

Synergistic removal of 1-(4-Hydroxy-3-Methoxyphenyl) ethenone (Apocynin) with enhanced immobilized and suspended Sr-doped LaNiO₃ based photo-catalytic membrane reactor under gamma irradiation

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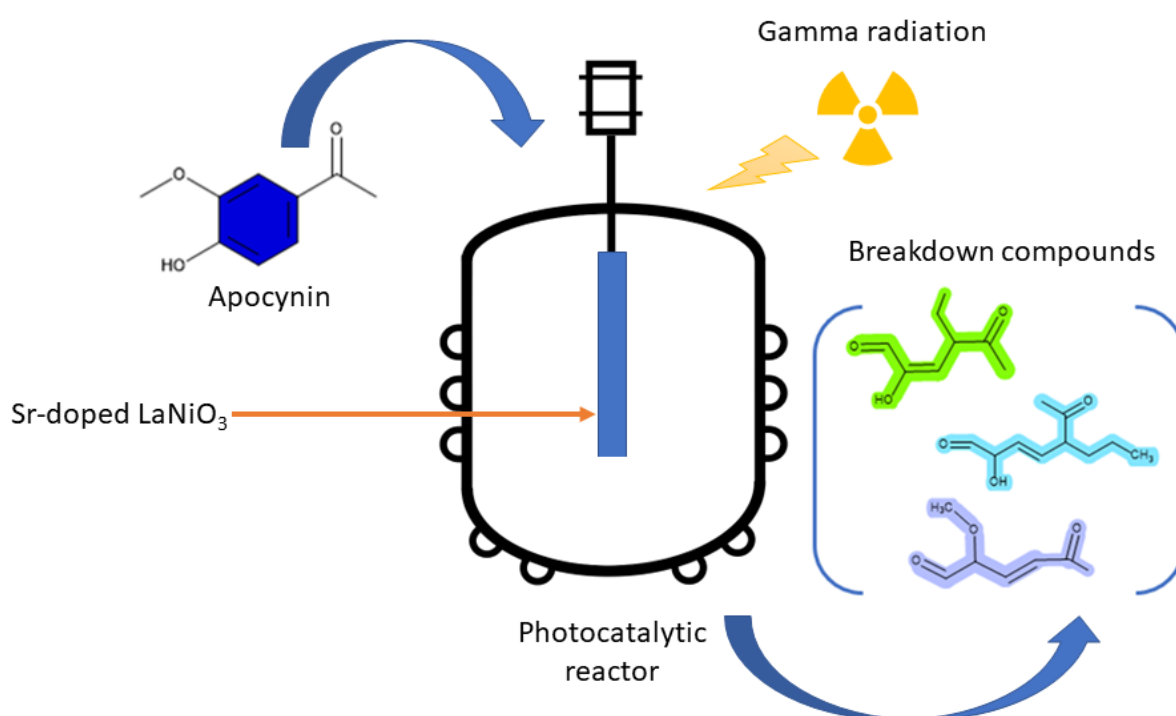
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GRAPHICAL ABSTRACT



Abstract

The electrolytic approach was combined with radiolysis employing gamma radiation as an enhanced oxidation process to study the decomposition of 1-(4-Hydroxy-3-Methoxyphenyl) ethenone (Apocynin (ACN)). For Apocynin breakdown from effluent water, a photo-catalytic membrane reactor (PMR) was utilized with suspended and immobilised Sr-LaNiO₃ with regarding to gamma irradiation. For phase procedures, the impacts of basic Sr-LaNiO₃ on the PMR containing residual Sr-LaNiO₃ was investigated. An RO (Reverse Osmosis) membrane was integrated by way of the PMR under continuous circumstances to improve standard of water. At such a moderate Sr-LaNiO₃ dosage (0.8 g/L), ACN deterioration by the PMR with attached and immobilised Pt-LaNiO₃ was similar in both methods, but improved with larger Sr-LaNiO₃ doses for the reactor with coupled Pt-LaNiO₃.

Keywords: *Apocynin, Electrochemical degradation, Photocatalytic reactor, gamma irradiatio*

1. Introduction

Several commodity-based industries that manufactures cork, pulp and paper uses a lot of electricity and freshwater, consequently generates large volumes of sewage water. The generated sewage water contains effective levels of biochemical oxygen demand (BOD) and chemical oxygen demand (COD), intense colour, particulates, and non-biodegradable chemicals [1,2]. The release of such unprocessed effluents might have a harmful influence on the environment because of the existence of these resistive chemicals, which should be reduced. Apocynin (ACN), a lignin degradation by-product found in pulp, paper and cork effluents. Microfiltration was used to remove apocynin from an aqueous mixture Gallic acid and Esculetin, according to some early researches that were published [3]. The persistence effectiveness of the membrane was shown to be dependent on the nature of the membrane, with poor persistence for ACN in the utilized membranes. Hence, it is essential to eliminate ACN from water prior to release into freshwater resources.

In treating sewage water, there seems to be a number of techniques for decontamination. For eliminating pharmaceutically active substances, reverse osmosis (RO) and nano-filtration (NF) are the effective methods. Furthermore, before even being released into freshwater resources, the re-circulated stream containing high level contaminants must be eliminated [4]. These chemicals have been successfully removed using advanced oxidation processes

(AOPs) such as ultraviolet (UV), semiconductor photo catalysis, Fenton and ozonation [5]. As a result, AOPs have a great deal of capability for decomposing such compounds in sewage water. LaNiO_3 photo-catalysts has gotten the greatest recognition because of their enhanced pursuit, thermal, chemical and photo-erosion tolerance, inexpensive, and eco-friendly [6,7]. Even so, anatase LaNiO_3 with a broad band gap primarily accumulates gamma light (400 nm), that reports for 5–8% of solar energy and moreover gives rise to non-metals or metals being on condition that with visible light without damage of gamma action, extending the operational span and increasing productivity [8]. The replacement Sr doped LaNiO_3 is the most efficient among some of the non-metal doped- LaNiO_3 , because of the bonding force amid the N 2p state and the O 2p state, therefore developing band-gap and shrinking Sr doped LaNiO_3 was capable in absorbing the light waves [9].

Molinari and his co-workers [10] were the first to create a hybrid method, which integrates membranes with catalysts. For the first time, a hybrid method was used to remove organic contaminants. Membrane serves as a barrier not just for molecule partition but also for catalyst isolation in this method. Photocatalytic membrane reactors (PMRs) can also conserve energy and minimize the complexity of an establishment [10,11]. The PMR has also been used in treatment processes of sewage water [12,13], as well as to remove bacteria like *Aspergillus fumigatus* [14] and medications like oxytetracycline [15]. Submerged photocatalytic membrane reactor (SPMR) is a form of PMR with cheap setting costs, reduced energy consumption, and ease of management [16]. The downsides of photo-catalytic immobilised in or above membrane, including a minor ratio of area-to-volume and lowered gamma light application efficacy [16,17], as well as the loss and prolonged usage of catalyst particles in assembly lines, can be determined with SMPR [16,18].

As per the knowledge of authors there were not many studies reported on Sr doped membranes on wastewater treatment especially on apocynin degradation. SPMRs with suspending and immobilised Sr- LaNiO_3 were examined for ACN elimination in continuous flow settings under gamma irradiation utilizing two photo-catalytic methods either with or without H_2O_2 . For high effluent quality, RO membranes were paired through the SPMR after ceramic microfiltration membrane in the constant cycle. The levels of ACN, RO and total organic carbon in the MF filtrate was confirmed.

2. Methods and Materials

2.1. Preparation of Sr-doped LaNiO_3

All the experimental reagents were analytical grade. A typical synthesis procedure was as follows: 0.9 mM of $\text{La}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, 0.2 mM of $\text{Sr}(\text{NO}_3)_2$, and 2 mM of $\text{Ni}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ were dissolved in 30 mL of N, N-Dimethylformamide and magnetically stirred for 4 h. Thereafter, 0.65 g of polyvinylpyrrolidone was added to the mixed to get a homogeneous solution by magnetically stirring it for 1.5 h. Subsequently, the homogeneous solution was heated under continuous stirring until the gel was formed. Then, the obtained gel was transferred to the crucible and placed in a muffle furnace for heating treatment at 730 °C for 2 h at a heating rate of 10 °C/min. The final synthesized product was nomenclatured as $\text{La}_{0.9}\text{Sr}_{0.1}\text{NiO}_{3-\delta}$ nanomaterials. Figure 1 illustrated the chemical structure of apocynin (ACN).

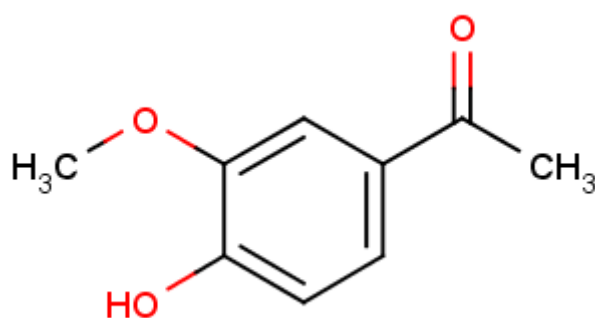


Fig 1. Chemical structure of apocynin (ACN)

2.2. Sr-LaNiO₃ immobilization on MF ceramic membrane

The overall surface of the MF membrane was 0.0150 m². To eliminate contaminants, the membrane area was rinsed numerous times with ethanol and desiccated at room temperature. In a mixer, 1.2 g Sr-LaNiO₃ particulates were added to 3.4 L of a sodium dioctyl sulfosuccinate solution (1.28 %), sonicated for 1.2 h to make the Sr-LaNiO₃ catalyst mixture. The MF membrane was gently immersed into the mixture, held in place for 2 min, and then steadily removed from the mixture. Lastly, the MF membranes were desiccated at 105°C in a kiln before being warmed to 550°C in a furnace at a heating rate of 10°C/min for 1.2 h. This technique was continued till the coating was spread to the appropriate thickness [19].

2.3. Experimental Setup

During investigations on synthetic ACN-treated effluent water, SPMRs with suspended and immobilised Sr-doped LaNiO₃ on the ceramic membrane with observable illumination was

utilized. It was also looked at combining H_2O_2 with the photo-catalytic method. The SPMR has a cylindrical photo-reactor container with a ceramic MF membrane embedded in the reactor's core and a conduit connecting it to a suction pump. A stainless-steel cylindrical photo-reactor tank with an operating capacity of 2.5 L was installed in a compartment. Around the exterior of the reactor, gamma source was mounted. A magnetic stirrer was used to guarantee that the ACN monomers and catalyst dispersion were homogeneous. The O_2 was transported through a canal suppressed below the MF membrane's base. There were batch and continuous procedures used.

For the batch tests, an SPMR with suspended and immobilised Sr-LaNiO_3 was employed for 200 min to examine the ACN extraction efficacy under Vis/ Sr-LaNiO_3 and Vis/ Sr-LaNiO_3 / H_2O_2 procedures. The impacts of the initial Sr-LaNiO_3 levels on the extraction efficacy were also looked at. The SPMR was connected to a RO membrane for constant circumstances. Outside the photo-catalytic reactor, the RO membrane was linked to the yield of pumping system. The Vis/ Sr-LaNiO_3 and Vis/ Sr-LaNiO_3 / H_2O_2 procedures were used for 72 h on SPMRs with suspended and immobilised Sr-LaNiO_3 . Through a hose, a feed tank was linked to the photo reactor. A regulating float was utilized to regulate the operating capacity of the photo-catalytic reactor, and a lever was employed to manage the RO membrane's discharge rate of flow. The RO permeate was mechanically reused into the reaction vessel. In this work, alternating performances of the SPMR units with a configuration of 8 min of operation (ON) and 2 min of halt were used to prevent membrane fouling (OFF). The MF and RO membranes' wastewater grade and pervade deadline were examined. Figure 2 illustrated the schematic representation of experimental setup.

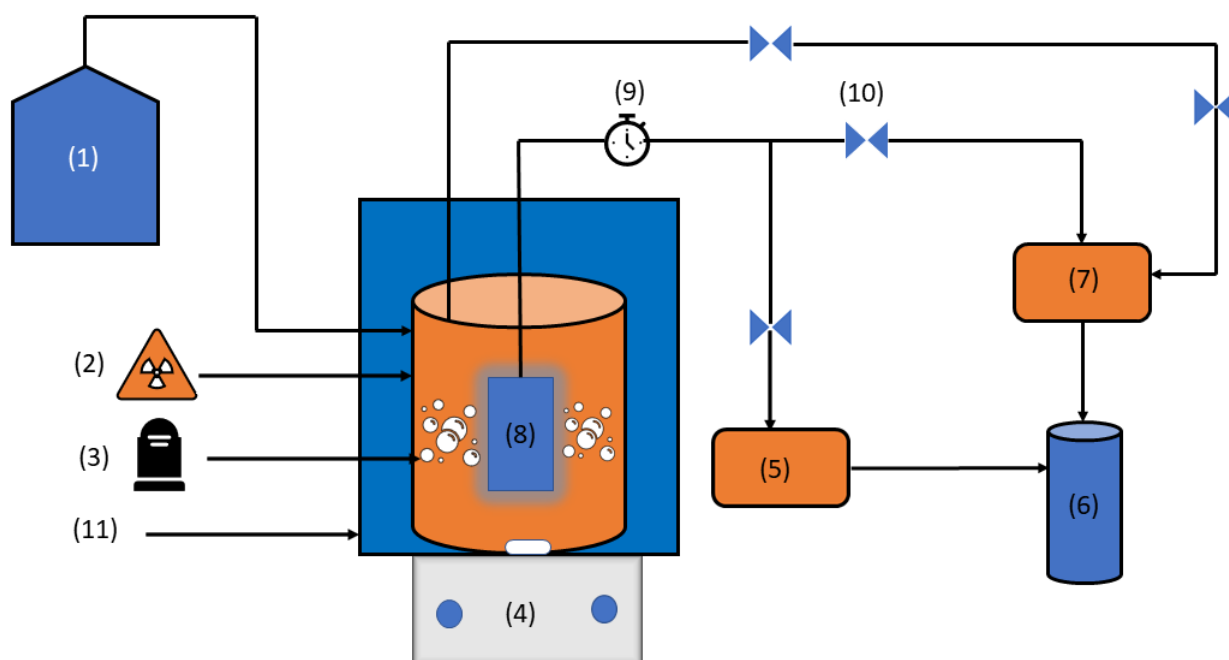


Fig 2. Schematic representation of experimental setup. 1) Feed tank, 2) Gamma irradiation, 3) Oxygen supply, 4) Magnetic stirrer, 5) Micro filtration (MF) permeates, 6) Reverse osmosis permeate, 7) Reverse osmosis membrane, 8) MF membrane, 9) Timer, 10) Suction pump, 11) Chamber.

2.4. Analytical Methods

At a wavelength of 300 nm, a UV-Vis spectrophotometer has been employed to assess the ACN contents. For the TOC assessment, a TOC analyzer model was used. Potassium phthalate norms were used to tune the instruments.

3. Result and Discussion

3.1. Batch photo-catalytic reactor

Batch experiments were conducted for each setting to assess ACN elimination through photocatalysis; RO membrane, MF membrane with and without immobilised Sr-LaNiO₃ on the exterior layer, and also H₂O₂. 60 mg/L ACN, pH 7, 2 g/L Sr-LaNiO₃ infusion into SMPR with dispersion, and 2 g Sr-LaNiO₃ deposited on MF membrane surfaces were used as the laboratory parameters. All of the trials were completed in 180 min. The research found that MF membrane, MF membrane coated Sr-LaNiO₃, and H₂O₂ alone elimination relatively little ACN at high concentrations (60 mg/L). The elimination of ACN by RO membrane has been estimated to be as high as 98%, showing that RO membrane has a great capability to exclude ACN molecule. The findings were in line with prior research, which shows that RO

membranes may exclude negatively charged ACN molecules up to 96% [20]. The difference in MF and RO extraction efficiency might be attributed to their pore diameters or molecular weight cut off (MWCO) [21]. For instance, MF membrane pore diameters vary from 0.2 to 5.4 μm or a MWCO larger than 1,500 kDalton, whereas RO membrane pore sizes vary from 0.0002 to 0.002 μm (MWCO less than 110 Dalton). Meanwhile, ACN has a molecular weight of 300 (g/mol). ACN with such a molecular weight can easily flow through the MF membrane, but it will be denied by the RO membrane. In comparison to MF membranes sans coated Sr-LaNiO₃ on the surface, MF membranes with Sr-LaNiO₃ immobilisation on the surface demonstrated somewhat greater ACN elimination. The discovery might be the cause of ACN monomers being deposited on the membrane surface by Sr-LaNiO₃ nanoparticles. Figure 3 illustrated the SEM micrographs of synthesized Sr-doped LaNiO₃

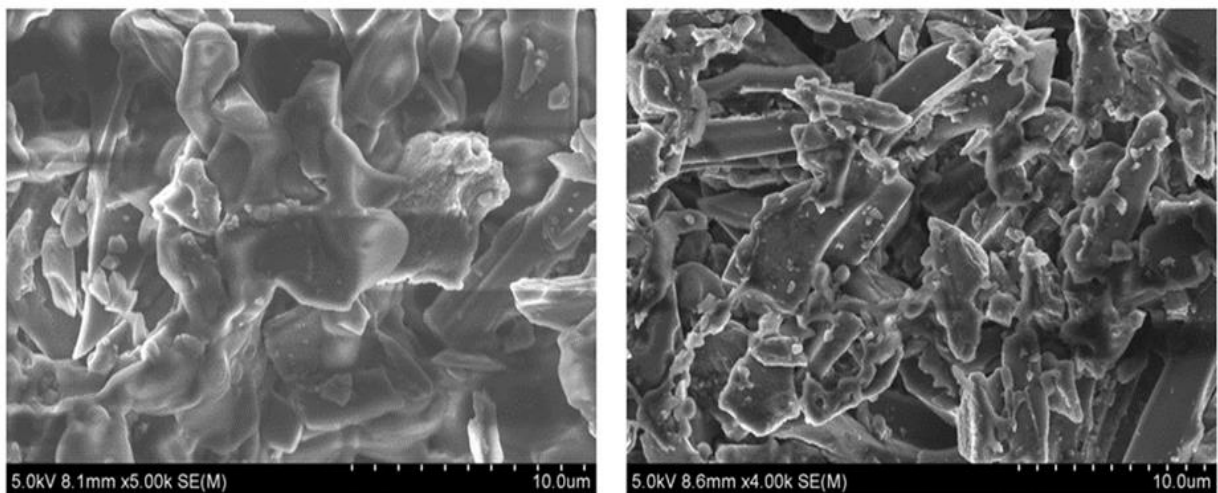


Fig 3. SEM micrographs of synthesized Sr-doped LaNiO₃

3.1.2 Impacts of Sr-doped LaNiO₃ accumulation in refinement process

In the batch settings, the influence of the starting Sr-LaNiO₃ level on the ACN extraction efficacy by the SPMR containing dispersed Sr-LaNiO₃ under Vis/ Sr-LaNiO₃ and Vis/ Sr-LaNiO₃ / H₂O₂ procedures is shown in Fig. 4. The ACN remaining in reactor (ACN level) measured by the SPMR with suspended Sr-LaNiO₃ during the photo-catalytic activity (Vis/N-TiO₂ procedure) is shown in Fig. 4a. For whatever starting Sr-LaNiO₃ level, ACN elimination did not achieve 100% after 200 min. With 0.6 g/L Sr-LaNiO₃, the lowest ACN and TOC elimination efficacy were achieved. At low catalytic dosages, there aren't enough photo-catalytic active sites in LaNiO₃, leading in a slow decomposition pace [22].

The ACN and TOC elimination efficiencies were lowered when the initial Sr-LaNiO₃ level was increased to 2 g/L compared to 1 and 1.5 g/L. Because the probing photon flux adsorbs to the catalyst surface when the catalyst dose is increased beyond saturation, the deterioration process may be slowed [23]. Furthermore, a high catalyst dose causes agglomeration of the catalyst, leading to reduced surface area [24]. With a starting Sr-LaNiO₃ level of 0.5 g/L, the highest ACN extraction efficacy was 54%. The deterioration rate rises at necessary catalyst doses since it avoids unwanted surplus catalyst and ensures full photon adsorption for effective photo-mineralization [25]. Because of the impacts on the catalyst surface and absorption of the photon flux on the exterior area of the catalyst, the catalyst dose has a positive or negative influence the photo-degradation rates in overall. The ACN elimination effectiveness using the photo-catalytic method combined with H₂O₂ is shown in Fig. 4b. The best initial Sr-LaNiO₃ level for ACN elimination using this procedure was 2 g/L, and the smallest ACN elimination appeared at 0.5 g/L, implying that the elimination efficacy for both photo-catalytic activity, with (Vis/ Sr-LaNiO₃ / H₂O₂) and without (Vis/ Sr-LaNiO₃ procedure), was maximized with the same initial Sr-LaNiO₃ level. Furthermore, as contrasted to exclusively photo-catalytic activity under visible irradiation, combining H₂O₂ with the photo-catalytic activity improved ACN and TOC elimination. H₂O₂ may readily interact with LaNiO₃ electrons to create hydroxyl radicals when combined with the photo-catalytic activity [26]. Furthermore, the interaction involving H₂O₂ and the oxygen species generates higher hydroxyl radicals in the photo-catalytic activity [27]. Additionally, the deposited H₂O₂ on the LaNiO₃ surface forms proxy compounds with the Sr ions, increasing the adsorption capability of LaNiO₃ when exposed to visible light [28]. The use of H₂O₂ in conjunction with the photo-catalytic method improves the elimination of ACN and TOC.

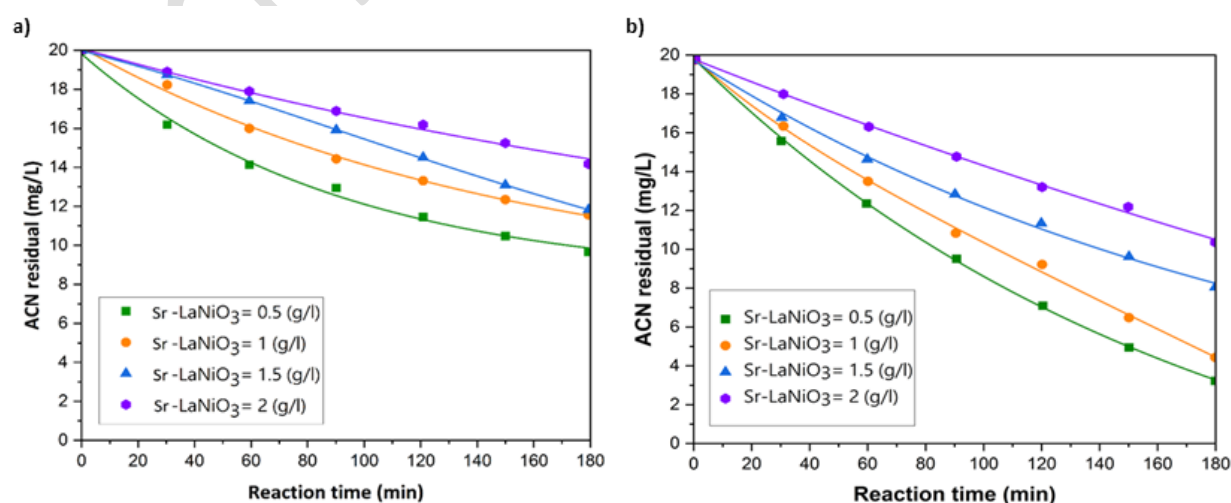


Fig 4. The impact of Sr-LaNiO₃ feeding on ACN deterioration by SPMR with stranded Sr-LaNiO₃; (a) Vis/ Sr-LaNiO₃ procedure, (b) Vis/ Sr-LaNiO₃ / H₂O₂ procedure. Levels of Sr-LaNiO₃ of 0.4, 1.0, 1.5, and 2.0 g/L were used in the experiments. [ACN] = 25 mg/L, observable light range = 300 W, and [H₂O₂] = 20 mM. pH = 7, [ACN] = 20 mg/L, observable light range = 300 W, and [H₂O₂] = 20 mM.

3.2 Regular parameters

3.2.1 Permeate quality

For SPMR with suspended and immobilised Sr-LaNiO₃, the longstanding constant operation capabilities of the Vis/ Sr-LaNiO₃ and Vis/ Sr-LaNiO₃ / H₂O₂ procedures were investigated. The Sr-LaNiO₃ loading was 2 g/L, the solution pH was 7, and the input ACN content was 25 mg/L. The RO permeate flow rate was maintained at 2.6 L/h, which corresponded to a hydraulic retention time (HRT) of 5 h. The ACN mixture was constantly delivered from the feed tank into the photo reactor. In the Vis/ Sr-LaNiO₃ / H₂O₂ procedure, H₂O₂ was combined with the ACN solution in the storage tank. The SPMR entity used sporadic membrane filtration and were monitored by a timer that interchanged amid 6 min of work (ON) and 2 min of stop (OFF). The long duration trials lasted for 72 h. At periodic times, samples taken from the MF and RO penetrates to determine the remaining ACN and TOC in the solution. To manage the SPMR's HRT, the RO permeate flow was monitored for 5 h. After 5 h, the MF permeate flow was determined using the MF membrane outflow. Under photo-catalytic operations with and without H₂O₂, the ACN levels in the MF and RO penetrates in the SPMR with suspended and immobilised Sr-LaNiO₃ are shown in Fig. 5. The findings demonstrate that during the initial 3 h, the ACN levels in MF permeates (ACN level in the photo reactor) tended to decrease for all operations. They also grew steadily after 4 h till the completion of the experiment. Throughout the study, the ACN contents in the RO filtrate (effluent) were constant. The ACN molecules flowing via MF membrane were restored to the SPMR via RO refusal as the ACN level in the photo-catalytic reactor increased. The SPMR treatment capability was overloaded due to rising ACN levels in the photo-catalytic reactor. As an outcome, the levels of ACN in the photo reactor rose. The ACN levels in the SPMR with immobilised Sr-LaNiO₃ were greater than those with suspended Sr-LaNiO₃ during the photo-catalytic process (Vis/ Sr-LaNiO₃). This implies that in the constant cycle, SPMRs with the immobilised Sr-LaNiO₃ catalyst had a poorer ACN extraction efficacy than SPMRs with suspended Sr-LaNiO₃. In comparison to the procedures without irradiation, combining H₂O₂ with the photo-catalytic activity improved ACN elimination under observable illumination

for both SPMR entities with suspended and immobilised Sr-LaNiO₃. The ACN elimination efficacy possibilities of the numerous SPMR samples with various photo-catalytic activity were in the following manner: SPMR with dispersed Sr-LaNiO₃ under the Vis/ Sr-LaNiO₃ / H₂O₂ process > SPMR with immobilised Sr-LaNiO₃ under the Vis/ Sr-LaNiO₃ / H₂O₂ procedures > SPMR with dispersed Sr-LaNiO₃ under the Vis/ Sr-LaNiO₃ procedures > SPMR with immobilised Sr-LaNiO₃ under the Vis/ Sr-LaNiO₃ procedures.

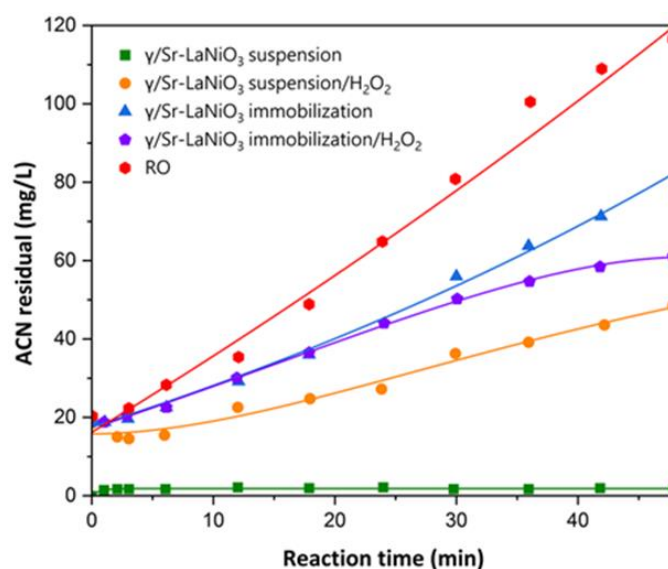


Fig 5. Periodic change of ACN leftover for 4 SPMR systems with RO membrane under long-term settings. Circumstances of the test: pH = 7; starting ACN = 25 mg/L; Sr-LaNiO₃ loading = 2g/L for suspended Sr-LaNiO₃; H₂O₂ = 20 mM for Vis/ Sr-LaNiO₃ / H₂O₂ process; intermittent membrane filtration: 10 minutes ON, 2 minutes OFF.

The majority of ACN molecules went through the MF membrane, whereas the RO membrane discarded them more frequently. Throughout the operating duration, the ACN contents in the RO filtrate for the SPMRs with suspended or immobilised Sr-LaNiO₃ and photo-catalytic procedures with or without H₂O₂ were less than 3 mg/L. The photo-catalytic membrane reactor, which integrated the MF membrane with an RO membrane in an PMR reduced RO membrane contamination as well as enhanced wastewater standard. Furthermore, the mixture resulted in a large rise in ACN levels in the photo reactors, causing an overflow in the SPMR system's processing capability.

The ACN discharges and the percent of elimination by the RO membrane are shown in Table 1. The proportion of ACN discarded by the RO membrane was around 97%, which corresponded to the findings, which showed that RO membranes may exclude negatively charged ACN up to 97% (20). Membrane characteristics, flow rate, and operation

circumstances are all parameters that influence RO membrane denial [29]. Furthermore, 3 processes might alter the solute's filtration by the RO membrane: physicochemical interactions between the solute, solvent and size confinement (sieving, steric effect) and membrane and charge confinement (electrical, Donnan) [30]. ACN has a pKa of 4.15, making it negatively charged, which lowers hydrophobicity and promotes solubility when pH = 6.5. As a result, a significant negative charge on the RO membrane, according to Donnan's theory, effectively denies the ACN, inhibiting diffusion and adsorption across the membrane. The often-employed measure for predicting filtering procedure's elimination effectiveness is molecular weight (21). ACN has a molecular weight of 300 g/mol and dimensions of $1.15 \times 0.95 \times 0.50$ nm, correspondingly. An RO membrane can eliminate molecules with molecular weights greater than 250 g/mol, implying that ACN, with a molecular weight of 300 g/mol, may be processed [31].

Table 1: ACN elimination through Reverse Osmosis membrane rebuff

SPMR process MF permeate	MF permeate = Reverse Osmosis feed (milligram per litre)	Reverse Osmosis permeate (milligram per litre)	Reverse Osmosis rebuff (%)
Vis/suspended Sr-LaNiO ₃	41.23±20.23	1.55±0.53	96.34±2.76
Vis/suspended Sr-LaNiO ₃ / H ₂ O ₂	27.23±11.34	1.45±0.32	95.23±2.45
Vis/immobilized Sr-LaNiO ₃	57.12±33.56	1.99±0.34	96.12±3.01
Vis/immobilized Sr-LaNiO ₃ / H ₂ O ₂	37.23±16.00	1.45±0.23	96.01±2.65

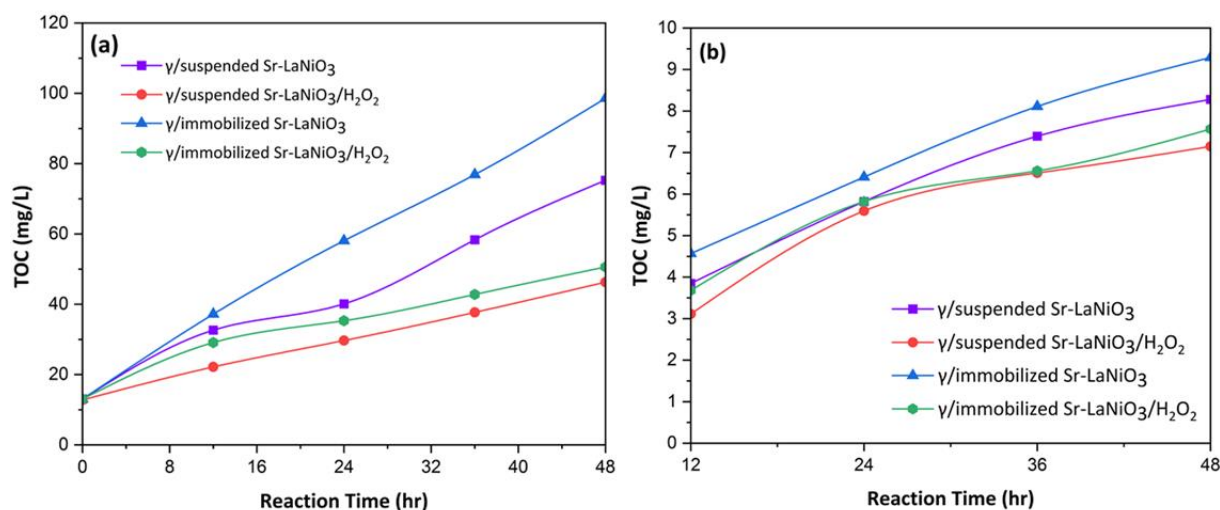


Fig 6. ACN mineralization throughout longstanding circumstances, (a) TOC of MF permeates, (b) TOC of RO permeates. pH = 7; starting ACN = 30 mg/L; Sr-LaNiO₃ loading = 2 g/L for suspended Sr-LaNiO₃; H₂O₂ = 20; periodic membrane filtration of 6 min ON and 2 min OFF.

At the time of 72 h of photo-catalytic oxidation, TOC concentration is eliminated by the MF (Fig. 6a) and RO permeates (Fig. 6b). The TOCs content in the photo-reactors and wastewater rose at the time of the reaction period for both SPMR structures and procedures, according to the findings. In the photo-reactor, the SPMR with immobilised Sr-LaNiO₃ under the Vis/ Sr-LaNiO₃ procedures showed the quickest rise in TOC level. The SPMR with suspended Sr-LaNiO₃ under the Vis/ Sr-LaNiO₃/ H₂O₂ procedure had the least rise in TOC content (Fig. 6a). The restoration of ACN monomers and compounds to the reactor via RO rebuff causes a rise in TOC levels in the photo-reactor. During the reaction period, the TOC contents in the wastewaters also increases (Fig. 6b). The lower molecular weight cut-off (MWCO) of the intervening products that may permeate via the pores of the membrane is connected to the increasing TOC levels in the RO penetrates (wastewater). These findings show that the RO membrane was critical in refusing not only the ACN molecules, but also their yields, and redirecting them to the photo-catalytic reactor for addition of minerals. The mixture of the RO membrane and the SPMR units led to enhanced wastewater standard and higher ACN breakdown into lower molecular weight compounds.

3.2.2 Permeate flux of MF and RO membrane

The penetrate fluxes across the MF and RO membranes was examined for long working term by regulating the proportion of flow, the MF and RO permeates later 5 h. An RO

permeates proportion of flow were monitored and controlled with a valve to keep the RO proportion at a constant flow rate of 2 L/h. With the multiple methods (Vis/ Sr-LaNiO₃ and Vis/ Sr-LaNiO₃ / H₂O₂), the MF permeate fluxes of the SPMR entities change. The data indicates that the MF permeate flow of the SPMR with suspended Sr-LaNiO₃ were lowered rapidly first, by way of a little drop later 30 h, but the MF permeate flux of the SPMR with immobilised Sr-LaNiO₃ were stable first, then fell significantly after 9h. Under the Vis/ Sr-LaNiO₃ method, the SPMR's MF permeate flux reduced faster than under the Vis/ Sr-LaNiO₃ / H₂O₂ operation. A decline in permeate flux, which can be caused by pore blockage, the creation of a cake layer on the surface of the membrane, smaller apertures, or foulant absorption of pores of the membrane, indicates membrane besmirching [32]. The initial decrease in MF permeate flow in the SPMR with scattered Sr-LaNiO₃ was due to nanoparticles of Sr-LaNiO₃ connected to the exterior layer of the membrane, whereas the flux in the SPMR with fixed Sr-LaNiO₃ held steady. When compared to disperse LaNiO₃, there were several benefits to immobilised LaNiO₃ on the membrane, including no need to remove and reuse catalysts, less membrane fouling, and improved wastewater quality owing to smaller membrane pores [33]. Nevertheless, there are certain drawbacks, including a poor catalyst area ratio, the inability to alter catalyst concentration, and the need for membrane renewal when the catalyst on the membrane is depleted. Whereas a LaNiO₃ layer forms on the membrane, LaNiO₃ nanoparticles can lead to the MF permeate flow to decrease [24]. The permeate flow is limited by elevating the LaNiO₃ concentration in the reactor. The naked eye was used to examine the surface of the MF membrane. The cake layer generated upon the MF membrane of the SPMR with dispersed Sr-LaNiO₃ under Vis/ Sr-LaNiO₃ was denser when compared to any other reactors, after 72 h of operation. Permeation was diminished because the combination of ACN and Sr-LaNiO₃ particles formed on the membrane had already become denser. The cake layer of LaNiO₃ with the organic component become denser at high organic levels [23].

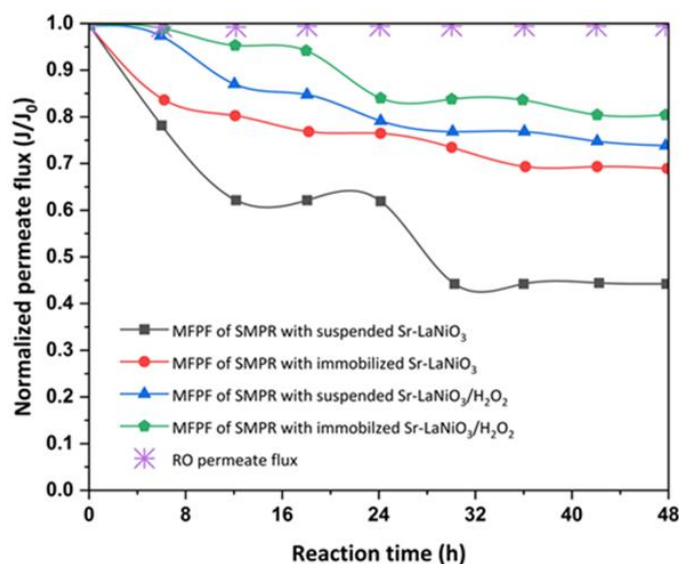


Fig 7. Changes in MF and RO permeate flow in constant settings. Initial ACN = 30 mg/L, Sr-LaNiO₃ loading = 2 g/L for suspended Sr-LaNiO₃; H₂O₂ = 30 mM for Vis/ Sr-LaNiO₃ / H₂O₂ procedure; pH = 7; and sporadic membrane filtration of 6 min ON and 2 min OFF.

Conclusion:

The efficacy of ACN exclusion in the photo-catalytic procedure with and without H₂O₂ was revised by the initial Sr-LaNiO₃ concentrations. Under visible irradiation, the greatest photo-catalytic ACN elimination efficacy was found at a Sr-LaNiO₃ level of 2 g/L. The use of H₂O₂ in conjunction with the photo-catalytic activity increased the effectiveness of ACN elimination. When the SPMRs with suspended and immobilised Sr-LaNiO₃ were compared, it was discovered that the SPMR with suspended catalyst had a higher ACN extraction efficacy because the Sr-LaNiO₃ particles increased ACN elimination. SPMRs with suspended and immobilised Sr-LaNiO₃ have both benefits and drawbacks. By raising the Sr-LaNiO₃ dose, the response proportion in the SPMR with dispersed Sr-LaNiO₃ may be increased. The SPMR with immobilised Sr-LaNiO₃ particle, on the other hand, exhibited less membrane fouling when compared to SPMR with dispersed Sr-LaNiO₃ molecules. When H₂O₂ was combined with the photo-catalytic activity, membrane fouling was reduced compared to when the photo-catalytic method was used separately. The pairing of the RO membrane and the SPMR produced high-quality wastewater. Nevertheless, since ACN and its by-products were restored to the photo reactor via RO repudiation, the ACN and TOC levels in the photo reactor raised. The Sr-LaNiO₃ particles combined with the organic compounds formed a cake layer on the MF membrane surface, which were principally capable in lowering MF permeate flow of the SPMR.

Conflicts of Interest

No Conflict of Interest

Reference

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