INTRODUCTION

Titanium dioxide (TiO₂) photocatalytic process is a promising technique to overcome water pollution due to the use of oxidation-reduction reactions triggered by excitation of electrons under photon irradiation to activate the band gap of TiO₂. Photogeneration process of electron pair (e⁻) and hole (h⁺) can be recombined, so if there is no recombination process, the pair of electrons can migrate to the surface of TiO₂ to participate in the process of reduction-oxidation reactions (redox), each of which allows the formation of superoxide anion radical (•O₂⁻) and hydroxyl radical (•OH)¹⁻³.

TiO₂ is selected as one of semiconductor materials due to it is inert to chemicals, non-toxic and unexpensive in its utilization⁴. TiO₂ has been used in various application fields such as for overcoming environmental pollution, self-clean,
paint industry, photoelectric sensors, anti-fungal material, photovoltaic cells as well as filtering materials. However, the disadvantages of TiO$_2$ compound are it is active only on irradiation of UV light with the band gap value of 3.2 eV, so various researches have been developed to change the TiO$_2$ band gap by inserting non-metal element, while trapping electrons by inserting metal elements. The process of doping on the TiO$_2$ semiconductor has an objective to change the properties of TiO$_2$ in order to make it works in visible light (low-energy) to be applied to the use of sunlight.

Various doping methods development of TiO$_2$ compounds lead to the utilization of very expensive, high toxicity and the difficulty prepared materials. So in this study, it led to the utilization of organic waste such as coconut shell which can encourage the development of new researches combined with P25 compounds using aqueous solvent, which have been developed based on TTIP material as a precursor of TiO$_2$. Coconut shell material can be made into charcoal that has excellent resistance since it has been widely used as harmful metal adsorbent media such as Mn, Zn, Cd and Pb, whereas the TiO$_2$ compounds can be used as water purification from bacteria because of its photoactive property resulting in the molecule breakdown of organic compounds in the water which makes it safe to consume.

Therefore, in this study preparation and characterization of activated carbon from coconut shell doped by P25 was performed to identify the utilization of conventional solvent, i.e distilled water to determine the interaction of both materials using distilled water which is one of the promising method developments in improving the doped TiO$_2$ semiconductor.

**EXPERIMENTAL**

**Preparation of Carbon from Coconut Shell**

Coconut shell, taken from the copra processed waste in the region of Southeast Sulawesi, was cleaned up of its fiber on shell using knife. Subsequently the shell was made into small pieces in size of 3-5 cm, sun-dried and aerated in order to reduce the water content of samples so it would produce good carbon.

**Carbonation and Activation of Coconut Shell**

Carbonation stage of coconut shell was done by conducting pyrolysis process at a temperature of £ 700ºC for 1 hour. Result of coconut shell charcoal carbonation was grinded and sieved using a 170 mesh sieve. Charcoal was then activated in termolite furnace with a temperature of 500ºC and simultaneously nitrogen gas was purged to produce coconut shell activated carbon.

**Synthesis of Carbon-doped TiO$_2$**

Synthesis of coconut shell activated carbon doped P25 was done by performing stoichiometrically ratio of 1:1, where the P25 was calcined in a furnace at a temperature of 500ºC to form anatase crystal then mixed in a beaker with the addition of distilled water. Microscopic of activated carbon doped P25 was observed using SEM, SEM-EDX, XRD and FTIR.

**RESULTS AND DISCUSSION**

**Sample Preparation of Coconut Shell**

The collecting of raw material, coconut shell, was taken from the copra processed waste in the region of Southeast Sulawesi. The collecting process was performed by selecting dried coconut shell and had good thickness to make shorter cleaning process, save time and to have good resistance to heat. Coconut shell was then cleaned thoroughly from coconut fiber attached to surface of shell. The final stage of raw material preparation was sun dried the coconut shell and left it overnight to produce dried shell to make it easy in the process of carbonation.

**Carbonation and Activation of Coconut Shell**

Carbonation was performed by pyrolysis process at a temperature of £ 700ºC for 1 hour and the results of the pyrolysis process produced quite hard charcoal. According to Jahirul et al. pyrolysis at temperatures >500ºC produces endurance and excellent charcoal crystal structure. Furthermore charcoal obtained was grinded to produce fine-grained charcoal, this technique can produce small particles, the smaller the particle, the greater surface area of particle. After the grinding process, then the next stage was sieving which aimed to produce charcoal powder with the same particles (170 mesh). Technique of charcoal activation was done using
physical methods performed by heating process within the termolitte furnace which simultaneously purged with nitrogen gas, expected to produce broader pore size so it can be used as binding medium between P25 compounds to be synthesized.

**Synthesis of Coconut Shell Activated Carbon doped TiO$_2$**

Activated carbon and P25 powder was weighed of 2.5 grams and put into a 100 mL beaker with the addition of a little distilled water, visually of this process, it produced colloid as shown in Fig. 1.

The process of activated carbon doped P25 was to determine the interaction between the activated carbon and P25 in water medium. This process was based on solid state reaction using easy conventional solvent. Moreover, the addition of TiO$_2$ in carbon served as pillarization of the structure, extending the carbon surface so that it has excellent adsorption$^{18}$ and improving the properties of TiO$_2$$^{19}$. Huang et al. reported the bond formation between the carbons doped TiO$_2$ tends to form carbon bond interstitially instead of substitutionally$^{20}$. Application of materials doped TiO$_2$ can be used in the adsorbent process in water treatment to eliminate bacteria contained in the water$^{11}$. This is because TiO$_2$ is a semiconductor which has photocatalysis property to solve various environmental problems among others are for water and air purification, destruction of microorganism such as bacteria and viruses, inactivation of cancer cell, degradation of dye and toxic chemical compounds and manufacture of hydrogen gas from water$^{21}$. TiO$_2$ semiconductor in photocatalytic oxidation process has an advantage in overcoming environmental pollution caused by dye$^7$.

**Characterization**

**Scanning Electron Microscope (SEM)**

Sample preparation of activated carbon doped P25 showed that there had been coating. The distribution of activated carbon to P25 indicated that TiO$_2$ filled the pores of activated carbon. Fig. 2 shows the particle size of activated carbon and P25 resulted 90 µm and 1 µm, respectively. So, it could be inserted between pores of activated carbon. It could be also assumed that the possibility of P25 can be pillarized and enlarge the pores of activated carbon so it enabled bond formation between C-O-Ti-O.

**Scanning Electron Microscopy – Energy Dispersive X-ray Spectroscopy (SEM – EDX)**

SEM-EDX data showed the morphology of synthesized materials from activated carbon doped P25 (Fig. 3) showed the composition of the presence of carbon compounds doped P25. The result of the spectrum, it showed the presence of Ti (Titanium) elements on energy bands, i.e 4.25 KeV and 0.20 KeV on the sample, it still showed a wide range of metals which enable material to be dispersed in samples of P25 which was extracted from ilmenite natural materials$^{22}$. This material showed three elements (C, Ti and O), the percentage of each element were 46.9%; 27.5% and 25.6%, respectively.
Preparation of activated carbon doped P25 was calcined at temperatures of 500°C aimed to form anatase crystals of TiO₂ which is photoactive, as well as to open the pores of activated carbon. From crystal analysis using XRD formed from activated carbon doped P25 (Fig. 4) the formation of carbon peaks on the field of 002 while TiO₂ of anatase crystal shown in fields of 101, 004, 200, 220, 105, 204, 116, 215 and the existence of the rutile crystal shown in fields of 110, 101, 210, 211, 002, 110 formed on the prepared sample. These indicated that the sample has been doped and interactions took place both on the surface of activated carbon and TiO₂.

Characterization using FTIR showed that based on the spectrum (Fig. 5) it was indicated the presence of –OH group stretching at wavenumber of 3500 cm⁻¹ (stretching vibration) with low signal. Similarly, at wavenumber of 2924 cm⁻¹, it showed C–H bonds of the aromatic ring with moderate intensity. Then the wavenumbers of 1645 cm⁻¹–1678 cm⁻¹ showed the C=C bonds of an alkene compound type, while 1516 cm⁻¹–1541 cm⁻¹ showed the C=C bonds of the aromatic ring compound type, this is supported by the presence of C–O at wavenumber of 1112 cm⁻¹. The presence of P25 at wavenumbers of 516 cm⁻¹–677 cm⁻¹.
CONCLUSION

By using conventional solvent (distilled water), it resulted in the formation of colloidal activated carbon from coconut shell doped P25 and had been characterized using SEM and the presence of interactions between activated carbon and P25 with a particle size were 90 µm and 1 µm, respectively. The composition of carbon, titanium and oxide by SEM-EDX are 46.9%; 27.5% and 25.6%, respectively. XRD data on the formation of P25 anatase crystals in the fields of 101, 004, 200, 220, 105, 204, 116, 215 and the carbon peak in the fields of 202. It was supported by FTIR data which was indicated –OH bond (3500 cm\(^{-1}\)); C–H (2924 cm\(^{-1}\)); C=C (1645 cm\(^{-1}\)–1678 cm\(^{-1}\)) and (1516 cm\(^{-1}\)–1541 cm\(^{-1}\)); C–O (1112 cm\(^{-1}\)) and the O-Ti-O (516 cm\(^{-1}\)–677 cm\(^{-1}\)).

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