



# INTERNATIONAL JOURNAL OF CREATIVE RESEARCH THOUGHTS (IJCRT)

An International Open Access, Peer-reviewed, Refereed Journal

## *Colocasia Esculenta*: From Traditional Medicine To Modern Applications

Bhakti D. Rahinj\* , Madhuri R. Kale

Dr. Naikwadi College of Pharmacy, Jamgaon, Sinner, Nashik, Maharashtra

**Abstract:** *Colocasia esculenta* (taro) is a staple food crop widely cultivated in tropical and subtropical regions and has been recognized for centuries for its nutritional and ethnomedicinal significance. Traditionally, various parts of the plant—including corms, leaves, and petioles—have been employed in the management of gastrointestinal disorders, inflammation, respiratory ailments, and skin diseases. Modern phytochemical investigations have revealed that *C. esculenta* is a rich source of bioactive compounds such as phenolics, flavonoids, alkaloids, and polysaccharides, which contribute to its broad pharmacological potential. Experimental studies have demonstrated antioxidant, antidiabetic, anticancer, antimicrobial, hepatoprotective, and immunomodulatory activities, making it a promising candidate for novel therapeutic development. Beyond its medicinal relevance, *C. esculenta* is emerging as an industrially valuable crop, with applications in starch production, biodegradable films, functional foods, and nutraceutical formulations. Its high starch content and physicochemical properties offer opportunities for innovation in the food, pharmaceutical, and packaging industries. This review provides a comprehensive synthesis of traditional knowledge, phytochemistry, pharmacological findings, and industrial applications of *C. esculenta*, critically analyzing current evidence and identifying research gaps. Future perspectives are also discussed, with an emphasis on molecular studies, clinical validation, and biotechnological approaches to maximize its therapeutic and commercial potential.

**Index Terms** - *Colocasia esculenta*, Taro, Ethnopharmacology, Phytochemistry, Antioxidant Activity, Antidiabetic Potential, Nutraceuticals, Industrial Applications, Therapeutic Development, Future Research.

### Introduction

Taxonomy and botanical description of *Colocasia esculenta*

*Colocasia esculenta* (L.) Schott is a perennial herb in the family Araceae.<sup>1</sup> The plant develops a substantial underground storage structure known as a corm, which is often globose or cylindrical, and produces large petiolate leaves.<sup>1</sup> Leaf blades are typically peltate, cordate or sagittate at the base, glabrous, dark green above and lighter beneath, with acute to obtuse apices.<sup>1</sup> Petioles may reach considerable length, supporting leaves often measuring up to ~40 × 25 cm.<sup>1</sup> Inflorescences consist of a spadix enclosed by a spathe; flowers are unisexual with male, female, and sterile zones arranged respectively.<sup>1</sup>



**FIG 1 - *C. ESCULENTA***

### **GLOBAL DISTRIBUTION AND CULTIVATION**

ORIGINALLY NATIVE TO SOUTHERN INDIA AND SOUTHEAST ASIA (THE INDOMALAYAN REGION), *C. ESCULENTA* IS NOW WIDELY NATURALIZED AND CULTIVATED ACROSS TROPICAL AND SUBTROPICAL REGIONS WORLDWIDE.<sup>1</sup> IT THRIVES IN DIVERSE CULTIVATION SYSTEMS—WETLAND PADDIES, FLOODED VALLEYS, RAINFED UPLANDS, AND SHADED OR SEMI-SHADED AREAS DEPENDING ON MOISTURE AVAILABILITY.<sup>2</sup> IN RECENT YEARS, GLOBAL PRODUCTION HAS RISEN SUBSTANTIALLY; FOR EXAMPLE, IN 2023, PRODUCTION REACHED APPROXIMATELY 18.07 MILLION METRIC TONS, WITH NIGERIA BEING THE LARGEST PRODUCER, FOLLOWED BY CHINA, CAMEROON, ETHIOPIA AND GHANA.<sup>2</sup>

### **HISTORICAL AND CULTURAL SIGNIFICANCE AS A STAPLE FOOD AND MEDICINAL PLANT**

ACROSS MANY CULTURES, *C. ESCULENTA* HAS SERVED BOTH AS A STAPLE FOOD AND MEDICINAL RESOURCE FOR CENTURIES.<sup>1</sup> ITS CORMS, LEAVES, AND PETIOLES ARE USED IN TRADITIONAL CUISINES, WHILE COOKED OR PROCESSED FORMS (BOILED, ROASTED, OR FERMENTED) ARE COMMON.<sup>1</sup> MEDICINALLY, VARIOUS PARTS ARE UTILIZED IN FOLK SYSTEMS FOR TREATING AILMENTS SUCH AS DIGESTIVE DISORDERS, SKIN AILMENTS, RESPIRATORY ISSUES, RHEUMATISM, AND AS REMEDIES FOR WOUNDS AND INFLAMMATION.<sup>3</sup>

### **IMPORTANCE OF COMPILING TRADITIONAL AND MODERN FINDINGS**

DESPITE WIDESPREAD TRADITIONAL USAGE, MANY CLAIMS REMAIN ANECDOTAL OR REGIONALLY CONFINED. BRIDGING ETHNOMEDICINE WITH MODERN SCIENTIFIC RESEARCH ENABLES VALIDATION OF PHARMACOLOGICAL ACTIVITIES, IDENTIFICATION OF BIOACTIVE COMPOUNDS, ASSESSMENT OF SAFETY, AND EXPLORATION OF INDUSTRIAL USES (E.G., NUTRACEUTICALS, FOOD PROCESSING, MATERIAL SCIENCE). BY SYNTHESIZING BOTH TRADITIONAL KNOWLEDGE AND CONTEMPORARY STUDIES, THIS REVIEW AIMS TO PROVIDE A COMPREHENSIVE FOUNDATION FOR FUTURE RESEARCH AND POTENTIAL COMMERCIAL APPLICATIONS OF *C. ESCULENTA*.

### **ETHNOBOTANY AND TRADITIONAL USES**

#### **ROLE IN AYURVEDA, TRADITIONAL CHINESE MEDICINE, AND OTHER INDIGENOUS SYSTEMS**

*COLOCASIA ESCULENTA* IS RECOGNIZED IN AYURVEDA AS *ALOOKAM*, *ALOOPAM*, USED TO PACIFY VITIATED VATA AND PITTA DOSHAS, AND TO TREAT AILMENTS SUCH AS GENERAL WEAKNESS, STOMATITIS, ALOPECIA, HEMORRHOIDS, AND CONSTIPATION.<sup>1</sup> ITS RASA (TASTE) IS DESCRIBED AS MADHURA (SWEET) AND KASHAYA (ASTRINGENT), GUNA (QUALITIES) AS GURU (HEAVY) AND SNIGDHA (UNCTUOUS), AND VIRYA (POTENCY) AS SEETA (COLD).<sup>4,5</sup> IN AYURVEDIC TRADITION BOTH LEAVES AND CORMS ARE EMPLOYED AS FOOD AS WELL AS MEDICINE.<sup>4</sup>

IN TRADITIONAL CHINESE MEDICINE (TCM), *C. ESCULENTA* IS USED TO STRENGTHEN THE SPLEEN AND STOMACH, IMPROVE DIGESTION, RELIEVE GASTROINTESTINAL DISCOMFORT, AND TO CLEAR HEAT AND RESOLVE SWELLING.<sup>6</sup> FRESH ROOT/PLASTER PREPARATIONS ARE USED EXTERNALLY FOR CONDITIONS SUCH AS BOILS, ABSCESSSES, AND PEST-RELATED BITES. FOLK AND INDIGENOUS SYSTEMS ALSO EMPLOY IT WIDELY: IN MANY RURAL INDIAN COMMUNITIES LEAF JUICE OR PASTES ARE APPLIED TO CUTS AND WOUNDS; THE CORM DECOCTION IS USED FOR DIARRHEA; LEAVES OR PETIOLE EXTRACTS ARE USED FOR COUGHS AND RESPIRATORY SYMPTOMS.<sup>7,8</sup>

### **FOLK USES: WOUND HEALING, DIARRHEA, COUGH, SKIN DISORDERS, INFLAMMATION**

SEVERAL TRADITIONAL PRACTICES ATTRIBUTE *C. ESCULENTA* WITH WOUND-HEALING PROPERTIES. FOR EXAMPLE, THE MUCILAGINOUS PULP OF THE PETIOLE IS APPLIED ON CUTS AND WOUNDS TO STOP BLEEDING IN KERALA, INDIA.<sup>4</sup> THE LEAF JUICE IS USED IN SNAKE BITES, SCORPION STINGS, AND ALSO FOR FOOD POISONING.<sup>4</sup> DECOCTIONS OF THE PEEL OR ROOTS ARE GIVEN FOR DIARRHEA.<sup>7,8</sup> LEAF JUICE AND LEAF STALK JUICE WITH SALT ARE APPLIED TO INFLAMED GLANDS (BUBOES), INTERNAL HEMORRHAGES, AND TO REDUCE INFLAMMATION AND SOOTHE SKIN DISORDERS.<sup>7</sup>

OTHER FOLK USES INCLUDE TREATING COUGH, EXPECTORANT APPLICATIONS, AND SOOTHING RESPIRATORY AILMENTS (E.G. SPUTUM IN ASTHMA).<sup>7,8</sup> THE PLANT IS ALSO USED FOR PILES, BODY ACHES, ALOPECIA, AND GENERAL DEBILITY IN VARIOUS TRIBAL OR FOLK MEDICINAL SETTINGS.<sup>7</sup>

### **NUTRITIONAL SIGNIFICANCE AS A STARCHY FOOD SOURCE**

*COLOCASIA ESCULENTA* IS VALUED FOR ITS CORMS AND LEAVES AS A SOURCE OF ENERGY AND MICRONUTRIENTS. THE CORMS ARE RICH IN CARBOHYDRATES, DIETARY STARCH (INCLUDING RESISTANT STARCH), AND PROVIDE SIGNIFICANT AMOUNTS OF PROTEIN, DIETARY FIBRE, AND MINERALS SUCH AS POTASSIUM, CALCIUM, IRON, MAGNESIUM, ZINC, AND COPPER.<sup>9,10</sup> LEAVES ARE ALSO RICH IN VITAMINS (ESPECIALLY B-COMPLEX, VITAMIN C), BETA-CAROTENE (IN YELLOW-FLESHED OR LEAF TISSUES), AND MINERALS.<sup>10,11</sup>

SPECIFIC NUTRITIONAL DATA: PER 100 G OF FRESH OR EDIBLE PORTION, TARO TUBERS MAY CONTAIN ~26-35 G CARBOHYDRATES, ~1-2 G PROTEIN, LOW FAT, ~3-5 G DIETARY FIBRE; MINERALS LIKE POTASSIUM IN THE ORDER OF HUNDREDS OF MG, CALCIUM, IRON ETC., AND VITAMINS SUCH AS THIAMINE, RIBOFLAVIN, NIACIN, VITAMIN C.<sup>9,10</sup> COOKED LEAVES SHOW HIGH VITAMIN C AND B COMPLEX VALUES COMPARED TO CORMS.<sup>6</sup>

### **PHYTOCHEMISTRY**

#### **PRIMARY PHYTOCONSTITUENTS: STARCH, DIETARY FIBERS, PROTEINS**

THE CORMS OF *COLOCASIA ESCULENTA* ARE PARTICULARLY RICH IN STARCH, FORMING ABOUT 70-80% OF DRY MATTER.<sup>14</sup> STARCH GRANULES ARE SMALL, AND THE CORM FLOUR ALSO CONTAINS CRUDE FIBRE, MODEST AMOUNTS OF PROTEIN, ASH, AND NEGLIGIBLE FAT.<sup>16</sup> LEAVES CONTRIBUTE DIETARY FIBER, VITAMINS, AND MINERALS, AND THE OVERALL PROTEIN CONTENT IN LEAVES AND CORMS VARIES WITH VARIETY AND ENVIRONMENTAL CONDITIONS.<sup>20</sup>

#### **SECONDARY METABOLITES: PHENOLS, FLAVONOIDS, SAPONINS, TANNINS, ALKALOIDS**

MULTIPLE STUDIES HAVE IDENTIFIED A WIDE SPECTRUM OF SECONDARY METABOLITES IN DIFFERENT PARTS OF *C. ESCULENTA*:

- PHENOLIC ACIDS AND THEIR DERIVATIVES (E.G. CHLOROGENIC ACID, CAFFEIC ACID, GALLIC ACID, BENZOIC ACID, ELLAGIC ACID) IN LEAVES AND EXTRACTS.<sup>12,13</sup>
- FLAVONOIDS INCLUDING VITEXIN, ISOVITEXIN, ORIENTIN, ISOORIENTIN, VINCENIN-2, LUTEOLIN AND THEIR GLYCOSYLATED FORMS.<sup>13,17,21</sup>
- TANNINS AND SAPONINS HAVE BEEN REPORTED IN LEAF, ROOT AND STOLON EXTRACTS.<sup>15,19</sup>
- ALKALOIDS AND GLYCOSIDES ARE ALSO PRESENT IN CERTAIN EXTRACTS.<sup>19,15</sup>

## BIOACTIVE COMPOUNDS RESPONSIBLE FOR MEDICINAL PROPERTIES

### SOME BIOACTIVE COMPOUNDS LINKED TO MEDICINAL EFFECTS:

- CATECHIN, (-) EPICATECHIN, RUTIN, AND OTHER FLAVONOIDS – POTENT ANTIOXIDANT AND ANTIMICROBIAL ACTIONS.<sup>12</sup>
- TRANS-FERULIC ACID, P-COUMARIC ACID, VANILLIC ACID ETC. – ANTIOXIDANT, POSSIBLY ANTI-INFLAMMATORY.<sup>15</sup>
- ANTHOCYANINS (CYANIDIN-3-GLUCOSIDE, CYANIDIN-3-RHAMNOSIDE, PELARGONIDIN-3-GLUCOSIDE) – CONTRIBUTE HEPATOPROTECTIVE AND ANTIOXIDANT ACTIVITIES.<sup>18,21</sup>

## PHARMACOLOGICAL ACTIVITIES OF *COLOCASIA ESCULENTA*

### ANTIOXIDANT ACTIVITY

**DESCRIPTION.** ANTIOXIDANT ACTIVITY REFERS TO THE CAPACITY OF PLANT EXTRACTS/COMPOUNDS TO SCAVENGE FREE RADICALS, REDUCE OXIDATIVE STRESS, AND PREVENT OXIDATIVE DAMAGE TO LIPIDS, PROTEINS, AND DNA.

**MECHANISM.** ACTIVITY IS MAINLY ATTRIBUTED TO PHENOLIC ACIDS, FLAVONOIDS AND ANTHOCYANINS THAT DONATE ELECTRONS OR HYDROGEN ATOMS TO NEUTRALIZE REACTIVE OXYGEN SPECIES (ROS), CHELATE TRANSITION METALS, AND UPREGULATE ENDOGENOUS ANTIOXIDANT ENZYMES (SOD, CAT, GPx).

**SUPPORTING STUDIES.** MULTIPLE STUDIES HAVE REPORTED STRONG DPPH, ABTS, FRAP AND REDUCING-POWER ACTIVITIES FOR METHANOLIC, ETHANOLIC AND AQUEOUS EXTRACTS OF TARO LEAVES AND CORMS; ULTRASOUND-ASSISTED AND OPTIMIZED SOLVENT EXTRACTION OFTEN YIELDS HIGHER FLAVONOID/PHENOLIC RECOVERY AND STRONGER ANTIOXIDANT READOUTS.<sup>22-24</sup>

### ANTI-INFLAMMATORY AND ANALGESIC EFFECTS

**DESCRIPTION.** ANTI-INFLAMMATORY ACTIVITY REFERS TO REDUCTION OF ACUTE OR CHRONIC INFLAMMATORY RESPONSES (EDEMA, CYTOKINE RELEASE), WHEREAS ANALGESIC EFFECTS RELIEVE NOCICEPTION AND PAIN BEHAVIORS.

**MECHANISM.** REPORTED MECHANISMS INCLUDE INHIBITION OF PRO-INFLAMMATORY MEDIATORS (TNF- $\alpha$ , IL-1 $\beta$ , IL-6), SUPPRESSION OF NITRIC OXIDE (NO) PRODUCTION VIA iNOS DOWNREGULATION, INHIBITION OF COX-2 OR PROSTAGLANDIN SYNTHESIS, AND STABILIZATION OF LYSOSOMAL MEMBRANES. MUCILAGE AND POLYSACCHARIDE FRACTIONS MAY ALSO MODULATE INFLAMMATORY SIGNALING.

**SUPPORTING STUDIES.** IN CARRAGEENAN-INDUCED RAT PAW EDEMA AND LPS-STIMULATED RAW264.7 MACROPHAGE MODELS, METHANOLIC ROOT AND CORM MUCILAGE EXTRACTS SIGNIFICANTLY REDUCED PAW SWELLING, NO PRODUCTION AND PRO-INFLAMMATORY CYTOKINE RELEASE; EFFECTS WERE DOSE DEPENDENT AND COMPARABLE TO STANDARD ANTI-INFLAMMATORY AGENTS IN SOME ASSAYS.<sup>25,26</sup>

### ANTIMICROBIAL & ANTIFUNGAL PROPERTIES

**DESCRIPTION.** ANTIMICROBIAL EFFECTS CONSIST OF BACTERIOSTATIC/BACTERICIDAL ACTIVITY AGAINST GRAM-POSITIVE/GRAM-NEGATIVE BACTERIA AND ANTIFUNGAL ACTIVITY AGAINST YEAST AND FILAMENTOUS FUNGI.

**MECHANISM.** LIKELY VIA MEMBRANE DISRUPTION, ENZYME INHIBITION, AND INTERFERENCE WITH MICROBIAL METABOLISM BY PHENOLICS, SAPONINS AND OTHER SECONDARY METABOLITES; SOME STUDIES ALSO DEMONSTRATE SYNERGISTIC EFFECTS WHEN TARO EXTRACTS ARE COMBINED WITH METALLIC NANOPARTICLES (GREEN SYNTHESIS AGNPs).

**SUPPORTING STUDIES.** ETHANOLIC AND AQUEOUS LEAF/CORM EXTRACTS SHOWED INHIBITION ZONES AGAINST *STAPHYLOCOCCUS AUREUS*, *ESCHERICHIA COLI*, *PSEUDOMONAS AERUGINOSA* AND *CANDIDA ALBICANS* IN AGAR DIFFUSION ASSAYS; MICs VARY BY EXTRACT AND SOLVENT. RECENT WORK ALSO USED TARO-MEDIATED SILVER NANOPARTICLES WITH ENHANCED ANTIMICROBIAL EFFICACY.<sup>27,28</sup>

### **ANTIDIABETIC AND ANTIHYPERLIPIDEMIC EFFECTS**

**DESCRIPTION.** ANTIDIABETIC ACTIONS LOWER BLOOD GLUCOSE (FASTING AND POSTPRANDIAL), IMPROVE INSULIN SENSITIVITY, AND/OR INHIBIT CARBOHYDRATE-DIGESTING ENZYMES; ANTIHYPERLIPIDEMIC EFFECTS REDUCE SERUM CHOLESTEROL, TRIGLYCERIDES AND IMPROVE LIPID PROFILE. **MECHANISM.** PROPOSED MECHANISMS INCLUDE INHIBITION OF  $\alpha$ -AMYLASE/ $\alpha$ -GLUCOSIDASE, DELAYED CARBOHYDRATE ABSORPTION (RESISTANT STARCH & DIETARY FIBER), ANTIOXIDANT-MEDIATED PRESERVATION OF PANCREATIC B-CELLS, AND MODULATION OF HEPATIC LIPID METABOLISM (REDUCED HMG-CoA REDUCTASE ACTIVITY OR INCREASED BILE ACID EXCRETION). POLYSACCHARIDES AND PHENOLICS ARE PRINCIPAL ACTIVE CLASSES.

**SUPPORTING STUDIES.** IN STREPTOZOTOCIN-INDUCED DIABETIC AND DIET-INDUCED HYPERLIPIDEMIC RAT MODELS, AQUEOUS/METHANOLIC CORM EXTRACTS DECREASED FASTING BLOOD GLUCOSE, IMPROVED ORAL GLUCOSE TOLERANCE, AND LOWERED SERUM TOTAL CHOLESTEROL, LDL AND TRIGLYCERIDES WHILE INCREASING HDL. SEVERAL IN VIVO ANIMAL STUDIES REPORT SIGNIFICANT ANTIHYPERGLYCEMIC AND ANTIHYPERLIPIDEMIC ACTIVITY AT ORAL DOSES OF PREPARED EXTRACTS.<sup>30-32</sup>

### **ANTICANCER POTENTIAL**

**DESCRIPTION.** ANTICANCER (CYTOTOXIC/ANTI-PROLIFERATIVE/ANTI-METASTATIC) EFFECTS INCLUDE INHIBITION OF CANCER CELL VIABILITY, INDUCTION OF APOPTOSIS, CELL-CYCLE ARREST, AND SUPPRESSION OF METASTATIC BEHAVIOR.

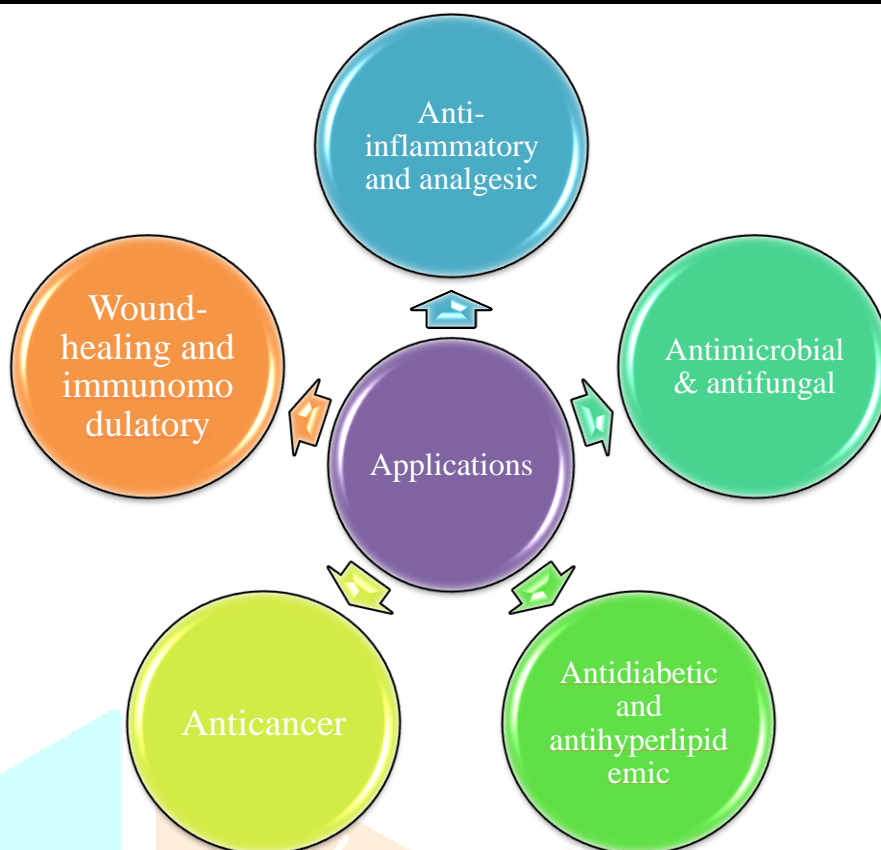
**MECHANISM.** MECHANISMS REPORTED INCLUDE ROS-MEDIATED APOPTOSIS INDUCTION, MITOCHONDRIAL PATHWAY ACTIVATION, CASPASE ACTIVATION, INHIBITION OF MATRIX METALLOPROTEINASES AND BLOCKADE OF ADHESION/MIGRATION PATHWAYS. POLYSACCHARIDE FRACTIONS HAVE SHOWN ANTI-METASTATIC ACTIVITY IN VITRO AND IN VIVO.

**SUPPORTING STUDIES.** SEVERAL IN VITRO CYTOTOXICITY STUDIES REPORT DOSE-DEPENDENT INHIBITION OF PROLIFERATION IN A549 (LUNG), HCT-116 (COLON), PC-3 (PROSTATE), PA-1 (OVARIAN) AND K562 (LEUKEMIA) CELL LINES BY LEAF AND CORM EXTRACTS; A WATER-SOLUBLE POLYSACCHARIDE FRACTION REDUCED METASTATIC POTENTIAL IN EXPERIMENTAL MODELS AND SHOWED IMMUNOMODULATORY EFFECTS.<sup>33,34</sup>

### **WOUND-HEALING AND IMMUNOMODULATORY EFFECTS**

**DESCRIPTION.** WOUND-HEALING EFFECTS ACCELERATE CLOSURE OF WOUNDS, IMPROVE COLLAGEN DEPOSITION AND RE-EPITHELIALIZATION. IMMUNOMODULATORY EFFECTS MODIFY INNATE OR ADAPTIVE IMMUNE RESPONSES (E.G., MACROPHAGE ACTIVATION, LYMPHOCYTE PROLIFERATION). **MECHANISM.** TARO MUCILAGE AND LEAF EXTRACTS PROVIDE A MOIST, PROTECTIVE MATRIX RICH IN POLYSACCHARIDES AND PROTEINS THAT PROMOTE FIBROBLAST PROLIFERATION, COLLAGEN SYNTHESIS AND ANGIOGENESIS; POLYSACCHARIDES MAY ALSO MODULATE MACROPHAGE ACTIVATION (NO, CYTOKINE RELEASE) AND STIMULATE IMMUNE CELL PROLIFERATION.

**SUPPORTING STUDIES.** IN VIVO EXCISION AND INCISION WOUND MODELS IN RODENTS DEMONSTRATED FASTER WOUND CONTRACTION, HIGHER HYDROXYPROLINE CONTENT (COLLAGEN MARKER), AND IMPROVED HISTOLOGICAL HEALING AFTER TOPICAL OR ORAL ADMINISTRATION OF TARO LEAF OR MUCILAGE PREPARATIONS. POLYSACCHARIDES ISOLATED FROM CORMS SHOWED IMMUNE-STIMULATORY ACTIVITY IN CELL-BASED ASSAYS.<sup>34</sup>



**FIG 2 - COLOCASIA ESCULENTA APPLICATION**

## INDUSTRIAL AND BIOTECHNOLOGICAL APPLICATIONS

### USE OF TARO STARCH IN BIODEGRADABLE FILMS AND PACKAGING MATERIALS

*COLOCASIA ESCULENTA* STARCH IS INCREASINGLY BEING USED TO DEVELOP BIODEGRADABLE FILMS (BIOPLASTICS) AS ALTERNATIVES TO PETROLEUM-BASED PLASTICS DUE TO THEIR RENEWABILITY, BIODEGRADABILITY, AND RELATIVELY LOW COST. SEVERAL WORKS HAVE FORMULATED FILMS FROM PURE TARO STARCH, COMPOSITE FILMS (REINFORCED WITH FILLERS OR BLENDED WITH POLYMERS), OR USING WASTE PARTS SUCH AS PEEL STARCH.<sup>35</sup>

- A STUDY PRODUCED BIODEGRADABLE FILMS FROM WILD TARO STARCH PLASTICIZED WITH GLYCEROL. AT STARCH LEVELS OF 5-10 G AND GLYCEROL 2-4%, FILMS SHOWED TENSILE STRENGTH UP TO ~9.51 MPa AND ELONGATION ~21.60%. ONE WEEK BIODEGRADABILITY TEST SHOWED ~64.5% DEGRADATION.
- TARO PEEL STARCH (TPS) WAS USED TO CAST FILMS WITH GLYCEROL. FILMS WITH 2.5-3.5% STARCH AND 25-35% GLYCEROL EXHIBITED THICKNESS 0.058-0.088 mm, WATER VAPOUR PERMEABILITY 0.06-0.09 g·m/m<sup>2</sup>·kPa·h, ETC. ALL FILMS FULLY DEGRADED IN RIVER WATER AND COMPOSTING SOIL WITHIN 5 DAYS.
- TO IMPROVE MECHANICAL STRENGTH AND OTHER PROPERTIES, REINFORCEMENT HAS BEEN USED. FOR EXAMPLE, BENTONITE WAS INCORPORATED INTO TARO STARCH FILMS; INCREASING BENTONITE CONCENTRATION IMPROVED TENSILE STRENGTH AND ACID/SALT RESISTANCE.
- ANOTHER COMPOSITE: TARO STARCH BLENDED WITH ALOE VERA GEL (PLUS GLYCEROL PLASTICIZER) WAS OPTIMIZED TO BALANCE TENSILE STRENGTH AND WATER-VAPOUR TRANSMISSION. THESE BIOCOMPOSITE FILMS SHOW PROMISE FOR FOOD PACKAGING APPLICATIONS.
- ALSO, TARO STARCH PLUS PVA POLYMER AND CaCO<sub>3</sub> FILLER WAS STUDIED TO PRODUCE BIODEGRADABLE PLASTIC, SHOWING EFFECTS OF STARCH CONTENT AND FILLER ON PROPERTIES IN AQUATIC ENVIRONMENT.

**MECHANISMS / REASONS FOR SUITABILITY:**

- TARO STARCH HAS HIGH STARCH CONTENT ( $\approx 70$ -80% IN MANY CULTIVARS) AND SMALL GRANULE SIZE, WHICH CAN CREATE DENSE FILM MATRICES.
- THE AMYLOSE/AMYLOPECTIN RATIO INFLUENCES FILM MECHANICAL AND BARRIER PROPERTIES; HIGHER AMYLOSE OFTEN GIVES GREATER TENSILE STRENGTH, LESS WATER PERMEABILITY.
- PLASTICIZERS (E.G. GLYCEROL), FILLERS (BENTONITE, ETC.), BLENDING WITH POLYMERS (E.G. PVA, ALOE VERA GEL) HELP ADJUST FLEXIBILITY, BARRIER, MECHANICAL STRENGTH, BIODEGRADATION RATE.<sup>36,37</sup>

**CHALLENGES / CONSIDERATIONS:**

- FILMS MAY HAVE LOWER DURABILITY, ESPECIALLY IN HIGHLY MOIST OR ALKALINE ENVIRONMENTS.
- COST AND PROCESS OPTIMIZATION (STARCH EXTRACTION, FILM CASTING, DRYING ETC) ARE IMPORTANT.
- BALANCE NEEDED BETWEEN MECHANICAL STRENGTH AND BIODEGRADABILITY / WATER BARRIER PROPERTIES.<sup>38</sup>

**POTENTIAL USE IN PHARMACEUTICAL FORMULATIONS (BINDER, DISINTEGRANT)**

**TARO STARCH HAS ALSO BEEN EXPLORED AS AN EXCIPIENT IN PHARMACEUTICAL FORMULATIONS, ESPECIALLY AS DISINTEGRANT AND BINDER.**

- A STUDY COMPARED NATIVE TARO STARCH AND CITRATE MODIFIED TARO STARCH AS TABLET DISINTEGRANTS (IN PARACETAMOL TABLETS BY WET GRANULATION). THE RESULTS SHOWED THAT BOTH NATIVE AND MODIFIED TARO STARCH PERFORMED WELL, WITH DISINTEGRATION EFFICIENCY BETTER THAN THAT OF STANDARD CORN STARCH IN SOME TESTS.
- TARO STARCH HAS SMALL GRANULE SIZE AND GOOD SWELLING PROPERTIES; MODIFIED VERSIONS (E.G. VIA CITRIC ACID) MIGHT FURTHER ENHANCE DISINTEGRATION AND DISSOLUTION CHARACTERISTICS.

**MECHANISMS:**

- AS A DISINTEGRANT, TARO STARCH ABSORBS WATER RAPIDLY, SWELLS, AND CAUSES TABLET MATRIX TO BREAK APART TO ENHANCE DISSOLUTION.
- AS A BINDER (THOUGH LESS DATA), STARCH CAN SERVE TO HOLD GRANULES/PARTICLES TOGETHER DURING TABLET FORMATION, PARTICULARLY WHEN MODIFIED OR COMBINED WITH BINDING AGENTS.

**ADVANTAGES:**

- READILY AVAILABLE, EDIBLE, NON-TOXIC.
- POSSIBLY MORE SUSTAINABLE / LOCAL THAN IMPORTED STARCHES IN MANY REGIONS.
- POTENTIAL FOR COST SAVINGS OR IMPROVED PROPERTIES DEPENDING ON MODIFICATION.<sup>39</sup>

**BIOETHANOL PRODUCTION FROM TARO STARCH**

**BIOETHANOL PRODUCTION FROM TARO STARCH/WASTE IS A PROMISING BIOENERGY USE, ESPECIALLY USING PARTS OF PLANTS OR CULTIVARS NOT MEETING FOOD STANDARDS.**

- A STUDY USED TARO WASTE (MAINLY PEELS WITH STARCH RESIDUES) AS CARBON SOURCE, WITH *KLUYVEROMYCES MARXIANUS* K21 (THERMO-TOLERANT YEAST). IN FLASK STUDIES SSF (SIMULTANEOUS SACCHARIFICATION AND FERMENTATION) AT OPTIMAL CONDITIONS (170 g/L TARO WASTE, ETC.) YIELDED  $\sim 48.98$  g/L ETHANOL, PRODUCTIVITY  $\sim 2.23$  g/L/H AFTER 22 HOURS; IN A 5-L BIOREACTOR,  $\sim 43.78$  g/L ACHIEVED,  $\sim 94.2\%$  THEORETICAL YIELD.
- ANOTHER RECENT STUDY EVALUATED TARO (BOTH WITH PEEL AND WITHOUT) USING ULTRASOUND-ASSISTED ENZYMATIC HYDROLYSIS ( $\alpha$ -AMYLASE + GLUCOAMYLASE), THEN FERMENTATION BY *SACCHAROMYCES CEREVISIAE*. ULTRASOUND PRETREATMENT SIGNIFICANTLY IMPROVED REDUCING SUGAR RELEASE; ETHANOL YIELD INCREASED BY ABOUT 35% IN TREATED SAMPLES COMPARED TO UNTREATED.<sup>40,41</sup>

## **MECHANISMS / PROCESS STEPS:**

- STARCH IN THE SUBSTRATE IS GELATINIZED / HYDROLYZED TO SUGARS VIA ENZYMES (AND/OR PHYSICAL PRETREATMENTS LIKE ULTRASOUND) → FERMENTATION BY YEAST → ETHANOL.
- PRETREATMENT (ULTRASOUND ETC.) HELPS IN DISRUPTING STARCH GRANULES, INCREASING SURFACE AREA FOR ENZYMATIC ATTACK.
- USE OF WASTE/TARO PEELS IMPROVES RESOURCE EFFICIENCY AND REDUCES WASTE.<sup>42</sup>

## **SAFETY, TOXICOLOGY, AND ANTINUTRITIONAL FACTORS**

### **PRESENCE OF OXALATES AND THEIR EFFECTS**

**TARO PLANTS, PARTICULARLY THE LEAVES AND STEMS, CONTAIN SIGNIFICANT LEVELS OF OXALATES, PRIMARILY IN THE FORM OF CALCIUM OXALATE CRYSTALS. THESE COMPOUNDS CAN POSE HEALTH RISKS IF CONSUMED IN LARGE QUANTITIES WITHOUT PROPER PREPARATION.**

- **OXALATE CONTENT:** THE TOTAL OXALATE CONTENT IN TARO LEAVES CAN RANGE FROM 433.8 TO 856.1 MG PER 100 G OF WET MATTER, WITH SOLUBLE OXALATES CONTRIBUTING TO A PORTION OF THIS TOTAL
- **HEALTH IMPLICATIONS:** OXALATES CAN INTERFERE WITH CALCIUM ABSORPTION AND MAY CONTRIBUTE TO THE FORMATION OF KIDNEY STONES. ADDITIONALLY, THE PRESENCE OF CALCIUM OXALATE CRYSTALS CAN CAUSE IRRITATION IN THE ORAL MUCOSA AND DIGESTIVE TRACT IF THE PLANT MATERIAL IS CONSUMED RAW OR INADEQUATELY PROCESSED.<sup>43,44</sup>

### **METHODS FOR DETOXIFICATION**

**SEVERAL TRADITIONAL AND SCIENTIFIC METHODS HAVE BEEN IDENTIFIED TO REDUCE THE OXALATE CONTENT IN TARO, MAKING IT SAFER FOR CONSUMPTION.**

#### **SOAKING**

- **PROCEDURE:** SOAKING TARO LEAVES IN WATER FOR EXTENDED PERIODS CAN LEACH OUT SOLUBLE OXALATES.
- **EFFECTIVENESS:** SOAKING FOR 18 HOURS HAS BEEN SHOWN TO REDUCE SOLUBLE OXALATE CONTENT BY APPROXIMATELY 26%, WITH NO SIGNIFICANT CHANGE IN INSOLUBLE OXALATES.
- **ADDITIONAL FINDINGS:** COMBINING SOAKING WITH DRYING FURTHER DECREASES OXALATE LEVELS; FOR INSTANCE, SOAKING AND DRYING TARO LEAVES REDUCED OXALATE CONTENT FROM 43 MG/100 G TO 4.2 MG/100 G.<sup>45</sup>

#### **BOILING**

- **PROCEDURE:** BOILING TARO LEAVES OR STEMS CAN SIGNIFICANTLY REDUCE OXALATE CONTENT.
- **EFFECTIVENESS:** BOILING FOR 20 MINUTES HAS BEEN REPORTED TO DECREASE SOLUBLE OXALATE CONTENT BY UP TO 79.6% IN TARO STEMS.
- **CONSIDERATIONS:** WHILE BOILING IS EFFECTIVE, IT IS ESSENTIAL TO ENSURE THAT THE COOKING TIME IS ADEQUATE TO ACHIEVE SIGNIFICANT OXALATE REDUCTION.<sup>46,47</sup>

#### **CALCIUM SALT TREATMENT**

- **PROCEDURE:** SOAKING TARO CORM CHIPS IN CALCIUM SALT SOLUTIONS (E.G., CALCIUM CHLORIDE) CAN FACILITATE THE FORMATION OF INSOLUBLE CALCIUM OXALATE, WHICH IS LESS BIOAVAILABLE.
- **EFFECTIVENESS:** SOAKING IN A 5% CALCIUM CHLORIDE SOLUTION FOR 60 MINUTES REDUCED SOLUBLE OXALATE CONTENT FROM 294.3 MG/100 G TO 35.1 MG/100 G.
- **ADDITIONAL INSIGHTS:** THIS METHOD NOT ONLY REDUCES OXALATE LEVELS BUT ALSO ENHANCES THE NUTRITIONAL QUALITY BY INCREASING CALCIUM BIOAVAILABILITY.<sup>48,49</sup>

**FERMENTATION**

- **PROCEDURE:** FERMENTATION INVOLVES THE USE OF MICROORGANISMS TO DEGRADE OXALATES OVER TIME.
- **EFFECTIVENESS:** FERMENTATION HAS BEEN IDENTIFIED AS AN EFFECTIVE METHOD TO REDUCE SOLUBLE OXALATE CONTENT, PARTICULARLY WHEN COMBINED WITH OTHER PROCESSING TECHNIQUES.
- **CONSIDERATIONS:** THE SUCCESS OF FERMENTATION DEPENDS ON FACTORS SUCH AS MICROBIAL STRAIN, FERMENTATION DURATION, AND ENVIRONMENTAL CONDITIONS.<sup>50</sup>

**TABLE 1 — SUMMARY OF OXALATE REDUCTION METHODS FOR TARO**

METHOD	TARGET OXALATE TYPE	REDUCTION (%)	REFERENCE
SOAKING (18 HOURS)	SOLUBLE	26%	( <a href="#">PUBMED</a> )
BOILING (20 MINUTES)	SOLUBLE	79.6%	( <a href="#">SCIRP</a> )
CALCIUM SALT TREATMENT	SOLUBLE	88%	( <a href="#">AJFS.JOURNALS.EKB.ENG</a> )
FERMENTATION	SOLUBLE	VARIABLE	( <a href="#">SCIENCE DIRECT</a> )

**CHALLENGES, RESEARCH GAPS, AND FUTURE PROSPECTS****NEED FOR MORE CLINICAL TRIALS TO VALIDATE PHARMACOLOGICAL CLAIMS**

WHILE NUMEROUS IN VITRO AND ANIMAL STUDIES HAVE DEMONSTRATED THE PHARMACOLOGICAL POTENTIAL OF TARO, THERE IS A SIGNIFICANT LACK OF HUMAN CLINICAL TRIALS TO SUBSTANTIATE THESE FINDINGS. FOR INSTANCE, WHILE TARO STARCH HAS SHOWN PROMISE AS A DISINTEGRANT IN TABLET FORMULATIONS AND AS A CARRIER FOR BIOACTIVE COMPOUNDS, THESE APPLICATIONS REMAIN LARGELY UNVERIFIED IN CLINICAL SETTINGS. THERE IS A PRESSING NEED FOR WELL-DESIGNED HUMAN STUDIES TO CONFIRM THE EFFICACY AND SAFETY OF TARO-BASED PHARMACEUTICAL FORMULATIONS.

**GENETIC IMPROVEMENT FOR YIELD, STARCH QUALITY, AND PEST RESISTANCE**

DESPITE ITS WIDESPREAD USE, TARO CULTIVATION FACES CHALLENGES RELATED TO LOW YIELD, POOR STARCH QUALITY, AND SUSCEPTIBILITY TO PESTS. ADVANCEMENTS IN GENETIC IMPROVEMENT ARE ESSENTIAL TO ADDRESS THESE ISSUES. RESEARCH INTO DEVELOPING TARO VARIETIES WITH ENHANCED STARCH CONTENT, IMPROVED QUALITY, AND GREATER RESISTANCE TO PESTS COULD SIGNIFICANTLY BOOST ITS AGRICULTURAL AND INDUSTRIAL VALUE.

**EXPLORATION OF NOVEL DRUG DELIVERY SYSTEMS USING TARO STARCH**

TARO STARCH HAS GARNERED ATTENTION FOR ITS POTENTIAL IN PHARMACEUTICAL APPLICATIONS, PARTICULARLY IN DRUG DELIVERY SYSTEMS. STUDIES HAVE EXPLORED ITS USE AS A DISINTEGRANT IN TABLET FORMULATIONS, DEMONSTRATING ITS ABILITY TO FACILITATE TABLET DISINTEGRATION AND ENHANCE DRUG RELEASE PROFILES. ADDITIONALLY, TARO STARCH HAS BEEN INVESTIGATED FOR ITS ENCAPSULATING PROPERTIES, SERVING AS A CARRIER FOR PROBIOTICS AND BIOACTIVE COMPOUNDS, THEREBY IMPROVING THEIR STABILITY AND CONTROLLED RELEASE.

HOWEVER, THESE APPLICATIONS ARE PRIMARILY BASED ON IN VITRO STUDIES, AND THERE IS A NEED FOR FURTHER RESEARCH TO VALIDATE THESE FINDINGS IN CLINICAL SETTINGS. MOREOVER, EXPLORING THE MODIFICATION OF TARO STARCH TO IMPROVE ITS PROPERTIES, SUCH AS PREGELATINIZATION TO ENHANCE COMPRESSIBILITY AND FLOWABILITY, COULD FURTHER EXPAND ITS UTILITY IN PHARMACEUTICAL FORMULATIONS.

TABLE 2 — SUMMARY OF TARO STARCH IN PHARMACEUTICAL APPLICATIONS

APPLICATION AREA	STUDY FOCUS	KEY FINDINGS
<b>TABLET DISINTEGRANT</b>	<b>EVALUATING TARO STARCH AS A DISINTEGRANT IN TABLET FORMULATIONS</b>	<b>DEMONSTRATED EFFECTIVE TABLET DISINTEGRATION AND ENHANCED DRUG RELEASE PROFILES</b>
<b>ENCAPSULATION OF BIOACTIVE COMPOUNDS</b>	<b>INVESTIGATING TARO STARCH'S ENCAPSULATING PROPERTIES FOR PROBIOTICS AND BIOACTIVE COMPOUNDS</b>	<b>SHOWED POTENTIAL FOR IMPROVING STABILITY AND CONTROLLED RELEASE OF ENCAPSULATED COMPOUNDS</b>
<b>MODIFICATION FOR PHARMACEUTICAL USE</b>	<b>STUDYING PREGELATINIZED TARO STARCH AS A DILUENT IN TABLET FORMULATIONS</b>	<b>FOUND THAT INCREASING PREGELATINIZATION TEMPERATURE IMPROVED COMPRESSIBILITY AND COMPRESSIBILITY</b>

## CONCLUSION

*COLOCASIA ESCULENTA* (TARO) REPRESENTS A UNIQUE INTERSECTION OF TRADITIONAL MEDICINE AND MODERN INDUSTRIAL APPLICATIONS. HISTORICALLY VALUED AS A STAPLE FOOD AND ETHNOMEDICINAL PLANT, TARO HAS BEEN UTILIZED IN VARIOUS INDIGENOUS HEALTHCARE SYSTEMS FOR ITS NUTRITIONAL AND THERAPEUTIC PROPERTIES. MODERN RESEARCH HAS VALIDATED MANY OF THESE TRADITIONAL CLAIMS, DEMONSTRATING ANTIOXIDANT, ANTI-INFLAMMATORY, ANTIMICROBIAL, ANTIDIABETIC, ANTICANCER, AND WOUND-HEALING ACTIVITIES.

BEYOND PHARMACOLOGY, TARO STARCH AND OTHER PLANT COMPONENTS HAVE SHOWN SIGNIFICANT POTENTIAL IN PHARMACEUTICAL FORMULATIONS AS EXCIPIENTS, IN THE DEVELOPMENT OF NOVEL DRUG DELIVERY SYSTEMS, AND AS BIODEGRADABLE MATERIALS FOR INDUSTRIAL APPLICATIONS. FURTHERMORE, ITS APPLICATION IN FUNCTIONAL FOODS AND BIOETHANOL PRODUCTION HIGHLIGHTS THE VERSATILITY AND SUSTAINABILITY OF THIS CROP.

Future research focusing on clinical validation, genetic improvement, and innovative biotechnological applications can unlock the full potential of *C. esculenta*. Its integration into drug discovery, nutraceutical development, and sustainable biomaterials underscores its value as a multipurpose crop bridging ethnomedicine and modern scientific innovation.

## REFERENCES

1. *Colocasia esculenta* (L.) Schott — Botanical description, taxonomy, morphology, native and naturalised distribution. CABI Compendium.
2. “From starch to bioactives: emerging trends in taro (*Colocasia esculenta* L.) research on composition, functionality, health benefits, and sustainable food potential.” *Frontiers in Nutrition*. 2024; Volume, Issue (if known). ([Frontiers](#))
3. “An overview of traditionally used herb, *Colocasia esculenta*, as a phytomedicine.” *International Journal of Herbal Medicine*. Published as open access. ([Longdom](#))
4. AyurvedicMedicinalPlants.com. *Colocasia esculenta* (Linn.) Schott. [Internet]. Available from: Plant pacifies vitiated vata, pitta, constipation, stomatitis, alopecia, hemorrhoids and general weakness; corms and tender leaves used. Sanskrit synonyms. AYURVEDIC PROPERTIES etc.
5. *Colocasia esculenta* (Linn.) Schott. Indian Medicinal Plants. SpringerReference. Synonyms, folk & Ayurvedic uses including petiole juice as styptic, corm juice for alopecia etc.
6. MedicineTraditions.com. *Colocasia, Taro – Ethnomedicinal uses (Ayurveda, TCM, Unani)*. Fresh root plaster, decoctions; clears heat, resolves swelling, abscess treatment etc.
7. Longdom.org. “An overview of traditionally used herb, *Colocasia esculenta*, as a phytomedicine.” Folk uses: wound healing, snake-bite, diarrhea, inflammation etc.
8. International Journal of Nutrition, Pharmacology, Neurological Diseases. “*Colocasia esculenta* (CE) Linn... used for various ailments such as asthma, arthritis, diarrhea, internal hemorrhage, neurological disorders, and skin disorders.”

9. Mdpi: "Taro Roots: An Underexploited Root Crop." Vitamins and minerals composition: carbs, protein, fibre, B-complex, vitamin C, beta-carotene etc.
10. Nutritional & Anti-Nutritional Quality of Taro (Innovation Science Publishing). Mineral content, vitamin content etc.
11. NutriOnio. Raw taro leaves – nutritional value per 100 g (vitamins, minerals, fibre) etc.
12. Phenolic Profile: Antimicrobial Activity and Antioxidant Capacity of *Colocasia esculenta* (L.) Schott. *Egyptian Journal of Chemistry*. 2021;64(4):2165-72. ([ejchem.journals.ekb.eg](http://ejchem.journals.ekb.eg))
13. "Further knowledge on the phenolic profile of *Colocasia esculenta* (L.) Shott." *PubMed*. ([PubMed](https://pubmed.ncbi.nlm.nih.gov/))
14. Functional profile and encapsulating properties of *Colocasia esculenta* (Taro). *Food Science & Nutrition*. 2023. ([Wiley Online Library](https://onlinelibrary.wiley.com/))
15. Antioxidant and antimicrobial prospects of *Colocasia esculenta* stolon: a phytochemical perspective. *Discover Food*. ([SpringerLink](https://www.springerlink.com/))
16. International Journal of Nutrition, Pharmacology, Neurological Diseases. "*Colocasia esculenta* (A potent indigenous plant)". 2011. ([Lippincott Journals](http://www.lippincottjournals.com/))
17. Potential of *Colocasia* leaves in human nutrition: Review on nutritional and phytochemical properties. *PubMed*. ([PubMed](https://pubmed.ncbi.nlm.nih.gov/))
18. *Colocasia esculenta* (L.) Schott: Pharmacognostic and pharmacological review. *Journal of Pharmacognosy and Phytochemistry*. 2020;9(4):S11937. ([Phytojournal](http://www.phytojournal.com/))
19. Wudali SN, Barwad A, Banadka A, Shaikh A, Al-Khayri JM, Nagella P. Bioactive Compounds and Biological Activities of Taro (*Colocasia esculenta* (L.) Schott). Reference Series in Phytochemistry. 2024; Volume F2504:37-59. ([archives.christuniversity.in](http://archives.christuniversity.in/))
20. "Potential of *Colocasia* leaves in human nutrition: Review on nutritional and phytochemical properties." *PubMed*. (same as #6) ([PubMed](https://pubmed.ncbi.nlm.nih.gov/))
21. Phytochemistry finding in leaf extracts including anthocyanins etc. *Research Journal of Pharmacology and Pharmacodynamics*, 2023;15(3).
22. Christou A, Leclerc E, et al. Ultrasound-Assisted Extraction of Taro Leaf Antioxidants: improved flavonoid recovery and antioxidant activity. *Antioxidants (Basel)*. 2023;12(8). ([PMC](https://pubmed.ncbi.nlm.nih.gov/))
23. Baro MR, Biswas M, et al. Exploring the anti-inflammatory potential of *Colocasia esculenta* methanolic root extract on carrageenan-induced rat paw edema and LPS-stimulated RAW264.7 cells. *International Journal of Biological Macromolecules* (or Journal indicated). 2023. ([PubMed](https://pubmed.ncbi.nlm.nih.gov/))
24. Anonymous / Singh et al. A study of antibacterial and antifungal activity of the leaves of *Colocasia esculenta* Linn. *International Journal of Pharmaceutical Sciences and Research (IJPSR)*. 2017 Mar;8(3). ([IJPSR](http://www.ijpsr.in/))
25. Lad SS, et al. Evaluation of antihyperlipidemic potential of aqueous corm extract of *Colocasia esculenta* in experimental rat models. \* Journal\* (SciDirect). 2023. ([ScienceDirect](https://www.sciencedirect.com/))
26. Jyothi R, et al. Cytotoxic potentiality of *Colocasia esculenta* leaves extract against human cancer cell lines. *International Journal of Green Pharmacy (IJGP)*. 2020;14(4). ([Green Pharmacy](http://www.greenpharmacy.in/))
27. Eltanbouly ND, et al. *Colocasia esculenta* L. Schott corm mucilage: anti-inflammatory and wound healing properties. *Pharmacognosy Communications / Phcog Res*. 2021. ([phcog.com](http://www.phcog.com))
28. Pereira PR, Gonçalves RF, et al. Anticancer and immunomodulatory benefits of taro (*Colocasia esculenta*): a review. *Frontiers in Pharmacology / Nutrients* (or journal). 2020. ([PMC](https://pubmed.ncbi.nlm.nih.gov/))
29. Wang X, et al. Green synthesis of *Colocasia esculenta*-based silver nanoparticles and evaluation of antimicrobial/anti-inflammatory effects. 2025. ([PMC](https://pubmed.ncbi.nlm.nih.gov/))
30. [Antihyperglycemic and antihyperlipidemic study] — (BiomedSciDirect / IJBMRF) *Evaluation of antihyperglycemic and antihyperlipidemic potential of Colocasia esculenta corms in vivo* 2019. ([Biomed Sci Direct](https://www.biomedsci.com/))
31. Esposito T, et al. Activity of *Colocasia esculenta* corms against selected targets: fractionation and bioactivity profiling. *Scientific Reports / MDPI* 2023. ([PMC](https://pubmed.ncbi.nlm.nih.gov/))
32. Zubair MW, et al. Functional profile and encapsulating properties of taro (*Colocasia esculenta*) starch and fibers: implications for health and metabolic effects. *Food & Function / MDPI* 2023. ([PMC](https://pubmed.ncbi.nlm.nih.gov/))
33. Park HY, et al. Anti-metastatic effect of a polysaccharide isolated from taro corm — mechanistic and in vivo studies. *International Journal of Molecular Medicine*. 2013;31(6):1400–1406. ([Spandidos Publications](http://www.spandidospublications.com/))
34. (Additional wound-healing and local studies) — Various in vivo reports and reviews (ResearchGate 2019–2023). ([ResearchGate](https://www.researchgate.net/))
35. Morales T, Tuates A. Biodegradable film from wild taro *Colocasia esculenta* (L.) Schott starch. *Agricultural Engineering International: CIGR Journal*. 2020;22(1):4085. ([cigrjournal.org](http://www.cigrjournal.org))

36. Belhocine K, et al. Characterization of Biodegradable Films Made from Taro Peel (*Colocasia esculenta*) Starch. *Polymers (Basel)*. 2023;15(2):338. ([MDPI](#))
37. Sitanggang MJ, et al. Development of biodegradable bioplastic films from taro starch reinforced with bentonite. *Carbohydrate Polymer Technologies and Applications*. 2021. ([ScienceDirect](#))
38. Azevedo NF, et al. Optimization of biocomposite taro starch and aloe vera gel based film using response surface methodology. *Journal of Polymers* (or relevant journal). 2023. ([PubMed](#))
39. Jurusan & Syahbanu I, Adhitiyawarman A. The effect of taro tuber starch addition to biodegradable plastic with PVA polymer and CaCO<sub>3</sub> filler. *Jurnal Ilmu Dasar*. XX;24(1):33409. ([jid.jurnal.unej.ac.id](#))
40. Lee YL, et al. Taro starch (*Colocasia esculenta*) and citric acid modified taro starch as tablet disintegrating agents. *Journal of Pharmaceutical Sciences / Indian Journal* (as per publication). 2018. ([PubMed](#))
41. Wu WH, Hung WC, Lo KY, et al. Bioethanol production from taro waste using thermo-tolerant yeast *Kluyveromyces marxianus* K21. *Bioresource Technology*. 2016;201:27–33. ([scholars.lib.ntu.edu.tw](#))
42. Almeida-Neto RA, et al. Evaluation of *Colocasia esculenta* as a raw material for bioethanol production through ultrasound-assisted enzymatic hydrolysis. *Fermentation*. 2025;11(2):102. ([MDPI](#))
43. Thanh HD, et al. Oxalate content of taro leaves grown in central Vietnam. *Journal of Food Science and Technology*. 2017;54(4):1023–1029.
44. Savage GP, Dubois MD. The effect of soaking and cooking on the oxalate content of taro leaves. *Food Chemistry*. 2006;98(2):235–240.
45. Vulla KE, Mmanda FP. Optimizing oxalate reduction in taro leaves (*Colocasia esculenta*) through soaking and drying techniques for inclusion in a farmed fish feed. *European Journal of Nutrition & Food Safety*. 2025;17(5):302–310.
46. Saleh S, et al. Reducing the soluble oxalate and phytic acid in taro corm chips by soaking in calcium salt solutions. *African Journal of Food Science*. 2019;13(4):89–95.
47. Huynh NK, et al. Effects of processing on oxalate contents in plant foods. *Food Research International*. 2022;157:111255.
48. Canh NT, et al. Effect of processing on oxalate and calcium contents in taro stems. *Food Chemistry*. 2022;373:131572.
49. Zayed A, et al. Management strategies for the anti-nutrient oxalic acid in plant foods. *Food Research International*. 2025;137:109676.
50. Bradbury JH, et al. The acidity of raphides from the edible aroids. *Journal of the Science of Food and Agriculture*. 2000;80(13):1849–1853.