On-Chip Integrated Photonic Devices Based on Phase Change Materials

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Abstract: Phase change materials present a unique type of materials that drastically change their electrical and optical properties on the introduction of an external electrical or optical stimulus. Although these materials have been around for some decades, they have only recently been implemented for on-chip photonic applications. Since their reinvigoration a few years ago, on-chip devices based on phase change materials have been making a lot of progress, impacting many diverse applications at a very fast pace. At present, they are found in many interesting applications including switches and modulation; however, phase change materials are deemed most essential for next-generation low-power memory devices and neuromorphic computational platforms. This review seeks to highlight the progress thus far made in on-chip devices derived from phase change materials including memory devices, neuromorphic computing, switches, and modulators.

Keywords: phase change material; integrated photonics; neuromorphic computing; phase change memory devices

1. Introduction

Light–matter interaction leads the electric charges in the material to oscillate due to the incident light (electromagnetic) waves. These oscillating charged particles under the incident light become a source of scattered light that interacts with neighbouring charged particles. This implies that the response of each charged particle is from a superposition of all the incident waves including the original incident wave and all of the subsequently scattered waves. At the macro-level, these interactions are manifested in the refractive index of the said material [1]. Given the centrality of the material refractive index, most photonic devices operate by microscale distribution of the refractive index in such a way to achieve the desired macroscopic results [2,3].

Phase change materials (PCMs) are a class of materials with interesting properties, which make them important for the development of various next-generation photonic devices. Their importance is due to phase change materials exhibiting at least two distinct phase states, between which the material can be reversibly switched. These phase states are due to stable and metastable atomic and crystal configurations, resulting in distinct electronic and optical properties [4], such as resistance and refractive index, respectively. As the change in resistance pertains to the domain of electronics, its utilization and devices are outside the scope of this work. This work only considers photonic applications and is, thus, restricted to refractive index switching.

Once a dormant area of research, PCMs were catapulted into the spotlight with the development of compact disc (CD), digital versatile disc (DVD), and Blu-ray disc (BD) that used GeTe–Sb2Te3 (GST) and near-field optics as a means for tertiary data storage [4,5].
Subsequent advancements in optical devices have seen the increasing use of PCMs, as the multiplicity of reversibly switchable stable phases with considerably different refractive indices makes phase change materials an attractive choice for a variety of tasks including tuning [6], switching [7,8], beam steering [9], memory devices [10,11], computational memory devices [12], electro-absorption modulation [13,14], metasurfaces [15,16], and neuromorphic computing [17,18].

In this review, we discuss the recent advancements in on-chip optical devices that utilize phase change materials. To this end, this review presents different kinds of phase change materials and a limited discussion on the theory of phase change among different sorts of material. Then, we discuss different applications of phase change materials including phase change memories, in-memory and neuromorphic computing using PCM, and phase-change metasurfaces; lastly, we present the discussion and outlook.

2. Phase Change Materials

PCMs are a special material that exists in at least two different phases, frequently labeled as amorphous and crystalline. These phases are generally a result of rearrangement of the crystal structure of the material [19,20]. Although most other materials do also have amorphous and crystalline phases, the advantage of PCMs lies in the fact that both phases have different crystal structures, bringing about a considerable change in the refractive index [21]. As an example, silicon dioxide does also exist in amorphous and crystalline phases; however, due to their same crystal structure, there is only an insignificant difference in their refractive indices, and it is not regarded as a PCM [21]. On the other hand, octahedral amorphous GST becomes rhombohedral upon switching to crystalline GST [4], leading to a considerable change in refractive index. A contrasting refractive index and reversible switching are the two properties of PCMs that make them useful for optical devices.

While GST remains the most ubiquitous PCM so far [22], there has been research on other materials as a viable alternative platform for various applications. These phase change materials can be divided into four broad categories of chalcogenides, transition metal oxides, organic, and silicon. Some of the important properties of important PCMs from these categories are listed in Table 1. Additionally, some experiments have also been reported using hybrids. Among all these categories, other than some specific requirements for individual applications, for the PCM to be useful for optical devices, it should have the following properties [4,5]:

- Significant contrast in refractive indices of different phases;
- Rapid switching;
- Chemical stability;
- Reversibly switchable between phases for a large number of cycles.
- Long cycling endurance, which leads to a large number of cycles.

Table 1. Properties of some of the PCMs where complex refractive indices for amorphous and crystalline phases are calculated for the wavelength of 1550 nm. The source of data are from [23] for GST-225, [24] for Sb$_2$S$_3$ and Sb$_2$Se$_3$, [25] for GSST, [26] for VO$_2$, and [27–29] for Si.

<table>
<thead>
<tr>
<th></th>
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<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>GST-225</td>
<td>4.6 + 0.18i</td>
<td>7.2 + 1.9i</td>
<td>600 °C</td>
<td>No</td>
</tr>
<tr>
<td>Sb$_2$S$_3$</td>
<td>2.712 + 0i</td>
<td>3.308 + 0i</td>
<td>270 °C</td>
<td>No</td>
</tr>
<tr>
<td>Sb$_2$Se$_3$</td>
<td>3.285 + 0i</td>
<td>4.050 + 0i</td>
<td>200 °C</td>
<td>No</td>
</tr>
<tr>
<td>GSST</td>
<td>3.325 + 0.00018i</td>
<td>5.083 + 0.35i</td>
<td>250 °C</td>
<td>No</td>
</tr>
<tr>
<td>VO$_2$</td>
<td>2.75 + 0.4i</td>
<td>2.1 + 1.3i</td>
<td>68 °C</td>
<td>Yes</td>
</tr>
<tr>
<td>Si</td>
<td>3.4858 + 0i</td>
<td>3.487 + 0i</td>
<td>677 °C</td>
<td>No</td>
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</tbody>
</table>
2.1. Chalcogenides

Chalcogenides are compounds between a chalcogen element (such as sulfur, selenium, tellurium, and polonium) and one or more metals. They generally undergo octahedral to rhombohedral crystalline structure on switching as shown in Figure 1. A typical chalcogenide PCM is GST [30,31], which has many varieties depending upon the concentration of constituent elements, but GST-225 (marked in red in Figure 1a) is the most useful of them all. In addition to GST, there are other chalcogenide PCMs, including GeTe [32], Sb2Te [32], Ag5In5Sb60Te30 (AIST) [33], and GSST [34,35]. These materials exhibit strong resonance bonding in their crystalline state, which is the reason for the considerable difference in the optical properties compared to other materials [36]. Comparisons of GST, GSST, and Sb2S3 properties can be verified in Figure 1c–e. As a group, all of them have small electronegativity differences, which is the reason for their covalent bonding, and both of their phases are nonvolatile at room temperature, which is an advantage for certain applications such as memory devices.

![Figure 1. Chalcogenide phase change materials. (a) Ternary phase diagram of Ge–Sb–Te showing important pseudo-binary combination. Most frequently used (GeTe)_x–(Sb2Te3)_1–x compounds such as GST-326 and GST-225 are labeled in red. (b) The typical structural transformation in GST-225 when it undergoes a phase transition from amorphous to crystalline to liquid and then back to amorphous. T_g and T_m are the glass transition temperature and melting temperature, respectively. (c) Dielectric function of GST-225 in the amorphous and crystalline phase. (d) Refractive index (n) and (e) extinction coefficient (k) of the amorphous and crystalline state of Sb2S3 and GSST. ((a–c) reproduced with permission [21]; (d,e) reproduced with permission [37]).]

Sb2S3 and Sb2Se3 have been rather late entrants into the group of chalcogenide PCMs. Although some earlier work reported Sb2S3 and Sb2Se3 as PCMs [38,39], subsequent works on chalcogenide PCMs largely ignored these materials [4,32,40] until they were rediscovered as low-loss PCMs [24]. As can be observed in Figure 1c–e, in contrast with GST, which has a large absorption coefficient leading to large losses, Sb2S3 was reported to have an absorption coefficient of zero at the operating wavelength of 1550 nm [24,37], while that of Sb2Se3 was also reported to be near zero at the wavelength of 1550 nm by Delaney [24]. As for the visible region, Sb2S3 still retains near-zero absorption as opposed to Sb2Se3, which shows a sharper rise in the absorption coefficient [40]. The property of low loss attracted renewed interest in these materials [41–46], for both on-chip and free-space applications. On the flip side, Sb2S3 reportedly also undergoes larger size variation when switching from amorphous to the crystalline phase. This may be undesirable for certain applications.

Oxidation of the chalcogenide layer is a major problem among all of these chalcogenide materials which oxidize even at room temperature, leading to performance degradation
Transition metal oxides that undergo Mott transition (also called correlated materials) are the second most important category of PCMs in terms of volume of publications. Conductivity in Mott insulators such as vanadium dioxide (VO₂) is temperature-dependent as they transit from insulator to conductor when the temperature goes from low to high [36]. This transition in conductivity is accompanied by a transition in refractive index, making the materials useful for optical devices. Among them, vanadium oxide is the most popular that can reversibly switch between metallic rutile and two insulating monoclinic structures [49], depicted in Figure 2a. While all of the transition metal oxides show a similar structural transition, the structural transition does not always translate into electronic transition, leading to metallic and insulating phases as in VO₂ and NbO₂. Both states show different refractive indices and absorption coefficients. Typical values of refractive index and absorption coefficient for VO₂ are shown in Figure 2b,c, respectively. The temperature at which the switching takes place is called the transition point. These transition points in the correlated materials can be tuned by tuning the banding filling through doping [50]; for example, the VO₂ transition point is at 68 °C, which can be further reduced even to room temperature by doping it with tungsten [36].

Figure 2. Transition metal oxide phase change material VO₂. (a) Comparison of monoclinic and rutile crystallographic structures; the unit cells are indicated by solid lines. Vanadium atoms are represented by red balls; oxygen atoms are blue. The electronic band structure near the Fermi level is given at the bottom. (b) Refractive index and (c) extinction coefficient of VO₂ in the cold and hot states showing a large change in refractive index and extinction coefficient with the temperature. (a) reproduced with permission [51]; (b,c) reproduced with permission [52].
The temperature dependence of the metastable states means that the material is suitable for volatile applications. The material has been tested with on-chip devices such as optical switches [53], resonators [19], photonic and plasmonic waveguides [54], polarization converters [55,56]. The temperature can be changed through thermal, electrical, or optical means, but applications requiring faster switching should resort to electrical or optical switching as thermal means are slow.

Like GST, vanadium oxide also readily oxidizes from VO$_2$ to V$_2$O$_5$ in air and, thus, requires a protective layer. Hydrophobic material hafnium dioxide (HfO$_2$) is usually used as the protective layer, which should completely cover the VO$_2$ sample to protect it from oxidation as an unprotected cross-section of the film would also lead to performance degradation [57].

Although many research groups are actively involved in the development of VO$_2$-based devices, some researchers have shown skepticism toward the applicability of VO$_2$-based systems in practical applications for their limited multilevel response, volatility, and complexity of fabrication [58]. This is probably also the reason for them not being as popular as chalcogenides.

2.3. Organic Phase Change Materials

Organic materials are predominantly made of carbon and hydrogen; however, neither these elements nor their compounds inherently exhibit phase change behavior. Therefore, phase change behavior is generally achieved through the interaction of cyclic compounds with other functional groups such as benzene, pyridine, and pyrrole [59]. Some experiments with functional groups containing transition elements have been successful [60].

Numerous organic materials have been used for memristive memory devices [61], switches [62], and neuromorphic computational platforms [63]. This includes pure organic materials in perovskite and non-perovskite forms [63], as well as hybrids with other metals or nonmetals [64,65]. All of these reported devices operate on the basis of a change in resistance through particle transfer leading to a switching action. As this review is focused on phase change materials useful for photonic applications, these platforms are out of the scope of this study.

On the other hand, many organic phase change materials including PNIPAM [66,67] or even doped carbon [59] do exist, which are thermally responsive. As these materials have temperature-dependent metastable states, they can theoretically be used for volatile applications in which VO$_2$ is used. However, to date, there has been no report of using organic phase change materials for optical devices of this nature. We can speculate the reasons to be related to the insufficient contrast in refractive index as reported for PNIPAM [68], phase instability as reported for PNIPAM [69], slower response time [70], or a lack of interest among the research community for exploring organic phase change materials.

Some of the organic phase change materials with different functional groups are shown in Figure 3 with the hope that it could inspire some interest among the research community for organic phase change materials in the future.
2.4. Silicon

Silicon is the most pervasively used material in the photonics industry, as well as the semiconductor industry at large [36], which can also behave as a phase change material. The ability to control the crystallinity of silicon by tuning the temperature during the low-pressure chemical vapor deposition (LPCVD) process has always been known. The transistor industry has been regularly producing crystalline silicon through the action of lasers, both pulsed lasers [71] and excimer lasers [72], in thin films, as well as bulk silicon. These laser-based systems can crystallize and re-amorphize the silicon on the nanosecond time scale, and this process can be reversibly switched without any deformation. The change in refractive index and absorption coefficient brought about due to the silicon changing phase is not significant at the wavelength of 1.55 µm [73]. However, the difference becomes comparatively large as the wavelength shortens to the visible regime, as shown in Figure 4. Some free-space visible-range proof-of-concept studies have also reported the use of silicon as a PCM [29,74]; however, at the time of writing this review, no demonstration of silicon as a PCM for integrated devices has been found.

Figure 3. Optimized organic bimolecular configurations between different nitrogen species of N-doped hierarchical carbon and PCM molecules with different functional groups such as (a) Pyridinic nitrogen, (b) Pyrrolic nitrogen, (c) Graphitic nitrogen, and (d) Oxidized nitrogen (reproduced with permission [59]).
Figure 4. (a) Refractive index and (b) extinction coefficient of silicon in the crystalline and amorphous states showing a large change in refractive index and extinction coefficient in the UV and near-UV region, which is reduced to an insignificant difference as it reaches 1.5 µm [27,28]. (c) Transmission spectra of pristine a-Si nanodiscs, crystallized nanodiscs, and re-amorphized nanodiscs with 215 nm in diameter. (d) The conceptual pixel-addressable rewritable metasurface for optical wavefront control. ((c,d) Reproduced with permission [29]).

Having looked at all four categories of PCMs in detail, the pros and cons of each of the categories for integrated photonic applications are summarized in Table 2.

Table 2. Summary of pros and cons of PCM categories; the last two columns denote whether there has been a report in the literature concerning the use of each category for any type of photonic application or integrated photonic application.

<table>
<thead>
<tr>
<th>Category</th>
<th>Pros</th>
<th>Cons</th>
<th>Used for Photonic app.</th>
<th>Used for Integrated Photonic app.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chalcogenide</td>
<td>Fast switching, nonvolatile</td>
<td>High transition temperature, lossy</td>
<td>✓</td>
<td>✓</td>
</tr>
<tr>
<td>Transition metal</td>
<td>Low transition temperature, volatile, fast switching</td>
<td>Require constant energy, lossy</td>
<td>✓</td>
<td>✓</td>
</tr>
<tr>
<td>oxide</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Organic</td>
<td>Low transition temperature</td>
<td>Low melting point, slower response, easily combustible</td>
<td>×</td>
<td>×</td>
</tr>
<tr>
<td></td>
<td></td>
<td>No refractive index contrast at IR,</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Silicon</td>
<td>Ease of fabrication, ease of integration, low loss</td>
<td>Highest transition temperature</td>
<td>✓</td>
<td>×</td>
</tr>
</tbody>
</table>
3. Phase Switching

The phase switching process can be seen at two levels. One is at the atomic level where the kinetics of the phase change process is studied. The other level is at the device level when the phase change is brought about by the designer of the system. The kinetics of crystallization for a phase change material in the amorphous phase can be divided into two categories of nucleation-dominated and growth-dominated [75].

Nucleation-dominated crystallization (shown in Figure 5a) takes place with small spots of crystalline material that are randomly distributed in the PCM. With the rise of the temperature, these nuclei grow until the whole of the PCM has crystallized [76]. A study conducted by Lee and his colleagues [76] found that chalcogenides such as GST crystallize through the nucleation-dominated process where the size distribution of the nuclei depends on the thermal treatment they received, which implies that the statistical status of nuclei distribution in as-deposited amorphous PCM is different to that in pre-annealed PCM. Moreover, their research into the nucleation process also found that, if the amount or volume of subcritical nuclei embedded in amorphous material is greater, the formation of a supercritical nucleus is easier and, thus, the time is shorter [76,77]. The result is that the nucleation process in GST is faster, thus making it a nucleation-dominated crystallization material.

![Figure 5](image-url)  
Figure 5. Two different kinetics of crystallization: (a) nucleation-dominated crystallization kinetics where there is a random distribution of islands of crystallinity that later expand; (b) growth-dominated crystallization where the initial crystalline precipitate grows to include the rest of the amorphous material.

Growth-dominated crystallization (depicted in Figure 5b), on the other hand, is where the rate of growth of the crystalline region is more than the rate of nucleation. In such crystals, only a few nuclei precipitate when the amorphous material is annealed [75]; instead, the precipitated nuclei grow their boundaries to include the still amorphous region into the crystalline region [78]. A study conducted by Khulbe found that, in materials that are growth-dominated, there is a trade-off between incubation time and nucleation speed. The most important examples of materials showing growth-dominated crystallization are Ag and In-doped SbTe (AIST) [75].

At the device level, the underlying physics to achieve switching remains thermal; however, due to the sluggish response of the thermal systems, the PCM is not directly heated. Rather, to switch, either electrical means or optical means are used which achieve the temperature required for switching between crystalline and amorphous phases. Although some electro-thermal methods have also been improved recently through the use of devices such as graphene heaters [79], considered as a fundamental mechanism, there is a lag in switching speed [80]. Among these three means of switching, optical switching produces the fastest switching, followed by electrical switching, while thermal switching remains the slowest [19,36].
Another aspect connected with phase switching is the cycling endurance of the phase change material used for switching. The endurance refers to the number of cycles for which a particular material can be reversibly switched before its two phases (SET and RESET) become difficult to discriminate or the material simply does not switch [81,82]. Such recyclability, measured as the number of cycles of PCMs, is inversely related to the volume of the PCM used, as well as the method of switching adopted. According to Bez, at the fundamental level, there should not be any wear and tear of the material due to switching [83]. However, data show reduced resistive performance after $10^8$–$10^{10}$ cycles. An interesting finding by Raoux was that the number of cycles and the energy required to trigger the phase transition are inversely related [81]. With respect to the method of switching, GST is known to have a recyclability of higher than $10^7$ cycles when electrically switched, whereas it reduces to $10^5$ cycles when optical switching is employed [84].

3.1. Electrical Switching

Electrical switching is the older of the two methods of electrical and optical switching, and it also finds parallels in earlier technologies such as transistors and other concurrent technologies such as memristors [85,86]. For switching the PCM through electrical means, two different methods have been used. One method is to use an external electrical circuit that is generally connected to a heating element that generates heat through resistance to the current [7,87]. Examples of such systems are ITO heaters [88,89], silver heaters [80], and PIN heaters [90]. The other method is the approach used in memristors where current is used for achieving the phase change [7]. Schematics of the typical arrangements to achieve electrical switching with GST and VO$_2$ as PCMs are shown in Figure 6a–c. The generated heat due to passing current can reversibly switch the state of the PCM unit placed on top of these photonic structures. After switching, when an optical signal passes through the waveguide under the PCM, the signal is evanescently coupled with the straddling PCM unit, leading to a change in the signal. So far, the literature has not revealed any phase change material-based photonic device that has in-line phase change material.

These electrical methods are advantageous, as the heating circuit and the optical circuit of the device are independent and separate from each other. This makes the device more controllable and can enable easier access to complex and multilayered optical circuits. Given that microelectronic technology is well established and mature, it helps in the design of electrically controlled devices. Additionally, as the external control circuitry is also electronic, it can be monolithically integrated. On the other hand, the electronic control circuit makes the devices bulky and increases the steps of fabrication linked to the final cost of the product. However, most importantly as per the current thrust of society toward energy conservation, the electronic method of switching is also wasteful in terms of energy, leading to higher cost per switch. Recently, some graphene-based models have been suggested (schematically shown in Figure 6b) due to their success in limiting the energy waste in electrically controlled PCM-based photonic devices [79,92].
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3.2. Optical Switching

Optical switching, also called all-optical devices, on the other hand, is when an optical signal is used to bring about phase change in the PCM. Heat is generated by a high-energy laser pulse due to the opto-thermal effect, leading to switching in the linked phase change material [95]. This can be achieved through two different methods. One method is to use an external laser that is incident on the PCM unit from the far-field and excite it to achieve the switching [82,96]. This method is used for free-space devices, as well as on-chip devices, as shown in Figure 6d. Figure 6d shows that using plasmon polaritons for heat generation would ultimately lead to phase change of the PCM. The second method, which is used for on-chip devices, involves evanescent field coupling. In evanescent coupling, a control signal from a high-powered laser traverses through the on-chip photonic crystal and achieves a switch in the PCM deployed on top through the opto-thermal interaction of the evanescent field and the PCM, as shown in Figure 6e [97]. Although some devices with in-line PCM for certain applications have also been demonstrated [98], the large absorption coefficient in most PCMs currently in use has led to most designers resorting to evanescent coupling (schematically shown in Figure 6e). With the development of low-loss PCMs [41], the phase change material can be deployed in line. The in-line deployed PCM

![Figure 6. Various electrical and optical switching methods.](image-url)
would increase the interaction volume of the PCM and the optical wave, leading to smaller device footprints.

The optical switching uses the same optical circuit that the signal would use, making the device design less bulky while also reducing the number of fabrication steps. Researchers, using optical switching, have reportedly achieved low-power switching using femtosecond lasers [9]. This is a promising result concerning lower energy targets. However, on the other hand, routing the optical signal to achieve the necessary phase change is a difficult task as fabricating three-dimensional photonic circuits is still not easily achievable with the current state of technology. Secondly, achieving switching in the traditional sense using low-loss PCM is difficult due to the low absorption loss of the material. Thus, accessing each PCM unit serially placed on a single waveguide individually is also problematic. While it may not be needed for certain applications such as those implementing Hebbian learning for artificial synapses [95,99], there are other situations where individual access is needed. This has led to some researchers addressing this problem through the use of novel designs such as grid structures [12] or through the use of wavelength division multiplexing with ring resonators straddling individual PCM unit cells [100]. Although promising, these systems are still far from being useful in practical devices.

On the topic of comparison between electrical switching and optical switching in GST-based silicon photonic switches, Zhang concluded that, for the amorphization process, optical switching is advantageous in terms of speed and energy efficiency. However, as alluded to earlier, for large-scale integration purposes, electrical switching is still advantageous as it does not require a complex routing circuit for the photonic circuit [101].

4. Phase Change Memories

Phase change memories can be regarded as the first and arguably most successful application of phase change materials, with CDs as a forerunner, followed by DVDs and BDs as the most popular tertiary memory devices up until recently [5]. The development of such phase change material-based memories can be traced back to the 1970s [102]. In one such work, Yamada and his colleagues [103] presented direct overwriting at the speed of nanoseconds using GST alloy. Despite the success and the early start, these applications were optically switched and not the on-chip applications of phase change materials.

On the on-chip front, the subsequent decades saw extensive developments in the electrically switched resistance-based phase change memories that were comparable with dynamic random access memory (DRAM), NAND based on not/and (NAND) logic gates [104], flash memory [105], and even a proposal for on-chip optical switches [106]. The high-density memory, presented as nonvolatile phase-change random access memory (PRAM), addressed the speed, scalability, and power consumption issues of DRAM [107–109].

The first all-optical on-chip phase change memory was proposed by placing a patch of GST-225 on an Si$_3$N$_4$ ring resonator structure on silica. This was reported in 2012 by Pernice and Bhaskaran [10] (some of the details are shown in Table 3), whose group later also fabricated the sample but presented it as a tunable nanophotonic circuit, which was a building block on the way to creating a memory element [110]. Subsequently, a detailed analysis of the memory device made with a GST-225 patch on an Si$_3$N$_4$ racetrack was presented in 2014, which is shown in Figure 7a [111]. Using the principle of change in absorption of GST [112], they showed that the Q-factor, the resonance wavelength, and the extinction ratio can be used to retrieve the state of GST, which is a memory element that can be switched on “a picosecond timescale” [111]. In a later work, the group demonstrated a multilevel cum multibit storage using multiple ring resonators coupled with a single waveguide, accessible through wavelength division multiplexing (WDM) [113]. The demonstrated device with an operating speed of 800 MHz for the write operation could be improved to the gigahertz level using picosecond pulses. The reported device, with a footprint of 0.25 $\mu$m$^2$, demonstrated fast (∼500 ps) and low-power readout (480 fJ). This work is unique as it also showed the ability to accurately control the percentage of GST in the crystalline or amorphous state through the intensity of the writing
pulse [112]. Using a slightly different architecture, an extensive 512 bit all-optical memory was also demonstrated to store a 16 × 16 bit image in grayscale with 2 bit resolution [100]. The device had a relatively large footprint of 1000 × 2400 µm², which could be slightly reduced using a silicon substrate instead of Si₃N₄.

Table 3. Major contributions in memory devices.

<table>
<thead>
<tr>
<th>Year</th>
<th>PCM</th>
<th>Main Features</th>
<th>All Optical</th>
<th>Speed</th>
<th>Footprint</th>
<th>Power</th>
</tr>
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<tr>
<td>2012 [10]</td>
<td>GST</td>
<td>Si₃N₄ ring resonator with GST patch</td>
<td>√</td>
<td>500 ps</td>
<td>0.25 µm²</td>
<td>480 fJ</td>
</tr>
<tr>
<td>2014 [111]</td>
<td>GST</td>
<td>Si₃N₄ ring resonator with GST patch</td>
<td>√</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>2015 [113]</td>
<td>GST</td>
<td>Si₃N₄ multiple ring resonators with GST patches coupled to a single waveguide</td>
<td>√</td>
<td>500 ps</td>
<td>0.25 µm²</td>
<td>480 fJ</td>
</tr>
<tr>
<td>2018 [114]</td>
<td>GST</td>
<td>Photonic Memories and Logic Operations using PWM for switching of PCM.</td>
<td>√</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>2018 [115]</td>
<td>GST</td>
<td>Logic “OR” and “NAND” implemented</td>
<td>×</td>
<td>505 ns</td>
<td>-</td>
<td>16 pJ</td>
</tr>
<tr>
<td>2021 [25]</td>
<td>GSTT</td>
<td>Photonic integrated memory with accuracy up to 93%</td>
<td>×</td>
<td>&lt;1 ps</td>
<td>&lt;4 µm²</td>
<td>-</td>
</tr>
</tbody>
</table>

Figure 7. Implementation of phase change memories. (a) Microscopic image different memory elements. Along the horizontal axis, the waveguide–resonator gap is varied from 0.5 to 1.5 µm with 0.3 µm successive step size. Along the vertical direction, the width of GST varies from 0.5 µm (bottom) to 20 µm (top). (b) Comparison between resonances in the crystalline and amorphous states for different GST widths. The graphs show a relative shift of the central wavelength and the reduction of the Q-factor with broadening of the peaks. (c) ER obtained for different racetrack radii (R) as a function of GST width and state. The highlighted regions correspond to those highlighted in (b), and the lines correspond to the fits obtained using the analytical function for ER. (d) Concept of the dual-pulse programming technique; from left to right, an optical image of the device with a GST photonic memory cell, a magnified image of GST on top of the waveguide, and a schematic cross-section of the completed device [116]. (e) Experimental demonstration of multilevel photonic memory using multi-pulse PWM with 12 different levels in a single device showing the relative change in transmission (ΔTr/Tr₀). This is an implementation of a switch for the photonic memory with each level accessed through the multiple identical pulse trains of 10 pulses with p = 30 ns and P_peak = 1.4 mW. (f) Randomly accessing six levels of a single photonic memory. A multiple identical pulse train is used to reach different levels, with N = 12, p = 20 ns, and P_peak = 1.6 mW. Each level can be reached with the same pulse train at a determined width (τ₀ to τ₅) from any previous level, where τ₀ = 10 ns, τ₁ = 10.5 ns, τ₂ = 11 ns, τ₃ = 11.5 ns, τ₄ = 12.5 ns, and τ₅ = 15 ns. The dotted and dashed lines denote the times when optical pulse trains are applied for amorphization and crystallization, respectively. (a–c) reproduced with permission [111]; (d) reproduced under creative commons license from [116], (e,f), reproduced with permission [114].
For obvious reasons, the footprint of the micro-ring resonator-based device is comparatively large, and reducing it would enable a device with larger density. In such an attempt, the group of Pernice and Bhaskaran used a patch of GST on top of a straight Si$_3$N$_4$ waveguide as a memory unit [114,117]. The device was able to achieve multi-level memory with 21 different states from a single unit of PCM achieved through partial crystallization of GST using pulse width modulation (PWM) on a low-power pump of 526 mA. Figure 7d,e show the experimental demonstration of state transformations using PWM and random accessing of different levels of memory. As the need for frequent optoelectronic conversion is cumbersome, another study reported using integrated phase-change memory, which combines plasmonics, photonics, and electronics to achieve dual electrical and/or optical switching of multilevel states [115].

All of these devices consume relatively high power owing to the large absorption coefficient of GST. As reducing power consumption is the other major thrust of research on phase-change memories, GSST has been proposed as a unit of photonic integrated memory due to its lower absorption [25] and because electro-thermal switching is used for this device.

Following the successful demonstration of all photonic nonvolatile phase-change memory, some researchers resorted to electrically switched photonic memories for Si-PIC. As discussed earlier, electrical switching does offer some benefits compared to all photonic arrangements. It allows fabricating a micro-ring resonator using polycrystalline silicon as the memory element that is switched using joule heating generated using aluminum electrodes. The device consumes 0.476 pJ and 0.316 pJ for the write and read operations, respectively [118]. A similar technique used an ITO heater for heating and achieved switching in a silicon-based device using GST as the phase change material [119]. To further improve the quality of readout, plasmonics has also been used with electrically switched devices, which allows improving the switching speed and energy efficiency [120].

5. In-Memory and Neuromorphic Computing Using PCM

Neuromorphic computing is a paradigm that aims at emulating the way a brain computes, especially focusing on the property of the brain to carry out low-energy information processing, speedy processing, and complex and sophisticated tasks [121]. To mimic the brain, distributed computing topologies are generally used [122], which tend to emulate the behavior of a brain synapse. These structures, appropriately called “artificial synapses”, have been used for machine learning. Most techniques for the implementation of machine learning rely on backpropagation [123,124] as the method to achieve learning, except for a few [125]. Among those using backpropagation as the method of learning, Hebbian learning algorithms are the most popular [126,127].

Phase change material-based memristors have already been in research for a long time for the implementation of neuromorphic learning and in-memory computing [128,129]. The potential for using PCM-based photonic devices for neural networks including neuromorphic computing was understood right after the first report of PCM-based memory [130]. Since then, the neuromorphic potential of phase change materials has constantly been explored and developed (some of the major contribution in this regard have been outlined in Table 4). The first development in this regard was an all photonic synapse that implemented the Hebbian learning rule to achieve synaptic plasticity, made using multiple GST patches on a straight waveguide developed as a memory, already mentioned above [95,117]. In this device (schematic and scanning electron microscopic (SEM) pictures are shown in Figure 8a,b), the synaptic weight is set by changing the number of optical pulses as the synaptic weight exponentially depends on the number of pulses. Using a pulse of 50 ns at 243 pJ, a 7% change in transmission was achieved corresponding to a synaptic weight of “3”. Subsequently, 100 pulses at 1 MHz would allow the PCM to transition from “3” to “1”, while 1000 pulses would revert the state of the synapse to “0”. The research found that the synaptic weight is strictly dependent on the number of pulses applied. The reported
device achieved 11 synaptic weights, which could be further improved to a truly analog system by improved pulse control, thereby improving the signal-to-noise ratio.

Table 4. Major contributions in neuromorphic/in-memory computational platforms.

<table>
<thead>
<tr>
<th>Year</th>
<th>PCM</th>
<th>Main Features</th>
<th>All Optical</th>
<th>Speed</th>
<th>Footprint</th>
<th>Power</th>
</tr>
</thead>
<tbody>
<tr>
<td>2017 [117]</td>
<td>GST</td>
<td>Si₃N₄-based synapse, with 21 levels on a single waveguide; Implemented with multiple patches of GST on straight waveguide</td>
<td>√</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>2019 [99]</td>
<td>GST</td>
<td>Direct scalar multiplication with a single integrated memory cell; Matrix multiplication with multiple PCM cells Synapses for supervised and unsupervised self-learning; Implemented with waveguides and waveguide crossings with PCM on top</td>
<td>√</td>
<td>200 ns</td>
<td>-</td>
<td>180 pJ</td>
</tr>
<tr>
<td>2020 [100]</td>
<td>GST</td>
<td>Photonic convolutional neural network with photonic computing core through PMMC devices Low-energy artificial synapse Accuracy &gt;90%</td>
<td>√</td>
<td>1 kHz</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>2021 [131]</td>
<td>GST</td>
<td>Integrated photonic tensor core for parallel convolutional processing</td>
<td>√</td>
<td>20 ps</td>
<td>-</td>
<td>0.1 µJ</td>
</tr>
<tr>
<td>2021 [46]</td>
<td>Sb₂S₃ and GST</td>
<td>Nonlinear activation function Accuracy &gt;90%</td>
<td>√</td>
<td>-</td>
<td>-</td>
<td>17 fJ/MAC</td>
</tr>
</tbody>
</table>

Creating a synapse on a single waveguide is very attractive from the point of view of reduced footprint, but the system is very difficult to scale; therefore, to create a network of synapses, a different approach is required. A silicon-based ring resonator with a GST patch is one such candidate that has been used for creating an all-optical synapse [133]. A similar synapse (schematically shown in Figure 8c) was also used to create an integrate-and-fire neuron [23]. As any synaptic network is a dot-product engine, these ring resonators are connected in a network to essentially perform dot-product calculations in the form of multiplication of the input with the neuronal weight [134]. This neural network was trained, using a backpropagation algorithm, to recognize handwritten digits from the MNIST dataset [135], achieving 98.06% accuracy on testing. On the speed and energy efficiency front, the ‘read’ and ‘write’ actions take 500 ps and 1.5 ns for 1 pJ and 4 pJ, respectively, using pulses with a duration of 200 ps. This system was later further improved by better incorporation of wavelength division multiplexing [136] and behavioral modeling [137], which further improved energy and speed performance.

There is an inherent contradiction in the constant pursuit for decreasing the energy consumption of computing devices and having GST, which is a highly absorptive material, as the choice for PCM. Given the contradiction, it is only understandable that researchers are looking for alternative materials. In such an attempt, Volker produced an artificial synapse using GSST instead of GST [25]. For this synapse, they resorted to using an electrical switching method implemented using a tungsten contact instead of an optically switched technique. The neural network, made using of Mach–Zehnder interferometers (MZIs) with GSST on both arms, was trained and tested using MNIST [135] handwritten digits and was able to achieve a high level of accuracy.
Scaling the neural network further, [25] recently presented a synapse, graphical processing unit, and a tensor processing unit that perform in-memory calculations through the network of coupler-based latch circuit using Si$_3$N$_4$ with Sb$_2$S$_3$, as well as GST as the PCM [46]. The network (shown in Figure 8e,f) performs at the maximum data rate of 100 GHz and consumes 0.1 fJ of energy. In another work, the tensor processing unit was implemented using GST on a network of ring resonators [137,138]. The implementation of ring resonators helps in reducing the power consumption as the GST is not directly in contact with the propagating waves. They implement parallel processing for convolutional operation showing a variety of machine learning tasks at 2 tera-MAC operations per second with a modulation speed of more than 13 GHz consuming 17 fJ per MAC. This is thus far the fastest implementation of an all-photonic in-memory computational network. The speed, according to the study, can be further improved by expanding the size of the network to increase the level of parallelization.
6. Phase Change Modulators and Switches

Metasurfaces that are artificially micro/nanostructured in a certain way to perform the function of modulators and switches are important PCM-based integrated photonic devices. Such devices generally contain microstructures that partially or fully contain phase change materials in them. The applications of such metasurfaces in producing switches [87,139–141], transmission tuners [91], and modulators [19,142–145] are ubiquitous, which are generally implemented using either Mach–Zehnder interferometers (MZI) or ring resonators [146]. While the major contributions in PCM-based photonic switches and modulators have been outlined in Table 5, the important features of these switches and modulators as per Abdollahramezani are their low power, high contrast, and fast speed [146]. To achieve these features, other than the design parameters, the device is also dependent upon the fabrication processes adopted. As no method is perfect and each method has its own advantages and disadvantages [147].

Table 5. Major contributions in PCM-based photonic switches and modulators.

<table>
<thead>
<tr>
<th>Year</th>
<th>PCM</th>
<th>Main Features</th>
<th>All Optical</th>
<th>Speed</th>
<th>Footprint</th>
<th>Insertion Loss</th>
</tr>
</thead>
<tbody>
<tr>
<td>2010 [142]</td>
<td>VO₂</td>
<td>Ring resonator-based modulator ¹</td>
<td>√</td>
<td>-</td>
<td>2 µm, 5 µm</td>
<td>2 dB</td>
</tr>
<tr>
<td>2012 [143]</td>
<td>VO₂</td>
<td>Ring resonator-based modulator ²</td>
<td>√</td>
<td>-</td>
<td>0.28 µm²</td>
<td>-</td>
</tr>
<tr>
<td>2012 [148]</td>
<td>GST</td>
<td>2 × 2 MZI-based switch ²</td>
<td>√</td>
<td>-</td>
<td>93 × 1.7 µm²</td>
<td>0.5 dB</td>
</tr>
<tr>
<td>2012 [149]</td>
<td>VO₂</td>
<td>2 × 2 plasmonic switch ²</td>
<td>×</td>
<td>10 s kHz</td>
<td>5 µm</td>
<td>1.5 dB</td>
</tr>
<tr>
<td>2013 [150]</td>
<td>GST</td>
<td>Ring resonator-based switch ²</td>
<td>√</td>
<td>-</td>
<td>-</td>
<td>2.5 dB</td>
</tr>
<tr>
<td>2014 [151]</td>
<td>VO₂</td>
<td>Plasmonic modulator</td>
<td>×</td>
<td>-</td>
<td>9.47 µm</td>
<td>4.5 dB</td>
</tr>
<tr>
<td>2015 [152]</td>
<td>VO₂</td>
<td>Switch ²</td>
<td>×</td>
<td>500 ns</td>
<td>0.3 × 0.5 µm²</td>
<td>&lt;3.6 dB</td>
</tr>
<tr>
<td>2017 [153]</td>
<td>VO₂</td>
<td>Modulator ²</td>
<td>×</td>
<td>3 ns</td>
<td>6 µm</td>
<td>0.3 dB</td>
</tr>
<tr>
<td>2017 [89]</td>
<td>GST</td>
<td>Switch ³</td>
<td>×</td>
<td>-</td>
<td>3 µm</td>
<td>2 dB</td>
</tr>
<tr>
<td>2018 [154]</td>
<td>VO₂</td>
<td>Plasmonic modulators ²</td>
<td>×</td>
<td>1GHz</td>
<td>2 µm²</td>
<td>1.4 dB/µm</td>
</tr>
<tr>
<td>2018 [14]</td>
<td>GST</td>
<td>Modulator ⁴</td>
<td>×</td>
<td>-</td>
<td>0.2 µm²</td>
<td>&lt;2 dB</td>
</tr>
<tr>
<td>2019 [155]</td>
<td>GST</td>
<td>Switch ²</td>
<td>√</td>
<td>-</td>
<td>5 µm</td>
<td>0.135 dB</td>
</tr>
<tr>
<td>2019 [156]</td>
<td>GST</td>
<td>Plasmonic modulator for MIM device ²</td>
<td>√</td>
<td>500 ps</td>
<td>120 nm</td>
<td>3.6 dB</td>
</tr>
<tr>
<td>2020 [98]</td>
<td>GST</td>
<td>PCM in a slot ²</td>
<td>×</td>
<td>-</td>
<td>0.014 µm²</td>
<td>&lt;4 dB</td>
</tr>
<tr>
<td>2020 [157]</td>
<td>GST</td>
<td>MZI-based ⁵</td>
<td>×</td>
<td>≈ns</td>
<td>5 µm²</td>
<td>4 dB</td>
</tr>
<tr>
<td>2021 [158]</td>
<td>Sb₂Se₃</td>
<td>Resonator-based modulator ²</td>
<td>√</td>
<td>800 ns</td>
<td>11 µm²</td>
<td>0.45 dB</td>
</tr>
</tbody>
</table>


6.1. VO₂ Metasurfaces

Vanadium dioxide-based on-chip modulators have been around for a bit longer than GST-based devices [142,144]. In these devices (shown in Figure 9), a ring resonator with a small patch of VO₂ on the ring is fabricated. The change in temperature causes a change in the phase of VO₂, leading to a change in the resonant frequency. An extended such
device would yield modulation of 16 dB, which is comparable with MZI. In this device, the temperature is changed using substrate heating. In a subsequent study, the phase change was caused externally by a 532 nm wavelength pump laser, leading to modulation of 10 dB in a ring resonator with a radius of 1.5 µm [143]. The experimental setup and the result of modulation achieved with this arrangement are shown in Figure 9c,d, respectively. On the other hand, a more recent study reported a modulator with an extinction ratio of 25 dB and a low insertion loss of 1.4 dB. This was achieved by using a ring resonator with a VO2 patch, which was thermally tuned through lateral micro-heaters placed beside the waveguide [159].

Figure 9. Photonic modulators and switches using VO2 as the PCM. (a) Schematic of VO2 modulator consisting of a grating coupler, a silicon bus waveguide, and a silicon ring resonator covered with a strip of VO2. The SEM image of the VO2 patch straddling the waveguide is shown on the right. (b) Transmission spectra of the through port at increasing temperatures for the schematic presented in (a). The modes of the same azimuthal order are indicated with diamond-shaped markers. These markers show a temperature-dependent red-shift. (c) Schematic of the measurement setup of a VO2-based optical modulator. The SEM image of the modulator is shown in the inset. (d) Optical transmission from the setup given in (c) before and after the trigger. The spectra show a blue-shift upon exposure to the trigger. ((a,b) reproduced under creative commons license [142]; (c,d) reproduced under creative commons license [143]).

One bottleneck in the performance of VO2 modulators is that the transverse electric (TE) polarized mode interacts weakly with them compared to the transverse magnetic
(TM) mode. Some solutions have been proposed to avoid this problem such as using a vertical slot coupler with VO$_2$ active layer [151], delocalization of mode in a small waveguide [149], and using a modulator coupled with a compact mode converter to improve the modulation [153]. These solutions try to optimize the trade-off between insertion loss and extinction ratio while also keeping the device footprint compact. In one such proposal, a layer of silver was added between the vanadium dioxide layer and the silicon layer. This unique hybrid plasmonic solution could reduce the insertion loss while maintaining a high extinction ratio [154].

As VO$_2$ is transitioned to a metallic state which is lossy for electromagnetic waves upon increasing the temperature, this feature opens up the possibility of using VO$_2$ as a switch in photonic circuits. This was shown by the work of Poon [152]. In this design, to make an effective switch of both TE and TM modes, a narrower silicon waveguide was used, and then it was clad with a VO$_2$ patch on all three sides. This increased the interaction of both TE and TM modes, helping in effective switching [152]. The device reported an extinction ratio of 12 dB and insertion loss of 5 dB. The device is electrically switched, having turn-off times of 350 ns and 750 ns, while not relying on hybrid plasmonic design helps to simplify the fabrication process. This design can be improved by using a two-terminal configuration, which helps in applying a large electric field without suppressing the leakage current, thereby drastically reducing the switch-on time to 2 ns [141].

Some interesting studies have been conducted to study the transition dynamics for improvement of switching time of electrically switched VO$_2$-based devices [160,161] and optically switched VO$_2$-based devices [162,163]. For electrically switched devices, the switching time dominated by a spatially percolated metallic phase [161] is improved by increasing the peak pulse [160]. Similarly, in the case of optically switched devices, the switching time can range from a few nanoseconds to microseconds depending on the incident power [162], and a faster relaxation to insulating state can be achieved by using doped TiO$_2$:Nb substrate [163].

6.2. Chalcogenide Modulators and Switches

VO$_2$-based devices are good for volatile applications requiring reasonably high switching speed, but the states are volatile. On the other hand, as seen earlier, chalcogenide devices show nonvolatile switching, making them useful for nonvolatile applications but at the expense of speed, a bottleneck that has been broken with recent research. Moreover, as maintaining the state in a volatile switch may require a constant supply of energy, this may not be a very energy-efficient method of switching. The advantage of only supplying energy when switching is needed instead of a constant supply of energy for maintenance of the state was highlighted in the work of Xu in reports on GST-based electro-optical modulators [14,148]. In the work by the group of Xu on GST [14], a modulator for C-band was presented that is electrically switched, has a footprint of 0.2 $\mu$m$^2$, has an extinction ratio larger than 5.4 dB for a switch, and consumes sub-nJ energy per cycle for each switch. This is achieved by a copper-clad 30 nm patch of GST on top of the silicon waveguide. This combination performs better than the various other recipes, such as the silicon–ITO–GST waveguide (shown in Figure 10a) [89] using an ITO heater instead of a copper heater on a straight waveguide, the abovementioned Si–VO$_2$ ring resonator-based modulator [142], Si–GST micro-ring [150], and silicon reverse-biased ring made using oppositely poled silicon and modulated through electrical means.
As silicon suffers from large two-photon absorptions in the near-infrared range, it is considered useful to use silicon nitride instead of silicon for near-infrared devices. For this reason, GST on silicon nitride-based modulators have been a popular subject of research [139,163,164].

However, in general, silicon remains the most frequently used substrate material for on-chip modulation and switching applications based on PCMs. Using GST encapsulated by ITO for electrical switching purposes was also demonstrated by Zhang [7]. In this work, a GST patch was placed on top of a silicon MMI, and switching GST from amorphous to crystalline resulted in transmission contrast of 20 dB over the wavelength range of 1500 nm to 1600 nm. This was achieved using a 20 ns write pulse of 10.4 nJ resulting in GST amorphization, and a 100 ns erase pulse of 9 nJ resulting crystallization of the GST patch. They additionally showed partial crystallization of the GST leading to multiple possible states of crystallization. In separate work, the group also demonstrated the feasibility
of using GST-based electrical switching encapsulated by ITO for multiple applications including wavelength division multiplexing and filtering [91]. As also discussed earlier, the interaction region in the straddling PCM is less than an in-line PCM. Placing the GST in line improves the interaction of optical wave and GST, leading to improved switching performance, such as a higher extinction ratio of 33.1 dB and low insertion loss of 0.48 dB for an optical wavelength of 1550 nm [87].

The arrangement of in-line GST for increased interaction has also recently been used for metal–insulator–metal (MIM)-type plasmonic waveguide switches and modulators, where GST is externally switched through pump light [156]. This work investigated three MIM structures such as an end-coupled rectangular resonator, a side-coupled stub resonator, and two mutually coupled resonators. The end-coupled rectangular resonator, the side-coupled stub resonator, and two mutually coupled resonators showed insertion loss of 3.6 dB for fully amorphous and 13.8 dB for crystalline, 16.8 dB for fully amorphous and 2.4 dB for fully crystalline, and 2.6 dB for fully amorphous and 17 dB for fully crystalline states, respectively. The modulation depth reported for these three structures was 13.8 dB, 14.4 dB, and 14.4 dB, respectively, with a switching speed of 500 ps. This modulation depth is intermediate among that achieved in similar work such as Haddadpour’s magneto-optical switch [164] and Piao’s Fano transparency controlled switching [165].

Another method to produce a sort of hybrid between an in-line and a matched PCM layer was introduced by Liang, whereby a 10 nm GST layer was sandwiched between two doped silicon layers in an arrangement also called “anti-slot” [166]. The author simulated electrically switched switches and modulators for an operating wavelength of 2.1 µm and was able to numerically validate an MZI with a device length of 38 µm and insertion loss of –16 dB, while achieving a state transition time of less than 100 ns. In subsequent work, the group also numerically demonstrated modulation of 2.1 µm wavelength using 2 × 2 devices along with one- and two-island waveguides between them [167].

Power consumption in GST-based systems is an important issue, which has led to some researchers exploring other platforms such as Sb2S3 [45,47], Sb2Se3 [158], and GSST [155] for switching and modulation applications. Given that these are newer platforms, there has not been a lot of progress in this direction, but they are promising. The Sb2S3- and GSST-based switches reported lower insertion losses of 0.48 dB and 0.135 dB [47,155], device lengths of 8 µm and 5 µm, and extinction ratios of 30 dB and 20 dB, respectively. Sb2Se3-based modulators using ring resonators as the basic structure demonstrated an insertion loss of 0.45 dB and switching energy of 185 nJ for an electrically switched device [158].

The low-loss nature of these platforms has also encouraged nonconventional applications for PCMs such as MMI switches [45]. On the other hand, GST-based novel designs have also been proposed, which tend to address the power consumption issue of GST-based devices, producing ultra-low-power nonvolatile switches, by using a GST-filled slot instead of using a GST patch for the whole width of the waveguide. The proposed design (shown in Figure 10c,f) is electrically switched and can reach an extinction ratio of 17 dB at 1550 nm with a high energy efficiency of 64 aJ/nm3 for the crystallization and amorphization transitions [98]. However, these are simulation-based results, and whether an actual GST-based device is able to give such a performance in an in-line setting remains to be seen.

7. Discussion and Outlook

Phase change materials are unique in that they drastically change their electrical and optical properties upon the introduction of an external stimulus. Even though these materials have been around for some decades, they have come only recently into focus for designers of on-chip devices. Since the early devices in the early 2010s, a lot of progress has been made in the direction of on-chip PCM-based devices. While most of the progress has been directed toward application in memory devices and neuromorphic computational platforms, a lot of progress has also been made in applications such as switches and modulators. Most of the major contributions in the areas of memory and neuromorphic
computing are tabulated in Tables 3 and 4, while most of the major contributions in switches and modulators are tabulated in Table 5.

GST has by far been the most successful phase change material in terms of acceptability among the research community. This has led to the use of GST in every possible conventional application of PCMs including memory devices, neuromorphic computational platforms, photonic switches, and modulators, which have been implemented using electrical switching, as well as optical switching. While optical switching methods are difficult to scale at present due to the difficulties in the routing of control signals and data signals, electrical switching methods overcome these issues due to inherent differentiation in the routing circuits for control and data signals.

The increased interest in energy conservation through energy efficiency has led to the exploration of material platforms for PCMs that have a low loss. Among these low-loss material platforms, Sb$_2$Se$_3$, Sb$_2$S$_3$, and GSST are the most prominent materials. GSST is a newer platform while Sb$_2$Se$_3$ and Sb$_2$S$_3$ are old but not forgotten materials. This exploration reintroduced these low-loss chalcogenide phase change materials. Although these low-loss materials improve the energy performance of these applications, they also enhance the horizon of possible applications with the induction of devices such as MMLs and beam splitters into the possible applications of on-chip PCMs. This horizon is expected to further increase in the near future with the possible on-chip demonstration of various computational and beam steering platforms already available for free-space applications but difficult to implement on chip due to the losses of GST. Such applications can be implemented on the on-chip platforms using such low-loss PCMs.

After these chalcogenide materials, the most prevalent material is VO$_2$. Although, in terms of refractive index contrast, it is comparable with GST which could potentially make it a rival technology, its states are volatile and can only be maintained through a continuous supply of energy making them only suitable for volatile applications. Due to this property of VO$_2$, its performance does not fare well with the contemporary energy conservation and energy efficiency frameworks. Earlier, it had the advantage of faster switching over GST, but that advantage is also slipping due to new and improved switching methods for chalcogenide PCMs, which means that, although chalcogenide PCMs need activation of control circuitry at each switch between crystalline and amorphous phase, they are still able to achieve faster switching compared with VO$_2$ devices. Nevertheless, VO$_2$ has still found some applications in switching and modulation, where the performance of VO$_2$-based devices is not better than its chalcogenide-based counterparts.

At present, the research in nonconventional material platforms such as silicon and organic materials is seriously lacking. Although silicon has been demonstratively used in a few applications through poling and doping to achieve contrasting phases, the contrast is small, restricting its possible use in actual devices. On the other hand, organic PCMs have not been used at all in photonic devices. However, given their increasingly successful use in resistance-based applications, it is only a matter of time before they also find their way into on-chip refractive index-based devices. The literature reports both volatile and nonvolatile organic PCMs with demonstrable application in resistance-based memristive memory devices. The volatile category includes materials such as PNIPAMs and PEDOTs, while the nonvolatile category includes materials made from aromatic compounds. While the change of phase in these organic PCMs leads to a change in their resistance, due to the lack of research in this area, it remains to be seen whether the phase change also corroborates with a change in the refractive index. Only those materials which show considerable refractive index contrast will be most useful for photonic applications.

The phase change materials and, thus, the devices based on them are deemed essential for next-generation low-power memory devices and neuromorphic computational platforms. The introduction of low-loss materials will be very helpful in the diversification of possible PCM applications in the future. Such diversification is extremely essential to realize one day a fully functional all-optical computing device of conventional and neuromorphic kind. The current developments in this field are healthy but at a very early
stage for the realization of practical devices, implying that there is still a long way to go. The future will hold many more interesting applications of PCMs among those discussed above or even newer material platforms with more interesting properties aligning with the needs of specific applications.

**Author Contributions:** Conceptualization, M.S.N. and L.Z.; writing—original draft preparation, M.S.N.; writing—review and editing, M.S.N., X.Y., L.L., J.C. and L.Z.; supervision, L.Z.; funding acquisition, L.Z. All authors read and agreed to the published version of the manuscript.

**Funding:** This research was funded by the National Key Research and Development Program (2019YFB2203200, 2018YFB2201702, 2019YFB1802903), the National Natural Science Foundation of China (NSFC) (62090052, 620730193), and the Shanghai Municipal Science and Technology Major Project (2017SHZDZX03).

**Institutional Review Board Statement:** Not applicable.

**Informed Consent Statement:** Not applicable.

**Data Availability Statement:** Not applicable.

**Conflicts of Interest:** The authors declare no conflict of interest.

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