
Stacked Polymer Computing (SPOC) Architecture Using Light-Induced Molecular Switching:

A Concept Proposal for 3D, Low-Energy, Optical Molecular Logic System

Concept released publicly by [@kadzdown](#)

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Concept Overview

This concept outlines a 3D, multi-layer polymer computing architecture that leverages light-induced molecular switching to enable ultra-fast, massively parallel molecular-scale computation. The architecture integrates broad-spectrum and wavelength-selective polymers to achieve selective addressing, optical gating, and extremely high information density. Computation is fully optical, with electronic conversion occurring only at the input/output (I/O) interface stage.

1. Layer Structure

Top Layer – Broadband Light-Responsive Polymers

- React to a wide range of light wavelengths.
- Function as a dynamic optical gate, controlling whether light reaches lower layers.
- Can either allow light to pass or absorb/block it, enabling selective activation of bottom layers.

Bottom Layer(s) – Wavelength-Selective Polymers

- Engineered to respond to specific wavelengths of light.

- Enable multi-bit encoding and parallel computation, as different wavelengths trigger different molecular switches.
- Layers can be stacked to increase information density vertically, without expanding surface area.

Layer-Specific Addressing

- Different wavelengths of light independently address different layers, enabling true 3D parallel computation.

Key Enhancement (Integrated): Each compute layer includes embedded plasmonic nanoparticles (e.g., Au or Ag, 5–20 nm) to amplify local electromagnetic fields, reducing input light intensity by 10–100× and enabling sub-femtojoule switching energies.

2. Mechanism

Computation and Storage

- Performed through light-induced isomerization or conformational changes at the molecular level.

Write / Compute Cycle

- Targeted light of a specific wavelength illuminates the polymer layer.
- Molecules in the lower layers switch state only if the upper layer permits light passage.
- **Plasmonic enhancement ensures efficient switching even at low input power.**
- Direct fiber-optic readout extracts the light signal from each layer; no electronic conversion occurs inside the stack.
- States are interpreted electronically only at the I/O stage.

Parallelism

- Multiple wavelengths can operate simultaneously, enabling massively parallel 3D processing.
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3. Advantages

- **Ultra-Fast Switching:** Molecular state changes occur on picosecond to nanosecond timescales.
- **High Information Density:** Stacking layers multiplies capacity beyond 2D molecular arrays.

- **Energy Efficiency:** Light-based switching consumes far less energy than electron-based transistors.
- **Selective Addressing:** The top gating layer enables precise control of activation in lower layers, reducing interference.
- **Native 3D Computation:** Vertical addressing shortens signal paths, reduces congestion, and allows massive parallelism.
- **Ultra-Low Power via Plasmonics:** Estimated sub-fJ/bit switching energies could enable scalable, heat-efficient operation.

4. Key Challenges

- **Fabrication:** Creating thin, defect-free, stacked polymer layers with precise thickness and uniformity.
 - **Fiber Integration:** Embedding ultra-thin fibers without alignment errors, leakage, or damage.
 - **Light Attenuation & Cross-talk:** Ensuring top layers and fibers do not scatter or bleed light to unintended layers.
 - **Thermal Stability:** Maintaining molecular states without relaxation or degradation.
 - **Readout & I/O:** Accurate decoding at the output stage, including wavelength separation and signal amplification.
 - **Integration:** Connecting these molecular layers with existing electronic or photonic systems.
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5. Candidate Materials

Broadband (Top Layer) Candidates

| Category | Example Materials | Key Features |
|--------------------------------|-----------------------------------|-------------------------------------------------------------------------|
| Azobenzene-based photochromics | Poly(azobenzene methacrylate) | Reversible isomerization under UV/visible light; fast switching. |
| Spiropyran/merocyanine systems | Spiropyran–PMMA copolymers | Broad visible spectrum response; fatigue-resistant under low intensity. |
| Diarylethenes | Fluorinated diarylethene polymers | Excellent thermal stability; visible–UV response. |

Wavelength-Selective (Bottom Layer) Candidates

| Target Band | Example Polymers | Notes |
|----------------------|--------------------------------------------------|----------------------------------------------------|
| UV (200–400 nm) | Poly(fluorene) derivatives | Strong UV absorption, stable isomerization cycles. |
| Visible (400–700 nm) | Polythiophenes, PPV (poly(p-phenylene vinylene)) | Tunable bandgap for selective visible response. |
| NIR (700–1200 nm) | Cyanine-based or porphyrin–polymer hybrids | Ideal for deep-layer activation; low energy loss. |

These materials are cited as examples of polymers known to respond to light of varying spectra; actual suitability would require experimental validation.

6. Potential Impact

- Could inform next-generation molecular computing and optical memory architectures.
- Offers a pathway to ultra-dense, low-energy, high-speed computing beyond silicon.
- Even partial implementations (e.g., single gating layer + responsive array with fiber readout) could produce valuable experimental results.
- If successful, the semiconductor industry could pivot from etching smaller silicon circuits to producing thinner, shielded optical fibers embedded in polymer stacks, creating a new paradigm for 3D computing.

While highly theoretical, the approach provides a structured direction for exploring polymer-based optical computation.

7. Note from the Contributor

This idea emerged through exploratory discussion assisted by ChatGPT-5 and GROK, without prior specialization in materials science or photonics. It is released publicly in the spirit of open scientific inspiration.

This document is conceptual and not experimentally validated. It is shared publicly for discussion and exploration.