Antioxidant Materials Based on 2D Nanostructures: A Review on Recent Progresses

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Abstract: Counteracting reactive oxygen species (ROS, e.g., superoxide radical ion, \( \text{H}_2\text{O}_2 \) and hydroxyl radical) is an important task in fighting against oxidative stress-related illnesses and in improving product quality in industrial manufacturing processes. This review focuses on the recent advances on two-dimensional (2D) nanomaterials of antioxidant activity, which are designed for effective decomposition of ROS and thus, for reduction of oxidative stress. Some materials featured in this paper are of uni- or multi-lamellar structures modified with small molecular or enzymatic antioxidants. Others are enzyme-mimicking synthetic compounds (the so-called nanozymes) prepared without antioxidant additives. However, carbon-based materials will not be included, as they were extensively reviewed in the recent past from similar aspects. Given the landmark development around the 2D materials used in various bio-applications, sheet-like antioxidant compounds are of great interest in the scientific and technological communities. Therefore, the authors hope that this review on the recent progresses will be helpful especially for researchers working on novel developments to substantially reduce oxidative stress either in biological systems or industrial liquors.

Keywords: antioxidant activity; 2D nanomaterial; nanozyme; clay; enzyme; chalcogenide; metal organic framework; oxidative stress; reactive oxygen species; radical scavenge

1. Introduction

Nanomaterials with two-dimensional (2D) structure have attracted notable scientific interest due to their high surface area and intriguing properties arising from their large variety of composition [1–5]. Starting with their light element-based representatives (e.g., graphene, graphite, graphite oxides, carbon nitride, boron nitride and covalent organic frameworks), through metal chalcogenide sheets, layered double hydroxides (LDHs), montmorillonites (MMTs), metal organic frameworks (MOFs) to thin noble metal platelets, one of the main advantages is that their composition is tunable for specific applications. In addition, 2D materials possess unusual electronic, mechanical and optical properties and hence, they can be used as building blocks in a wide range of composites. Concerning the contemporary applications of 2D compounds, they serve as efficient catalysts for water splitting [6–15] or organic reactions [16–21], drug carriers [22–31], sensors [32–39], materials for energy production and storage [2,15,21,40–49] or adsorbers [50–55].

Biocompatible 2D materials gained considerable popularity in the biochemical community, as indicated by the rapidly growing number of their bio-medical applications [56–58]. Among them, 2D antioxidant composites were developed [59–61] and used in decomposition of reactive oxygen species.
species (ROS) [62] and subsequently, to reduce oxidative stress in living and industrial systems. Tremendous improvement has been accomplished in the number and variety of antioxidant materials obtained over this period and the following approaches were reported (Figure 1): (i) immobilized natural and artificial enzymes, (ii) 2D materials loaded with molecular antioxidants and (iii) 2D composites of radical scavenging activities (nanzymes). In the present contribution, the antioxidant characteristics of such 2D materials are summarized emphasizing metal-based nano-objects. The carbonaceous materials of 2D structure were greatly investigated in the recent past and collective reviews were published on their applications [57,63–79]. Therefore, carbon-based 2D materials will not be discussed here.

![Figure 1. Illustration of molecular and enzymatic antioxidants immobilized in/on layered 2D materials. Schematic representation of superoxide radical anions and H₂O₂ is also presented.](image)

2. Natural Defense Systems versus Oxidative Stress

The most effective defense systems against reactive oxygen species (ROS), which induce oxidative stress at higher concentrations, are the antioxidant enzymes [80]. ROS include, for instance, superoxide radical anion, H₂O₂, hydroxyl and alkoxy radicals of short life time, but of extremely high reactivity [62]. They damage cell constituents leading to the development of various diseases such as chronic inflammation, neurological disorders and cancer [81]. On the other hand, efficient defense against ROS is also required in industrial manufacturing processes (e.g., food, cosmetic and textile industry) to improve the quality of the products [82,83].

Superoxide anion is one of the most notable ROS, since its decomposition by the superoxide dismutase (SOD) enzyme leads to the formation of molecular oxygen and H₂O₂. Therefore, the dismutation reaction is also a ROS source. Enzymatic assays were developed to estimate the SOD activity. In the most popular tests, radicals are produced in enzymatic ways and indicator molecules change their color upon reduction by the superoxide anions [84]. Inhibition of the radical-indicator reaction by the enzyme is the measure to assess the dismutation ability.

The two common enzyme groups that can break up H₂O₂ are catalases and peroxidases [85,86]. During neutralization of harmful peroxides, catalases generate O₂ gas and H₂O, while peroxidases perform homolytic bond cleavage on H₂O₂ (Figure 1). The as-obtained hydroxyl radicals (OH⁻) either are consumed by an appropriate substrate near the active site or converted to water. The antioxidant enzymes decompose ROS in tandem reactions in the intracellular environment, e.g., H₂O₂ produced by SOD is subsequently decomposed by catalase (see graphical abstract).
Besides antioxidant enzymes, molecular antioxidants also play an important role in reducing oxidative stress. They involve vitamins, carotenoids and flavonoids, for instance [87]. Supplementation from certain foods is feasible, however, their activity in ROS decomposition lags behind the enzymatic antioxidants. Non-enzymatic routes are also established to estimate antioxidant capacity. Methods based on free radicals involve 2,2-diphenyl-1-picrylhydrazyl (DPPH) [88], nitrogen monoxide (NO) [89], radical form of 2,2′-azino-bis(3-ethylbenzthiazoline-6-sulfonic acid) (ABTS⁺) [90] and peroxynitrite ion (ONOO⁻, prepared by the direct, cold peroxidation of nitrite salts) [91], which are relatively stable and can be reduced by the antioxidant of choice.

In addition, Fenton reaction, which is a catalytic route to decompose H₂O₂ in the presence of metal ions, especially Fe²⁺ or Fe³⁺, generating aggressive OH⁻ and OOH⁻ radicals, therefore used for strong oxidations and thus, it is a radical source to probe enzymatic antioxidant compounds [92]. Two indirect radical-based tests include thiobarbituric acid (TBA) as substrate [93]. The degradation of 2-deoxyribose by Fenton reaction [94] and egg lipid peroxidation [95] produces small dialdehydes, mainly malondialdehyde that gives a colored condensation product with TBA. If OH⁻ radicals in Fenton kinetics or the lipid damaging radicals are consumed by antioxidants, the color in the reactions with TBA are less profound. Further, two assays take oxidative metal complexes as reagents with Fe²⁺ (Ferric reducing antioxidant power assay (FRAP) [96], with 2,4,6-tris(2-pyridyl)-s-triazine ligand) and Cu²⁺ (cuppic reducing antioxidant capacity (CUPRAC) assay [97], with 2,9-dimethyl-1,10-phenanthroline ligand) central ions. The latter test has a kinetic advantage as the Cu²⁺ complex is easier to reduce. For detailed antioxidant effect of various materials, the reader is referred to the works of Nimse and Pal [98] and Valgimigli et al. [99].

Due to the increased harmful environmental effects such as air pollution, food additives and radiations, the ROS level increased and thus, oxidative stress became an important issue nowadays. The supplementation of the above natural antioxidants is complicated due to their high sensitivity to the environmental conditions (enzymatic) or limited water solubility (molecular). Therefore, their heterogenization in composite materials, which provides sufficient protection and solubilization of the antioxidants, is a promising research direction.

3. Metal Oxide Structures

3.1. Cobalt-Based 2D Oxides

The intrinsic peroxidase and catalase activity of Co₃O₄ was first described in 2012 for cubic nanoparticles [100] (NPs) and was proven at several occasions for spherical NPs [101] and architectures consisting of Co₃O₄ nanorods [102]. Morphology-dependent study was also conducted, revealing an order of nanocubes < nanorods < nanoplates (Figure 2) in catalytic activity, pointing out that the efficiency depends on the most exposed crystal plane [103]. The same correlation was found by Zhang et al. who were able to produce polyhedrons of Co₃O₄, possessing approximately 55% of peroxidase and 70%–80% of catalase activity, compared to the 2D platelets [104]. Data of the quantitative assessment of the activity, together with other 2D antioxidant systems discussed in this feature article, are given in Table 1.

Table 1. Two-dimensional 2D nanomaterials of antioxidant properties and the quantitative assessment of their activities in test reactions.

| Materials                     | Mimicked Enzymes | Activity Assessmentᵃ | Ref.  |
|-------------------------------|------------------|----------------------|-------|
| Cu(4-hydroxythiophenol)       | Peroxidase       | H₂O₂, TMB            |       |
| Nanosheets                    | Kₘ (mM)          | 0.716, 0.431         | [19]  |
| MoS₂                           | Peroxidase       | H₂O₂                 |       |
| Nanosheets                    | Kₘ (mM)          | 0.003, 3.4           | [39]  |

SOD Fridovich methodᵇ IC₅₀ (mg/L)ᶜ
| System                          | Enzyme 1 | Enzyme 2 | Michaelis Menten Parameters | Others |
|--------------------------------|----------|----------|-----------------------------|--------|
| Mg/Al-CO$_3$-LDH-SOD           | SOD      | Fridovich method $^b$ | $IC_{50}$ (mg/L) | [61]   |
| Mg/Al-CO$_3$-LDH-SOD-Hep       | SOD      | Fridovich method $^b$ | $K_m$ (mM)     |        |
| Co$_3$O$_4$ Sisal-like Structures | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 0.056  |
| Co$_3$O$_4$ Nanoplates         | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 0.096  |
| Co$_3$O$_4$ Nanoplates         | Peroxidase | H$_2$O$_2$ | $v_{max}$ (10$^{-8}$ M/s) |        |
| Co$_3$O$_4$ Microbelts         | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 0.057  |
| Co$_3$O$_4$-CeO$_2$ Nanosheets | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 0.015  |
| Co$_3$O$_4$ Nanosheets         | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 132.2  |
| V$_2$O$_5$ Nanosheets          | Glutathione Peroxidase | GSH | $K_m$ (mM)     | 2.22   |
| VO$_2$ Nanosheets              | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 2.924  |
| HRP-Titanate Nanosheets        | HRP      | Guaiacol | $K_m$ (mM)     | 2.06   |
| SOD-Titanate Nanosheets        | SOD      | Hypoxanthine/XO Cytochrome c | $IC_{50}$ (U) | 0.036  |
| HPR-TiO$_2$ nanosheets-magnetic beads | HRP | Guaiacol | $K_m$ (mM)     | 1.8    |
| Au-MoS$_2$ Nanoribbons         | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 0.105  |
| MoS$_2$-Co(OH)$_2$ Nanoflakes  | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 0.127  |
| MoSe$_2$ Nanosheets            | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 0.155  |
| VS$_2$ Nanosheets              | Peroxidase | H$_2$O$_2$ | $K_m$ (mM)     | 3.49   |
|                             | TMB   | Km (mM) | v<sub>max</sub> (10<sup>-8</sup> M/s) | Reference |
|-----------------------------|-------|---------|-------------------------------------|-----------|
| **Cu-Boron Nitride Nanosheets** Peroxidase |       |         |                                     |           |
| H<sub>2</sub>O<sub>2</sub> | 25    | 1.775   | 12.5                                | [116]     |
| TMB                          | 0.175 | 3.76    |                                     |           |
| **Ni-MOF Nanosheets** HRP |       |         |                                     |           |
| H<sub>2</sub>O<sub>2</sub> | 2.49  | 3.65    | 130                                 | [117]     |
| TMB                          | 0.365 | 6.53    |                                     |           |
| **HRP-LDH** HRP Guaiacol |       |         |                                     |           |
| Km (mM)                      | 5.25  | 272     |                                     | [118]     |
| v<sub>max</sub> (10<sup>-8</sup> M/s) |       | 6.53    |                                     |           |

**Carnosine-MgAl-LDH**
- Scavenging activity 95.9%

**Gallic acid-MgAl-LDH**
- Scavenging activity 83.9%

**Gallic acid-ZnMgAl-NO<sub>3</sub>-LDH**
- Scavenging activity 93%

**Organic solvent treated EA-LDH**
- Scavenging activity up to 80-90%

**IrganoxCOO-H-MgAl-LDH**
- Scavenging activity 5.25

**Trolox-MgAl-LDH**
- Scavenging activity 0.24

**EC<sub>50</sub> (µg/mL)**
- DPPH
  - Olanzapine-CaAl-LDH
  - Olanzapine-NiAl-LDH
  - EC<sub>50</sub> 8.541
  - EC<sub>50</sub> 11.630

**SOD-Hep-LDH**
- SOD Fridovich Method
  - IC<sub>50</sub> (nM)
  - Cu(Hsm)<sub>2</sub>-PVPMAA-LDH
  - IC<sub>50</sub> 2.9
  - IC<sub>50</sub> 97

**DNA-CoAl-LDH** Peroxidase
- H<sub>2</sub>O<sub>2</sub>
  - Km (mM) 10.24
  - Km (mM) 1.775
  - v<sub>max</sub> (10<sup>-8</sup> M/s) 2.3
  - v<sub>max</sub> (10<sup>-8</sup> M/s) 4.09

**[Bael oil proteins]-MgAl-LDH**
- Scavenging activity 42-50%

**CoAl-LDH** Peroxidase
- H<sub>2</sub>O<sub>2</sub>
  - Km (mM) 22.13
  - Km (mM) 0.372
  - v<sub>max</sub> (s<sup>-1</sup>) 0.598
  - v<sub>max</sub> (s<sup>-1</sup>) 0.101

**NiFe-LDH** Peroxidase
- H<sub>2</sub>O<sub>2</sub>
  - Km (mM) 2.4
  - Km (mM) 0.5
| Material                          | Enzyme or Assay  | Substrate  | $K_m$ (mM) | $v_{max}$ $(10^{-8} \text{ M/s})$ | Ref. |
|----------------------------------|------------------|------------|------------|----------------------------------|------|
| Polyethylene Glycol-Fe-LDH      | Peroxidase       | H$_2$O$_2$ | 0.09       | 176                               | [129]|
| AEA-GSH-MMT                     | $-$               | ABTS      |            | Scavenging activity 59.88%       | [130]|
| Blue Berry Extract-MMT          | $-$               | DPPH      |            | Scavenging activity 38%          | [131]|
| PHB-TPS/eugenol-MMT             | $-$               | DPPH      |            | Scavenging activity 92%          | [132]|
| Hemin-Hsm-MMT                   | HRP              | Guaiacol   |            | Rate constant (A/min) 0.107      | [133]|
| TOC-CSNPs/CS/M MT               | $-$               | DPPH      |            | Scavenging activity 46.5%        | [134]|
| ZnS-MMT                         | Peroxidase       | H$_2$O$_2$ | 0.0254     | 0.750                             | [135]|
| CuS-MMT                         | Peroxidase       | H$_2$O$_2$ | 2.27       | 0.971                             | [136]|
| AgS-MMT                         | Peroxidase       | H$_2$O$_2$ | 1.874      | 2.286                             | [137]|
| CoNiS-3/MMT                     | Peroxidase       | H$_2$O$_2$ | 1.247      | 6.92                              | [138]|
| HRP-MMT                         | HRP              | Phenol    | 12.96      | 7150                              | [139]|
| Cu–Zn complex-MMT               | SOD              | Fridovich method$^b$ | IC$_{50}$ ($\mu$M) | 91.0   | [140]|
| Cu-histidine-MMT                | SOD              | Fridovich method$^b$ | IC$_{50}$ ($\mu$M) | 251    | [141]|
| Ni Foam-CoP Nanosheets          | Peroxidase       | H$_2$O$_2$ | 4.90       | 0.54                              | [142]|
| Co(OH)$_2$:CO$_3$-CeO$_2$ Nanneosheets | Peroxidase       | H$_2$O$_2$ | 10.01      | 10.21                             | [143]|
| Fe(PO$_4$)$_3$-H$_2$O Nanoflowers | Peroxidase       | H$_2$O$_2$ | 0.11       | 5.58                              | [144]|
| Porous Iron Oxide               | Peroxidase       | H$_2$O$_2$ | 150.47     | 3.12                              | [145]|

$^a$ The columns contain the applied substrate or assay and the quantitative measure of the activity.

$^b$ The method is described in reference [84].

$^c$ Concentration of enzyme or enzyme mimic necessary to decompose 50% of the radicals forming in the test reaction.

$^d$ General radical scavenging activity was assessed, no enzymatic assays was applied.

$^e$ Top values belong to olanzapine-CaAl-LDH and the bottom ones to olanzapine-NiAl-LDH.
Abbreviation list: TMB: 3,3',5,5'-tetramethylbenzidine, SOD: superoxide dismutase, GSH: glutathione, HRP: horseradish peroxidase, XO: xanthine oxidase, DPPH: 2,2-diphenyl-1-picrylhydrazyl, CUPRAC: cupric reducing antioxidant capacity, ABTS: 2,2'-azino-bis(3-ethylbenzthiazoline-6-sulfonic acid).

Figure 2. TEM image of a typical Co$_3$O$_4$ nanoplate (a), high-resolution transmission electron microscopy (HRTEM) images of site 1 (a), site 2 (c) and site 3 (d) of Co$_3$O$_4$ nanoplate, a HRTEM image of a Co$_3$O$_4$ nanorod (e), a HRTEM image of a Co$_3$O$_4$ nanocube (f) and exposed crystal planes on the nanoplate, nanorod and nanocube of Co$_3$O$_4$ (g). Reproduced from Reference [103] with permission from the PCCP Owner Societies.

Elongated 2D Co$_3$O$_4$ belts were obtained by electrospinning technique from Co(NO$_3$)$_2$ containing polyvinylpyrrolidone gel with superior affinity compared to the native horseradish peroxidase (HRP) enzyme and up to 100-fold higher activity was calculated from the Michaelis constants ($K_m$) determined in the test reactions involving H$_2$O$_2$ and TMB (3,3',5,5'-tetramethylbenzidine). Besides, the belts were found to be stable at significantly higher temperature than HRP [105]. The high-resolution transmission electron microscopy (HRTEM) images of the belts obtained reflect characteristic basal distances of 0.24 and 0.29 nm assigned to (311) and (220) interplanar spacings in Co$_3$O$_4$, respectively.

Platelets of Co$_3$O$_4$ were combined with CeO$_2$ sheets to a nanocomposite of relatively low affinity towards H$_2$O$_2$, as indicated by higher $K_m$ values, but excellent maximum reaction rate ($v_{max}$) was observed in the diffusion controlled regime [106]. Separately, both oxides are considered as peroxidase nanoenzymes that was reinforced in the composite, which was later used as a paper based analytical glucose sensor with smartphone software quantification.

In a recent finding, porous Co$_3$O$_4$ nanoplates of pH-switchable peroxidase-like (acidic medium) or catalase-like (basic medium) properties was synthesized [107]. Overall, the porous flakes possessed remarkable kinetic parameters and were used as a glucose sensor enabled by glucose oxidase (GOx)-glucose selective reaction (Figure 3).
Finally, a mixed-valance state lamellar cobalt oxide with non-stoichiometric composition was also used as a nanozyme to oxidize TMB substrate with molecular O$_2$ \cite{146}. This structure was synthesized by partial oxidation of Co(II)-trimesate and was applied as a glutathione (GSH) sensor.

### 3.2. Vanadium-Based Oxides

Ghosh et al. extensively examined the morphology-activity correlation of orthorhombic V$_2$O$_5$ structures of peroxidase activity \cite{108}. Unlike Co$_3$O$_4$ \cite{104}, they found that surface area is not a main factor in activity, but it is the accessibility of active sites on the dominant crystal faces. The GSH-mediated peroxidase activity and surface area of the nanomaterials obtained are depicted in Figure 4 (with V$_2$O$_5$ abbreviated as V), showing that V$_2$O$_5$ spheres reached the highest maximum rate, followed by flowers, sheets and wires, out of which only wires and 2D sheets are saturated at the lowest H$_2$O$_2$ concentration.
Figure 4. SEM and TEM (inset) images of V$_2$O$_5$ nanowires (V$_{NW}$) (a), V$_2$O$_5$ sheets (V$_{Sh}$) (b), V$_2$O$_5$ nanoflowers (V$_{Nf}$) (c) and V$_2$O$_5$ spheres (V$_{Sp}$) (d). Reduction of H$_2$O$_2$ by GSH in the presence of nanomaterials, glutathione reductase (GR) and NADPH (e). The GSH peroxidase-like activity using H$_2$O$_2$, tert-butyl hydroperoxide (t-BuOOH) and cumene hydroperoxide (CuOOH) (f). Michaelis–Menten plot for the nanozymes (g). Trends in $v_{max}$ and surface area (h). Copyright (2020) Wiley. Used with permission from (Ghosh, S.; Roy, P.; Karmodak, N.; Jemmis, E.D.; Mugesh, G. Nanoisozymes: Crystal-facet-dependent enzyme-mimetic activity of V$_2$O$_5$ nanomaterials. Angew. Chem. Int. Ed. 2018, 57, 4510–4515) Reference [108].

Aside from its peroxidase mimic, a layered V$_2$O$_5$ 2D structure was also used to detect GSH via an oxidase-type mechanism [147]. Lower valence state vanadium oxide, VO$_2$ was investigated in various morphologies [109]. Fibrillar, lamellar and elongated NPs were tested against TMB substrate and it was pointed out that VO$_2$ sheets exhibited the best kinetic overall parameters in regard of $K_m$ and $v_{max}$. Additionally, the aforementioned VO$_2$ particles were applied as H$_2$O$_2$ sensors with the nanosheets providing linear range up to 62.5 mM H$_2$O$_2$, four times higher than the second best performing nanorods.
3.3. Titania-Based Composites

The prominent photocatalytic activity of pure and doped TiO$_2$ is known for decades [4]. Despite Ti(IV) is moderately redox active, TiO$_2$ is not an antioxidant nanozyme. However, its negative surface charge in a broad range of pH makes it a suitable vehicle for HRP immobilization, for instance. Xie et al., constructed hollow titania spheres based on sheets as building blocks, capable storing 20 mass% HRP owing to the porous structure [148]. This highly loaded material was used for electrochemical H$_2$O$_2$ sensing with linearity over three orders of magnitudes.

Moreover, titania materials were proved as excellent solid supports for other antioxidant enzymes due to their biocompatibility, high abundance of surface functionalities and chemical inertness. Surface modifications with polyelectrolytes resulted in stable structures capable of hosting various enzymes. Accordingly, lamellar titania nanosheet (TNS)-based nanocomposite with this setup had peroxidase activity nearly identical to the bare HRP (Figure 5) [149]. An adsorbed positively charged poly(diallyldimethylammonium chloride) (PDADMAC) polyelectrolyte layer improved the enzyme’s structural integrity over longer timeframes and maintained a good colloidal stability for the platelets.

![Figure 5](image-url)

**Figure 5.** Illustration of HRP (a) and SOD (b) immobilization on titania nanosheet (TNS) of saturated poly(diallyldimethylammonium chloride (PDADMAC) polyelectrolyte layer on the surface. Reprinted with permission from (Rouster, P.; Pavlovic, M.; Saringer, S.; Szilagyi, I. Functionalized titania nanosheet dispersions of peroxidase activity. *J. Phys. Chem. C* **2018**, *122*, 11455–11463) Reference [149]. Copyright (2020) American Chemical Society (a). Copyright (2020) Wiley. Used with permission from (Rouster, P.; Pavlovic, M.; Szilagyi, I. Immobilization of superoxide dismutase on polyelectrolyte functionalized titania nanosheets. *ChemBiochem* **2018**, *19*, 404–410) Reference [61] (b).

The same TNS was used to immobilize SOD enzyme [61]. Surface functionalization was carried out first with PDADMAC to obtain positively charged sheets and the enzyme was attached through electrostatic and hydrophobic interactions. The same superoxide anion radical scavenging activity of the free and attached SOD indicated that the enzyme kept its structural integrity upon immobilization.

Kamada et al. demonstrated that nanometric titanate nanosheets enhanced the catalytic properties of aqueous HRP and SOD [110]. It was claimed that TNS served as a surfactant and helped to decrease the number of HRP and SOD aggregates in the solution, exposing more catalytic centers for the reactions to occur. This was possible due to electrostatic attraction, thus immobilization of the enzymes on TNS particles, as indicated by dynamic light scattering measurements.

Stable suspensions were obtained by immobilizing HRP and magnetic beads on positively charged layered titania in slightly acidic samples [111]. Although steric hindrance and possibly slight conformational changes led to reduced HRP activity, the magnetic composite was separable with strong magnets at the end of catalytic cycles. Furthermore, the reused material maintained 80% of its original activity after 5 repetitions.
4. Chalcogenide Structures

Among the other lamellar chalcogenides, molybdenum compounds were inspected thoroughly. Pure MoS$_2$ has multienzymatic activity under physiological conditions. It was reported to have the ability to scavenge superoxide radicals generated by the xanthine/xanthine oxidase system and to degrade H$_2$O$_2$ in catalase and peroxidase-type manners [59]. Although the generation of OH$^\cdot$ radicals was evident by using DMPO (5,5-dimethyl-pyrroline N-oxide) as a radical trap for electron paramagnetic resonance (EPR) spectroscopy measurements, the OH$^\cdot$ radical products were transformed to H$_2$O, proven by the scavenging of Fenton-type developed radicals. Furthermore, the 2D MoS$_2$ was used as a scavenger for other reactive species, e.g., NO and DPPH radicals, meaning that the nanozyme was an excellent oppressor of oxidative stress during in vitro measurements (Figure 6). Moreover, in vivo experiments revealed that MoS$_2$ sheets protected *E. coli* and *S. aureus* bacteria from H$_2$O$_2$-induced oxidative stress. While the nanozyme was fully biocompatible with *E. coli*, it expressed a moderate toxicity on *S. aureus*.

Figure 6. Illustration of the multi-fold antioxidant activity of MoS$_2$ nanosheets. Reprinted with permission from (Chen, T.M.; Zou, H.; Wu, X.J.; Liu, C.C.; Situ, B.; Zheng, L.; Yang, G.W. Nanozymatic antioxidant system based on MoS$_2$ nanosheets. *ACS Appl. Mater. Interfaces* 2018, 10, 12453–12462) Reference [59]. Copyright (2020) American Chemical Society.

Later, Nandu et al. showed that out of many proteins, lipase selectively masked the peroxidase activity of 2D MoS$_2$ [39], as indicated by the increasing $K_m$ values, i.e., decreasing affinity towards H$_2$O$_2$. The $v_{max}$ value was also decreased by more than 94% in the presence of lipase. Based on this principle, the MoS$_2$ NPs were used as a colorimetric lipase sensor.

The peroxidase activity of MoS$_2$ sheets can be improved by depositing Au NPs on the surface to drastically increase reaction rate (100-fold increment in $v_{max}$) and the affinity to TMB substrate (350-fold decrease in $K_m$) [112]. This system was also used as a probe in detection of free cholesterol.

In another study, 2D Co(OH)$_2$ flakes were combined with MoS$_2$ ions and treated hydrothermally to get a mainly amorphous CoMo(OH)$_x$S$_y$ or simply CoMo hybrid, perceived as MoS$_2$ doped Co(OH)$_2$ [113]. This novel material acted as a tri-enzymatic mimic with catalase, peroxidase and oxidase activity (Figure 7).
Figure 7. Absorbance intensity of TMB and 50 mM H$_2$O$_2$ upon the addition of different materials (a). Time-dependent absorbance of TMB at 652 nm varied with different catalysts used (b). O$_2$ generation from H$_2$O$_2$ decomposition with different catalysts (c). Catalytic activity comparison of CoMo hybrids and the supernatant coexisting with the CoMo hybrids (d). Visualized reaction scheme to obtain the CoMo hybrids (e). Reprinted with permission from (Ding, Y.Q.; Wang, G.; Sun, F.Z.; Lin, Y.Q. Heterogeneous nanostructure design based on the epitaxial growth of spongy MoS$_2$ on 2D CO(OH)$_2$(2) nanoflakes for triple-enzyme mimetic activity: Experimental and density functional theory studies on the dramatic activation mechanism. ACS Appl. Mater. Interfaces 2018, 10, 32567–32578) Reference [113]. Copyright (2020) American Chemical Society.

The molybdenous modification improved the Michaelis–Menten parameters of lamellar Co(OH)$_2$. Exfoliated MoSe$_2$ sheets with few hundreds nm lateral size also possessed superb peroxidase mimicking properties [114]. Although this structure’s maximum rate in H$_2$O$_2$
decomposition with TMB as chromogen is lower than that of native HRP's, the $K_m$ values observed are outstanding, making the material an effective sensor for xanthine and H$_2$O$_2$ detection (Table 1).

Huang et al. proved that VS$_2$ sheets catalyzes homolytic cleavage of H$_2$O$_2$ in a faster reaction than HRP and the nanozyme also possesses higher affinity to both H$_2$O$_2$ and TMB compared to the native enzyme [115]. The same mechanism was implied for a BN@CuS composite [116], suitable for detection of cholesterol and for an HRP@WS$_2$ hybrid [150], evidently. Several different sulfides were used as a composite building block with montmorillonites, which are detailed later.

5. Metal Organic Frameworks (MOFs)

MOFs are a subgroup of coordination polymers built up by metal ions coordinated and interlinked by multidentate bridging organic ligands [48]. In MOFs, the coordination network contains voids leading to enlarged surface area. Utilizing their favorable surface properties, Huang et al. constructed ca. 4 nm thick Cu-based MOFs with Fe or Co complexes of a porphyrin ligand (tetrakis(4-carboxyphenyl)porphyrin, TCPP) decorated with 2 nm spherical Au NPs [151]. While the bare MOF was attested to be a peroxidase mimic (in acidic buffers near pH 3−4, similar for all MOFs with peroxidase activity), Au deposition turned the material into a GOx mimic, after which the concentration of gluconic acid was measured by a photometric assay. Similarly, Qin et al. used the same Fe-TCPP in Zn, Co and Cu-MOFs as peroxidases with altered activity in the presence of numerous phosphates of biological relevance [152]. Based on their activity change, the peroxidase probe reaction was utilized to detect the biomolecules mixed with the MOFs. Ultrathin (2 nm) variants of the same Cu and Co-MOFs, alongside a Ni-containing one, were also synthesized and used as a modifier on glassy carbon electrodes fused with carbon nanotubes or graphene oxide [49].

Although MOFs are noteworthy peroxidase nanozymes under pH 5, this value makes it hard for researchers to utilize them during in vivo experiments. Bridging this gap, a cascade system was fabricated using the well-known Cu-MOF applying Fe-TCPP in the synthesis [153]. In the first step, GOx was immobilized on the framework that oxidized glucose in the presence of molecular O$_2$. During this reaction, the pH dropped (Figure 8), therefore the generated H$_2$O$_2$ was decomposed by a self-activated peroxidase mechanism. The resulting OH$^-$ radicals are strong potent antibacterial agents, decimating $S$. aureus bacteria after injecting onto mice wound using a band-aid containing the biocompatible GOx@MOF and glucose.
Figure 8. TEM micrograph of Cu-TCPP(Fe) 2D MOF nanosheets (a). AFM image of Cu-TCPP(Fe) 2D MOF nanosheets and their thickness distribution (b). Darkfield TEM image of typical Cu-TCPP(Fe) 2D MOF nanosheets and the corresponding TEM element mappings (c). TEM image of Cu-TCPP(Fe) 2D MOF/GOx (d). Time-dependent color changes of TMB reaction solutions catalyzed by the Cu-TCPP(Fe) 2D MOF/GOx every 30 min in pH 7.4 PBS buffer (0–240 min, e) or pH 7.4 PBS buffer (0–240 min, f). Methyl red pH indicator (0.001%) was used to signal the developing acidic conditions. Reprinted with permission from (Liu, X.P.; Yan, Z.Q.; Zhang, Y.; Liu, Z.W.; Sun, Y.H.; Ren, J.S.; Qu, X.G. Two-dimensional metal-organic framework/enzyme hybrid nanocatalyst as a benign and self-activated cascade reagent for in vivo wound healing. ACS Nano 2019, 13, 5222–5230) Reference [153]. Copyright (2020) American Chemical Society.

Chen et al. obtained Ni-MOF with p-benzenedicarboxylate ligands [117]. In acidic acetate buffer, TMB was oxidized via the peroxidase intrinsic activity of the material. The nanosheets had impeccable affinity towards the substrates, while maintaining high maximum rate. Since the selectivity of the catalyst was high, it was applied in the detection of H2O2 in human serum samples with good accuracy. Furthermore, the material possessed remarkably low limit of detection.

6. Layered Double Hydroxides (LDHs)

The main structural motifs of LDHs are the positively charged metallic lamellae containing metal ions coordinated by hydroxide ions [5]. Generally, they are based on divalent and trivalent cations, although many exceptions are known. Since the surplus layer charge is neutralized by interlamellar (or intercalated) anions, the family of LDHs is suitable for immobilization of enzymatic [118,154] and molecular [119,155,156] antioxidants. Alternatively, large antioxidants may be anchored to the outer surface of LDHs through electrostatic attraction [60]. In this section, LDHs modified with simpler, then more complex drugs via intercalation will be introduced first. Thereafter, surface modified LDHs of antioxidant effect will be discussed, followed by some details about LDH nanozymes.

Among antioxidant drugs, the main target molecules for intercalation are phenolic compounds, e.g., ascorbic acid, ferulic acid and gallic acid. Gallate, the anionic form of gallic acid, preserved its
antioxidant character in MgAl-LDH [119] and ZnAl-LDH [120]. Release kinetics were also scoped and for ZnAl-LDH, parabolic model (governed by intraparticle diffusion) provided the best fit, while for MgAl-LDH, a fast, diffusion-controlled stage (Freundlich model) was followed by the slow parabolic release. The antioxidant capacity was evaluated in DPPH assays. MgAl(gallate)-LDH scavenged 83.9% of initial DPPH radicals and the same method yielded a maximal 95% scavenging for ZnAl(gallate)-LDH. A plausible reason of the difference may arise from the synthetic procedure: the ZnAl variant was obtained from the delamination-reconstruction process, which can lead to LDHs of higher surface area, contrary to the coprecipitation method used for the synthesis of MgAl(gallate)-LDH.

Beside the gallate-containing MgAl-LDH, Kong et al. also crafted carnosine-intercalated MgAl-LDH through ion-exchange, which provided nearly complete scavenging (95.9%) of DPPH radicals [119]. Interestingly, although gallic acid is a strong antioxidant [87,157], it can act as a prooxidant [157], especially without oxidants present. This effect was exploited by Arratia-Quijada et al., who prepared ZnAl(gallate)-LDH that was used as cytotoxic agent against lung cancer cells [158]. Lima et al. modified ZnAl-LDH with ferulate anions [159]. The intercalant is known for its high antioxidant effect as a pure substance, but the authors proved that ferulate keeps its advantageous scavenging properties in the LDH as well and thus, successfully preventing H2O2-induced oxidative damage generation in fibroblast cultures. The prolonged release (10 hours) from the carrier meant that the protective effect could be maintained for longer time frames. Similar multi-hour release kinetics was measured for two of the most known antioxidant, ascorbic acid (from MgAl-, MgFe- [160] and CaAl-LDH host [161]) and epigallocatechin gallate (from CaAl-LDH [162]).

Recently, polyphenolic ellagic acid (EA) was used to prepare an antioxidant LDH hybrid. Although the cleavage of lactone bonds in EA occurred, the material preserved its antioxidant activity after intercalation. Additionally, the LDH shell served as a tool to drastically improve the dispersibility of the originally hydrophobic EA. It was also proved that the release of EA from the mixed material is minimal. Organic solvents were also used to modify the surface properties of the hybrids and the originally ca. 60% DPPH scavenging activity of untreated EA-LDH increased to ca. 90% after treatment with ethanol and acetonitrile [121]. The materials also showed high TEAC (trolox equivalent antioxidant capacity) values. Trolox is the common name of 6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid.

Regarding larger, but still molecular antioxidants, the most common target molecules are polyphenols. In this family, BHPPA (3-(3, 5-di-tert-butyl-4-hydroxy-phenyl)-propionic acid) is a model compound with good antioxidant effect, expressed in DPPH assay after intercalation in various LDHs [156,163,164]. The DPPH scavenging capability of such LDHs ranged from ca. 50% to 70%, which are comparable to the 60%–70% effectivity of pure BHPPA, depending on the reaction conditions. Even though the values obtained are well below 100%, the nanocomposites were excellent polypropylene (PP) stabilizers against accelerated oxidation (Figure 9) [163], which probably arose from i) the BHPPA-protective effect of the LDH structure and ii) the general polymer stabilizing characteristics of LDHs originated from endothermic water loss at higher temperatures [165] and the reduced gas permeability through plastic [166]. The BHPPA-LDH hybrid surpassed Irganox 1010 [164], a frequent polymer antioxidant used in the industry.
Figure 9. Radical-scavenging activity of BHPPA (denoted as AO) and BHPPA-LDH in a 100 mL of 100 μM DPPH ethanolic solution. The top inset represents the color change and the bottom inset shows UV-Vis absorption curves after radical scavenging for 30 min in a dark box (a). FT-IR absorbance spectra of pure PP in the range of 3700–1250 cm$^{-1}$ as a function of accelerated thermal aging time at 150 °C. Inset shows a magnified region of the FT-IR spectra in the wavenumber range of 1810–1660 cm$^{-1}$, thus the carbonyl region indicating oxidation of the plastic (b). Reprinted with permission from (Feng, Y.J.; Jiang, Y.; Huang, Q.; Chen, S.T.; Zhang, F.B.; Tang, P.G.; Li, D.Q. High antioxidative performance of layered double hydroxides/polypropylene composite with intercalation of low-molecular-weight phenolic antioxidant. Ind. Eng. Chem. Res. 2014, 53, 2287–2292) Reference [163]. Copyright (2020) American Chemical Society. BHPPA: (3-(3, 5-di-tert-butyl-4-hydroxy-phenyl)-propionic acid; LDH: layered double hydroxides

In 2013, Lonkar et al. also reported the improved thermal behavior of PP containing BHPPA-LDH as a filler [167]. Gomez Amaro et al. investigated BHPPA and Trolox in intercalated from [122], revealing up to 90% DPPH scavenging for both antioxidant-modified LDHs with 14 and 13 mass% organic content, respectively. Irganox 1425, an ionic antioxidant consisting of a benzylphosphonate-type anion in Ca$^{2+}$ salt, a common building block in the LDH structures, was used as precursors to prepare CaAl-LDH intercalated with its anion [155]. The polymer additive kept its DPPH scavenging activity after intercalation with ca. 55% maximum effect, which is very similar, if not slightly higher than bare Irganox 1425. The LDH hybrid of olanzapine (2-methyl-4-(4-methylpiperazin-1-yl)-10H-thieno [2,3-b][1,5]-benzodiazepine) was also developed to overcompensate its low water solubility [123]. This drug molecule is a popular antipsychotic agent and due to the correlation between oxidative stress and brain damage [168], its antioxidant potential may be a reason for its pharmaceutical efficacy. It was demonstrated that the LDH hosts (CaAl- and NiAl-LDH) did not have negative effects in the assays investigated, e.g., DPPH assay, ABTS assay, NO scavenging and TBARS (TBA reactive substances) test. Moreover, the LDHs had synergistic effects with the olanzapine molecules. This was more pronounced for CaAl-LDH, which was often significantly more active than NiAl-LDH, as a carrier for olanzapine.

As mentioned earlier, larger, or more importantly, positively charged agents can be immobilized only via surface modifications. These examples tend to find enzymatic applications, e.g., the histamine (Hsm) containing Cu(Hsm)$^{2+}$ complex fixed on a polymer modified MgAl-LDH surface [124]. Poly(vinylpyridine-b-methacrylic acid (PVPMAA) copolymer of negative charge was used to reverse the originally positive surface charge of LDH enabling the adsorption of the positively charged Cu$^{2+}$ complex. Charge reversal also achieved higher colloidal stability, i.e., the LDH precipitated at higher background electrolyte concentrations. The complex-LDH composite was then used as a SOD mimic with comparable activity to bare SOD or a SOD-coated MgAl-LDH [60]. The latter material was obtained by modifying the surface of SOD-MgAl-LDH composite by heparin (Hep) polyelectrolyte giving rise to highly stable antioxidant dispersions. It was apparent that both
SOD functioning LDHs were highly active, but unlike natural enzymes, Cu(Hsm):2+ was not heat sensitive and kept 90% of its activity at 80 °C, while SOD quickly denatured at higher temperatures (Figure 10).

The intrinsic peroxidase activity was enhanced using deoxyribonucleic acid (DNA) from herring sperm [125]. The hybrids were constructed by surface electrostatic force between the building blocks. Furthermore, DNA-induced delamination of the carrier LDH (proved by X-ray powder diffraction and light scattering) facilitated the antioxidant reaction in the TMB test reaction, which showed best activity at 35–45 °C and at pH 3–4 with sudden decrease afterwards. Combined with GOx-glucose pair, Chen et al. used their DNA-LDH hybrid as a colorimetric glucose sensor.

Natural HRP bears gross positive charge in its active form, thus a Hep coating was used on the surface of MgAl-LDH before its immobilization [118]. The kinetic parameters in the guaiacol assay of the as-prepared composite LDH were slightly different compared to the free enzyme, representing that adsorption forces were insufficient to overcome the protein rigidity and to considerably influence the native structure of the enzyme. Beforehand, HRP-containing LDHs were synthesized using transition metal LDHs that were applied in the electrochemical detection of H2O2 [154,169].

Mimicking the active site of HRP, hemin was immobilized in/on FeNi- and CuAl-LDH by Zhang et al. [170] and Qiao et al. [171], respectively. Despite its size, hemin is a good target for intercalation owing to the planar structure and negative charge through the carboxylic groups. It was demonstrated that FeNi(hemin)-LDH was an excellent peroxidase, surpassing free hemin. This is attributed to the dimerization of hemin outside the layered material, while in intercalated form, the hemin rings are mostly found as monomers. Finally, Zhang et al. developed an electrochemical sensor based on this material [170]. The CuAl(hemin)-LDH designed by Qiao et al. was tested as a scavenger of ONOO− ions [171] (Figure 11). They also found that the composite was more active than hemin alone, which is interpreted by the electrostatic attraction between the anionic ONOO− and the LDH particles, and also due to the π donor property of LDHs to the Fe center of hemin, as stated by the authors. The composite also inhibited the reaction between tyrosine and ONOO− up to 80%.

Figure 10. Schematic representation of formation of Cu(Hsm):2+ on PVPMAA-coated LDH (a) and its activity in SOD assay compared to bare SOD, LDH-Hep-SOD and Cu(Hsm):2+ at 80 °C as a function of reaction time (b). Reprinted from (Highly stable enzyme-mimicking nanocomposite of antioxidant activity, 543, Pavlovic, M.; Nafradi, M.; Rouster, P.; Murath, S.; Szilagyi, I., Journal of Colloid and Interface Science 2019, 543 174–182) Reference [124] Copyright (2020), with permission from Elsevier.
A novel protein-LDH interaction was recently revealed when protein extracted from bael (*Aegle marmelos*) were anchored to the outer surface of MgAl-LDH [126]. The natural antioxidant proteins were scavengers of DPPH radicals on the LDHs (ca. 50% effectivity) and possessed antimicrobial effect towards *S. coliform*. In addition, the composite adsorbed Pb$^{2+}$ ions according to Freundlich model.

The earliest report on LDHs with intrinsic nanozyme activity was published by Zheng et al. [172]. For this purpose, like for all the others, redox active metal ions are represented in the lamellae. In this instance, CoFe-LDH was used as a peroxidase mimic. The characteristics of the nanozyme were like those of HRP, regarding optimal pH and temperature dependence, but the platelets were active in a larger H$_2$O$_2$ concentration regime. Thus, H$_2$O$_2$ detection is possible in a range of reaction conditions. The obtained LDH was also used in the GOx mediated selective sensing of glucose via TMB test reaction. Afterwards, CoAl-LDH sheets exfoliated in L-asparagine solution were proven to have peroxidase mimic with pH optimum at 6 [127]. A similar approach with L-asparagine was applied to get exfoliated 2D NiFe-LDH with 2 nm thickness [128]. Both asparagine exfoliated LDHs showed Tyndall effect in aqueous suspension, indicating the formation of stable colloids. While the nanosheets of CoAl-LDH showed relatively lower affinity to H$_2$O$_2$ ($K_m = 22.1$ mM), NiFe-LDH and bare HRP had similar values with 2.4 and 3.7 mM, respectively (Table 1). The detection limit of NiFe-LDH was estimated to be 4.4 mM H$_2$O$_2$. The peroxidase-like activity of NiCo-LDH was described by Su et al. using homovanillic acid as a fluorescent dye upon reacting with H$_2$O$_2$ [173]. It was found that the catalysis was strongly pH dependent and reached optimum above pH 8.3 with a linear response at pH between 8.4 and 8.9, before plateauing at 9.6. Another peroxidase mimic, CuAl-LDH was grown onto carbon fibers (Figure 12) after atomic layer deposition of Al$_2$O$_3$ and hydrothermal conversion to the end-product [174]. Interestingly, the optimum temperature for this material was 70 °C, but at 60 and 80 °C, it showed similar efficiency. Activity also grew when using more catalyst, indicating that the reaction was not diffusion controlled. It was also shown that radical scavengers inhibit the catalytic reaction in a great manner. Peroxidase nanozyme CoFe-LDH was used as catalyst to obtain fluorescent polydopamine (PDA) from its monomer [175]. This procedure avoided using concentrated H$_2$O$_2$ and yielded shorter reaction times. PDA is strongly fluorescent in aqueous solutions which is quenched by Fe$^{3+}$ ions through coordination. However, intensity is restored when strong iron complexing pyrophosphate is introduced, with good linearity and selectivity. Another detection is possible through the addition of pyrophosphatase, which decomplexes the quenching Fe$^{3+}$ ions and lower fluorescent signal is observed.
Recently reported MnFe-LDH possessed catalase mimicking activity and exhibited photothermal effect [176]. Methylene blue loading enhanced the efficacy of O$_2$-dependent photothermal phenomenon. This way, the phototherapeutic effect in hypoxia cancer cells were gapped, as demonstrated in human cells and mice, in which the relative size of tumor was also reduced. We mention that even though it was not used as an antioxidant, rather a prooxidant, but Fe$^{2+}$-bearing FeAl-LDH was used to decompose H$_2$O$_2$ with a Fenton-type mechanism inducing apoptosis in cancer cells through the generated OH$^\cdot$ and OOH$^\cdot$ radicals [129].

7. Montmorillonites (MMTs)

Montmorillonites are sheet-like silicate minerals with anionic layers and feasible, water-exchangeable cations between the individual sheets [31,130,177]. Their charging features, therefore, are the reverse of the already discussed LDHs. Hence, MMTs may possess antioxidant capacity by their own through fixing antioxidant compounds in their structures (e.g., intercalation or physical adsorption) or by the preparation of composite materials with synergistic nanomaterials.

Regarding intercalated MMTs, Baek et al. constructed GSH-containing MMT, coated with poly(vinylacetel diethylaminoacetate) (AEA) [130]. GSH is the most common antioxidant material in the body, but with low bioavailability, in case it is taken in orally. Carriers can overcome this barrier and it was found that GSH and GSH-MMT had an ABTS radical scavenging activity of ca. 60%. Diethyl aminoacetate coating was also introduced as a protective layer, which did not interfere with the antioxidant effect. After oral administration for mice, the concentration of GSH significantly grew
in liver, to which delivery of free GSH is minimal. Naturally occurring ionic species may also be intercalated from plant extracts, e.g., antioxidants from blueberry (*Vaccinium corymbosum*) that provided an intelligent potential food packaging nanomaterial, which is color-sensitive to pH [131]. Compared to the negligible DPPH scavenging activity of pure MMT (which probably arose from adsorption), the blueberry extract-modified clay showed exceptional activity with ca. 40% scavenging.

An antioxidant activity, close to the one of ascorbic acid, was assigned to the eugenol-MMT hybrid that was used as a filler in poly(3-hydroxybutyrate)-thermoplastic (PHB-TPS) starch composite that resulted in a functional material with evolved heat resistance and mechanical properties that also possessed antioxidant and antifungal properties [132]. The peroxidase nanzyme activity of hemin was demonstrated after adsorbing its highly active monomeric form in 7 mg/g dose on MMT [133]. Earlier, it was demonstrated that histidine amino acid residues activate the catalytic activity of HRP [178], which was substituted with intercalated Hsm in the case of hemin-MTT. This way, the relative catalytic activity of the MMT composite jumped to over 350% at 60 °C, where the performance of natural HRP decreased.

A chitosan (CS) coated MMT@(Ca-alginate) nanoparticle synthesis was recently reported with covalently bonded HRP enzyme on its outer surface to degrade aflatoxin B1 [179], a dangerous poison produced by molds. The role of negatively charged MMT in this structure was to promote aflatoxin adsorption and to attract more chitosan chains. Multiple chitosan/MMT composites were prepared in Fernando’s group as a biocompatible substitute of classic polymer food wrappers [177,180,181]. These materials had adequate physical characteristics (tensile strength, optical properties, etc.) and were incorporated with rosemary (*Rosmarinus officinalis*) and ginger (*Zingiber officinale*) essential oils. This incorporation provided the composite with good antimicrobial activity and antioxidative protection for applications in poultry packaging. Yan et al. used α-tocopherol (TOC) in a chitosan-MMT matrix as preservative for wrapping sliced ham with film of the material [134]. The film coating showed good antioxidant activity in DPPH assay even after 16-day storage, while in lipid oxidation tests, the remarkable antioxidant activity of the MMT composite was expressed after 120 days.

Regarding MMT composites with other inorganic materials of overall antioxidant capacity, sulfide-MMT structures were under investigation previously. A facile way was worked out to construct ZnS/MMT composite via coprecipitation [135]. This material acted as a peroxidase mimic in a rapid fashion with merely a few minutes to form oxidized TMB. The activity measured was similar in the pH range 4–7 and from 20 to 70 °C. Besides, outstanding substrate affinity was achieved, compared to bare HRP (148-fold and 8-fold increase towards H2O2 and TMB, respectively). A CuS/MMT composite was also considered as a peroxidase mimic with short timeframe (< 1 min) to reach colorimetric visualization with poorer kinetic parameters [136] (Table 1). Reaction kinetics for ZnS/MMT were also proved by EPR, trapping and detecting the OH· radicals formed from H2O2 by DMPO spin trap. A very similar coprecipitation method, starting from thioacetamide, was used to obtain Ag2S/MMT composite [137]. The pH and temperature dependence profiles were fairly similar to other MMT systems and it was found that pure MMT also possessed peroxidase activity. In this case, terephthalic acid was used as fluorescent OH· radical trap to confirm the mechanism. Recently, the composite of MMT and CoNiSx was prepared, and remarkable peroxidase activity was observed [138]. It exhibited good affinity towards H2O2 and TMB substrates, often surpassing bare HRP. Contrary to other MMT composites, elevated temperatures had negative impact on its activity above 35 °C, but at ambient temperatures showed linear response to H2O2.

MMT was also found to be the ideal clay mineral for soil organic matter modification linked with HRP adsorption [139]. Furthermore, MMT retained 91.3% activity of bare HRP and was reusable for phenol removal, tested for six cycles with slight activity loss.

Heterobinuclear [140] and mononuclear [141] Cu2+ complexes containing amine and amino acid ligands were intercalated between the layers of MMT to obtain a heterogeneous catalysts for dismutation of superoxide radicals. The composites showed significant SOD-like function in the test reactions, in which the superoxide radical anions were generated by a photochemical reaction. Finally, montmorillonite-intercalated Cu2+-tris(2-aminoethyl) amine complexes were highly active in
decomposition of H₂O₂, i.e., they acted as an excellent catalase mimicking antioxidant composite [182].

8. Miscellaneous Structures

Structures listed in this paragraph either did not fall under the categories listed before or were borderline (e.g., CeO₂@CO₂(OH)₂CO₃) or were the only example of the material (see Fe₂O₃).

He et al. published the synthesis of flaky CoP 2D particles on Ni foam substrate [142]. First, Co(OH)₂ was built on the metal carrier, which was phosphorylated with NaH₂PO₂ in a hydrothermal manner. The precursor Co(OH)₂ had mediocre activity in the peroxidase test reaction, the end product possessed outstanding performance, while Ni foam (NF) was completely inert. More importantly, the same piece of catalyst showed the same activity for 12 cycles, which was unaffected by pH (2–14) and temperature (0–95 °C). The material was successfully used to detect uric acid in
serum samples (with overall 5% error compared to clinical method) and in real urine samples as well with good selectivity to the substrate (Figure 13).

A facile detection method based on CeO$_2$ nanoparticles on 2D CO$_2$(OH)$_2$CO$_3$ was introduced by Alizadeh et al [143]. After the material was characterized as an excellent peroxidase mimic with higher maximum rate than HRP’s, a paper based colorimetric immunoassay method was developed with smartphone image analysis selective to carcinoembryonic antigen.

Deshapriya et al. used aminated and artificially cationized BSA (bovine serum albumin) as an overcharging agent on negatively charged, exfoliated ZrP (standing for -Zr(HPO$_4$)$_2$) sheets [183]. This led to the possible immobilization of numerous enzymes, including catalase. For GOx and tyrosinase, loading up to 640 and 380 mass% was achieved, respectively. Moreover, the bound enzymes kept their native structure and at least 80% of their activity, excluding tyrosinase with a value of ca. 40%.

Iron containing simple compounds were also tested as peroxidases. Fe$_3$(PO$_4$)$_2$ × 8 H$_2$O sheets in a flower-like array were utilized as a potent nanozyme to detect H$_2$O$_2$ with over 30 ionic and non-ionic abundant substances tested as an interfering agent, but the detection was considerably not distorted [144]. Glucose sensing was also achieved through the glucose-GOx reaction with good selectivity amongst other bioactive compounds. Tanaka et al. started from Fe$^{3+}$-glycerate precursor and applied thermal treatment to obtain porous Fe$_3$O$_4$ flakes [145]. It was found that lower pyrolysis temperatures resulted in better Fenton-type activity for the synthesized Fe$_3$O$_4$, since those products had higher surface area and higher hexagonal $\alpha$-Fe$_3$O$_4$ content with more active sites compared to cubic $\gamma$-Fe$_3$O$_4$ (Table 1).

9. Summary and Outlook

As discussed above, antioxidant compounds of 2D structure has attracted widespread contemporary interest in the scientific and technological communities, wherever the goal is to decrease oxidative stress or to develop ROS sensing devices. The main advantages of the 2D structures are the high specific surface area and that processable samples for homogeneous film formation (e.g., for electrode modifications) can be easily prepared. Given the fact that these compounds are excellent candidates as building blocks of sensing materials, the ease of synthesis even in larger amount will definitely help in the commercialization of sensors and detectors.

Enzyme-loaded 2D NPs represent a very efficient ROS consuming class, however, the high sensitivity of the proteins to the environmental conditions often remains upon immobilization. Such a low resistance against pH, pressure and temperature can be improved by applying enzyme mimicking metal complexes as catalytic center in the 2D NP-based hybrids, but their efficiencies usually lag behind the one of the native enzymes. A returning problem in case of many molecular antioxidants is their low water solubility and thus, low bioavailability. The insufficient solubility may be increased by loading the 2D particles of high specific surface area with the antioxidant molecules, provided the carrier can be dispersed in liquid media. The new wave in the field of 2D antioxidants are the so-called nanozymes, which are able to decompose ROS without embedding enzymatic or molecular antioxidants in their structure. Metal-containing compounds hold great promise towards development of such 2D nanozymes.

Apart from the 2D antioxidant materials reported to date, novel hybrid materials are continuously being developed in the academic and industrial laboratories. Achieving excellent radical scavenging activity is the main priority, however, other important issues must also be considered. These include the physiological effect on living organisms, i.e., cell toxicity assessment is a must prior to the biomedical application. In addition, dispersions of antioxidant 2D particles are applied in most of the systems, therefore, the stability of the dispersions has to be precisely controlled. Unwanted particle aggregation processes lead to significant loss in the antioxidant activity and to non-processable samples. The stability issue can be solved by choosing appropriate stabilizing agents. Comprehensive studies in this topic are missing in the relevant literature.

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