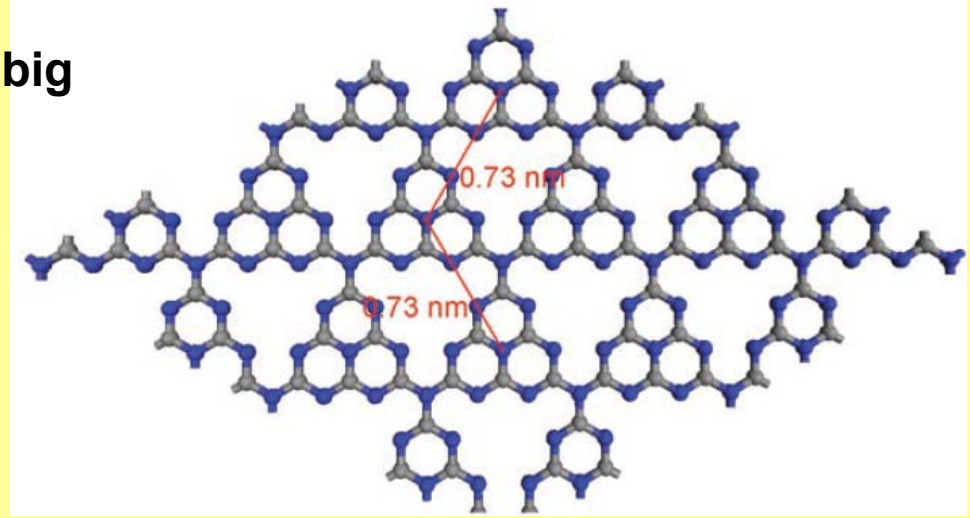


- **Predicted**
- **“Non-derivatives“ of graphene**
- **Semiconductors**
- **Movement of electrons as in graphene but only in one direction**

Graphitic Carbon Nitride



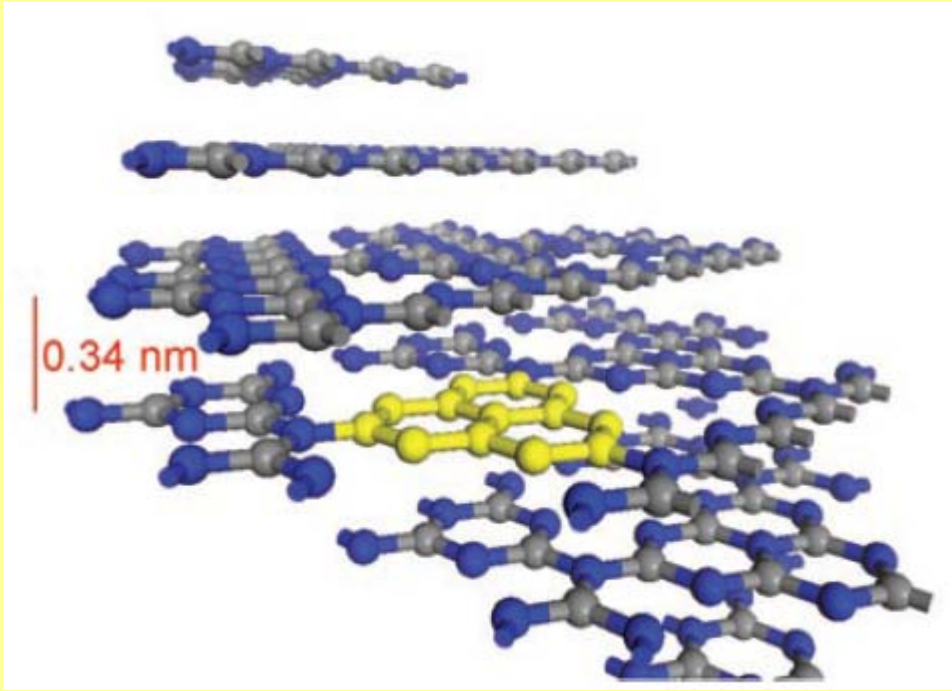
1834 Berzelius, Liebig



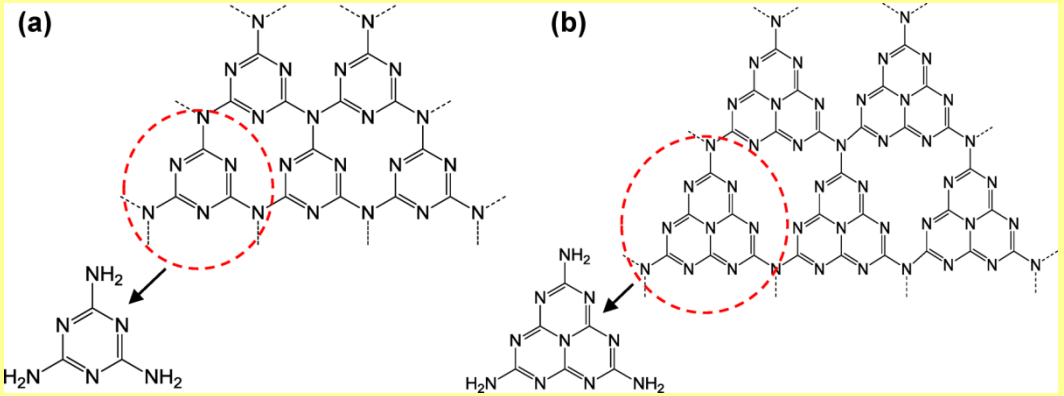
Temperature-induced condensation

dicyandiamide $NH_2C(=NH)NHCN$

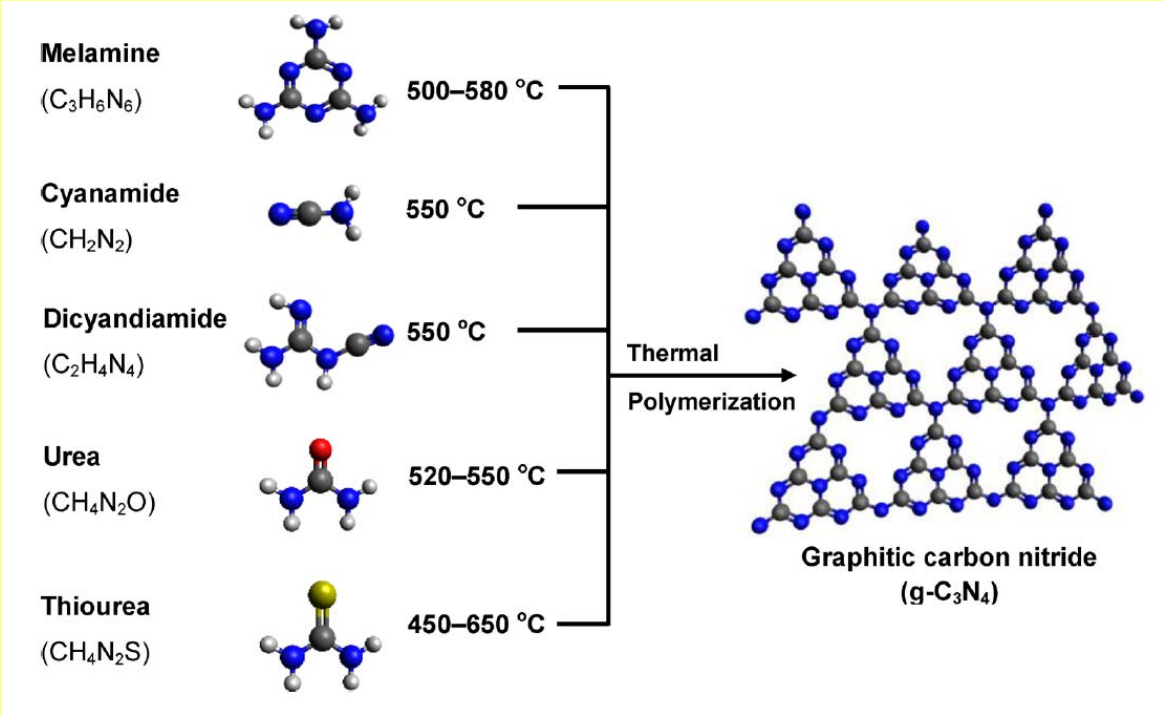
In a LiCl/KCl melt



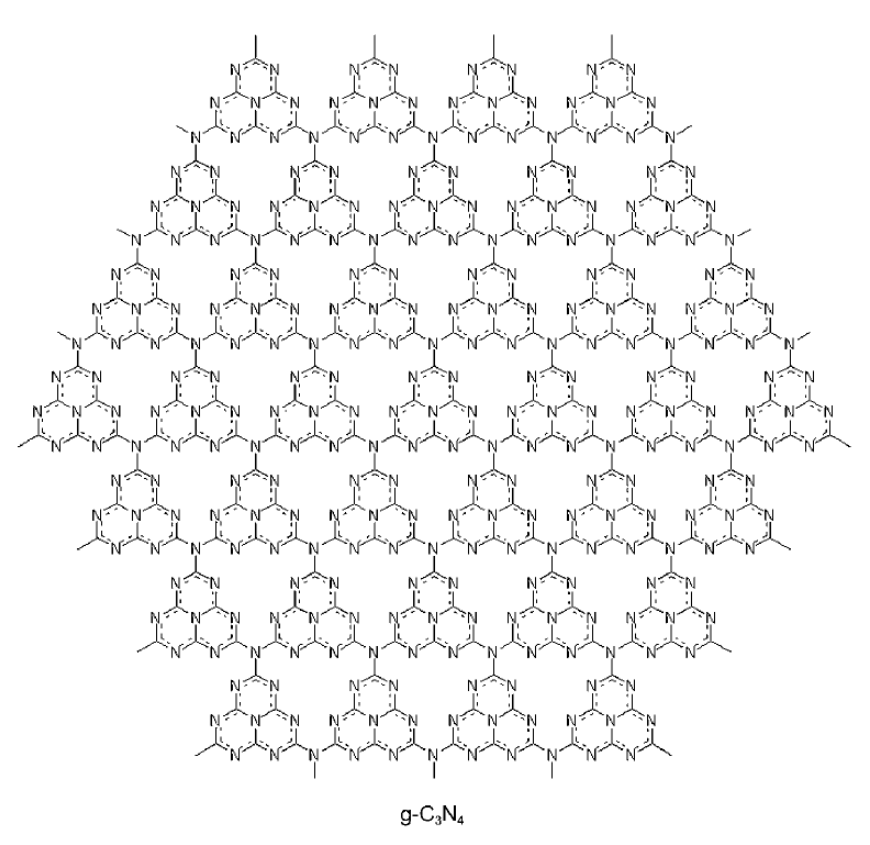
Graphitic Carbon Nitride



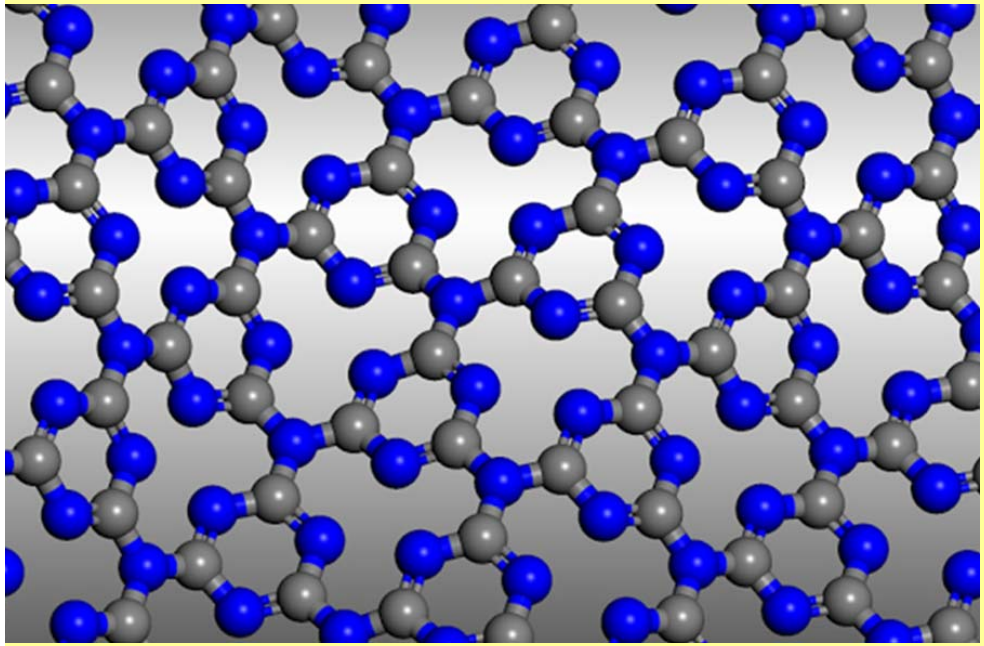
(a) triazine and (b) tri-s-triazine (heptazine)



Graphitic Carbon Nitride

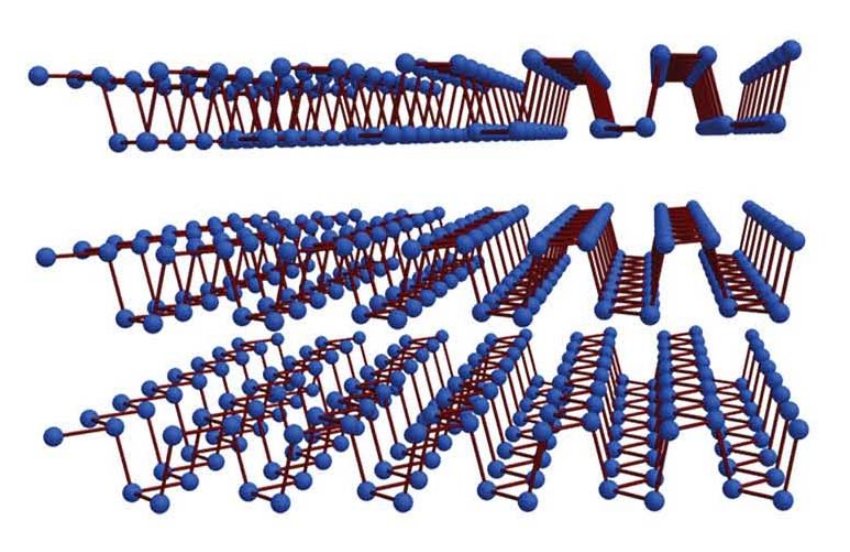


("g-C_3N_4")



band gap 1.6 - 2.0 eV
small band gap semiconductors
Si (1.11 eV), GaAs (1.43 eV), and GaP (2.26 eV)

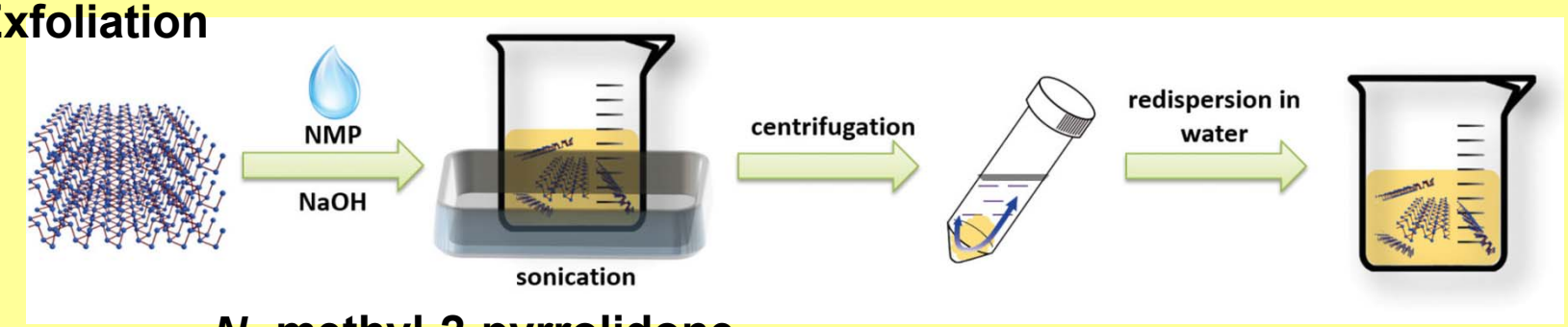
Phosphorene



Black phosphorus

Orthorhombic
a = 3.31 Å, b = 4.38 Å, c = 10.50 Å
$\alpha = \beta = \gamma$ = 90°
Space group *Bmab*

Exfoliation

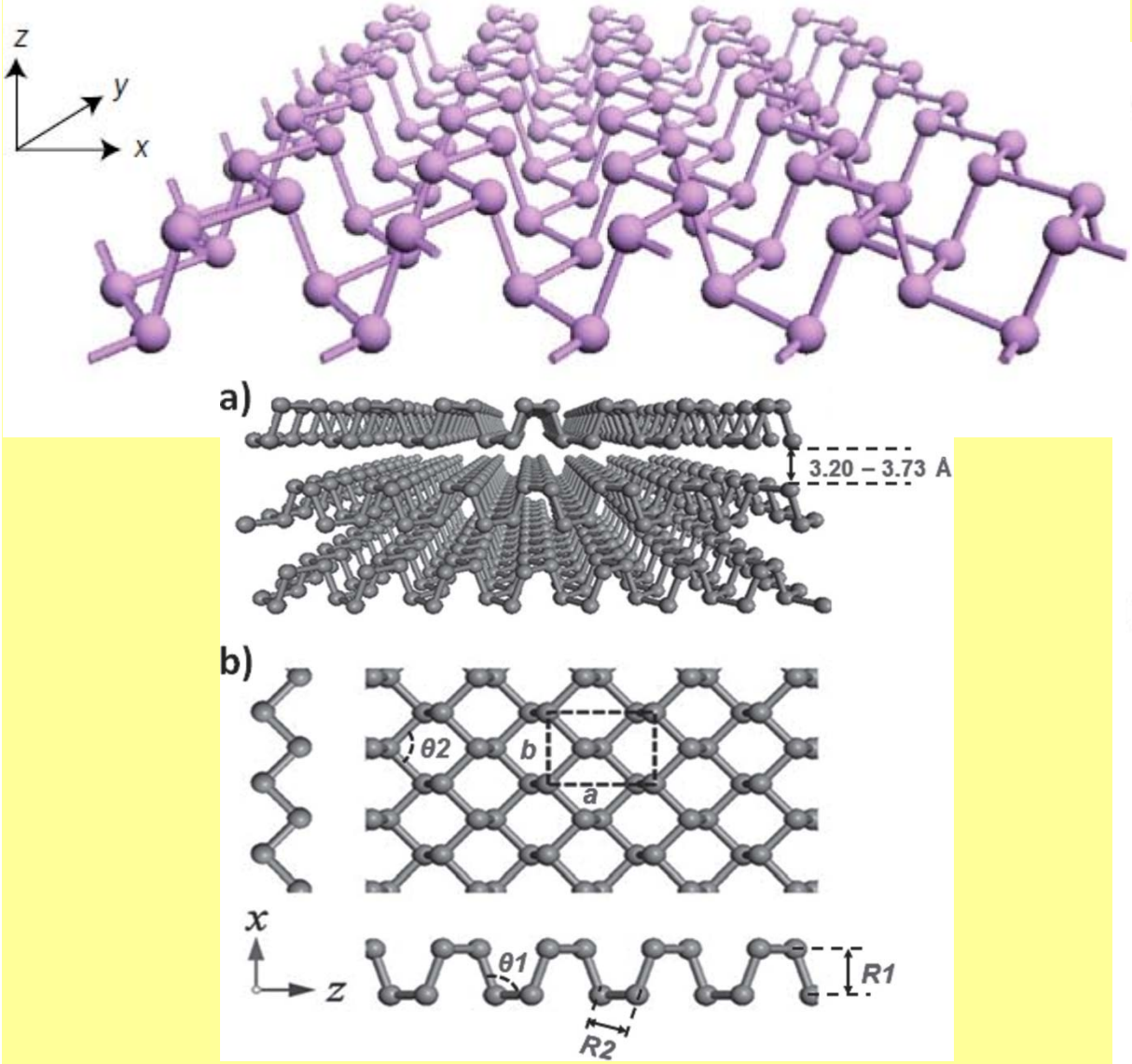


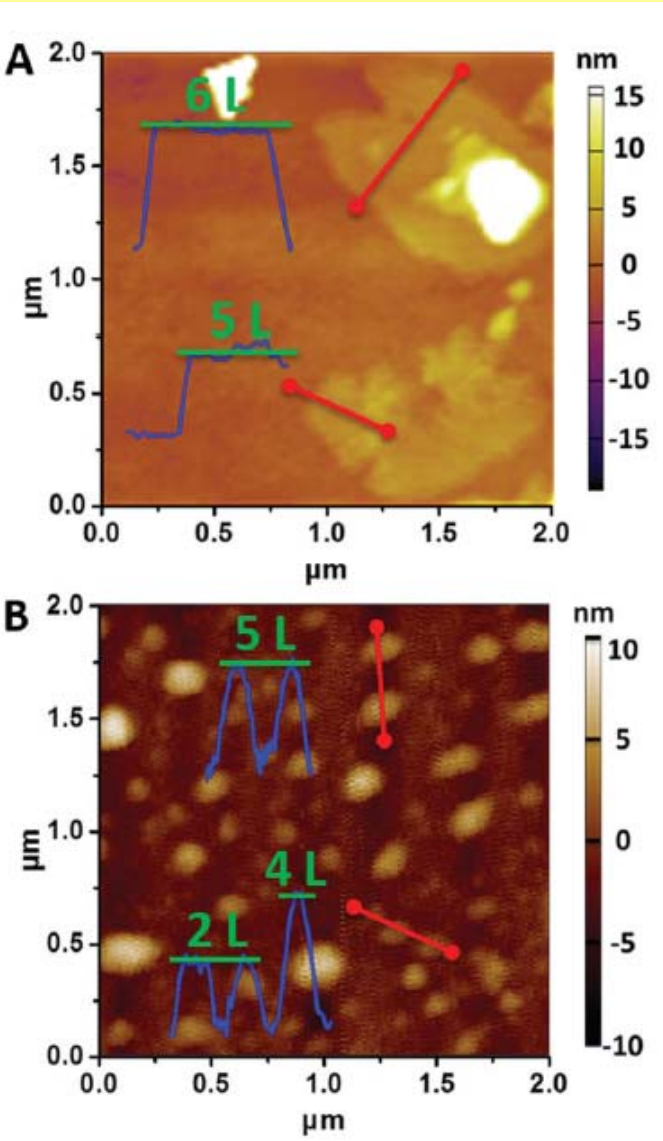
***N* -methyl-2-pyrrolidone**

Semiconductor - direct band gap
bulk BP 0.3 eV
monolayer phosphorene 1.5 eV

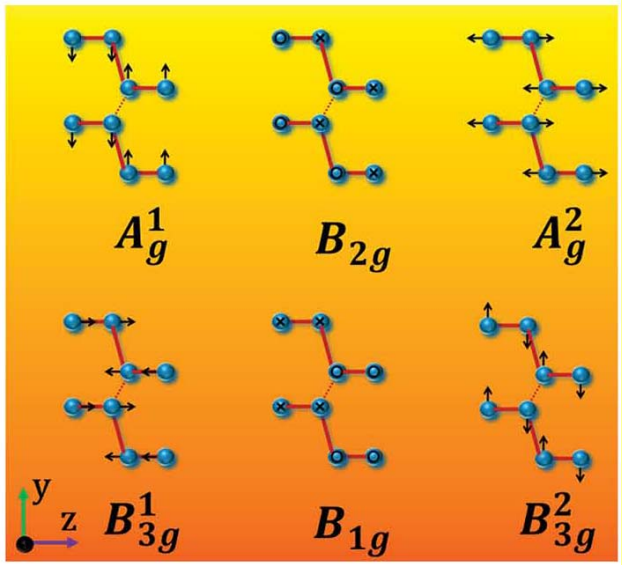
Phosphorene

Height-mode AFM images
single-layer phosphorene
ca. 0.9 nm



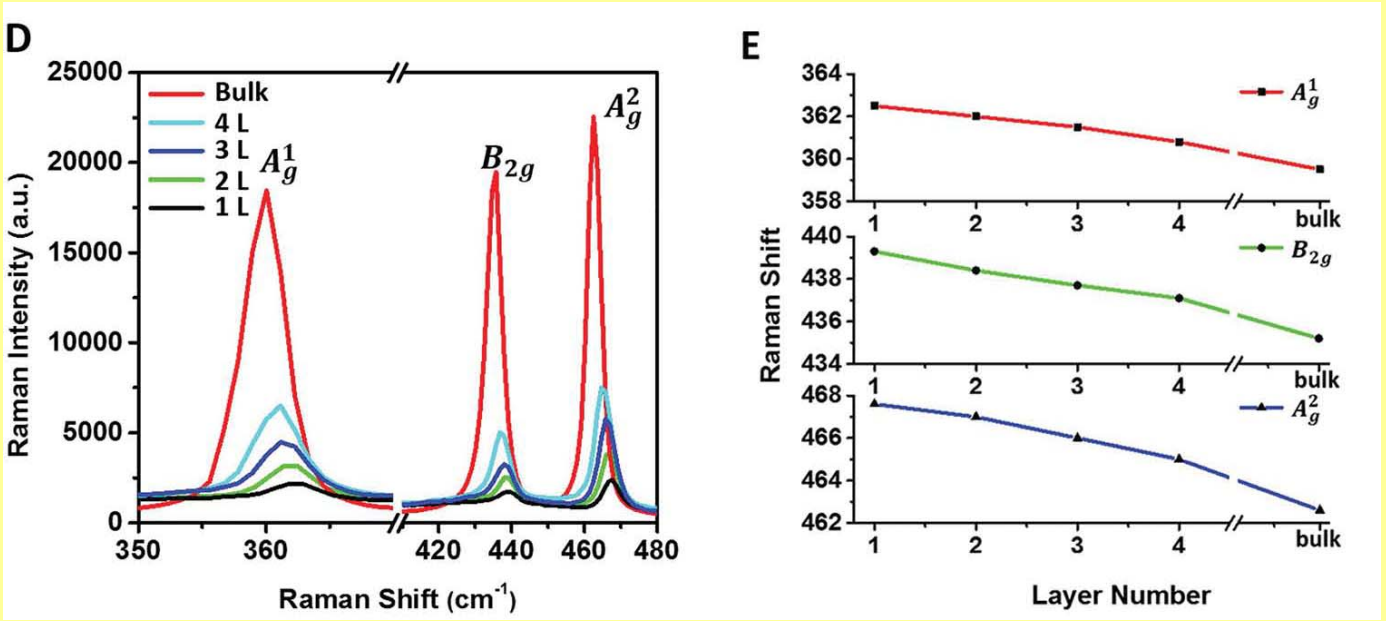


Phosphorene

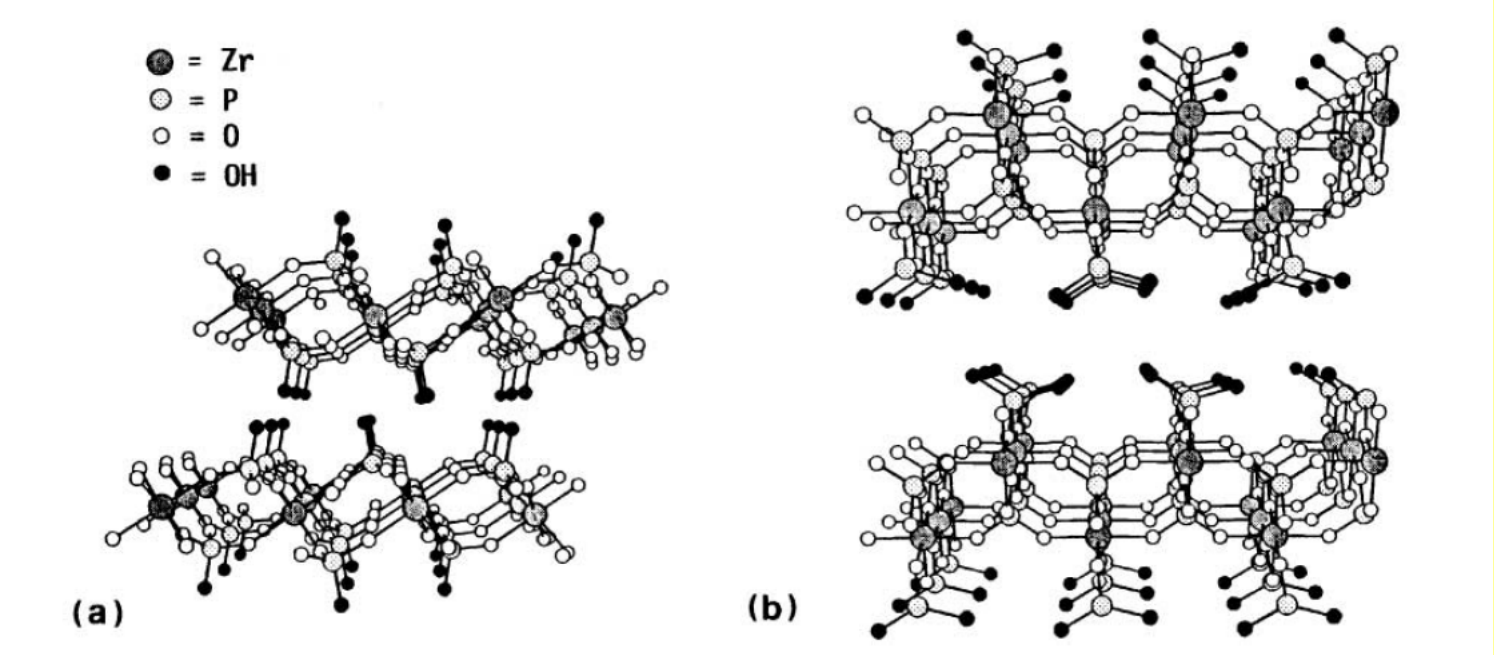


Black phosphorus
12 lattice vibrational modes
6 Raman active modes
3 vibrational modes A^1_g , B_{2g} , and A^2_g can be detected when the incident laser is perpendicular to the layered phosphorene plane: 361 cm^{-1}, 438 cm^{-1}, 465 cm^{-1}

As the number of phosphorene layers increases, the three Raman peaks red-shift



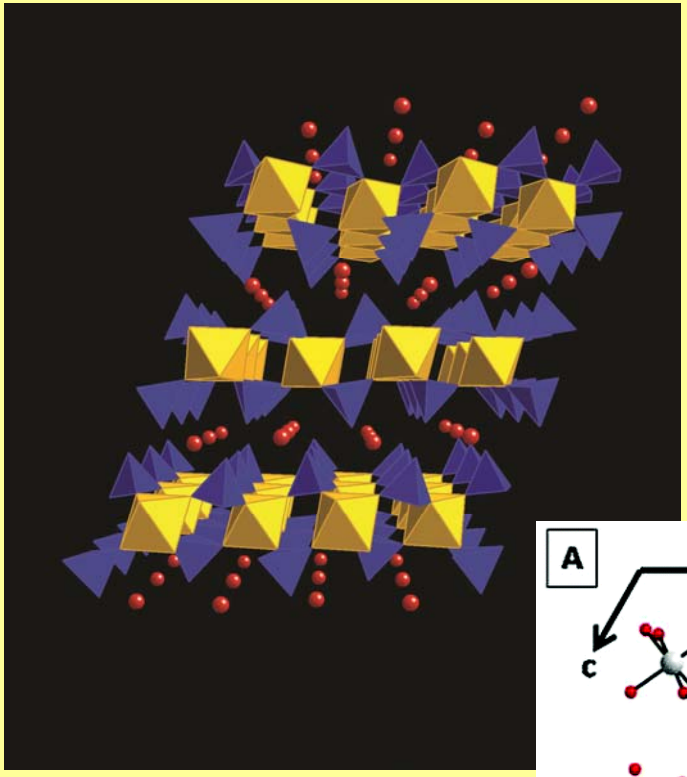
Zirconium Phosphates



(a) α-zirconium phosphate = $Zr(HPO_4)_2.H_2O$
interlayer spacing 7.6 Å

(b) γ-zirconium phosphate = $Zr(PO_4)(H_2PO_4)2H_2O$
interlayer spacing 12.2 Å

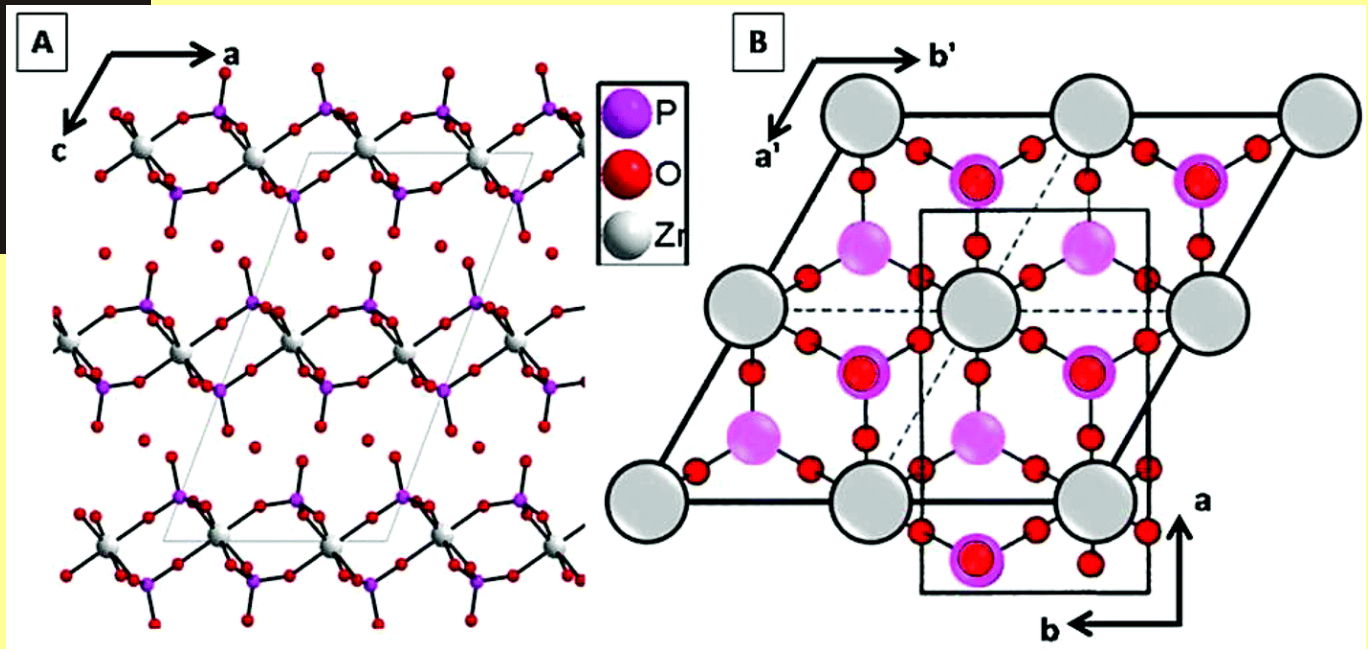
Zirconium Phosphates



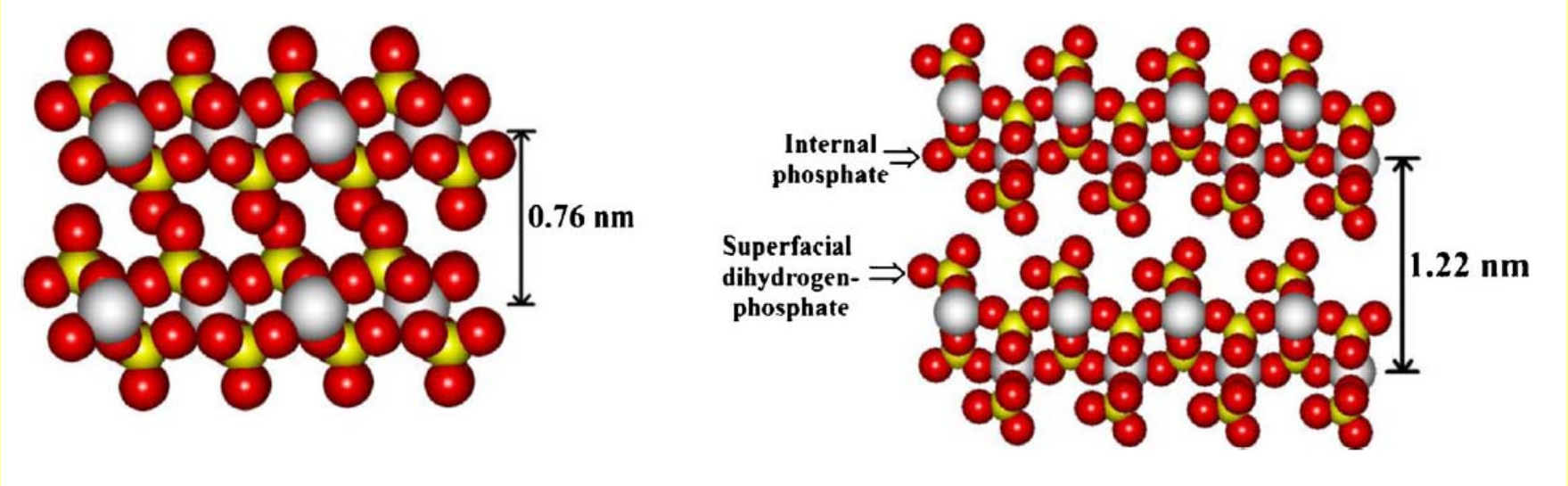
α-zirconium phosphate

$Zr(HPO_4)_2.H_2O$

interlayer spacing 7.6 Å



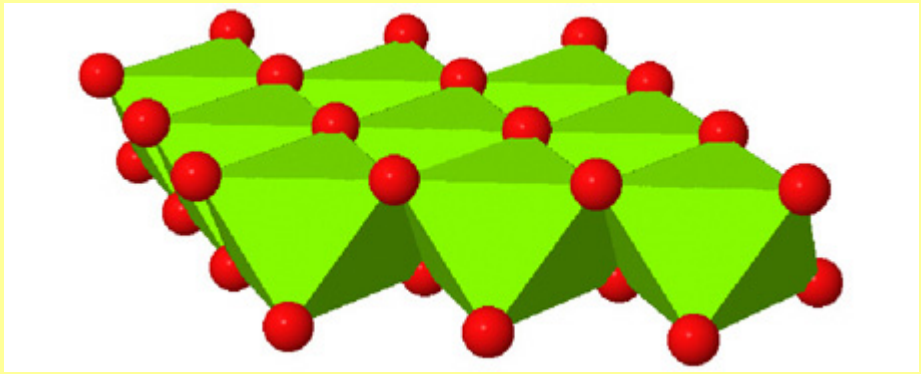
Zirconium Phosphates

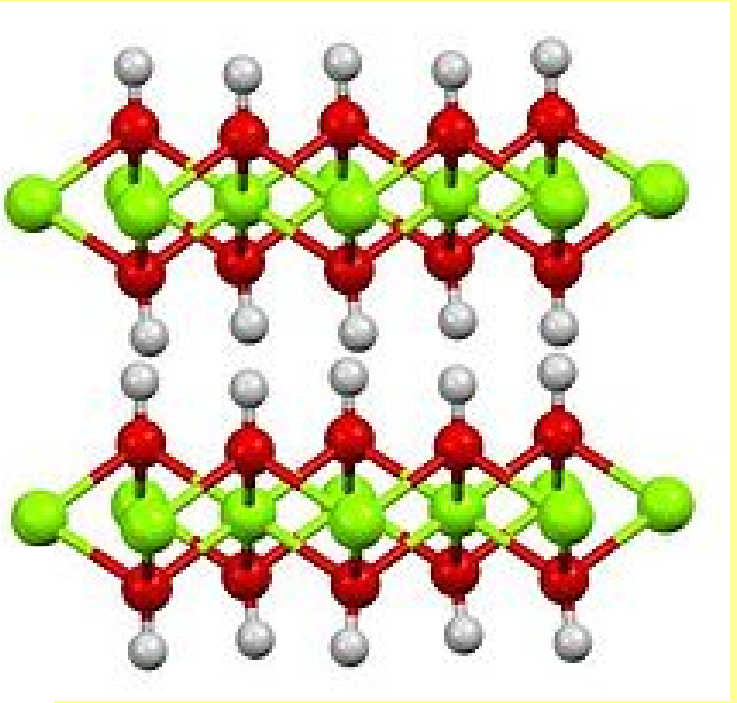


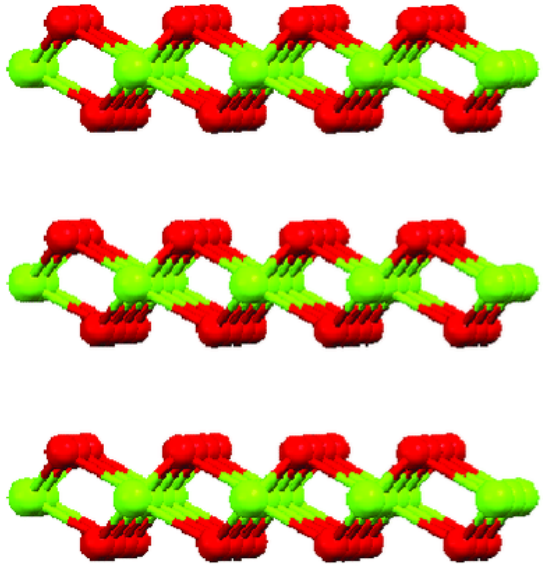
(a) α-zirconium phosphate = $Zr(HPO_4)_2.H_2O$
interlayer spacing 7.6 Å

(b) γ-zirconium phosphate = $Zr(PO_4)(H_2PO_4)2H_2O$
interlayer spacing 12.2 Å

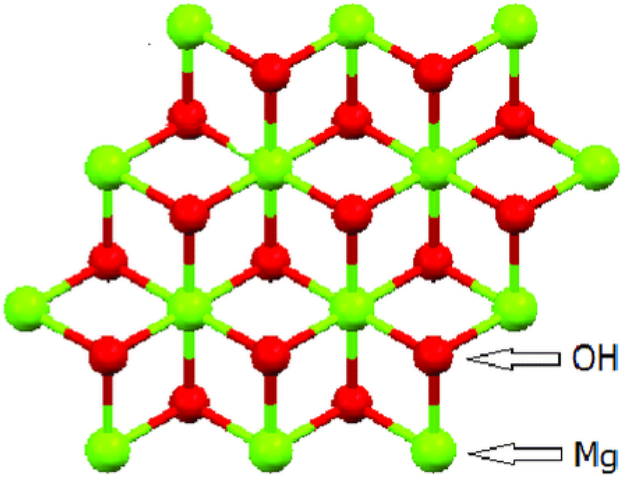
Brucite - $Mg(OH)_2$





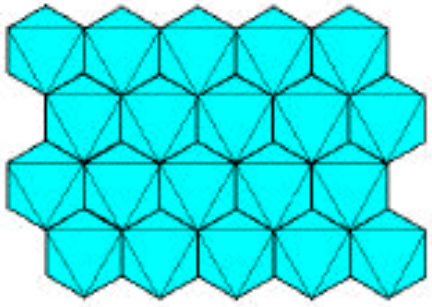


Side view

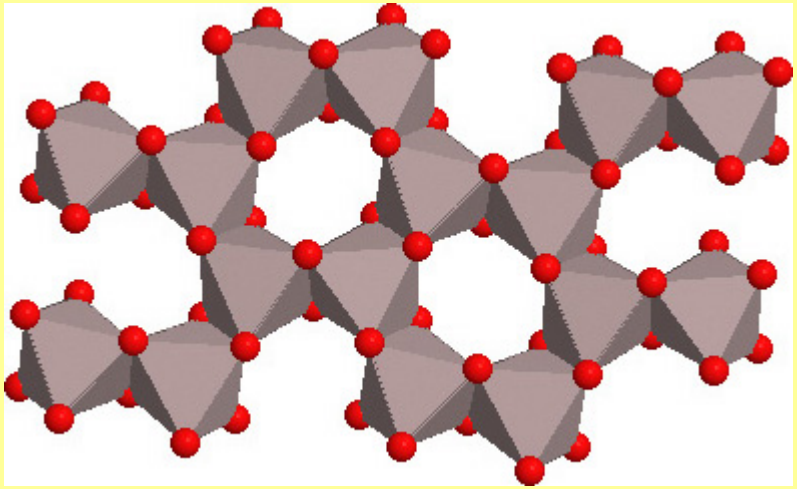


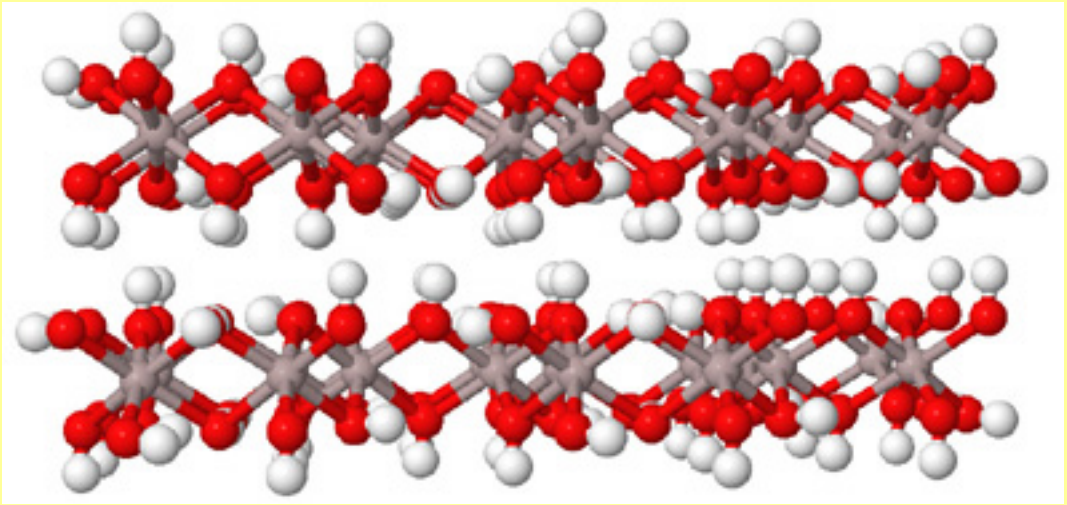
Top view

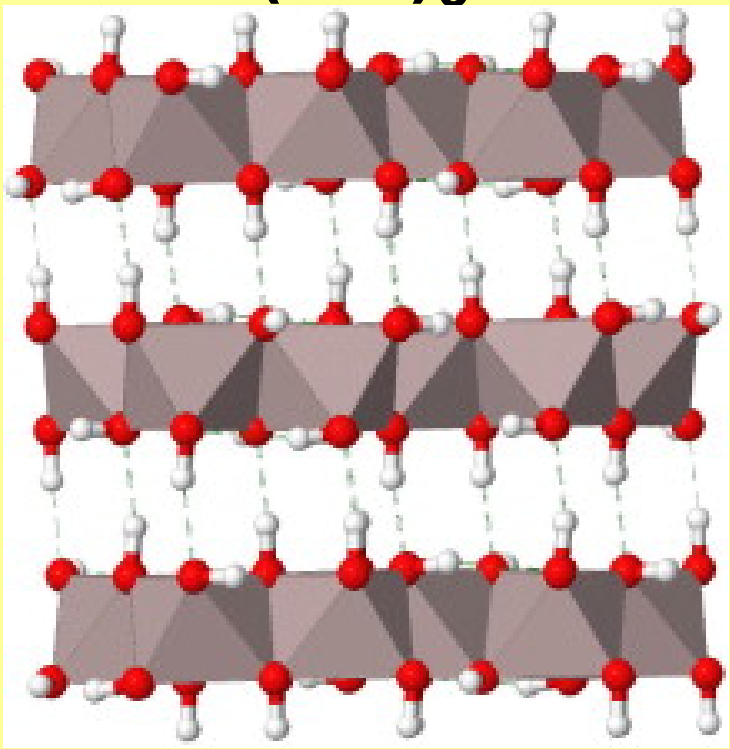
trioctahedral



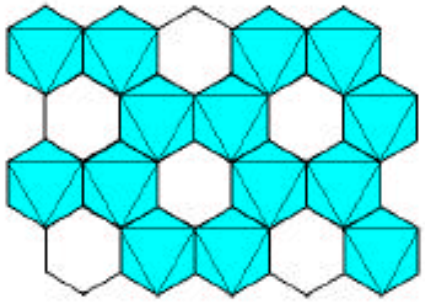
Bayerite and Gibbsite - $Al(OH)_3$



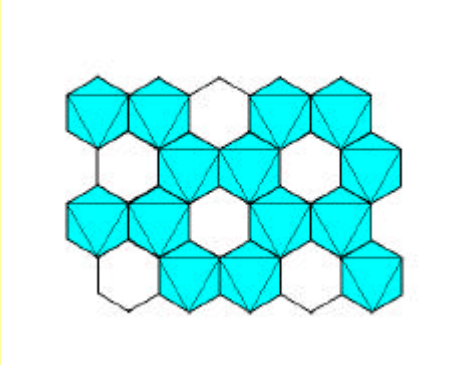




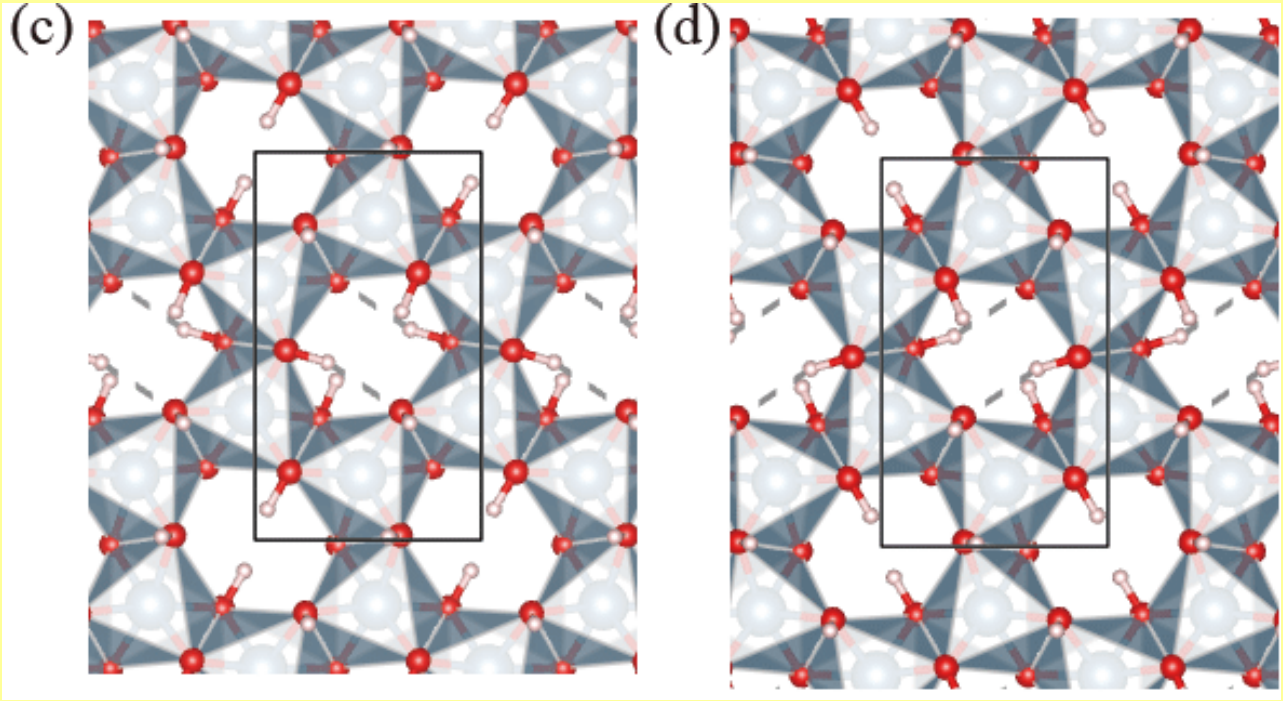
dioctahedral



Bayerite and Gibbsite - $Al(OH)_3$



Opposite faces of a single layer $Al(OH)_3$ (A and B sides, respectively)

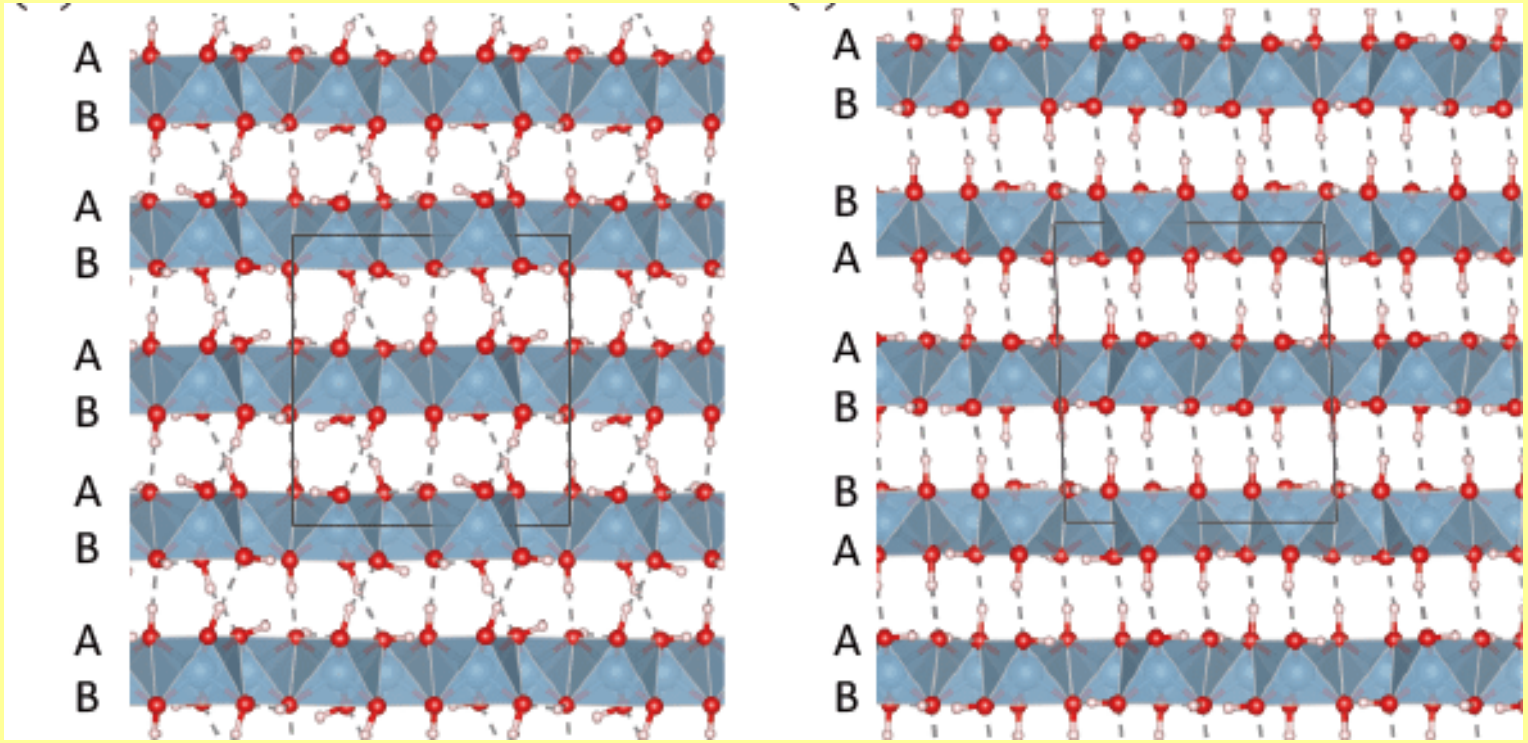


Bayerite and Gibbsite - $Al(OH)_3$

Bayerite and Gibbsite phases have an identical single layer as the building block

Bayerite is stacked by AB-AB sequence HCP of oxides

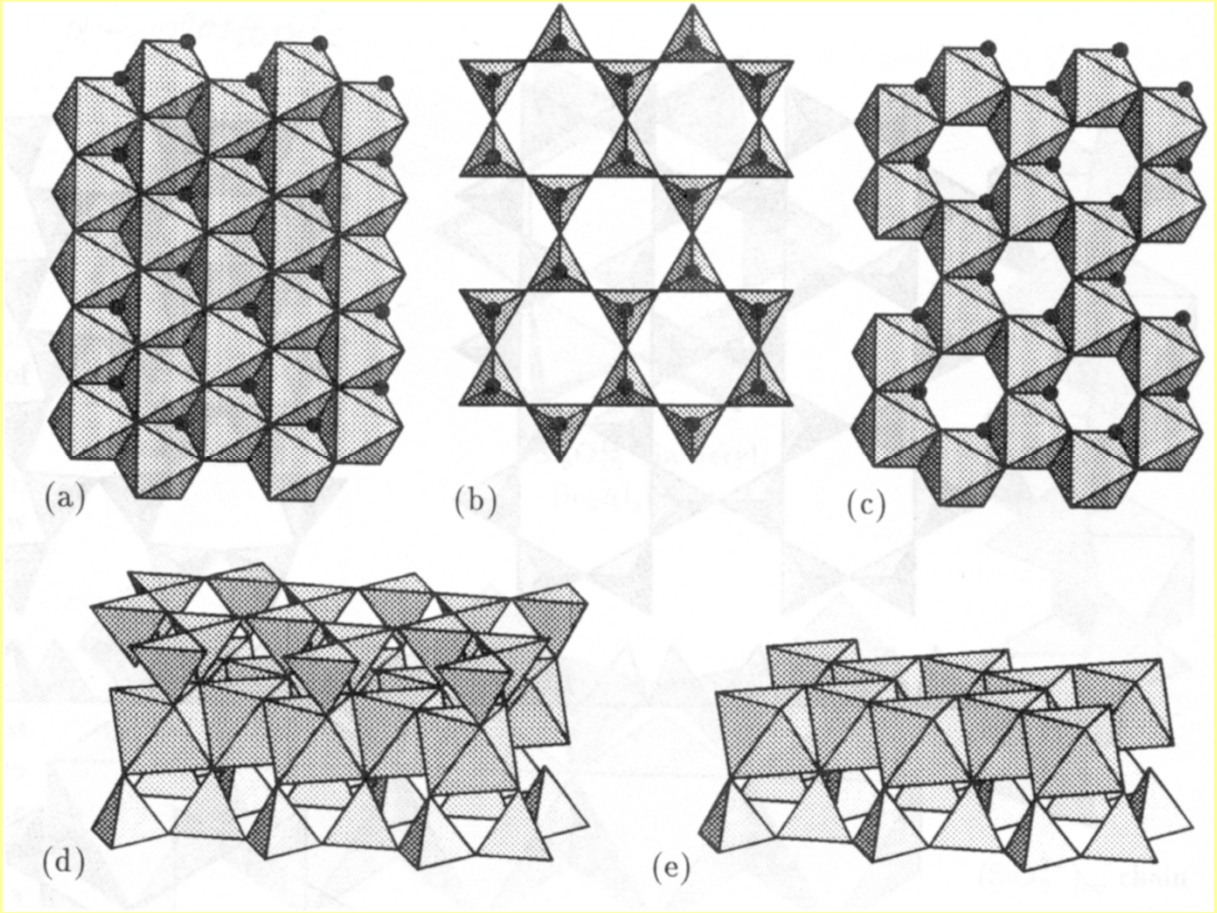
Gibbsite is stacked by AB-BA sequence CCP of oxides



Clay Minerals

$[Si_4O_{10}]^{4-}$ tetrahedral sheet

$[Mg_6O_{12}]^{12-}$ trioctahedral sheet of octahedral units



$[Al_4O_{12}]^{12-}$ dioctahedral sheet

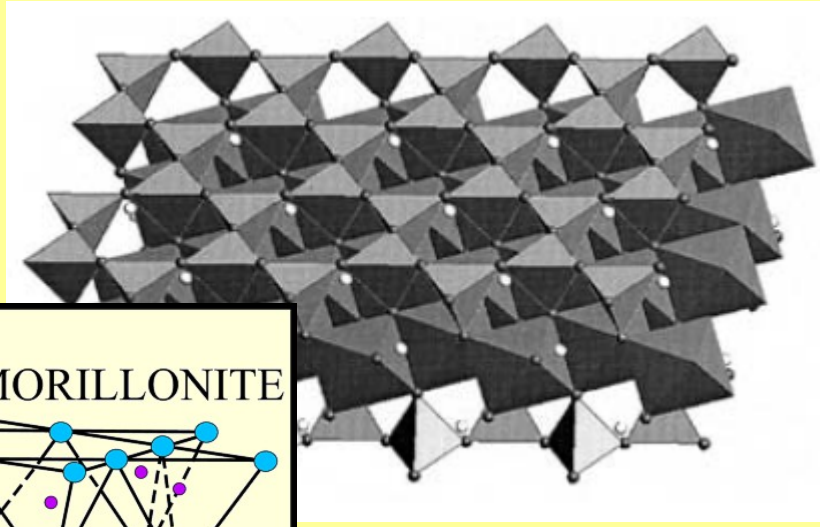
2:1

montmorillonite

1:1

kaolinite

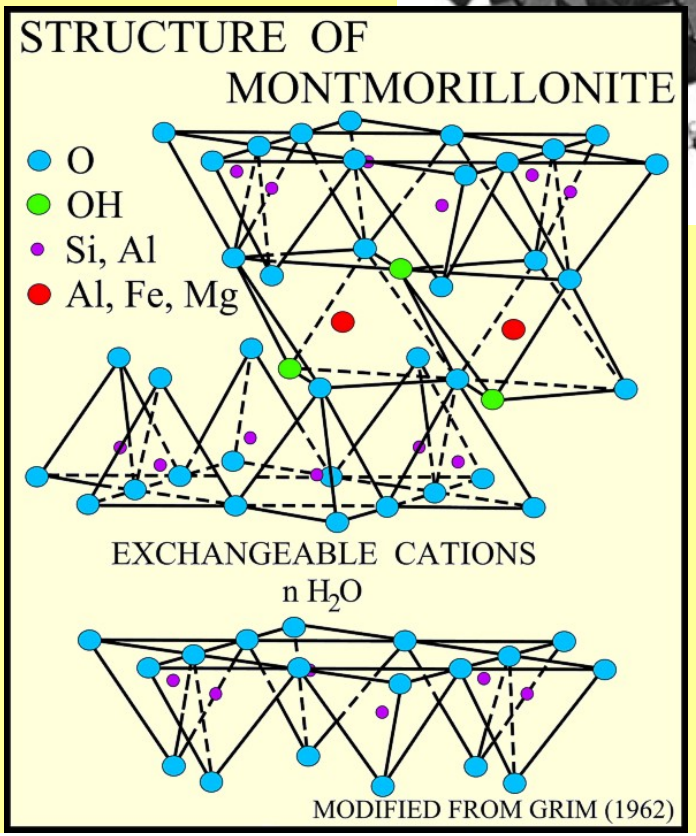
Montmorillonite



- *Dioctahedral clay mineral*
- T_d-O_h-T_d sandwich
- Isomorphous substitution

O_h: Al^{3+} by Mg^{2+}
T_d: Si^{4+} by Al^{3+}

→ **Net negative charge**
→ **Interlayer cations**

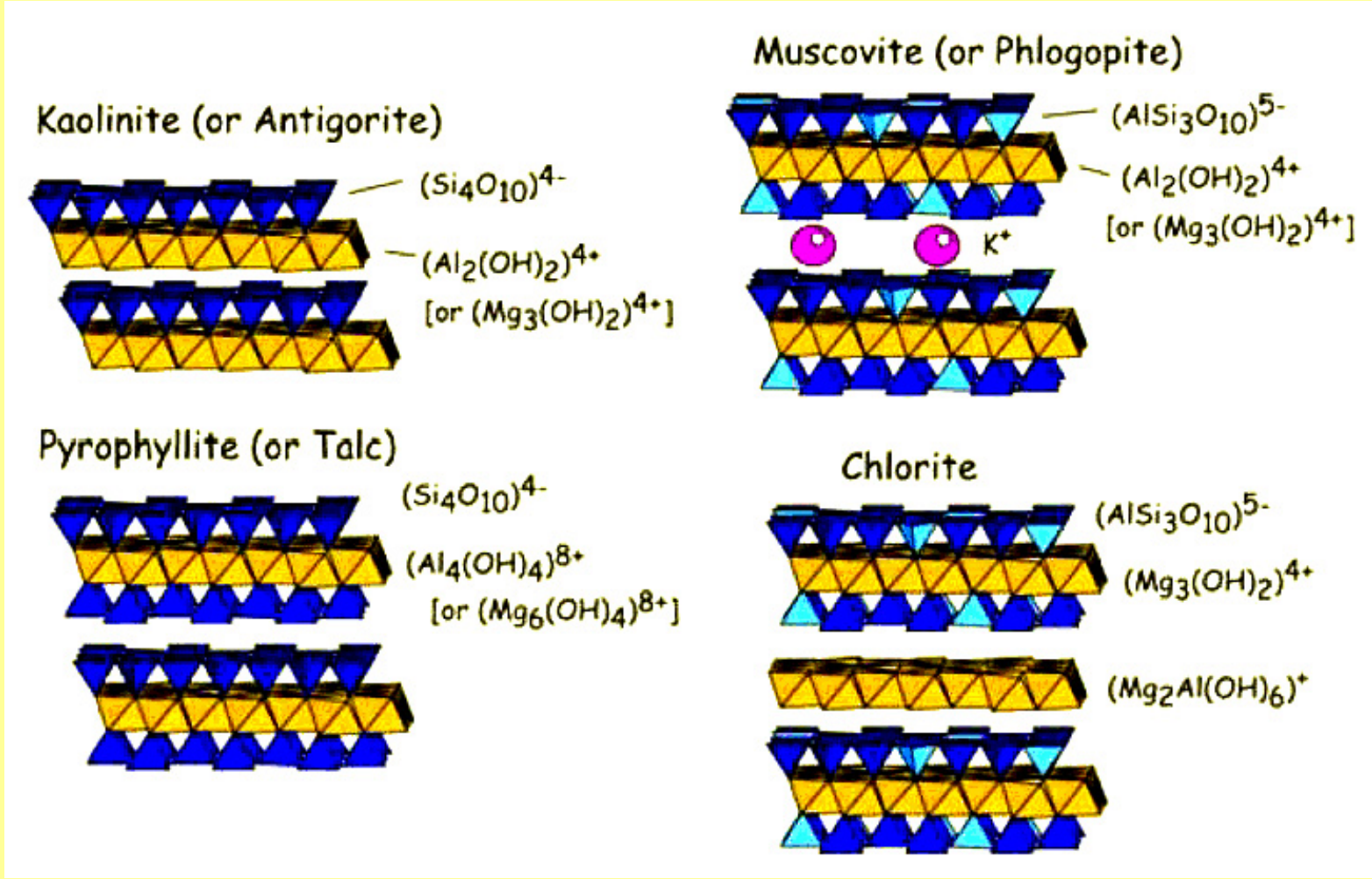


$(Na,Ca)_{0.33}(Al,Mg)_2(Si_4O_{10})(OH)_2 \cdot 10H_2O$

Phyllosilicate Minerals

Structure	Interlayer Charge	Trioctahedral (Y^{2+})	Dioctahedral (Y^{3+})
O	~0	Brucite	Gibbsite
TO	~0	Serpentine	Kaolinite
TOT	~0	Talc	Pyrophyllite
TOT O TOT	~0	Chlorite	
TOT (X^+, X^{2+}, H_2O) TOT expandable clay	~0.2-0.6	Saponite (smectite)	Montmorillonite (smectite)
	~0.6-0.9	Vermiculite	
TOT (X^+, X^{2+}) TOT non-expandable clay	~0.5-0.75	-	Illite
TOT X^+ TOT true mica	1	Phlogopite, Biotite	Muscovite, Paragonite
TOT X^{2+} TOT brittle mica	2	Clintonite	Margarite

Phyllosilicate Minerals



Clay Minerals

N_2 sorption isotherms

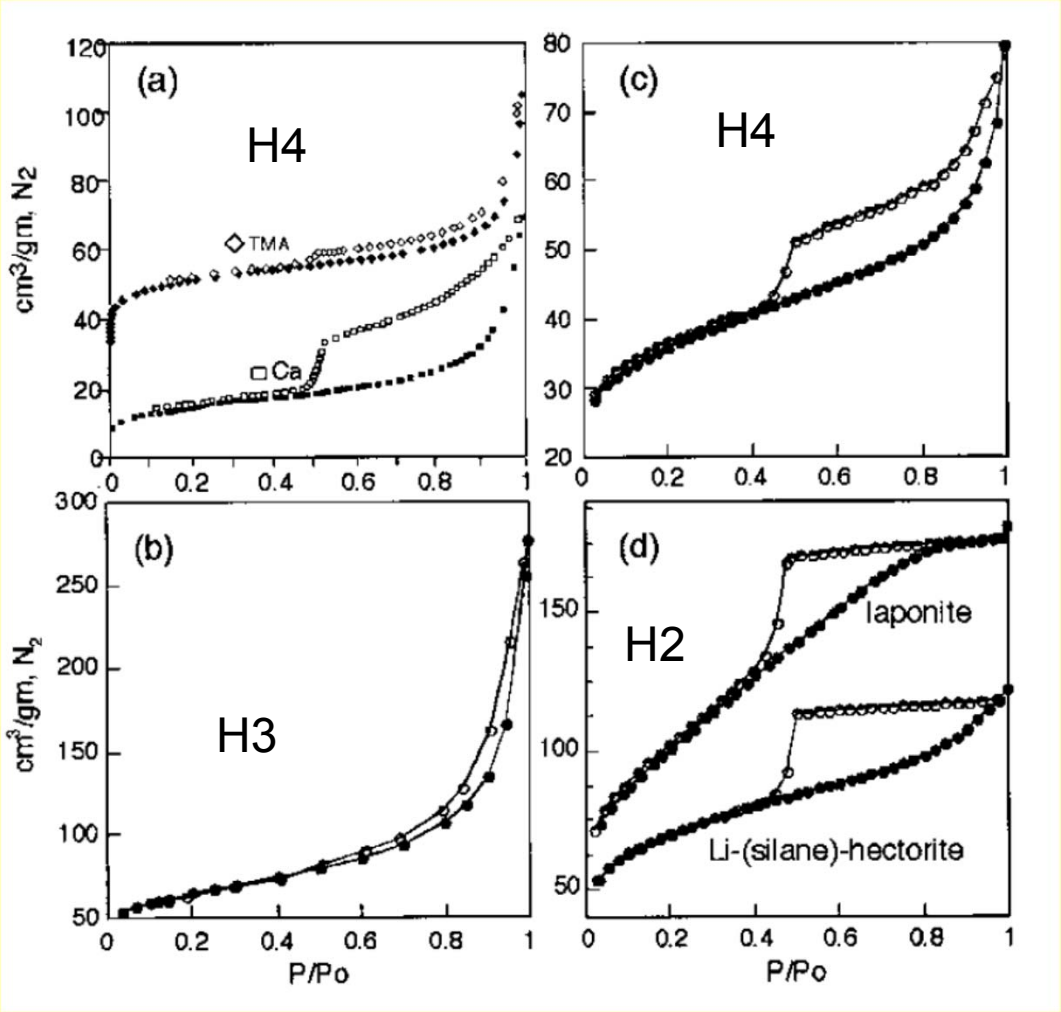
(a) TMA- and Ca-montmorillonite

(b) An Italian sepiolite

(c) Natural SHCa-1 Na-hectorite

(d) synthetic laponite and Li-(silane)-hectorites

Closed symbols = adsorption
Open symbols = desorption



Surface Area

the most important parameters of clays with respect to catalytic applications

TABLE 3 N_2 BET Surface Areas of Various Clay Minerals

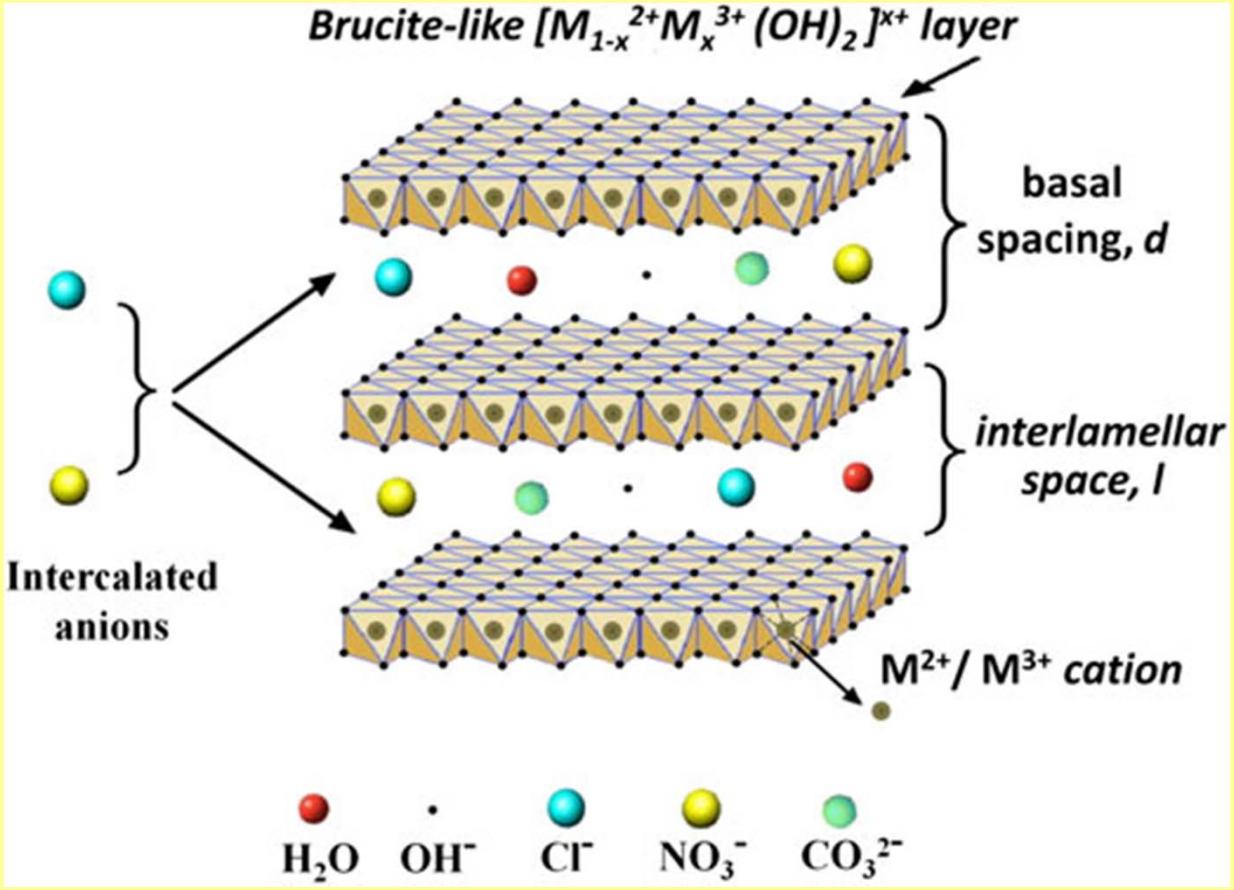
Clay	Outgassing conditions	S. A., m^2/g
Kaolinite[a,b]	200 °C, overnight, $<10^{-2}$ torr	8.75
Na,Ca-montmorillonite[a,c]	same	31.0
Ca-montmorillonite[a,d]	same	80.2
Ca-montmorillonite[a,e]	same	93.9
Na-hectorite[a,f]	same	64.3
Laponite[g]	105 °C, overnight, 10^{-3} torr	360
Sepiolite[h]	96 °C, 3 h	378
Palygorskite[h]	95 °C, <70 h	192

nonpolar guest molecules N_2 do not penetrate the interlayer regions

Na^+ forms of smectites and vermiculites – no penetration
larger ions (Cs^+ and NH_4^+ keep the basal planes far enough) - limited penetration

Layered Double Hydroxides

LDH = layered double hydroxides
HT = hydrotalcites

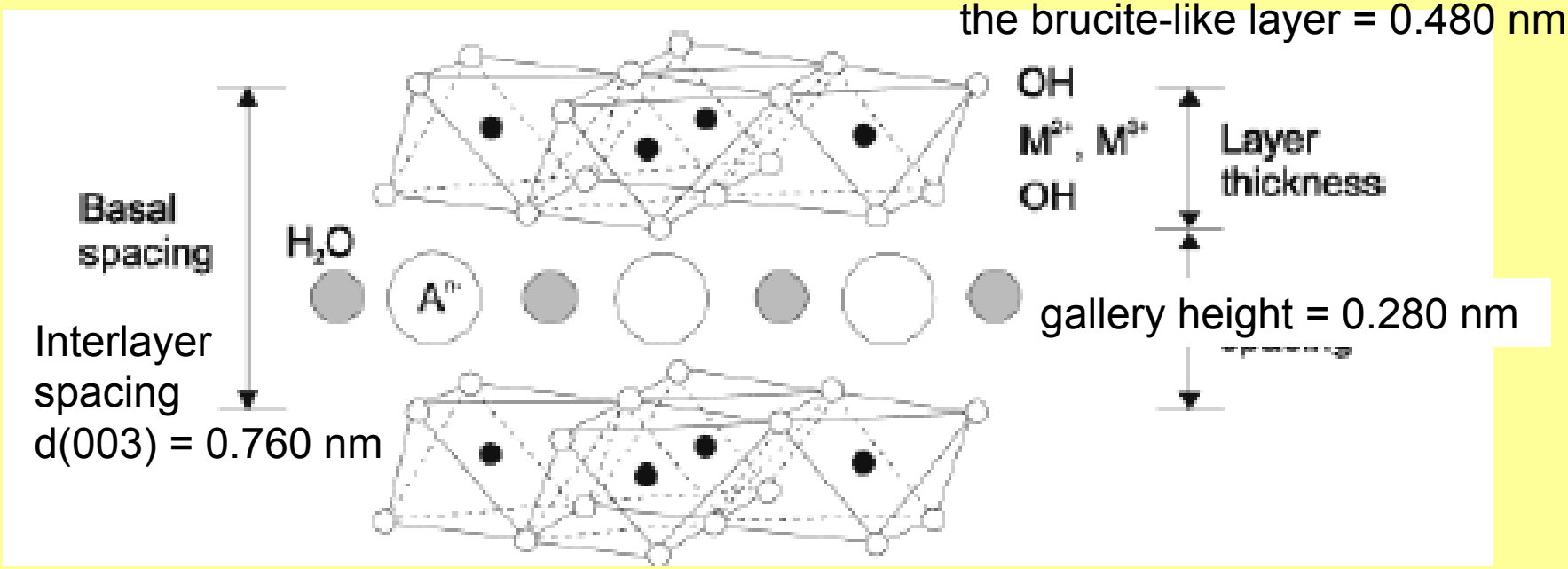


Layered Double Hydroxides

LDH = layered double hydroxides
HT = hydrotalcites
Natural mineral hydrotalcite $Mg_6Al_2(OH)_{16}CO_3.4H_2O$

Brucite layers, Mg^{2+} substituted partially by Al^{3+}

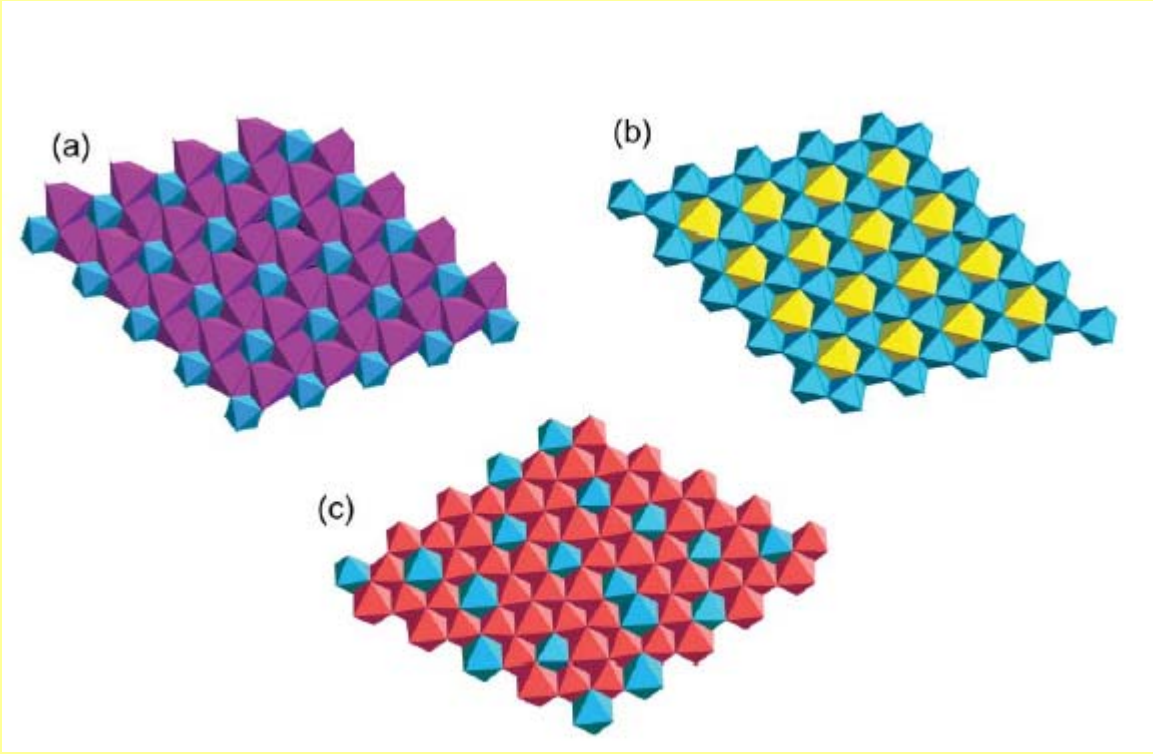
Layers have positive charge



Hydrotalcite $Mg_6Al_2(OH)_{16}CO_3.4H_2O$

Hydrotalcites

Brucite layers, Mg^{2+} substituted partially by Al^{3+}
Layers have positive charge



(a) $[Ca_2Al(OH)_6]_2SO_4.6H_2O$ (b) $[LiAl_2(OH)_6]Cl$ (c) $[Mg_{2.25}Al_{0.75}(OH)_6]OH$

Hydrotalcite

The layered structure of LDH is closely related to brucite $Mg(OH)_2$

a brucite layer, Mg^{2+} ions octahedrally surrounded by six OH^-
the octahedra share edges and form an infinite two-dimensional layer
the brucite-like layers stack on top of one another
either rhombohedral (3R) or hexagonal (2H) sequence

Hydrotalcite $Mg_6Al_2(OH)_{16}CO_3.4H_2O$ - 3R stacking

$$[M^{II}_{1-x}M^{III}_{x}\ (OH)_2]^{x+}(A^{m-})_{x/m}]\cdot nH_2O$$

x = 0.25 $Mg_6Al_2(OH)_{16}CO_3$

x = 0 $Mg(OH)_2$

Hydrotalcite

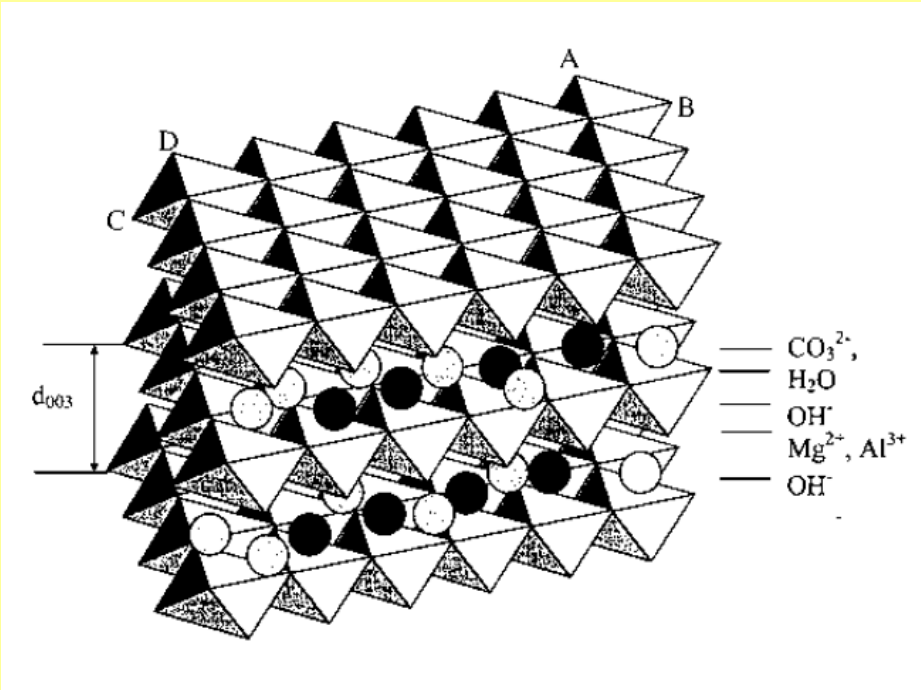
The interlayer spacing *c′* is equal to *d*003, 2*d*006, 3*d*009, etc.;

$c' = (d003 + 2d006 + \ldots + nd00(3n)) / n$

The cell parameter *c* is a multiple of the interlayer spacing *c′*

$c = 3c'$ for rhombohedral (3R)

$c = 2c'$ for hexagonal (2H) sequences



Hydrotalcite

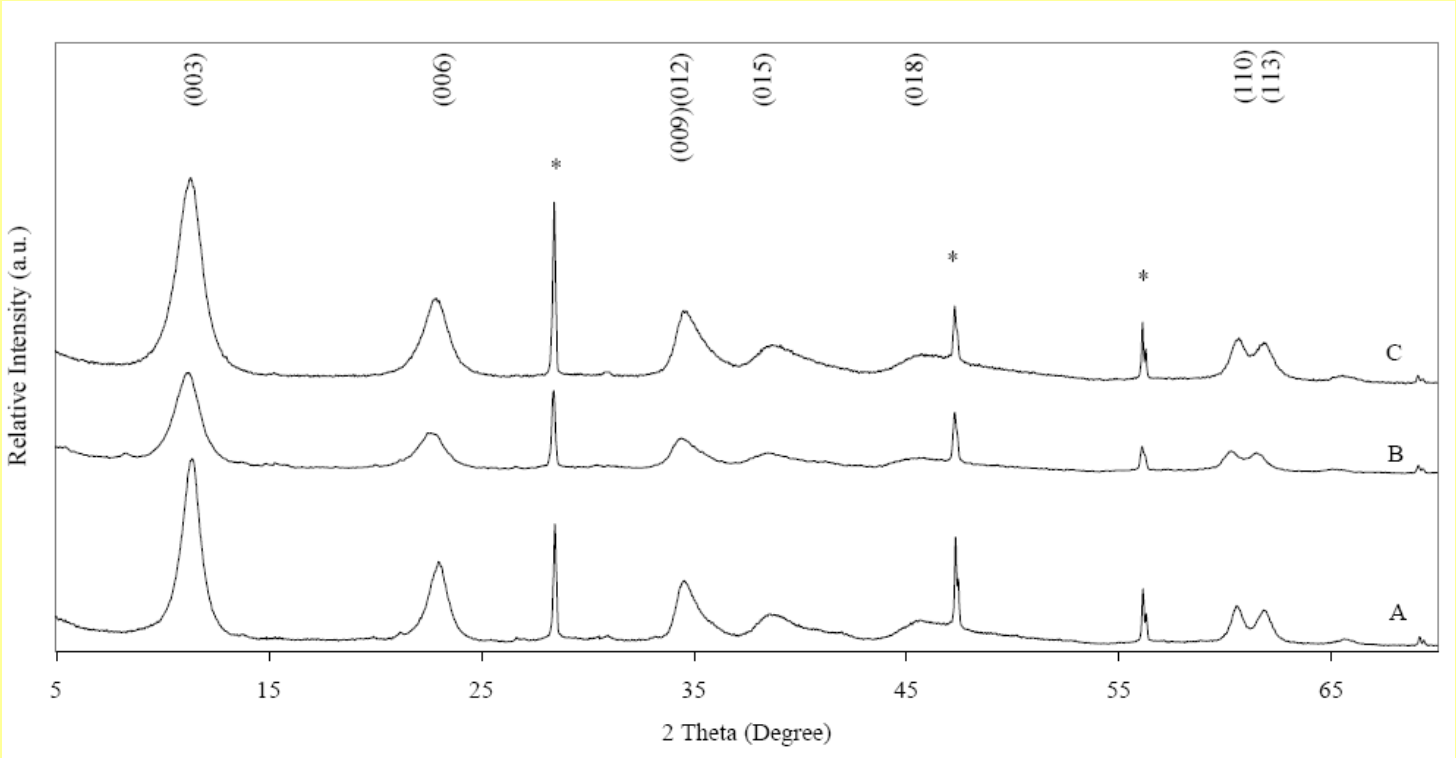
Hydrotalcite $Mg_6Al_2(OH)_{16}CO_3.4H_2O$ - 3R stacking

unit cell parameters

a = 0.305 nm c = 3d(003) = 2.281 nm

the interlayer spacing: d(003) = 0.760 nm
the spacing occupied by the anion (gallery height) = 0.280 nm
a thickness of the brucite-like layer = 0.480 nm
the average M—O bond = 0.203 nm
the distance between two nearest OH^- ions in the two opposite side layers = 0.267 nm shorter than a (0.305 nm) and indicative of some contraction
along the c-axis.

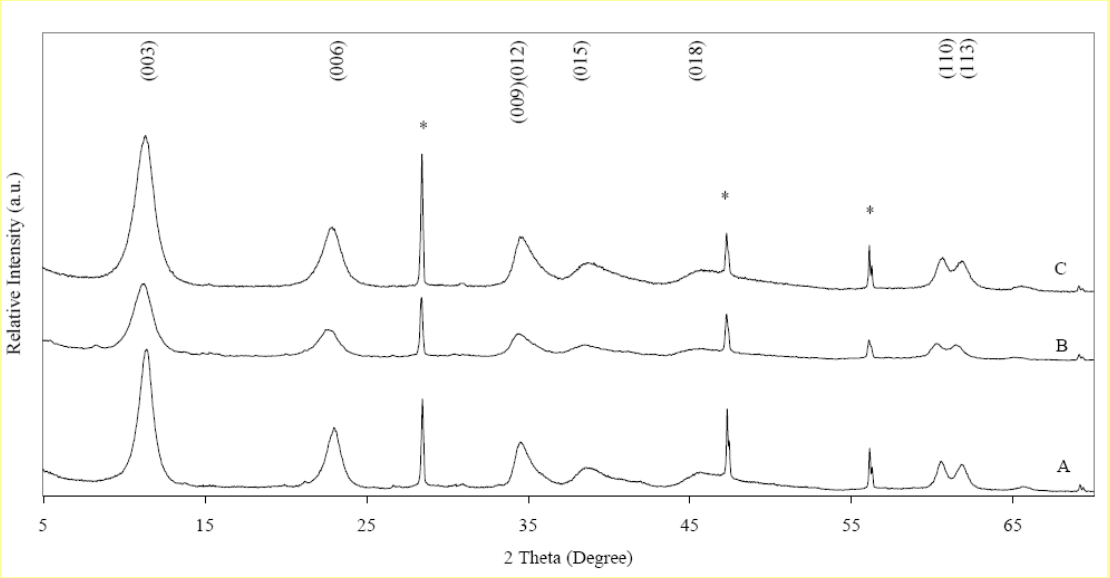
XRD Patterns of LDH



XRD patterns of layered double hydroxides synthesized by coprecipitation method with various cations composition:
A – Mg/Al; B- Mg/Co/Al; C- Mg/Ni/Al

*** = Reflections from Si crystal used as a reference**

XRD Patterns of LDH



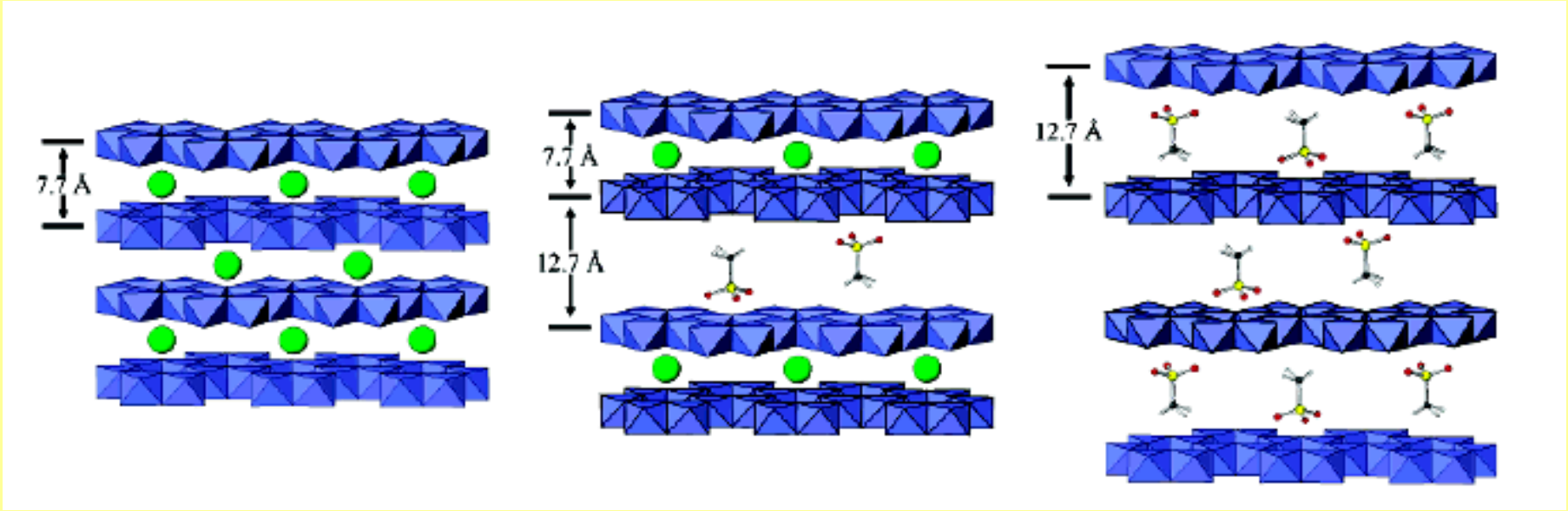
rhombohedral structure
the cell parameters *c* and *a*

The lattice parameter *a* = *2d*(110) corresponds to an average cation–cation
distance

The *c* parameter corresponds to three times the thickness of *d*003

***c* = 3/2 [*d*003+2*d*006]**

Intercalation to LDH



the intercalation of methylphosphonic acid into Li/Al LDH

(a) $[LiAl_2(OH)_6]Cl.H_2O$
(b) second-stage intermediate, alternate layers occupied by Cl and MPA anions
(c) first-stage product with all interlayer regions occupied by MPA.

Intercalation to LDH

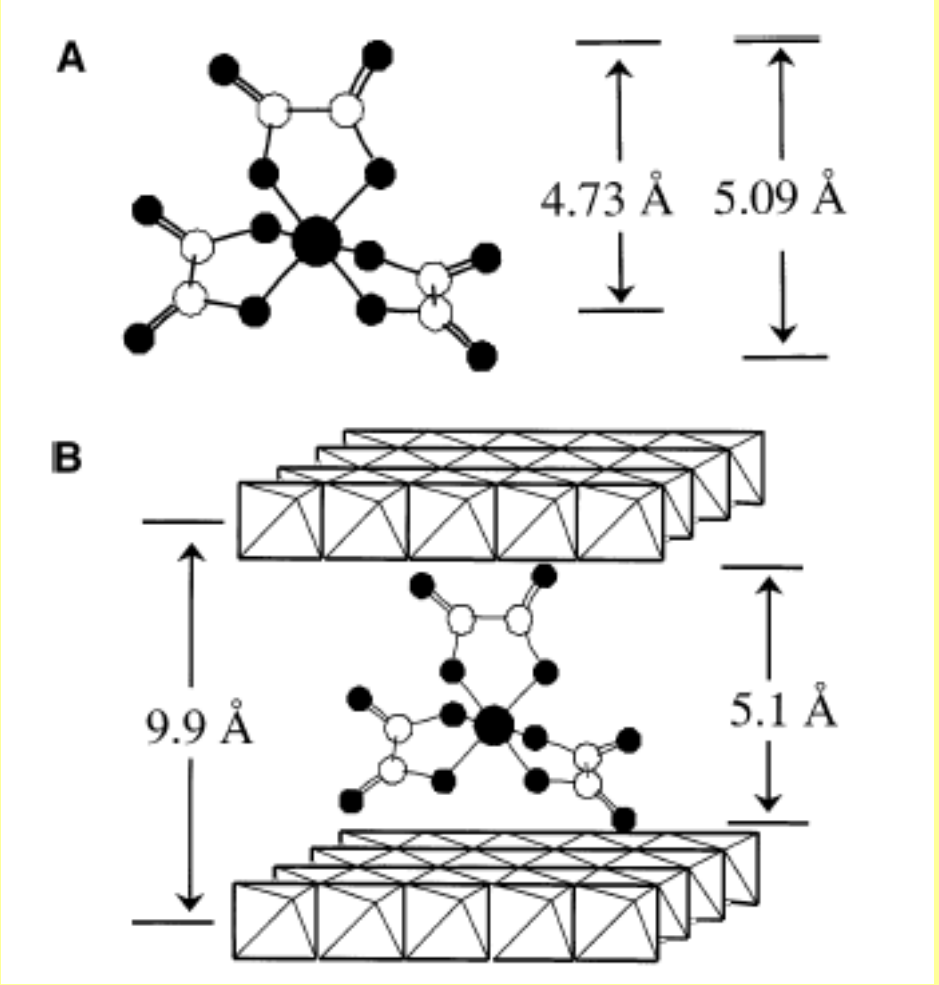
LDH = layered double hydroxides

hydrotalcites
mineral $Mg_6Al_2(OH)_{16}CO_3.4H_2O$

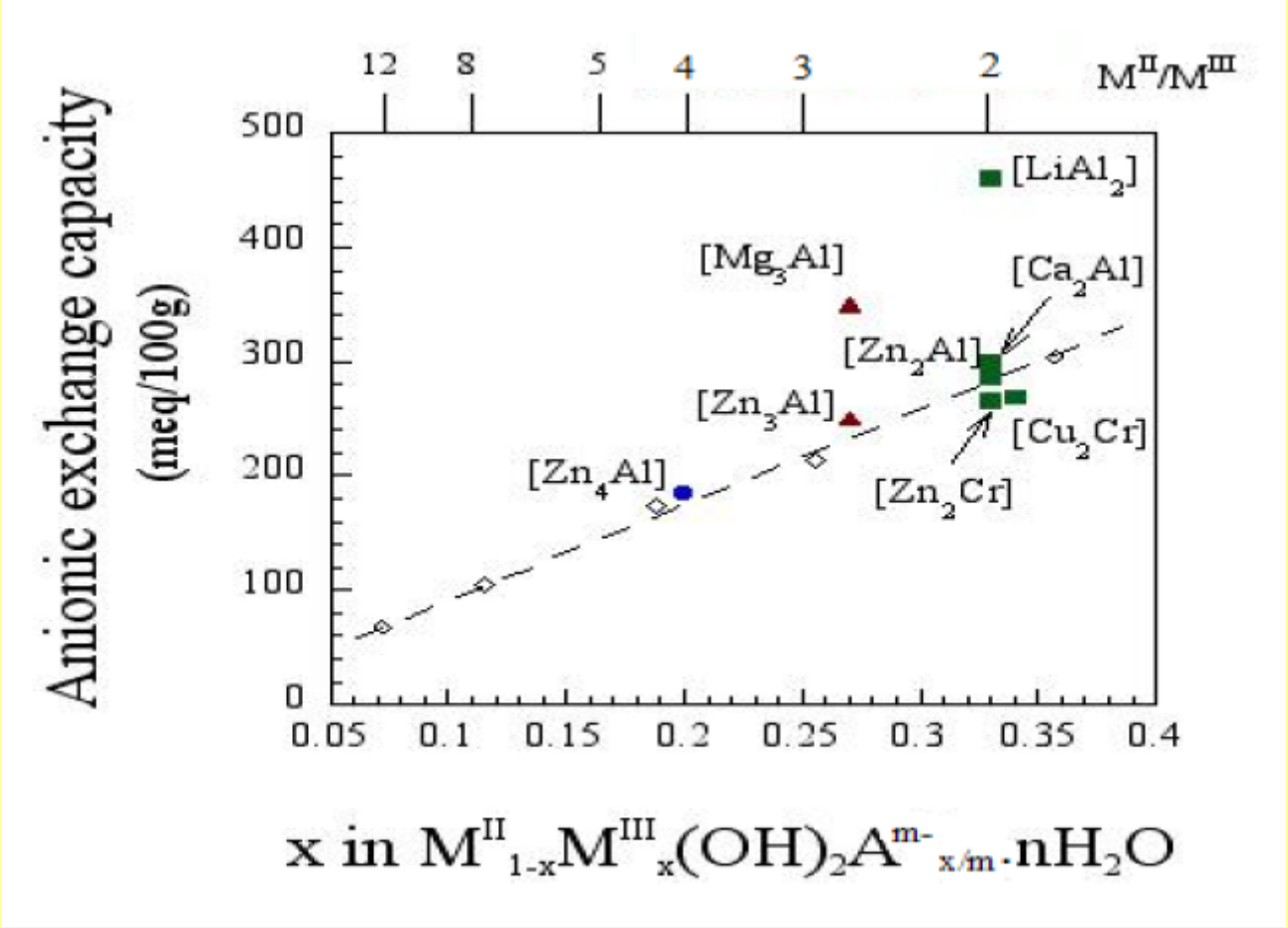
Brucite layers, Mg^{2+} substituted partially by Al^{3+}

Layers have positive charge

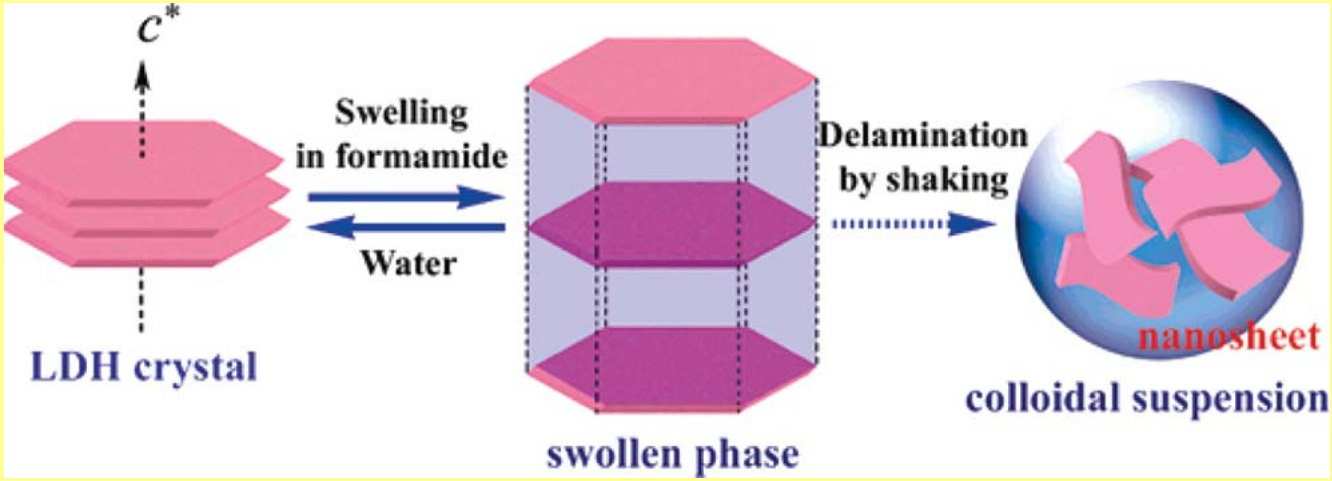
Intercalate anions $[Cr(C_2O_4)_3]^{3-}$

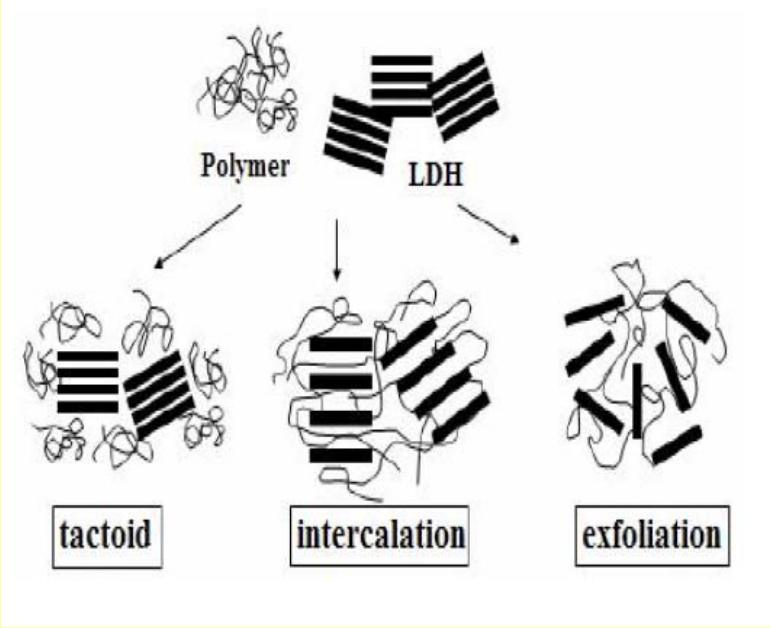


The Anionic Exchange Capacity (AEC)

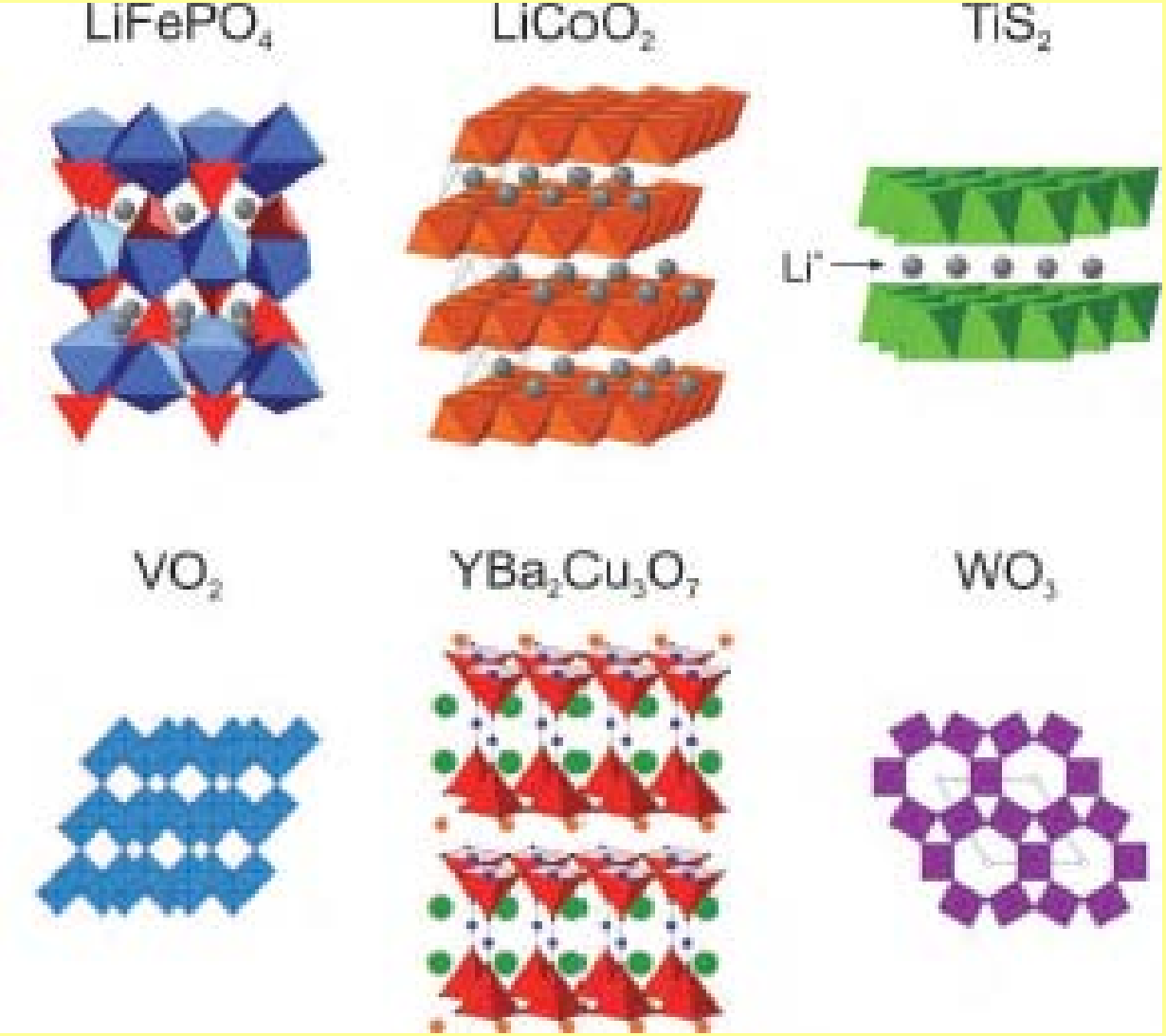


LDH Composite Structures



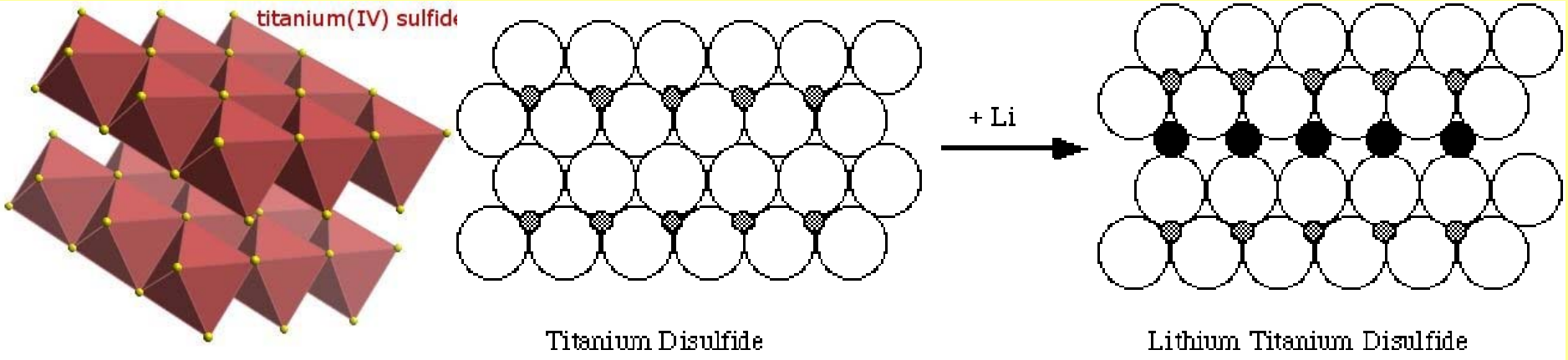


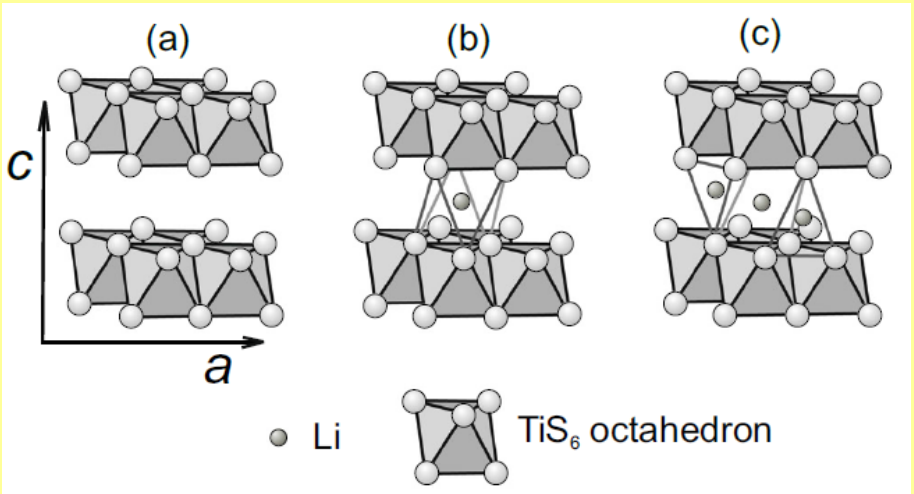
Li Intercalation Compounds

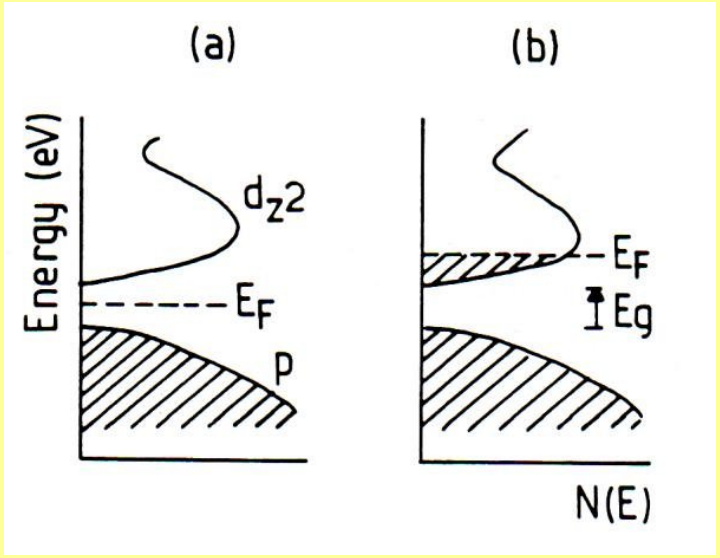


Li Intercalation

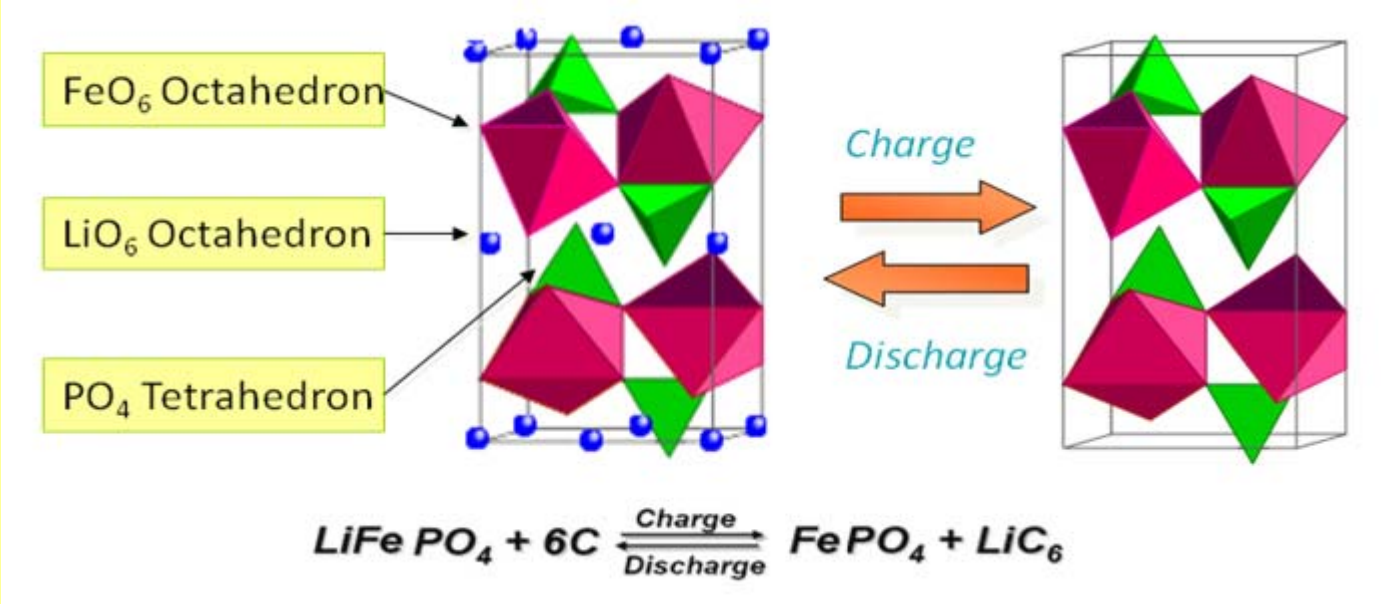
$$x\,Li + TiS_2 \rightarrow Li_xTiS_2$$





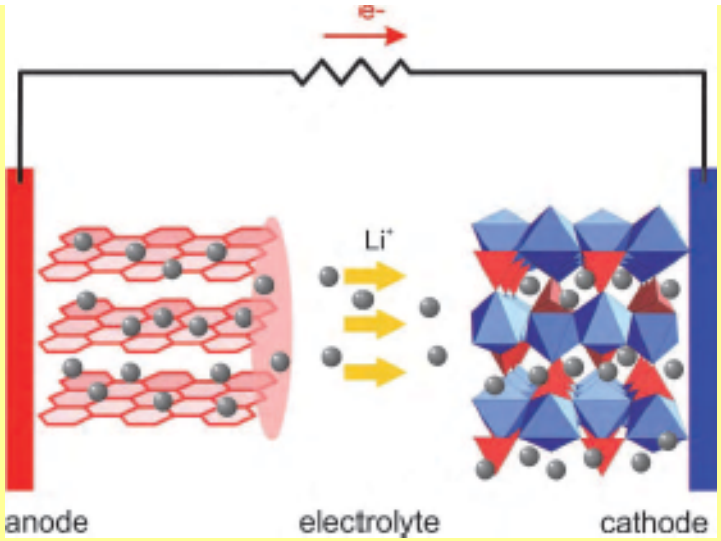


Li Intercalation



$Li/C \rightarrow e + Li^+ + C$

$Li^+ + e + FePO_4 \rightarrow LiFePO_4$

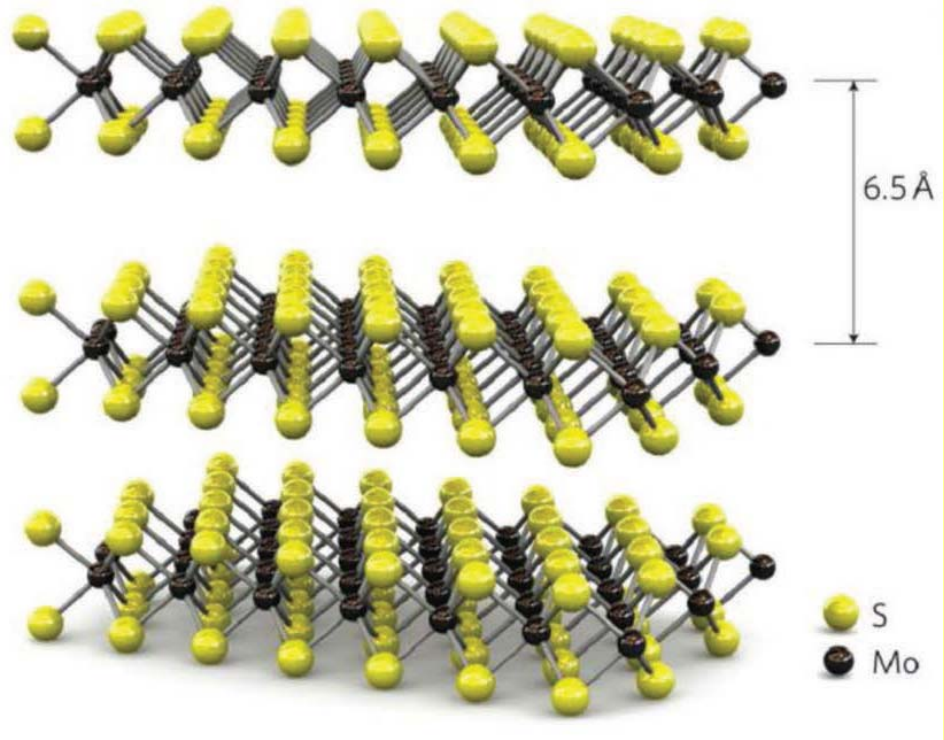


Molybdenum Disulfide (MoS_2)

Mineral molybdenite

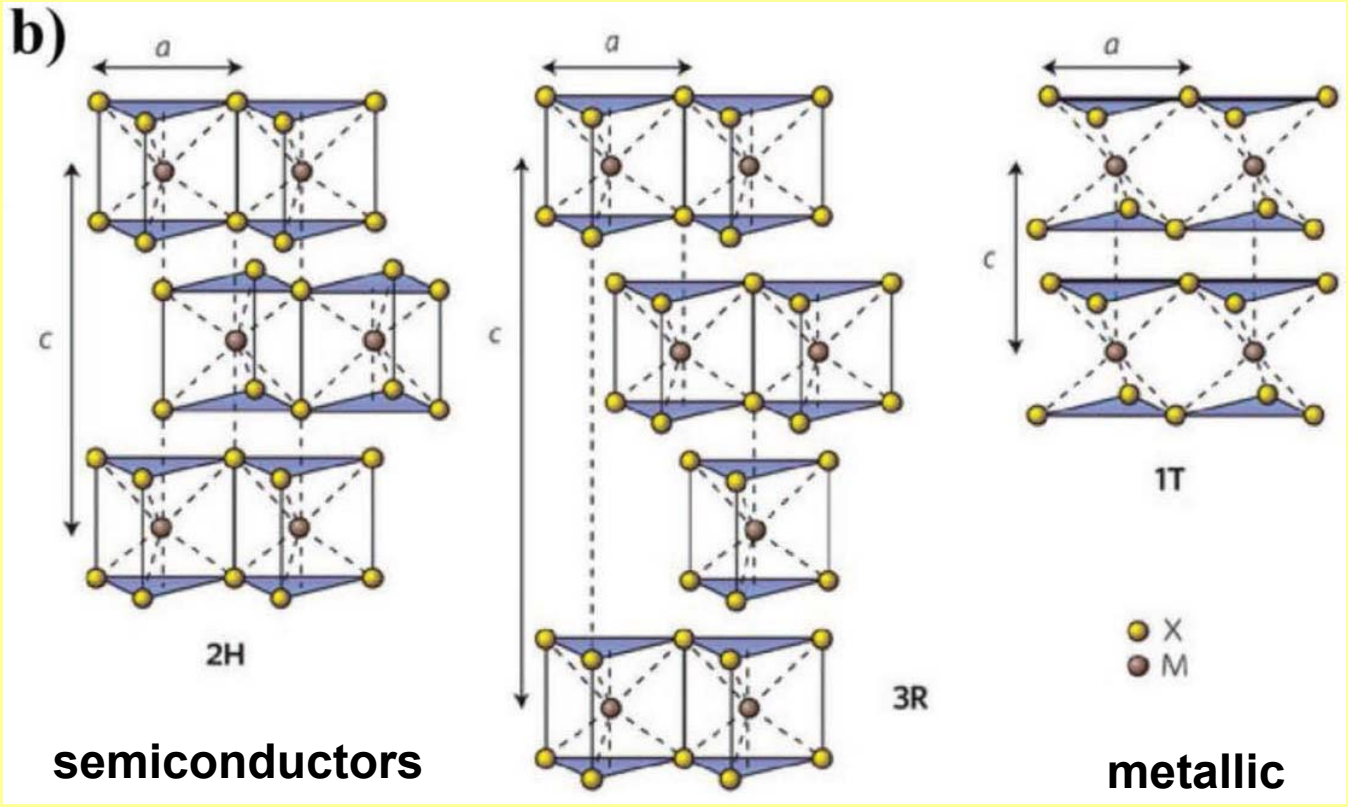
Hydrodesulfurization catalyst at edges

Lubricant

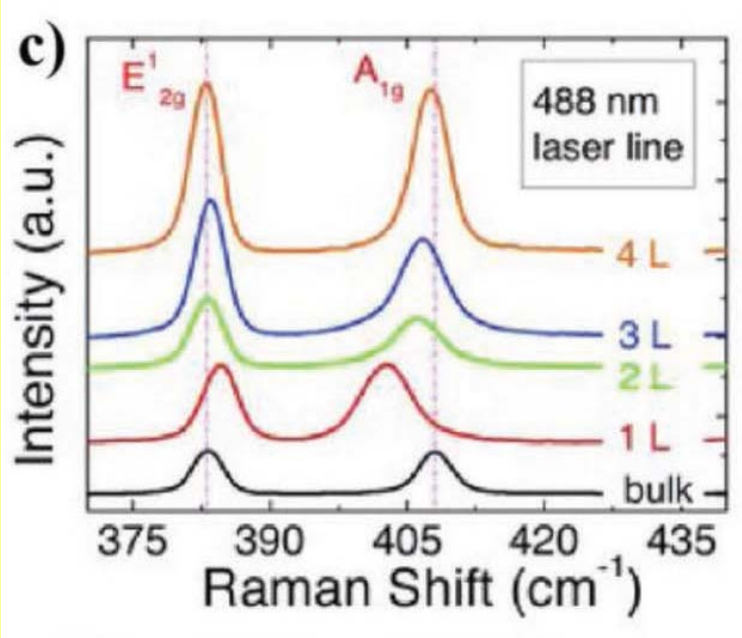


Molybdenum Disulfide (MoS_2)

Polymorphs of MoS_2

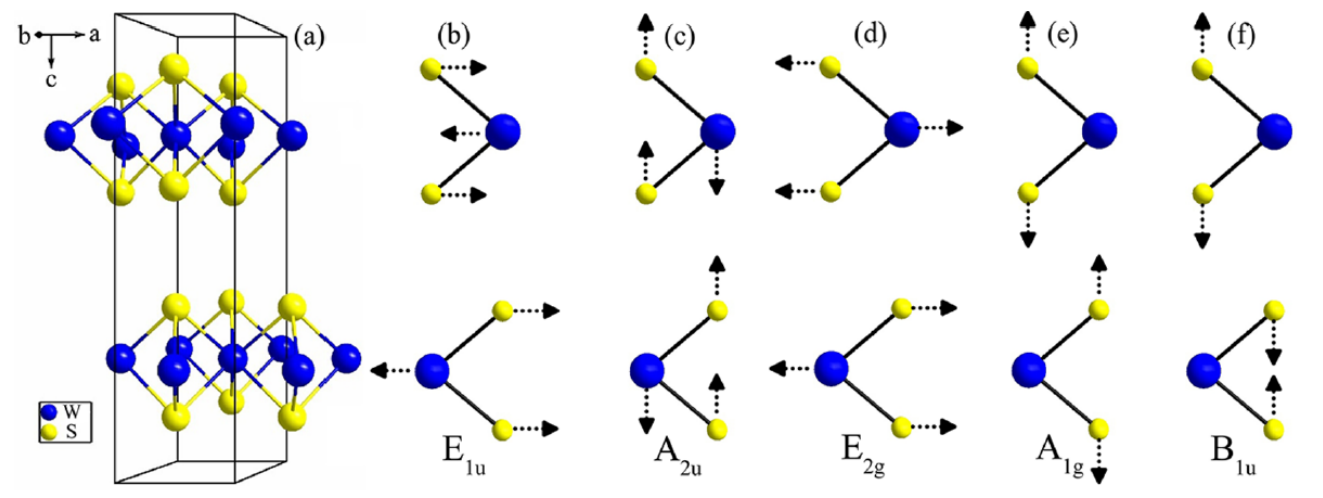


Molybdenum Disulfide (MoS_2)

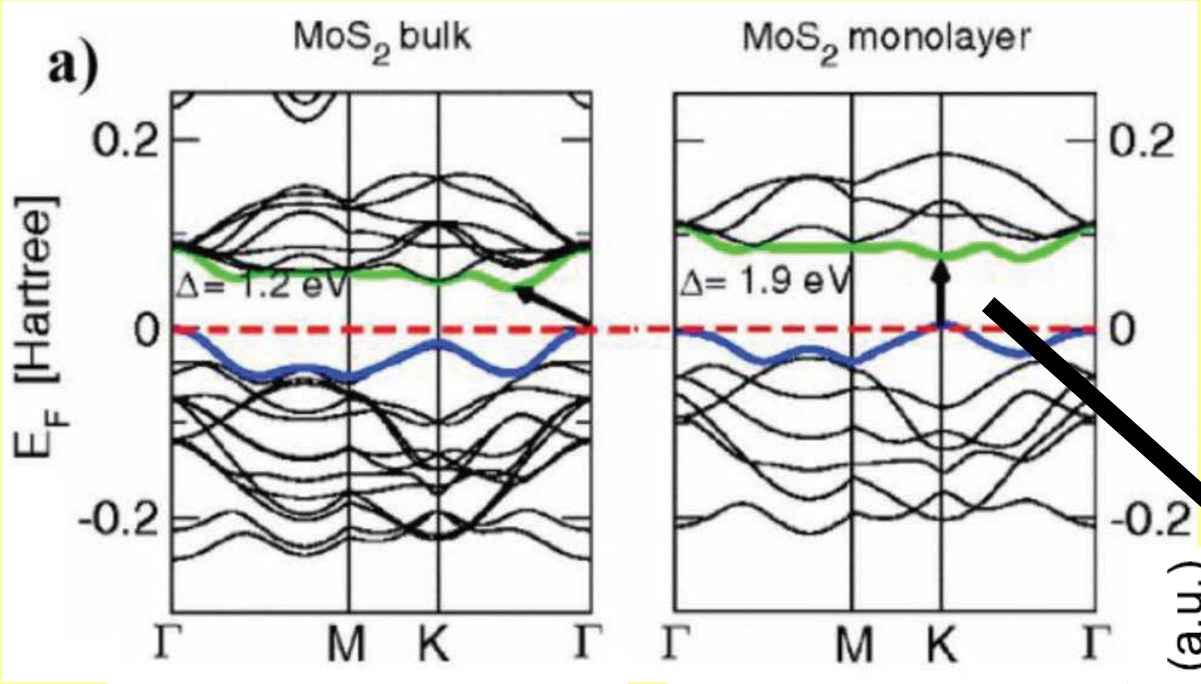


Frequency of A_{1g} band is increasing while that of E^1_{2g} is decreasing with increase in number of layers

(b,c) infraredand (d−f) Raman-active



Molybdenum Disulfide (MoS_2)



An indirect band gap
1.2 eV

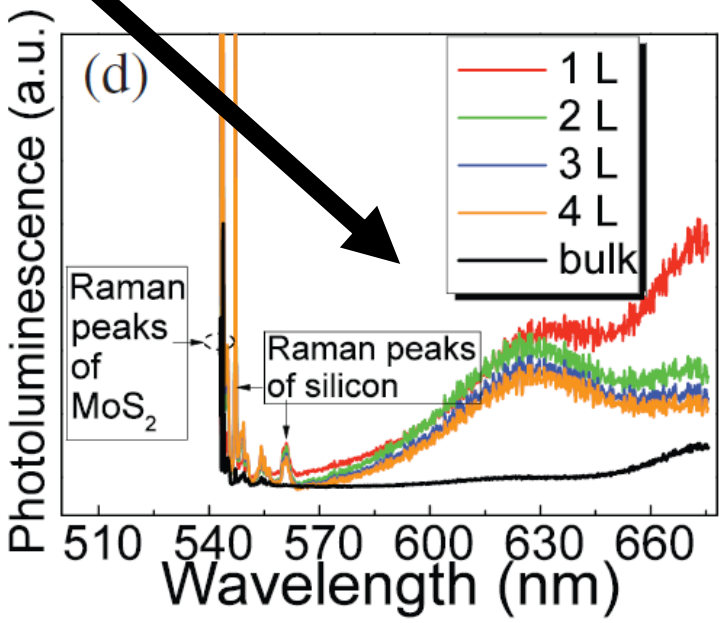
A direct band gap
1.8 eV

Conduction band

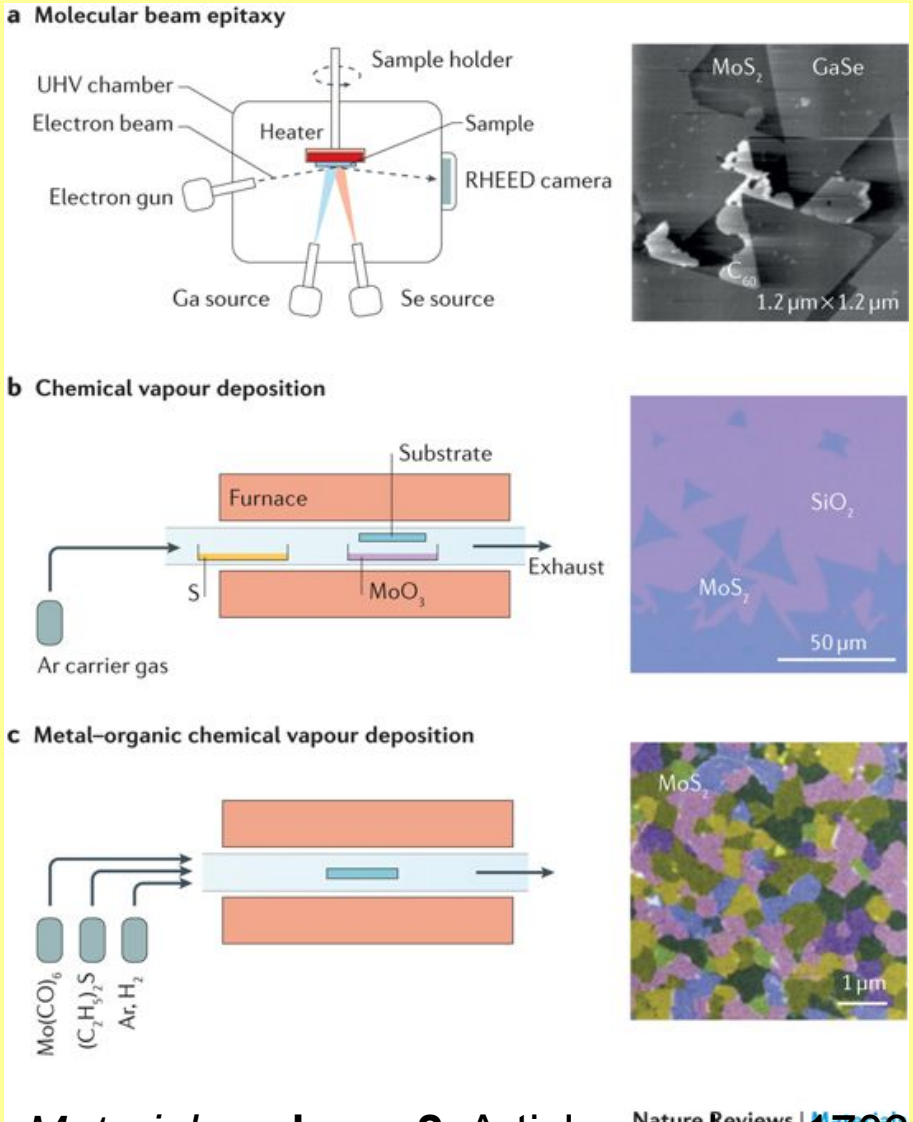
Fermi level

Valence band

photoluminescence



Molybdenum Disulfide (MoS_2)



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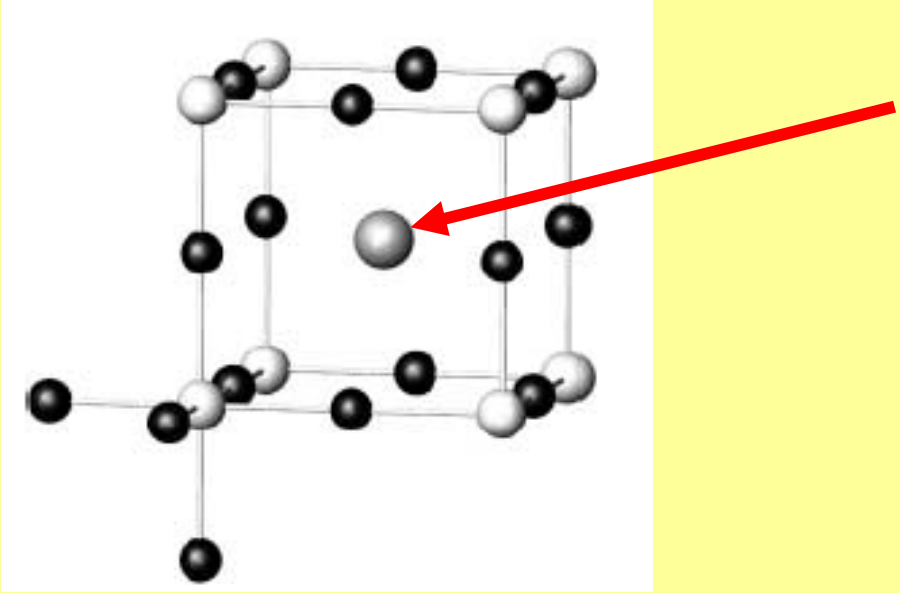
3D Intercalation Compounds

Cu_3N and Mn_3N crystallize in the (anti-) ReO_3-type structure



the large cuboctahedral void in the structure can be filled

By Pd to yield (anti-) perovskite-type $PdCu_3N$

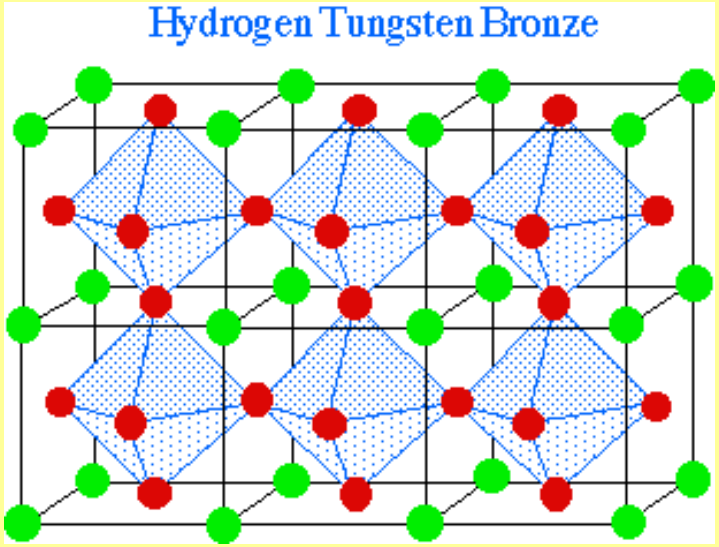
By M = Ga, Ag, Cu leading to MMn_3N

3D Intercalation Compounds

Tungsten trioxide structure
= WO_6 octahedra joined at their corners
= the perovskite structure of $CaTiO_3$ with all the calcium sites vacant

$$Zn + 2\ HCl \rightarrow 2\ H + ZnCl_2$$

$$WO_3 + x\ H \rightarrow H_xWO_3$$



The color and conductivity changes are due to the intercalation of protons into the cavities in the WO_3 structure, and the donation of their electrons to the conduction band of the WO_3 matrix. The material behaves like a metal, with both its conductivity and color being derived from free electron behavior.
The coloration reaction used in electrochromic displays for sun glasses, rear view mirrors in cars

0D Intercalation Compounds

C_{60} = FCC

$K_3 C_{60}$