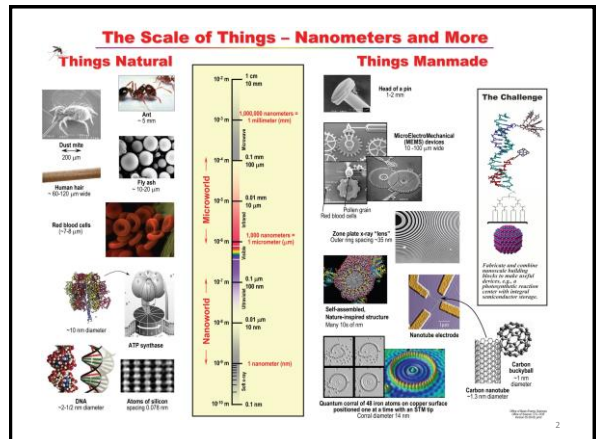


Atomic and molecular interactions; applications

Tamás Bozó
30 October 2014

Damjanovich, Fidy, Szöllősi: *Medical Biophysics*, pp. 44-51; 579-583
Lab Manual, Chapter 18.c., Resonance

1



2

Atomic interactions

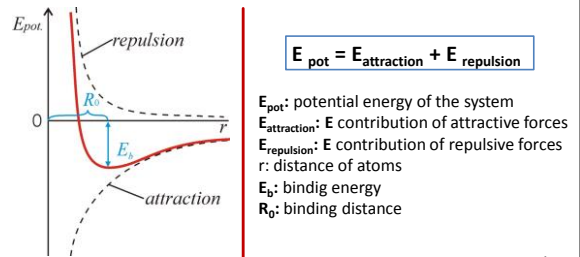
Noble gases: found in their atomic form in nature (*He, Ne, Ar, Kr, Xe, Rn*)

Other elements: form molecules held together by chemical bonds. (e.g. H_2 , HCl , H_2O , ...) Each atom has a (more or less) fixed position in the molecule.

3

Atomic interactions

Bringing two atoms (ions/molecules/...) close to each other attractive and repulsive forces emerge between them resulting in a net potential energy function with an energy minimum at a certain distance:



4

Bond Types

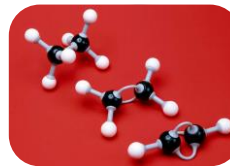
Classification systems:

- Intra/intermolecular bonds
- According to the strength of bonds:
 - Strong/weak
 - Primary (covalent, metallic, ionic)
 - Secondary (dispersion, dipole-dipole, hydrogen)
- Other Bond types...



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I. Covalent bond



| | R_0 | E_b |
|----------------------------|----------|---------------------------|
| H_3C-CH_3 ethane | 154 pm | -331 kJ·mol ⁻¹ |
| $H_2C=CH_2$ ethylene | 139,9 pm | -590 kJ·mol ⁻¹ |
| $HC\equiv CH$ acetylene | 120,3 pm | -812 kJ·mol ⁻¹ |

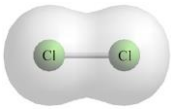
- Atoms are held together by electrons of common orbitals, called molecular orbitals.
- Valence electrons interact the most.
- Single, double or triple bond depending on the number of shared electrons.
- Some electrostatic component may be present (see later), but not predominant.
- Strong interaction: $E_k > 1 \text{ eV}$
- $(1 \text{ eV} = 1,6 \cdot 10^{-19} \text{ J} = 96 \text{ kJ/mol} \sim 100 \text{ kJ/mol})$

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I. Covalent bond

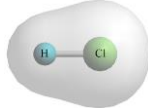
Apolar (nonpolar or homeopolar) bond

Center of (+) és (-) charges of the molecule coincides
Binding electron(s) shared equally
„purely covalent“
E.g. H_2 , Cl_2 , O_2 , ...



Polar (heteropolar) bond

Centers of (+) and (-) charges do not coincide
Electrostatic contribution appears
Electron cloud is polarized
Electric dipole is formed
E.g.: HCl , HF , H_2O , ...



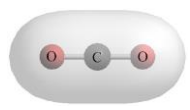
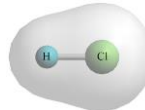
7

I. Covalent bond

Electric dipole moment: measure of the separation of positive and negative electric charges.

$$p = Qd$$

p : electric dipole moment
 Q : charge (positive and negative)
 d : distance of separation
[D, debye] ($1D = 3,34 \cdot 10^{-30} \text{ Cm}$)



8

I./b Metallic bond

- Atoms are held together by common electrons (delocalized orbital)
- Valence electrons form continuous electron band
- Strong interaction: $E_k > 1 \text{ eV}$
- Multi atomic systems

Periodic Table of the Elements

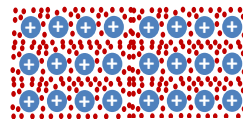
| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
|---|---|---|---|---|---|---|---|---|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 | 21 | 22 | 23 | 24 | 25 | 26 | 27 | 28 | 29 | 30 | 31 | 32 | 33 | 34 | 35 | 36 | 37 | 38 | 39 | 40 | 41 | 42 | 43 | 44 | 45 | 46 | 47 | 48 | 49 | 50 | 51 | 52 | 53 | 54 | 55 | 56 | 57 | 58 | 59 | 60 | 61 | 62 | 63 | 64 | 65 | 66 | 67 | 68 | 69 | 70 | 71 | 72 | 73 | 74 | 75 | 76 | 77 | 78 | 79 | 80 | 81 | 82 | 83 | 84 | 85 | 86 | 87 | 88 | 89 | 90 | 91 | 92 | 93 | 94 | 95 | 96 | 97 | 98 | 99 | 100 | 101 | 102 | 103 | 104 | 105 | 106 | 107 | 108 | 109 | 110 | 111 | 112 | 113 | 114 | 115 | 116 | 117 | 118 |
|---|---|---|---|---|---|---|---|---|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|

Legend:

- hydrogen
- alkali metals
- alkali earth metals
- transition metals
- poor metals
- nonmetals
- noble gases
- rare earth metals

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I./b Metallic bond

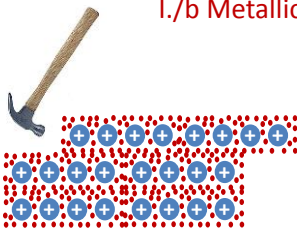


Metallic lattice: positively charged metal ions in a crystall lattice surrounded by a cloud of delocalized electrons.



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I./b Metallic bond



Physical properties:

Metals are:

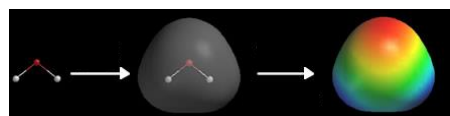
- Non transparent
- Ductile (shapeable)
- Good electric conductors
- etc.

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II. Bond types involving electrostatic interactions

Electronegativity

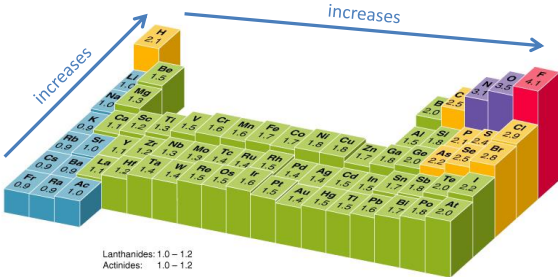
- is a chemical property that describes the tendency of an atom or a functional group to attract electrons towards itself.
- The higher the associated electronegativity number, the more an element or compound attracts electrons towards it.
- Approximately proportional to the sum of ionization energy and electron affinity.
- Calculated with different methods (*Pauling*, *Mulliken*, *Sanderson...*)



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II. Bond types involving electrostatic interactions

Electronegativity according to L. Pauling (dimensionless units)

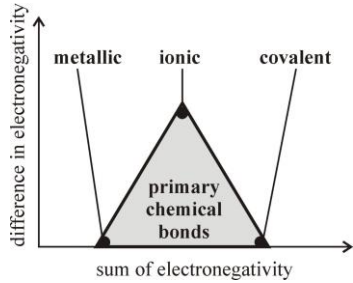


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II. Bond types involving electrostatic interactions

Classic bonds classified according to electronegativity:

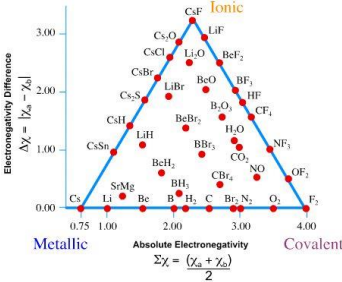
ΔEN : $< 0,6$ (apolar covalent) $0,6 - 2,1$ (polar covalent) $2,1 <$ (ionic)



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II. Bond types involving electrostatic interactions

Classic bonds classified according to electronegativity: an example



(This model utilizes Norman (and not Pauling) EN values.)

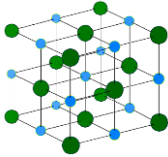
15

II./a Ionic bond

- Atoms are held together by Coulombic forces between (+) és (-) point charges
- „Limiting case of heteropolar bonds”
- Formed between atoms of significantly different electronegativity (e.g.: NaCl , $\Delta EN = 3,0 - 0,9 = 2,1$)
- Can form between two atoms, but ions are usually multi-atom systems.
- Long range interaction - attraction is inversely proportional to the distance (decreases slowly with it).
- Electrostatic interaction can be largely affected by other charged components (eg. dissociation in water!)
- Strong interaction ($E_b > 1 \text{ eV}$)

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II./a Ionic bond



Ionic crystals: stoichiometric ratio of positive and negative ions are structured into a periodic crystalline structure. (e.g.: NaCl)



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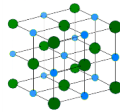
II./a Ionic bond



$$E_i = 496 \text{ kJ/mol}$$



$$E_{\text{aff}} = -349 \text{ kJ/mol}$$



$$E_{\text{lattice}} = -788 \text{ kJ/mol}$$

Ionization energy: is the energy required to remove electrons from gaseous atoms or ions.

Electron affinity: amount of energy released when an electron is added to a neutral atom or molecule to form a negative ion (measured in the gaseous state).

Lattice energy: measure of the strength of bonds in an ionic compound. Energy required to completely separate one mole of a solid ionic compound into gaseous ionic constituents.

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II./b Dipole-dipole interaction

- Constant charge distribution is present in a (given part of a) molecule
- Partially (+) and (-) segments are held together by electrostatic interactions (Coulombic forces)
- Intra/intermolecular interaction.
- Weak interaction ($E_b = 0,003-0,02$ eV)
- Energy of attracting interaction between dipoles:

$$E_{\text{attraction}} = p E$$

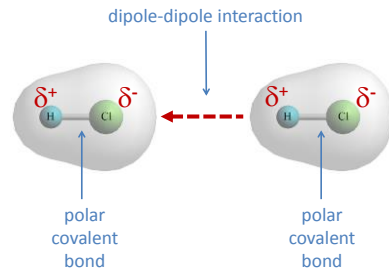
p : dipole momentum

E : electric field strength generated by the surrounding partners

($E_{\text{repulsion}}$: stems from the repulsion of the participant's electron cloud)

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II./b Dipole-dipole interaction



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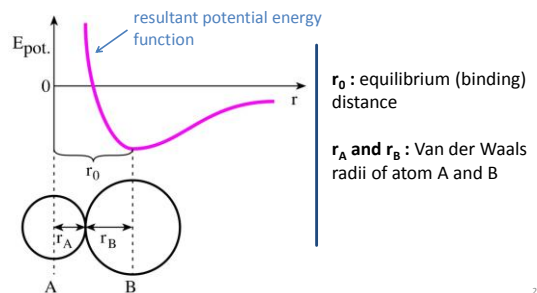
III. Van der Waals-interactions

- Sum of attractive and repulsive interactions between two apolar atoms, molecules or apolar molecular parts.
- The **attractive contribution** (also called *London-*, or *dispersion force*) is a result of temporarily created dipoles that can induce the polarization of other apolar molecule or molecular part.
- Intermolecular or intramolecular interaction.
- Important biological role: formation of organic structures.
- Weak connection ($E_b \sim 0,02$ eV)
- [according to other classifications Van der Waals interactions involve all types of weak electrostatic connections (permanent dipole-permanent dipole; permanent dipole-induced dipole, induced dipole-induced-dipole)]

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III. Van der Waals-interactions

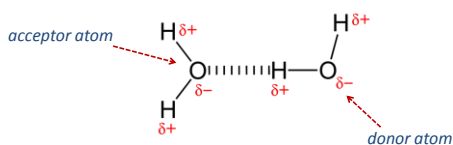
Equilibrium distance of a Van der Waals interaction can be considered as one definition of atomic size.



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IV. Hydrogen bond

- Two atoms of high electronegativity are held together with a Hydrogen-bridge.
- Primarily between **F, N, O** atoms (pillar atoms).
- Intermolecular/intramolecular interaction
- Typical bond distance: 0,23 – 0,35 nm
- Well defined geometry.
- Important role in structural biology and biochemical reactions.
- Medium strong interaction (typical $E_b = 0,2$ eV)



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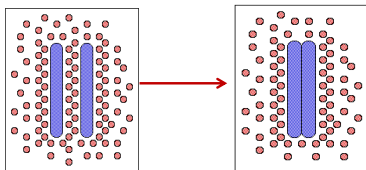
V. Hydrophobic interaction

- Appears in aqueous systems (like biological environments!)
- Apolar, hydrophobic molecules tend to aggregate (form bonds with each other) in aqueous environment in order to exclude the polar water molecules and minimize their surface area exposed to water.
- Intra/intermolecular interaction
- It has mostly entropic origin (see later at Thermodynamics) through reduction of highly structured water cage around the apolar surfaces. (see Organisation of Water later)
- Important role in structural biology and biochemical reactions.
- Weak interaction.



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V. Hydrophobic interaction



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Scanning Probe Microscopy (SPM)

Family of instruments used for studying surface properties of various materials.

How do they work?

Etimology and function:

Microscopy: a method being able to form image of small objects. How small? Size of resolvable objects spans from few pm-s to several μm -s.

They are not „scope“-s in the classic sense of the word: They do not „see“ the object, they „touch“ it.

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Scanning Probe Microscopy (SPM)

Probe:

- A tiny, specifically designed component particularly sensitive to *atomic interactions*.
- The probe is brought very closed to the sample surface.
- The sensed interactions can be correlated with the distance between the probe and the sample.
- Various interactions can be observed depending on the design of the probe.
- SPM methods are named after the type of atomic interaction sensed by a certain probe.

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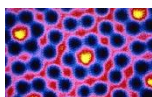
Scanning Probe Microscopy (SPM)

Scanning:

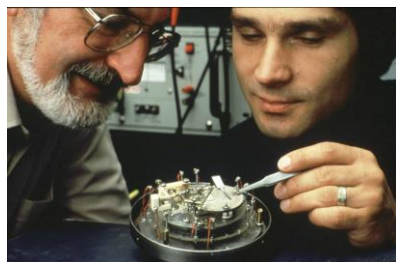
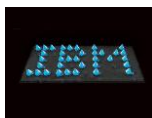
- A scanner controls the precise *position* (X; Y) of the probe and its *distance* (Z) from the surface to be imaged.
- The probe can be moved with pm sensitivity in X; Y; Z directions.
- The surface of region of interest (ROI) is scanned point by point during a measurement.
- (The material that enables such precise positioning is *piezoelectric ceramic*. If voltage is applied on it, the ceramic changes its geometry.)

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Scanning Tunneling Microscope (STM) 1981



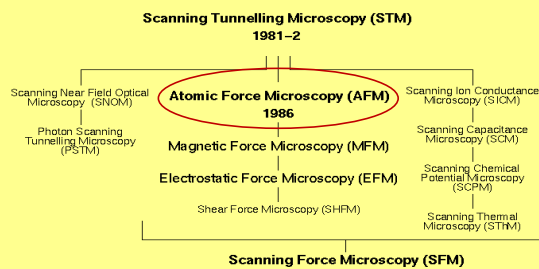
Atoms in a silicon chip



Heinrich ROHRER and Gerd BINNING
Nobel prize: 1986

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Scanning Probe Microscopy "Family Tree" (SPM)



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Atomic Force Microscopy (AFM)

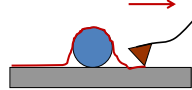
- The probe is a tiny, sharp tip, usually few μm -s tall, and only few nm-s in diameter at the apex.
- The tip is located at the free end of a cantilever that is usually 100-500 μm long.
- Topographic image is collected with ~ 10 pm vertical and somewhat worse horizontal resolution.
- Any surfaces (conductors, insulators and semiconductors) can be imaged.
- Works in air and in fluid environment as well.
- Usually does not require fixation or staining of the sample.
- Biological samples can be examined in their native state and physiological environment.

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Atomic Force Microscopy

Contact mode:

- The tip is in perpetual contact with the sample surface.
- The deflection of the cantilever, i.e. the force exerted on the sample by the tip is held constant.
- A Z feedback system is utilized to maintain the deflection at a constant value (setpoint) by lifting or lowering the cantilever.
- Topography data (i.e.: height) in each X;Y point is calculated from these Z movements



Disadvantage:

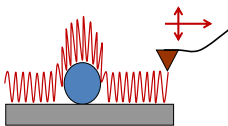
- Lateral forces exerted by the scanning tip may damage softer samples.

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Atomic Force Microscopy

Oscillating mode: (Tapping mode, Non-contact mode)

- Cantilever is oscillated close to its resonant frequency.
- The tip taps the surface gently
- The amplitude of cantilever oscillation changes with surface topography.
- A Z feedback system is utilized to maintain the amplitude at a constant value (setpoint) by lifting or lowering the oscillating cantilever.
- Topography data (i.e.: height) in each X;Y point is calculated from these Z movements

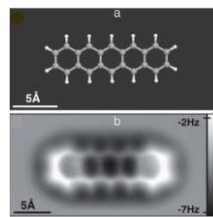


Advantage:

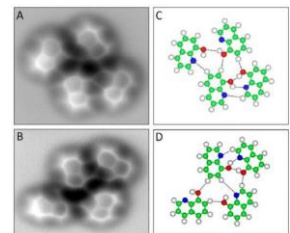
- Virtually eliminated lateral forces.
- Allows more gentle imaging.
- Applicable for soft samples.

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Scanning Probe Microscopy (SPM)



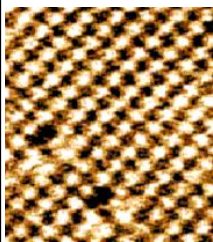
Pentacene molecule
imaged with AFM



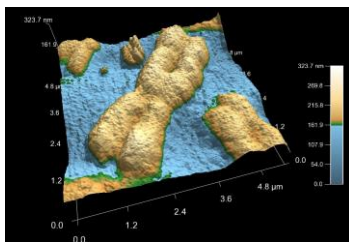
Hydrogen bonds between 8-
hydroxyquinoline molecules scanned
with AFM

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Atomic Force Microscopy Images



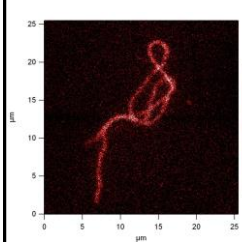
NaCl crystal surface



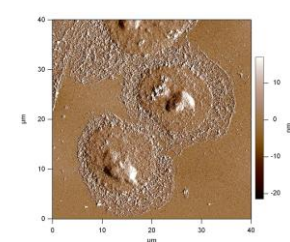
Human metaphase chromosomes

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Atomic Force Microscopy Images



„The thinker“
a single actin polymer



HeLa cells on glass

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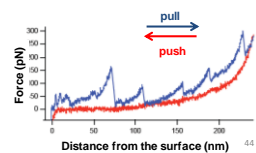
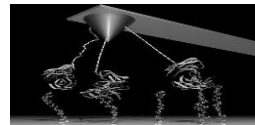
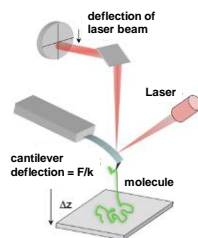
Atomic Force Microscopy

<http://www.youtube.com/watch?v=BrsoS5e39H8>

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Atomic Force Microscopy

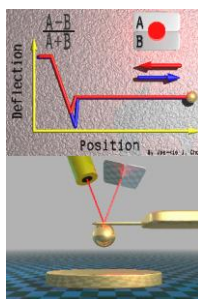
Force spectroscopy: Force-distance traces registered upon push-pull cycles of the AFM tip (movement only in Z direction)
~10 pN sensitivity



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Atomic Force Microscopy

Force spectroscopy:



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Thank you for your attention!



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