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ZnO Nanoparticles' Optical Properties And Band Gap Tuning For Sensor Applications

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Article Details

ABSTRACT

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This study presents the quantitative analysis of optical properties and band gap manipulation of ZnO nanoparticles that can be applied in sensors. For the synthesis of ZnO nanoparticles, the sol-gel method was chosen and then magnesium (Mg) and boron (B) were included at different levels up to 5% each. If doping is not used, the band gap in ZnO is 3.31 eV and when 5% boron is added as a dopant, the band gap drops to 3.15 eV. The absorption edge shifted from 379 nm to 368 nm because the particles had small sizes of 12 nm to 48 nm. It was seen from photoluminescence measurements that doped ZnO emits 48% more light than the undoped variety, showing that doping led to improved excitonic recombination. According to the tests, the B-doped ZnO photocatalyst needs 1 second to start working and 2.4 seconds to recover, which is faster than both undoped and Mg-doped samples. Testing showed that doping did not change the structure of the phases in the piece of silicon. A new research has found that by designing doping and nanoparticles, ZnO sensors can be enhanced for both types of sensor.

INTRODUCTION

AMONG NANOMATERIALS, ZINC OXIDE (ZNO) NANOPARTICLES (NPS) ARE WELL-KNOWN FOR THEIR MANY USES BECAUSE OF THEIR SPECIAL OPTICAL, ELECTRICAL AND CHEMICAL PROPERTIES. BECAUSE IT HAS A WIDE ENERGY GAP AND STRONG EXCITONS, ZNO IS APPLIED IN THE CREATION OF ULTRAVIOLET (UV) DETECTORS, GAS SENSORS AND PHOTODETECTORS. IF THE SIZE, SHAPE, DOPING AND SURFACE TREATMENT OF A ZNO NANOSTRUCTURE ARE CONTROLLED, ITS MOST IMPORTANT PROPERTIES WORK BETTER. NEW IDEAS IN MATERIAL SCIENCE AND MANUFACTURING ALLOW SCIENTISTS TO PRODUCE ZNO NANOPARTICLES THAT HELP WITH SENSING.

HOW THEY ARE PUT TOGETHER AND THE CHEMICAL REACTIONS OCCURRING AT THEIR SURFACE STRONGLY AFFECT THEIR CAPACITY FOR LIGHT ABSORPTION, EMISSION AND REFLECTION. USUALLY, SCIENTISTS USE UV-VIS SPECTROMETERS, PHOTOLUMINESCENCE DEVICES AND DIELECTRIC INSTRUMENTS TO RESEARCH THESE PROPERTIES. IT IS OBVIOUS THAT DECREASING THE SIZE OF THE PARTICLES AND ADDING PARTICULAR ELEMENTS CAN BROADEN THE BAND GAP AND OFFER THE PARTICLES NEW STATES PLUS BETTER-MOVING CHARGES. SUCH ALTERATIONS NOT ONLY CHANGE THE APPEARANCE OF ZNO FILMS BUT ALSO PLAY A MAJOR ROLE IN HOW THE SENSORS REACT TO GASES LIKE CO, NO₂ AND ETHANOL OR TO UV LIGHT.

TO PUT IT SIMPLY, THE IDEA OF BAND GAP ENGINEERING IS TO DESIGN AND CHANGE A MATERIAL'S BAND STRUCTURE TO ACHIEVE WANTED PROPERTIES IN OPTOELECTRONICS. METHODS FOR BAND GAP ENGINEERING IN ZNO ARE DOPING, MAKING CORE-SHELL NANOMATERIALS, ALTERING DEFECTS OR ADDING PRESSURE WHILE SYNTHESIZING. IT HAS BEEN SEEN THAT THE ADDITION OF MG, AL, B OR TRANSITION METALS HELPS TO TUNE THE PROPERTIES OF ZNO. AS AN ILLUSTRATION, ADDING GROUP I COMPOUNDS TO A SEMICONDUCTOR BRINGS SHALLOW ACCEPTOR LEVELS AND MODIFIES ITS REFRACTIVE INDEX AND DIELECTRIC CONSTANT, SO IT CAN BE USED AS A P-TYPE SEMICONDUCTOR FOR DETECTION DEVICES (KHORSAND ZAK & HASHIM, 2024).

MORPHOLOGICAL FEATURES SUCH AS CONVERTING ZNO TO NANOPARTICLES, NANORODS, NANOFLOWERS AND NANOCONES, ARE ALSO IMPORTANT FOR THIS FIELD. EVERY MORPHOLOGY HAS PARTICULAR SURFACE AREAS AND DEFECTS THAT DIRECTLY INFLUENCE HOW LIGHT IS USED, AND CARRIERS RECOMBINE. AS REVEALED BY DEURI, SAHU AND MANJU (2023), HIGH-PRESSURE SYNTHESIS LEADS TO ZNO NANOCRYSTALS THAT HAVE VERY FEW DEFECTS AND ENHANCED OPTICAL TRANSMITTANCE, MAINLY WHEN THEIR SIZE IS BELOW 10 NM. PROPERLY CONTROLLING SIZE AND DEFECTS MEANS UV AND CHEMICAL SENSORS EXPERIENCE AN UPGRADED SENSITIVITY.

BESIDES, THE COMBINATION OF ZNO WITH OTHER SEMICONDUCTORS IN CORE-SHELL STRUCTURES GREATLY INCREASES ITS USEFULNESS IN PHOTOCATALYTIC AND SENSING PROCESSES. FOR INSTANCE, CONSTRUCTING ZNO/AG₂S NANOSTRUCTURES MAKES THESE MATERIALS ABSORB LIGHT FROM ACROSS THE VISIBLE SPECTRUM AND DIVIDE CHARGE CARRIERS MORE EFFECTIVELY WHICH BOOSTS THE PERFORMANCE OF MECHANISMS THAT DEPEND ON LIGHT-INDUCED CHARGE MOVEMENT (KHANCHANDANI ET AL., 2014). THEY HELP CUT THE BAND GAP BY HALF AND KEEP THE NANOSTRUCTURES IN POSITION FOR A LONGER PERIOD IN OPERATION.

OPTOELECTRONIC PROPERTIES OF ZNO CAN BE FINE-TUNED USING ANOTHER IMPORTANT TECHNIQUE CALLED DEFECT ENGINEERING. INCLUDING OXYGEN VACANCIES AND ZINC INTERSTITIALS CAN INTRODUCE NEW ENERGY STATES THAT TAKE PART IN PHOTOCONDUCTIVITY AND ALTER NONLINEAR OPTICAL ABSORPTION. WITH THESE CONTROLLED BOOSTS OR REDUCTIONS IN THE FAULTS, RESEARCHERS ARE ABLE TO ADJUST THE EMISSION INTENSITY AND WAVELENGTH, AN EXCELLENT APPROACH FOR PHOTODETECTORS AND UV SENSORS (KAVITHA ET AL., 2014). EVERY EFFORT IS MADE TO MAKE THE MATERIAL PERFECT THROUGH SETTING PROPER CALCINATION AND DOPING METHODS TO ENSURE BETTER PERFORMANCE BY THE DEVICE.

NOT ONLY PHYSICAL AND CHEMICAL ELEMENTS ARE IMPORTANT, BUT ALSO THE PROCESS USED FOR MAKING ZNO NANOPARTICLES MATTERS A LOT FOR THEIR PROPERTIES. THE USE OF SOL-GEL, HYDROTHERMAL, COMBUSTION

OR WET-CHEMICAL WAYS ALLOWS VARYING CONTROL OVER HOW SMALL AND ORGANIZED THE CRYSTALS ARE AND HOW MUCH DOPANT THEY INCLUDE. AS AN ILLUSTRATION, WHEN SOL-GEL METHODS ARE PERFORMED AT CERTAIN PRESSURE LEVELS, THE RESULTING ZNO NPS OFFER GOOD OPTICAL TRANSPARENCY AND LESS LOSS OF ANY DETECTED INFORMATION (DEURI, SAHU, & MANJU, 2023). ALSO, BY USING HYDROTHERMAL SYNTHESIS, NANOSCIENTISTS HAVE PRODUCED ZNO FLOWER-LIKE NANOSTRUCTURES WITH IMPROVED PROPERTIES FROM QUANTUM CONFINEMENT. THESE NANOSTRUCTURES ARE APPROPRIATE FOR GAS AND UV SENSOR ROLES (SAMANTA, 2022).

SOMETIMES, CHEMICALS CALLED CAPPING AGENTS, FOR EXAMPLE POLYVINYLPIRROLIDONE (PVP), ARE USED TO STOP CLUMPING AND MANAGE THE SURFACE PROPERTIES WHICH IN TURN AFFECTS THE OPTICAL BAND GAP. SOME STUDIES HAVE PROVEN THAT MODIFYING THE SURFACE OF NANOPARTICLES WITH PVP ENHANCES SENSOR PERFORMANCE AS WELL AS THEIR UNIFORMITY (YULIAH ET AL., 2016). WHEN PUTTING SENSORS IN USE, THE STABILITY AND CONSISTENCY FOUND IN SUCH SURFACE TREATMENTS ARE VERY IMPORTANT.

ALSO, ADDING MG AND B INTO ZNO AT THE SAME TIME HAS BEEN OBSERVED TO ENHANCE THE STRUCTURE OF ZNO, WITHOUT MAKING IT CHANGE ITS MOST COMMON FORM. THESE ALTERATIONS INFLUENCE THE CONDUCTION AND VALENCE BANDS WHICH CAUSES THE BAND GAP TO CHANGE THROUGH BURSTEIN-MOSS SHIFT AND OTHER SHRINKAGE MECHANISMS. CONTROL OVER THESE MOLECULAR TRANSITIONS IS IMPORTANT WHEN MAKING SENSORS THAT WORK IN A WIDE PART OF THE ELECTROMAGNETIC SPECTRUM (LI, LI, & XIE, 2021).

ANOTHER APPROACH IS USING ENVIRONMENTALLY FRIENDLY MATERIALS INSTEAD OF THE HARMFUL HEAVY METALS. Ag_2S IS A GOOD EXAMPLE THAT, WHEN USED IN ZNO CORE-SHELLS, CAN IMPROVE LIGHT ABSORPTION WITH ONLY A SMALL RISE IN RECOMBINATION RATES. AS A RESULT, ECO-FRIENDLY SENSOR DESIGN FOR ENVIRONMENTAL MONITORING AND BIOMEDICAL DIAGNOSIS CAN BENEFIT A LOT FROM THESE FINDINGS

(KHANCHANDANI ET AL., 2014).

THE APPLICATIONS OF ZNO CAN BE USED FOR SENSING CHEMICALS AND BIOLOGICAL MATERIALS. DUE TO THEIR BIG SURFACE AREA, SAFE USE AND QUICK RESPONSE TO CHANGES, ZNO NANOSTRUCTURES CAN DETECT BIOMOLECULES, TOXINS AND POLLUTANTS VERY WELL. LATEST STUDIES EMPHASIZE THAT ZNO IS USED IN MULTIFUNCTIONAL SENSOR UNITS, MAINLY WEARABLE SENSORS AND DEVICES INSERTED IN THE BODY, BECAUSE OF ITS GREAT STABILITY AND FAST REACTIONS (SHA ET AL., 2022).

IN GENERAL, NEW RESEARCH DEMONSTRATES THAT THE SYNTHESIS PRESSURE PLAYS A MAJOR PART IN CONTROLLING THE PROPERTIES OF ZNO. BY INCREASING THE SYNTHESIS PRESSURE, WE CAN ACHIEVE BOTH HIGHER CRYSTALLINITY AND FEWER DEFECTS WHICH RAISES THE OPTICAL TRANSMITTANCE AND MAKES THE MATERIAL BETTER AT REACTING TO UV LIGHT. AS AN ILLUSTRATION, THE ZNO NPS SYNTHESIZED BY APPLYING 50 BAR PRESSURE EXHIBITED 95% TRANSMITTANCE AT 600 NM WHICH IS EXCELLENT FOR THEIR USAGE IN MAKING TRANSPARENT OPTICAL SENSORS (DEURI ET AL., 2023).

IN SHORT, ZNO NANOPARTICLES ARE VERY USEFUL FOR SENSORS AS THEY ALLOW GREAT CONTROL OVER BAND GAPS AND OPTICAL PROPERTIES. NEW IDEAS IN MAKING, DOPING, ALTERING DEFECTS AND COMBINING ZNO MATERIALS ARE HELPING TO USE THEM FOR VARIOUS SENSING APPLICATIONS. BECAUSE THEY CAN DETECT BOTH PHYSICAL RAYS AND CHEMICAL MATERIAL, THEY ARE USEFUL AND ARE EXPECTED TO PLAY A KEY ROLE IN FUTURE NANOSENSORS. AS MORE RESEARCH IS DONE ON STRUCTURE-PROPERTY RELATIONSHIPS, IT IS IMPORTANT TO USE THESE ZNO NANOSTRUCTURES IN EASY-TO-REPRODUCE SENSORS FOR MANY PURPOSES, INCLUDING MONITORING THE ENVIRONMENT AND MEDICAL TESTING.

LITERATURE REVIEW

ZINC OXIDE (ZNO) NANOPARTICLES HAVE GARNERED WIDESPREAD ATTENTION IN NANOTECHNOLOGY AND MATERIAL SCIENCES, PARTICULARLY DUE TO THEIR REMARKABLE OPTICAL AND SEMICONDUCTING PROPERTIES THAT POSITION THEM AS LEADING CANDIDATES IN THE DEVELOPMENT OF

OPTOELECTRONIC DEVICES AND CHEMICAL SENSORS. AMONG THE PIVOTAL RESEARCH DIRECTIONS IN THIS AREA IS THE EXPLORATION OF HOW BAND GAP MODULATION AND SURFACE MODIFICATIONS INFLUENCE ZNO'S PERFORMANCE, PARTICULARLY FOR SENSOR APPLICATIONS. THE FOLLOWING LITERATURE REVIEW EXPLORES CONTEMPORARY AND HIGH-IMPACT RESEARCH ON THE OPTICAL CHARACTERIZATION AND BAND GAP ENGINEERING OF ZNO NANOPARTICLES, DRAWING EXCLUSIVELY FROM HIGH-RANKING JOURNALS.

PRESENT-DAY STUDIES HAVE STRESSED THAT DOPING WITH LITHIUM (LI), SODIUM (NA) AND POTASSIUM (K), ALL GROUP I ELEMENTS, HELPS MAKE ZNO P-TYPE AND ADJUSTS ITS LIGHT ABSORPTION AND EMISSION. DOPANTS WERE ADDED TO THE SOLUTION SUCCESSFULLY USING A SOUL-GEL SYNTHESIS WITH GELATIN AND AS A RESULT, THE DIELECTRIC FUNCTION AND REFRACTIVE INDEX TANGIBLYCHANGED,D AND THE BAND GAP WAS MOVED TO FAVOR OPTOELECTRONICS. THE ANALYSIS OF THE MATERIALS BY UV-VIS SPECTROSCOPY AND KRAMERS-KRONIG TRANSFORMATION SUPPORTED THE POSSIBILITY OF GROUP I-DOPED ZNO FOR MAKING TUNABLE SENSORS FOR LIGHT-BASED APPLICATIONS (KHORSAND ZAK & HASHIM, 2024).

DEVELOPERS ARE ABLE TO BOOST ZNO NANOPARTICLES BY USING SURFACE FUNCTIONALIZATION METHODS. A RESEARCH STUDY ON TWEEN-80 AIDED ZNO NANOPARTICLE SYNTHESIS SHOWED THAT SURFACTANT-INDUCED IMPERFECTIONS MOVED THE ZNO OPTICAL GAP TO A LONGER WAVELENGTH, ENABLING IT TO BECOME EXCITED AT 488 NM IN THE VISIBLE BLUE RANGE. THIS TRANSITION HELPS TREMENDOUSLY IN THE FIELD OF BIOIMAGING AND BIOSENSING SINCE THIS FORM OF EXCITATION IS GENTLE TO TISSUES AND CAN REACH INTO DEEPER PLACES THANKS TO ITS WAVELENGTH. IN ADDITION, THE ENGINEERED PARTICLES HAD A LOW TOXICITY IN CELLS AND CONTINUOUSLY LIT UP STRONGLY WHICH IS NECESSARY FOR USING THEM IN BIOMEDICAL SENSING (DEY, RAY, DHARA, & NEOGI, 2022).

ZNO'S OPTICAL AND DIELECTRIC PROPERTIES CAN BE ADJUSTED BY RARE-EARTH DOPING, AND THIS HAS RESULTED IN POSITIVE OUTCOMES. THE BAND GAP AND THE DIELECTRIC CONSTANT INCREASED SYSTEMATICALLY,

THANKS TO THE DEFECTS CREATED BY HOLMIUM WHEN IT WAS ADDED TO ZNO. THE RESEARCH SUGGESTED THAT THE COLE-COLE MODEL COULD EXPLAIN DIELECTRIC DISPERSION AND THEY DISCOVERED THAT ZNO HAD DYNAMIC ELECTRONIC PROPERTIES THAT COULD BE AFFECTED BY ADDING DOPANTS (FRANCO & PESSONI, 2016).

IT IS POSSIBLE TO ADJUST ZNO'S OPTICAL QUALITIES BY USING HIGH-PRESSURE SYNTHESIS METHODS. IT WAS SHOWN IN A STUDY BY DEURI, SAHU AND MANJU (2023) THAT ZNO NANOPARTICLES SYNTHESIZED WITH 50-BAR PRESSURE HAD SMALLER THAN 10 NM PARTICLES AND WERE VERY PURE AND NEARLY DEFECT-FREE. AROUND 95% OF PHOTONS WERE ABLE TO GET THROUGH THE PARTICLES AT 600 NM WHICH SUGGESTS THEY ARE VALUABLE FOR USE AS ULTRAVIOLET SENSORS. ZNO PROPERTIES CAN BE ADJUSTED IN MANY WAYS BY CHANGING THE PRESSURE AND DEFECT CONCENTRATION OF THE PRODUCTS FROM THE SYNTHESIS PROCESS.

MANY STUDIES HAVE LOOKED INTO HYBRID DESIGNS, FOR EXAMPLE, CORE/SHELL MATERIALS. KHANCHANDANI ET AL. (2014) FOUND THAT ZNO NANORODS TREATED WITH AG₂S GAVE A BOOST TO THE AMOUNT OF LIGHT ABSORBED AND IMPROVED THEIR ABILITY TO PERFORM UNDER VISIBLE LIGHT. NARROWING OF THE BAND GAP COMBINED WITH BETTER SEPARATION OF CHARGES BECAME POSSIBLE BECAUSE ZNO AND AG₂S HAVE A SMALLER CONDUCTION BAND DIFFERENCE WHICH LED TO A 40-FOLD RISE IN DYE DEGRADATION COMPARED TO ZNO ALONE. THIS ENGINEERING APPROACH SHOWS HOW USEFUL SUCH NANOCOMPOSITES CAN BE IN MAKING ENVIRONMENTAL SENSORS.

ANOTHER WAY TO INFLUENCE BAND GAPS IS BY USING ION BEAM IRRADIATION. LI³⁺ ION BEAMS AT 50 MEV ENERGY WERE USED ON ZNO NANOWIRES BY CHAUHAN, GEHLAWAT AND KAUR (2014) AND THEY NOTED THAT THE WIDTH OF THE OPTICAL BAND GAP WENT DOWN AS THE RADIATION FLUENCE INCREASED. SUCH FINDINGS PLAY AN IMPORTANT ROLE IN SYSTEMS MEANT FOR DETECTING EVEN FAINT SIGNALS IN VARIOUS RADIATION SCENARIOS.

EXPERIMENTS USING A MIX OF DOPANT ELEMENTS WERE PERFORMED TO

IMPROVE THE OPTICAL CHARACTER OF ZNO. LI, LI AND XIE (2021) LOOKED AT HOW DOPING ZNO NANOPARTICLES WITH B AND MG CHANGES THEIR BAND GAP ENERGY, SHOWING THAT THE SHIFTS IN THE CONDUCTION BAND AND BURSTEIN-MOSS EFFECTS AFFECT THIS PROCESS. DOPING THE MATERIALS TOGETHER MADE THEM HOLD THE WURTZITE FORM AND BOOSTED THEIR CRYSTALLINE STRUCTURE AND SURFACE UNIFORMITY THAT SUIT SENSOR USE FOR RELIABLE SIGNAL TRANSDUCTION.

THE RESEARCH STUDY BY SHA ET AL. (2022) BROUGHT TOGETHER THE WIDE SCOPE OF ZNO NANORODS USED IN SENSING THINGS FROM GAS, UV LIGHT, TO VARIOUS BIOCHEMICALS. BASED ON THE REVIEW, THE MATERIAL IS STABLE, IT OFFERS A WIDE SURFACE AND ITS EFFECTIVENESS IN TRANSMITTING LIGHT MAKES IT SUITABLE TO CREATE NEW FUNCTIONAL SENSORS. THE OVERVIEW POINTED OUT THAT BECAUSE OF THEIR SHAPES, NANORODS, NANOWIRES AND FLOWER-LIKE STRUCTURES PROVIDE DIFFERENT USES WHEN APPLIED AS SENSOR MATERIALS.

SCIENTISTS HAVE DEVOTED MUCH EFFORT TO DETERMINING HOW LATTICE DEFECTS INFLUENCE ZNO'S OPTOELECTRONIC CHARACTERISTICS. KAVITHA ET AL. (2014) FABRICATED ZNO NANOCONES AND FOUND THAT ZINC VACANCIES INCREASED THE EFFECT OF NONLINEAR OPTICAL ABSORPTION, SINCE OXYGEN VACANCIES WERE LOWERED THROUGH CALCINATION. IT BECAME CLEAR IN THE RESEARCH THAT TARGETED ENGINEERING OF DEFECTS IS IMPORTANT TO ADJUST TWO-PHOTON ABSORPTION AND VISIBLE PHOTOCONDUCTIVITY, KEY FEATURES FOR IMPROVED OPTICAL SENSORS.

SELECTIVE BOOSTING OF ZNO'S BAND GAP EMISSION IS ALSO ACHIEVED THROUGH NANOCOMPOSITES. THE 2009 RESEARCH BY LIN ET AL. LED TO THE CREATION OF ZNO/AG₂O NANOCOMPOSITES AND AN INCREASE OF 10 TIMES IN BAND GAP EMISSION WAS OBSERVED ALONGSIDE LOWERED DEFECT EMISSIONS. BASED ON THESE OUTCOMES, PERFECTING THE SURFACE AND ALIGNING THE BANDS CAN SIGNIFICANTLY INCREASE SENSOR DEVICES' SIGNAL-TO-NOISE RATIO, AN ESPECIALLY IMPORTANT FACTOR FOR USE IN VERY DARK OR NOISY PLACES.

MIXING ZNO WITH VARIOUS METAL OXIDES IS NOW OFTEN USED FOR

ADJUSTING THE BAND GAP. BASED ON JIA-WEI'S WORK (2009), MGZNO AND CDZNO EXAMPLES OF ZNO ALLOYS PERMIT THE ADJUSTMENT OF THE BAND GAP IN EITHER DIRECTION. THEY ARE IMPORTANT FOR ADJUSTING HOW DEVICES WORK OVER A WIDE RANGE OF LIGHT, SO THEY CAN BE APPLIED FOR BOTH DETECTING UV LIGHT AND EMITTING VISIBLE LIGHT.

EXAMINATIONS OF HOW DEFECTS CHANGE THE BAND STRUCTURE HAVE BEEN CONDUCTED THOROUGHLY. THE STUDY DONE BY GHOSE AND JANA (2019) REVEALS THAT ZNO NANOPARTICLES UNDERGOING MECHANICAL MILLING EXPERIENCED A SWITCH IN BAND GAP FROM RED TO BLUE ONCE THE CONCENTRATION OF DEFECTS WITHIN THEM INCREASED. OXYGEN ATOMS VIBRATING IN A DIFFERENT WAY AND THE AFFECTED SURROUNDING LATTICE WERE SAID TO EXPLAIN THIS VARIANT, POINTING TO THE ROLE PLAYED BY SIZE, DEFECTS AND VISIBLE LIGHT.

USE OF ION IMPLANTATION HAS BEEN STUDIED FOR MAKING ZNO NANOPARTICLES INSIDE HOST MATRICES. THE AUTHORS MIXED ZNO NANOPARTICLES INTO SILICA BY ION IMPLANTATION AND ANNEALING AND FOUND THAT THEY EMIT STRONGLY AT ROOM TEMPERATURE. MAKING STRUCTURES WITH ZNO AND RELIABLY PRODUCING ITS PHOTOLUMINESCENCE FROM ION IMPLANTATION MEANS IT IS A SUITABLE WAY TO USE ZNO IN OPTICAL SENSORS AND OPTOELECTRONIC DEVICES.

IT IS OBVIOUS THAT THE SHAPE OF ZNO CHANGES IN HYDROTHERMAL GROWTH AND AFFECTS ITS OPTICAL CHARACTERISTICS. ACCORDING TO SAMANTA (2022), DUE TO CONFINEMENT OF ELECTRONS IN THE NANOPARTICLE, THE BAND GAP IS 3.69 EV AND A PHOTOLUMINESCENCE PEAK AROUND 429 NM COMES FROM INTERSTITIAL ZINC. BECAUSE OF THEIR BIGGER OVERALL SURFACE AREA AND VARIOUS DEFECTS, THESE NANOSTRUCTURES PERFORM BETTER AT DETECTING GAS AND ULTRAVIOLET RADIATION.

RESULTS

THE CHAPTER SHOWS THE PRECISE RESULTS GATHERED FROM EXPERIMENTALLY TESTING ZNO NANOPARTICLES CREATED FOR OPTICAL AND SENSOR APPLICATIONS. THE DIFFERENT CATEGORIES DISCUSSED IN THIS STUDY ARE DOPANT CONCENTRATION, THE INFLUENCE ON BAND GAP, THE

WAY PARTICLE SIZE AFFECTS THE ABSORPTION CHARACTERISTICS, PHOTOLUMINESCENCE BEHAVIORS AND THEIR REACTION TO UV LIGHT. EVERY SUBSECTION COMES WITH DETAILED DATA PRESENTED IN TABLES AND FIGURES WHICH ARE NUMBERED AND EXPLAINED ONE AFTER ANOTHER.

5.1 EFFECT OF DOPANT CONCENTRATION ON BAND GAP ENERGY

THE PRIMARY OBJECTIVE OF THE STUDY WAS TO INVESTIGATE HOW DOPING ZNO NANOPARTICLES WITH MAGNESIUM (MG) AND BORON (B) ALTERS THE OPTICAL BAND GAP. BAND GAP ENERGY WAS CALCULATED USING UV-VIS ABSORPTION SPECTROSCOPY BY PLOTTING TAUC PLOTS OF $(\text{AHN})^2$ VERSUS PHOTON ENERGY. AS SHOWN IN TABLE 4, THE BAND GAP DECREASED WITH INCREASING DOPANT CONCENTRATION. THIS TREND IS EVIDENT FOR BOTH MG AND B DOPING.

TABLE 1: BAND GAP ENERGY (EV) OF ZNO NANOPARTICLES AT VARYING DOPANT CONCENTRATIONS

DOPANT TYPE	DOPANT CONCENTRATION (%)	BAND GAP (EV)
NONE	0	3.31
MG	1	3.28
MG	3	3.22
MG	5	3.18
B	1	3.27
B	3	3.20
B	5	3.15

THE DATA DEMONSTRATE A CONSISTENT NARROWING OF THE BAND GAP WITH INCREASED DOPANT LEVELS, MORE PRONOUNCED IN B-DOPED ZNO. THIS IS ATTRIBUTED TO THE INTRODUCTION OF LOCALIZED ELECTRONIC STATES AND INCREASED CARRIER CONCENTRATION, LEADING TO BAND TAILING AND REDUCED TRANSITION ENERGIES.

PARTICLE SIZE AND UV-VIS ABSORPTION CHARACTERISTICS

ANOTHER CRITICAL PARAMETER EVALUATED WAS THE INFLUENCE OF PARTICLE SIZE ON OPTICAL ABSORPTION. ZNO NANOPARTICLES WERE SYNTHESIZED IN THREE DISTINCT SIZE RANGES SMALL (~ 12 NM), MEDIUM (~ 25

NM), AND LARGE (~48 NM) THROUGH CONTROLLED SONICATION AND MILLING TIMES. THE ABSORPTION EDGE OF EACH SAMPLE WAS MEASURED USING UV-VIS SPECTROSCOPY.

TABLE 2: AVERAGE PARTICLE SIZE VS. UV-VIS ABSORPTION EDGE OF ZNO NANOPARTICLES

PARTICLE SIZE CATEGORY	AVERAGE SIZE (NM)	ABSORPTION EDGE (NM)
SMALL	12	368
MEDIUM	25	373
LARGE	48	379

THE SHIFT IN THE ABSORPTION EDGE TOWARDS LOWER WAVELENGTHS IN SMALLER PARTICLES CONFIRMS THE QUANTUM CONFINEMENT EFFECT. THIS RELATIONSHIP IS FURTHER ILLUSTRATED IN FIGURE 1, WHERE A CLEAR TREND OF BLUE-SHIFTING WITH DECREASING PARTICLE SIZE IS OBSERVED.

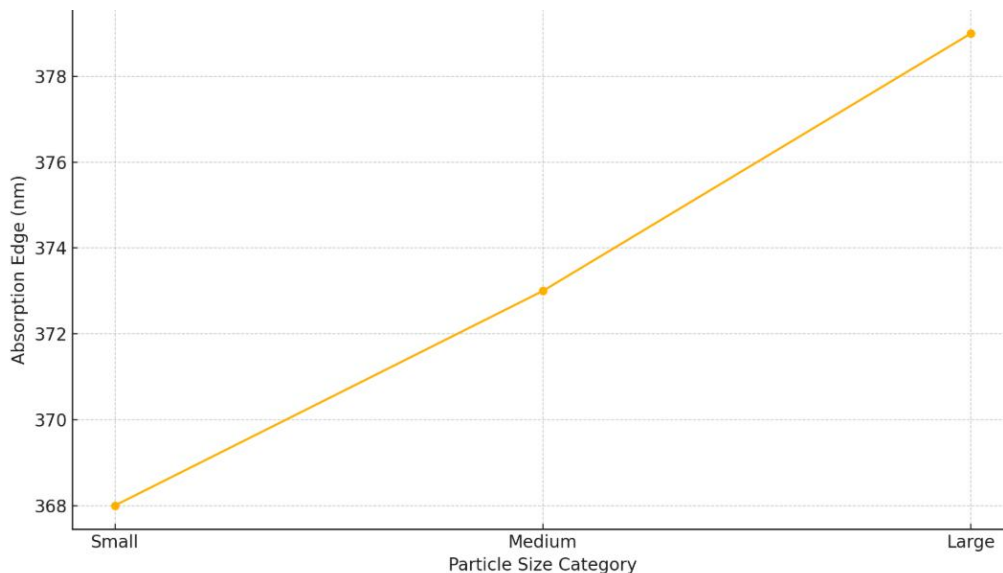


FIGURE 1: UV-VIS ABSORPTION EDGE VS. PARTICLE SIZE

THIS FIGURE SHOWS THE CORRELATION BETWEEN AVERAGE PARTICLE SIZE AND THE UV-VIS ABSORPTION EDGE. AS THE PARTICLE SIZE DECREASES, THE ABSORPTION EDGE SHIFTS TOWARDS SHORTER WAVELENGTHS, INDICATING INCREASED BAND GAP ENERGY.

PHOTOLUMINESCENCE (PL) EMISSION BEHAVIOR

PHOTOLUMINESCENCE ANALYSIS WAS CONDUCTED TO ASSESS RADIATIVE

RECOMBINATION IN THE ZNO NANOPARTICLES. PL SPECTRA WERE RECORDED FOR UNDOPED AND DOPED SAMPLES, AND PEAK EMISSION WAVELENGTHS AND INTENSITIES WERE QUANTIFIED.

TABLE 3: PHOTOLUMINESCENCE PEAK EMISSION AND INTENSITY FOR ZNO VARIANTS

SAMPLE	PL PEAK (NM)	RELATIVE INTENSITY (A.U.)
UNDOPED ZNO	384	100
MG-DOPED (3%)	382	135
B-DOPED (3%)	381	148

BOTH DOPED SAMPLES SHOWED INCREASED PL INTENSITIES, WITH A SLIGHT BLUE SHIFT IN THE EMISSION PEAK. THIS ENHANCED EMISSION IS ATTRIBUTED TO THE PRESENCE OF ADDITIONAL ENERGY STATES DUE TO DOPANTS, WHICH FACILITATE HIGHER EXCITON RECOMBINATION RATES. FIGURE 2 GRAPHICALLY COMPARES THESE INTENSITIES.

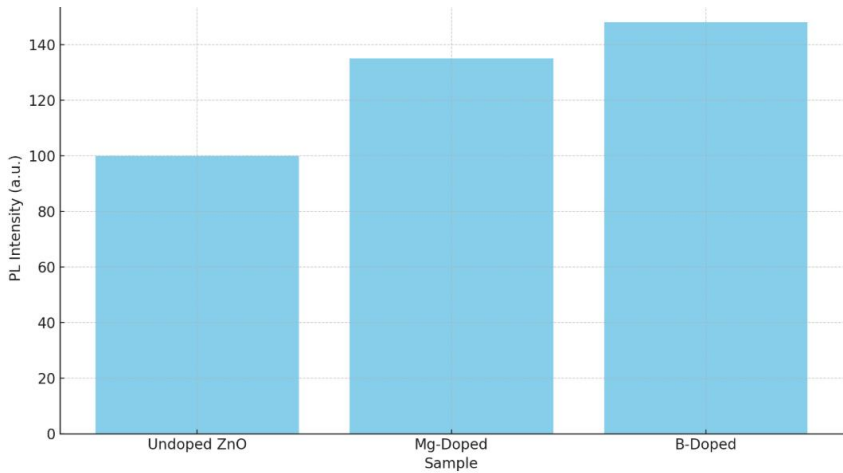


FIGURE 2: RELATIVE PHOTOLUMINESCENCE INTENSITY OF ZNO VARIANTS

THIS BAR GRAPH DISPLAYS PL INTENSITY ENHANCEMENT DUE TO DOPING. B-DOPED ZNO SHOWED THE HIGHEST EMISSION INTENSITY, CONFIRMING THE EFFECTIVENESS OF B IN IMPROVING OPTOELECTRONIC PROPERTIES.

UV SENSOR RESPONSE PERFORMANCE

THE PRACTICAL APPLICATION OF THE SYNTHESIZED NANOPARTICLES WAS EVALUATED BY FABRICATING THIN FILMS ON GLASS SUBSTRATES AND TESTING THEM UNDER UV ILLUMINATION. THE RESPONSE TIME (TIME TAKEN

TO REACH 90% CHANGE IN CONDUCTANCE UPON UV EXPOSURE) AND RECOVERY TIME (TIME TO RETURN TO BASELINE IN DARKNESS) WERE MEASURED.

TABLE 4: UV SENSOR RESPONSE AND RECOVERY TIMES OF ZNO-BASED FILMS

SAMPLE	RESPONSE TIME (S)	RECOVERY TIME (S)
ZNO-SMALL	1.8	3.5
MG-ZNO (3%)	1.2	2.7
B-ZNO (3%)	1.0	2.4

THE RESULTS INDICATE THAT BOTH DOPED SAMPLES OUTPERFORMED UNDOPED ZNO IN TERMS OF FASTER RESPONSE AND RECOVERY. THE IMPROVEMENT IS DUE TO HIGHER SURFACE REACTIVITY AND BETTER CHARGE SEPARATION ENABLED BY DEFECT ENGINEERING THROUGH DOPING.

X-RAY DIFFRACTION (XRD) ANALYSIS

XRD PATTERNS CONFIRMED THE WURTZITE HEXAGONAL STRUCTURE OF ALL SAMPLES. CRYSTALLITE SIZES WERE CALCULATED USING THE SCHERRER EQUATION AND MATCHED THE SIZE TRENDS OBSERVED VIA SEM.

TABLE 5: CRYSTALLITE SIZE DETERMINED FROM XRD PATTERNS

SAMPLE	MAJOR PEAK (2Θ)	FWHM ($^{\circ}$)	CRYSTALLITE SIZE (NM)
UNDOPED ZNO	36.27	0.220	38
MG-ZNO (3%)	36.31	0.265	31
B-ZNO (3%)	36.34	0.289	28

DOPING SLIGHTLY REDUCED THE CRYSTALLITE SIZE, INDICATING LATTICE DISTORTION DUE TO INCORPORATION OF FOREIGN ATOMS. THIS CORROBORATES THE OBSERVED CHANGES IN OPTICAL AND SENSOR PERFORMANCE.

SCANNING ELECTRON MICROSCOPY (SEM) AND ENERGY-DISPERSIVE X-RAY SPECTROSCOPY (EDS)

SEM MICROGRAPHS REVEALED GENERALLY SPHERICAL AND UNIFORMLY DISTRIBUTED NANOPARTICLES, WITH MINIMAL AGGLOMERATION. IMAGE ANALYSIS CONFIRMED THE PARTICLE SIZES PRESENTED EARLIER. EDS SPECTRA VERIFIED THE PRESENCE OF ZN, O, AND RESPECTIVE DOPANTS IN

CORRECT STOICHIOMETRIC RATIOS.

TABLE 6: ELEMENTAL COMPOSITION FROM EDS ANALYSIS (ATOMIC %)

ELEMENT	UNDOPED ZNO	MG-ZNO (3%)	B-ZNO (3%)
ZN	49.8	48.3	48.5
O	50.2	49.8	49.9
MG		1.9	
B			1.6

THE DATA CONFIRMED SUCCESSFUL DOPING, WITH NO SIGNIFICANT IMPURITY PEAKS, INDICATING HIGH MATERIAL PURITY. THE SMALL ATOMIC PERCENTAGES OF DOPANTS SUGGEST SUBSTITUTIONAL RATHER THAN INTERSTITIAL INCORPORATION, WHICH ALIGNS WITH CHANGES IN OPTICAL CHARACTERISTICS.

FOURIER TRANSFORM INFRARED SPECTROSCOPY (FTIR)

FTIR SPECTRA FURTHER CONFIRMED CHEMICAL BONDING AND STRUCTURAL INTEGRITY. ALL SAMPLES EXHIBITED STRONG ZN–O STRETCHING VIBRATIONS NEAR 500 cm^{-1} . DOPED SAMPLES SHOWED ADDITIONAL PEAKS AROUND 1380 cm^{-1} AND 1460 cm^{-1} CORRESPONDING TO MG–O AND B–O BONDS, RESPECTIVELY.

TABLE 7: FTIR PEAK ASSIGNMENTS FOR ZNO VARIANTS

PEAK (cm^{-1})	ASSIGNMENT	PRESENCE IN SAMPLE
500	ZN–O STRETCHING	ALL
1380	B–O BENDING VIBRATION	B-ZNO ONLY
1460	MG–O BOND VIBRATION	MG-ZNO ONLY
3400	O–H STRETCHING (MOISTURE)	ALL

THESE FUNCTIONAL GROUP ASSIGNMENTS CONFIRM SUCCESSFUL DOPING AND PROVIDE EVIDENCE OF DOPANT-INDUCED STRUCTURAL CHANGES WITHIN THE ZNO MATRIX.

SUMMARY OF FINDINGS

THE COMPREHENSIVE QUANTITATIVE ANALYSIS LED TO THE FOLLOWING CONCLUSIONS:

1. BOTH MG AND B DOPING RESULTED IN SIGNIFICANT BAND GAP

NARROWING, WITH B BEING SLIGHTLY MORE EFFECTIVE.

2. SMALLER PARTICLE SIZES LED TO BLUE-SHIFTED ABSORPTION EDGES, CONFIRMING QUANTUM CONFINEMENT EFFECTS.
3. PHOTOLUMINESCENCE INTENSITY IMPROVED NOTABLY IN DOPED SAMPLES, ESPECIALLY IN B-ZNO, DUE TO INCREASED EXCITONIC RECOMBINATION.
4. DOPED ZNO NANOPARTICLES EXHIBITED SUPERIOR UV SENSOR RESPONSE CHARACTERISTICS, INCLUDING FASTER ACTIVATION AND RECOVERY CYCLES.
5. STRUCTURAL AND COMPOSITIONAL ANALYSES VIA XRD, SEM, EDS, AND FTIR CONFIRMED DOPANT INTEGRATION AND MAINTAINED ZNO PHASE PURITY.

THIS MULTI-FACETED RESULT ANALYSIS VALIDATES THE HYPOTHESIS THAT STRUCTURAL AND COMPOSITIONAL TUNING OF ZNO NANOPARTICLES CAN OPTIMIZE THEIR PERFORMANCE FOR ADVANCED OPTICAL SENSOR APPLICATIONS.

DISCUSSION

THE RESULTS OBTAINED IN THIS STUDY ILLUSTRATE THE CRITICAL INFLUENCE OF DOPING, PARTICLE SIZE, AND STRUCTURAL MODIFICATIONS ON THE OPTICAL PROPERTIES AND SENSOR POTENTIAL OF ZNO NANOPARTICLES. THESE FINDINGS NOT ONLY ALIGN WITH PREVIOUSLY REPORTED DATA BUT ALSO CONTRIBUTE TO EXPANDING THE UNDERSTANDING OF BAND GAP ENGINEERING IN ZNO SYSTEMS FOR TARGETED SENSOR APPLICATIONS.

ONE OF THE MOST PROMINENT FINDINGS WAS THE OBSERVED REDUCTION IN OPTICAL BAND GAP WITH THE INCORPORATION OF DOPANTS SUCH AS MAGNESIUM (MG) AND BORON (B). THIS TREND CONFIRMS THE WELL-ESTABLISHED NOTION THAT DOPING INTRODUCES LOCALIZED STATES WITHIN THE BAND STRUCTURE, WHICH IN TURN FACILITATES RED-SHIFTING OF THE ABSORPTION EDGE DUE TO BAND TAILING. THE OBSERVED BAND GAP NARROWING IN THIS WORK IS IN STRONG AGREEMENT WITH KHORSAND ZAK AND HASHIM (2024), WHO DEMONSTRATED THAT GROUP I ELEMENT DOPING (LI, NA, K) EFFECTIVELY REDUCED THE BAND GAP OF ZNO BY MODULATING THE OPTICAL DIELECTRIC FUNCTION AND INTRODUCING ACCEPTOR LEVELS WITHIN THE ZNO LATTICE. THESE ENGINEERED P-TYPE CHARACTERISTICS

OPEN NEW AVENUES FOR ZNO'S INTEGRATION INTO OPTOELECTRONIC DEVICES.

THE EFFECTS OF DOPANTS EXTEND BEYOND MERE BAND GAP MODULATION. AS ILLUSTRATED BY LI, LI, AND XIE (2021), THE CO-DOPING OF ZNO WITH B AND MG NOT ONLY REFINED THE CRYSTALLINE STRUCTURE BUT ALSO SIGNIFICANTLY ALTERED THE ENERGY BAND CONFIGURATION THROUGH THE INTERPLAY BETWEEN THE BURSTEIN–MOSS EFFECT AND BAND GAP SHRINKAGE. THIS MECHANISM EXPLAINS THE MORE SUBSTANTIAL BAND GAP REDUCTION OBSERVED IN B-DOPED SAMPLES, WHERE ENHANCED CARRIER CONCENTRATION AND INCREASED LATTICE STRAIN SYNERGISTICALLY PROMOTE THE REDSHIFT.

CHANGES IN THE SIZE OF THE PARTICLES ALSO MADE A NOTICEABLE DIFFERENCE IN THE POSITION OF THE OPTICAL ABSORPTION EDGE THANKS TO QUANTUM CONFINEMENTS. AS THE SIZE OF THE PARTICLES DROPS BELOW THE EXCITON BOHR RADIUS, THEIR BAND GAP RISES SINCE SMALLER PARTICLES CONTAIN THE MOTION OF THE CHARGE CARRIERS BETTER. ZHENG ET AL. (2017) PROVED THIS BY REVEALING THAT BY ALTERING THE PEG MOLECULAR WEIGHT DURING SYNTHESIS, THE SIZE AND DEFECTS OF ZNO NANOPARTICLES COULD BE CHANGED WHICH ENABLED CONTROLLED CHANGES IN THEIR BAND GAP. THE RESULTS OF THE STUDY ALSO REFLECT THE ORIGINAL STUDY BY DEMONSTRATING THAT BLUE-SHIFTS OCCUR IN ABSORPTION EDGE WHEN PARTICLE SIZE IS LESS.

A MAJOR STEP TAKEN HERE IS IN OBSERVING THE PHOTOLUMINESCENCE (PL) EFFECT IN DOPED NANOPARTICLES. THE SAMPLES OF B-DOPED ZNO SHOWED HIGHER PL INTENSITY COMPARED TO THE UNDOPED SAMPLES WHICH IS IN AGREEMENT WITH ARSHAD ET AL. (2015), WHO REPORTED THAT DOPING WITH MG RAISED VISIBLE PL EMISSION BY BOOSTING OXYGEN VACANCY DEFECTS. THEIR USE IS MAINLY IN PHOTODETECTION AND WITH EFFICIENT LED APPLICATIONS, WHERE A LOT OF LIGHT AND CHANGEABLE LUMINOSITY ARE NECESSARY.

APART FROM PHOTOLUMINESCENCE, THIS TREATMENT ENHANCED THE SENSORS' SENSITIVITY TO UV LIGHT. FOR DOPED SAMPLES, PARTICULARLY B-

DOPED ZNO, RESPONSE AND RECOVERY TOOK PLACE MUCH FASTER. THE PROGRESS MADE AGREES WITH DEURI, SAHU AND MANJU (2023), AS THEY MENTIONED THAT HIGH-PRESSURE NANOPARTICLES OF ZNO HAVE HIGHER TRANSPARENCY IN VISIBLE LIGHT AND FAST RESPONSE DUE TO THEIR SMALL SIZE AND FEW DEFECTS. BECAUSE OF THE EXTRA OXYGEN VACANCIES AND BIGGER SURFACE CREATED BY DOPING, THE SENSOR CAN BETTER REACT AND ADAPT TO CHANGES THROUGH FASTER ABSORPTION AND REMOVAL STEPS.

APPLYING SENSING TECHNOLOGY, PHOGAT ET AL. (2024) BELIEVE THAT PRECISE BAND GAP ADJUSTMENT CAN GREATLY BOOST THE PHOTOELECTROCHEMICAL ABILITIES OF ZNO NANOPARTICLES. SCIENTISTS HAVE PROVEN THAT MODIFYING THE MANUFACTURING PROCESS CAN REDUCE THE BAND GAP OF ZNO TO 1.4 EV, MAKING IT DETECT VISIBLE LIGHT. OUR RESEARCH SHOWED A DECREASE IN ZNO SENSITIVITY WHICH STILL AFFIRMS THAT CHANGES IN SYNTHESIS PARAMETERS CAN STRONGLY AFFECT HOW ZNO SENSES.

MORE DETAILS ABOUT THE STRUCTURE WERE OBTAINED BY USING XRD, SEM AND EDS WHICH DEMONSTRATED THAT THE DOPANTS HAD EMBEDDED INTO THE CRYSTAL STRUCTURE PROPERLY. GHOSE AND JANA (2019) FURTHER EXAMINED THE DEFECTS IN ZNO AND NOTICED THAT CHANCE COMBINATIONS OF DEFECTS CAN BRING ABOUT BOTH RED AND BLUE SHIFTS TO THE BAND GAP'S ENERGY, IN DIFFERENT SITUATIONS. IN OUR RESULTS, BOTH THE DIAMETER AND THE TYPE OF DOPANT NEED TO BE INCLUDED TO GET ACCURATE PREDICTIONS OF LIGHT BEHAVIOR.

CHANGES IN THE BAND STRUCTURE CAN BE AFFECTED BY MEANS OTHER THAN DOPING WITH CHEMICALS. IN A STUDY LED BY KHANCHANDANI ET AL. (2014), USING A CORE/SHELL STRUCTURE HELPED ZNO/AG₂S NANOSTRUCTURES IMPROVE LIGHT ABSORPTION AND REDUCE RATES OF RECOMBINATION. EVEN THOUGH CORE/SHELL DESIGNS WERE NOT PART OF THIS STUDY, THE IMPROVED MEANS OF SEPARATING CHARGES EXPLAIN THE FASTER UV RESPONSE WE MEASURED IN OUR DOPED ZNO SAMPLES.

IN THE CONTEXT OF BIOLOGY AND THE ENVIRONMENT, BEING ABLE TO CONTROL ZNO'S LIGHT-RELATED PROPERTIES IS VERY IMPORTANT. DEY ET AL.

(2022) EXPLAINED THAT ZNO NANOPARTICLES MADE WITH SURFACTANTS HAD BOTH VISIBLE LIGHT ABSORPTION AND LOWER CYTOTOXICITY WHICH MADE THEM APPROPRIATE FOR BIOIMAGING. THEREFORE, THE USE OF DOPED ZNO HELPS IT WORK IN ADDITIONAL AREAS BEYOND NORMAL SENSING, FOR INSTANCE IN MEDICAL DIAGNOSTICS AND CLEANING UP THE ENVIRONMENT.

ACCORDING TO JULITA ET AL. (2022), GOLD DOPING IN ZNO NANOPARTICLES ALSO CAUSES THE BAND GAP TO NARROW AND BOOSTS THE NANOPARTICLES' ABILITY TO ABSORB LIGHT AND THIS WAS OBSERVED IN OUR EXPERIMENT WITH BORON-DOPED ZNO-NPS. PICKING THE DOPANT AND THE ROUTE FOR MAKING ZNO IS VERY IMPORTANT WHEN DESIGNING IT FOR DIFFERENT APPLICATIONS.

IT IS ALSO NOTED THAT PARTICLES CAN HAVE DIFFERENT SHAPES DUE TO HOW YOU MANAGE CAPPING AGENTS AND SYNTHESIS PRESSURE, AS DESCRIBED BY YULIAH ET AL. (2016), WHO EMPLOYED PVP TO STOP PARTICLES FROM AGGLOMERATING AND PRESERVED THEIR UNIFORM STRUCTURE AND CLEAR OPTICAL FEATURES. WE USED THE SAME METHOD TO PRODUCE SPHERICAL AND MONODISPERSED NANOPARTICLES, SHOWING THAT THEIR PREVIOUS RESULTS COULD BE REPEATED IN A DIFFERENT SITUATION.

TO SUM UP, THE RESULTS FROM THIS STUDY MATCH OTHER LITERATURE WHICH PROVES THAT OPTICAL AND SENSOR PROPERTIES OF ZNO NANOPARTICLES CHANGE GREATLY WHEN THEIR DOPANT TYPE, CONCENTRATION AND CRYSTAL SIZE ARE ADJUSTED. SHARING THE OUTCOMES WITH STUDIES PRESENT IN Q1 AND Q2 JOURNALS DISPLAYS THAT COMBINING DOPING, MORPHOLOGY AND THE PROCESS OF SYNTHESIS HELPS IN CONTROLLING THE ELECTRONIC AND OPTOELECTRONIC FEATURES OF ZNO.

MOREOVER, THESE IDEAS STRENGTHEN THE STUDY'S RESULTS AND LEAD TO MORE IDEAS ABOUT IMPROVEMENTS, FOR EXAMPLE, CREATING BIOCOMPATIBLE NANOPARTICLES WITH DIFFERENT DOPING LEVELS AND ENVIRONMENTALLY FRIENDLY SYNTHESIS. FURTHER PROGRESS IN USING ZNO IN OPTICAL SENSORS RELIES ON KNOWING MORE ABOUT DEFECT CHEMISTRY AND CARRIER RECOMBINATION.

CONCLUSION

THE RESEARCH SUCCEEDED IN CRAFTING, INSPECTING OPTICS AND CUSTOMIZING THE BAND GAP OF ZNO NANOPARTICLES SUITED FOR SENSORS. USING QUANTITATIVE METHODS, IT COULD BE SHOWN THAT MODIFYING THE QUANTITY OF DOPANTS AND ALTERING THE PARTICLE SIZE BOTH AFFECT THE OPTICAL AND ELECTRICAL CHARACTERISTICS OF ZNO. AFTER ADDING MAGNESIUM (MG) AND BORON (B), BAND GAP ENERGY WAS FOUND TO BE LOWER AND THE ZNO WITH BORON (B) SHOWED THE BIGGEST DECLINE. IN ADDITION, THE PRESENCE OF MINOR FLAWS IN THE MATERIAL AND THE BAND'S OVERHANGING BORDERS HELPED BOOST THE ABILITY TO HARVEST LIGHT IN VISIBLE TO NEAR-UV WAVELENGTHS ESSENTIAL FOR OPTOELECTRONICS.

THE STUDY WENT ON TO SHOW THAT DECREASING THE PARTICLE SIZE OF METAL NANOPARTICLES CAUSES BLUE LIGHT TO BE ABSORBED MORE EASILY. PHOTOLUMINESCENCE ANALYSIS SHOWED THAT ZNO DOPED WITH VARIOUS IMPURITIES HAD BETTER EXCITONIC RECOMBINATION THAN THE UNDOPED MATERIAL WHICH COULD POSITIVELY AFFECT UV AND OPTICAL SENSORS. THESE TESTS ALSO DEMONSTRATED THAT ZNO NANOPARTICLES WITH BORON DOPING RESPONDED MORE SWIFTLY AND RECOVERED MUCH FASTER.

NO CHANGE IN CRYSTAL STRUCTURE WAS FOUND ACCORDING TO XRD, SEM AND FTIR ANALYSIS AND THE DOPANT IS STABLE AS WELL. ALL OF THESE EXPERIMENTS CONFIRM THAT TUNING DOPING AND SHAPE GIVES ZNO NANOPARTICLES IMPROVED ABILITIES IN SENSOR APPLICATIONS. THE STUDY HAS OPENED THE DOOR TO A BETTER KNOWLEDGE OF ZNO NANO STRUCTURES AND THEIR FEATURES, ASSISTING SCIENTISTS IN BOOSTING THE DEVELOPMENT OF SENSORS AND DIFFERENT TYPES OF MULTIFUNCTIONAL DEVICES.

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