

# Modeling Advanced Technology of Biodegradable Mulch Films

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**Abstract** - Growing environmental concerns have accelerated the development of biodegradable plastics, which are applied widely in organic waste collection bags, disposable food containers, and agriculture mulch films. Mulch films are widely used in agriculture to suppress weeds, retain soil moisture, regulate temperature, protect soil structure, and control pests, thereby enhancing crop yields. However, conventional polyethylene (PE) mulch films accumulate as plastic fragments, causing significant soil pollution. In contrast, biodegradable mulch films are completely assimilated by soil microorganisms as a carbon source, safely reintegrating into the ecosystem. The major critique for existing biodegradable films is that they perform well for short-growth-cycle crops (<90 days; e.g., potatoes, strawberries) but disintegrate prematurely in long-growth-cycle crops like cotton (requiring 120–150 days of coverage), leading to substantial yield losses. This study aims to enhance biodegradable film properties with advanced technologies, such as grafting biopolymers with inorganic fillers to improve tensile strength, UV resistance, and water barrier properties while maintaining biodegradability to align with cotton growth timelines to ensure functionality during cultivation and complete disintegration before the next planting season.

The results obtained from laboratory aging tests (100 hours) and biodegradation tests (180 days) confirmed the biodegradable mulch film's durability and decomposition profile. Subsequently, the field tests in cotton fields demonstrated that the optimized film with advanced technology sustained the support for cotton growth during 120–150 days, degraded on schedule, and delivered yields comparable to PE films.

**Keywords:** Biodegradable mulch film, disintegration, grafting, tensile strength, aging test, biodegradation test, field test

## 1. Introduction

Polyethylene (PE) mulch film is widely accepted agriculturally due to its myriad of agronomic benefits. Current PE mulch research demonstrates that mulch film can increase soil temperatures to accelerate plant growth and protection from various types of erosion damage, as well as reducing weed levels by up to 98%, ultimately supporting crop growth [1]. All of these characteristics are especially beneficial in arid and temperate climates, where it is necessary to optimize soil and water conditions. PE mulch film enhances crop growth through maximization of the root zone microclimate, which enhances productivity and farmers' yields [1]. PE mulch film is also appealing because of its low manufacturing costs, resilience in fields, and ease of application, and so it is widely used on both small-scale and industrial farms. However, the application of PE mulch film bears environmental consequences. PE mulch is made from petrochemical-derived plastics that cannot biodegrade, and postharvest film recovery is labor intensive and imperfect. Large quantities of mulch films are left in fields, where they fragment into microplastics over time. Trapped in the soil, they degrade soil structure and reduce water infiltration, among other negative consequences. Additionally, recycling the mulch is ineffective as it is already contaminated, and inappropriate disposal methods greatly damage the environment through pollution and toxic emissions.

To reduce the long-term environmental footprint of plastic mulching, biodegradable mulch films (BDMs) are a strong candidate for a more sustainable alternative. These BDMs mainly use polylactic acid or starch-based polymers that degrade through microbial activity under field conditions. BDMs are able to replicate the major functions and benefits of PE films such as preserving moisture, controlling temperature, and suppressing weeds at an even higher effectiveness than PE film—all without the need for removal and cleaning up after harvest [2, 3]. One example from China utilizing on-field studies found that under the same conditions, BDMs can match PE mulch in terms of yield and water use efficiency in a multitude of plants [2]. On top of environmental benefits, biodegradable films may also be reinstalled in fields after harvests without extra labor costs to remove old ones, thus assisting farmers in their work. Research has shown that especially in arid and semi-arid regions, the use of BDMs can increase crop yields by 28%, demonstrating levels of applications for BDMs [3]. Viability is further supported by tests conducted in Spain, in which target levels of degradation and yields over multiple years were achieved and followed European regulations, namely Regulation CEE 2092/91 [1].

While BDMs are effective, they face a critical problem: most commercial biodegradable mulch films disintegrate prematurely in long-growth cycle crops like cotton (requiring 120-150 days of coverage). However, this issue can be minimized through further materials development. Time-to-event decomposition models and field experimentation demonstrate the possibility of customizing the BDM degradation timeline to align with crop cycles. Thus, with careful tuning, these biodegradable mulch films can remain unimpaired during the plant growth stage, then start the process of degradation during or post-harvest—an extremely functional and sustainable advantage that can be customized to different agricultural environments [4]. Understanding this customization process is rooted in understanding the scientific mechanisms behind biodegradation and the ability to control it via material formulation. Researchers engineered a BDM via reactive grafting of poly(butylene adipate terephthalate) (PBAT/Ecoflex®) with talc filler to enhance tensile strength, to improve UV resistance, and to maintain hydrophobicity within cotton's 120–150-day growth window while ensuring full biodegradability after harvest. We will demonstrate the results of field validation after laboratory biodegradable confirmation.

## 2. Materials Preparation

In previous literature, grafting reactive extrusion has already been applied and used with polymers to generate copolymers [5]. Moreover, the usage of silane to graft polymers via reactive extrusion processes has also been previously accomplished [6]. To enhance the compatibility and mechanical properties of PBAT blended with talc filler, our study employs a film manufactured using a reactive extrusion process. Silane-functionalized PBAT is grafted to talc, which is hydroxyl-rich, using a free-radical initiator (Luperox 101). This forms covalent Si-O-Si bonds that bridge the organic polymer and inorganic filler. This method aims to overcome interfacial weakness in PBAT/talc composites while reducing material costs. Afterwards, Soxhlet extraction, Fourier-transform infrared spectroscopy (FTIR) tests of the PBAT/talc/silane blend, as well as mechanical tests have also confirmed the structural change of PBAT, as well as the increased tensile properties in machine and transverse direction. Ultimately, the study confirmed the PBAT mulch film under such grafting refinement has a significant improvement on mechanical properties, making it comparable to traditional PE mulch film in terms of utility.

### 2.1. Material Synthesis and Grafting Confirmation

PBAT (Ecoflex® FBX 7011, BASF) was reactively grafted with talc (Luzenac 20M00S, ~6  $\mu\text{m}$ , dried) using vinyl-trimethoxysilane (VTMOS) or vinyl-methyldimethoxysilane (VMDMOS) (Gelest) as coupling agents and 2,5-Bis(tert-butylperoxy)-2,5 dimethylhexane (Luperox 101, Sigma-Aldrich) as a free-radical initiator. PBAT (700g) was hand-mixed sequentially with the initiator (0.1 wt% of PBAT), silane (0.5 or 1.0 wt% VTMOS or VMDMOS of PBAT), and talc (300g). The mixture was extruded in a co-rotating twin-screw extruder (Century ZSK-30, L/D=40:1) at 150 rpm. The temperature profile ranged from 90°C (feed) to 175°C (die). Strands were cooled and pelletized.

Grafting was confirmed via solvent extraction (dichloromethane, 24hr) and FTIR. Unmodified PBAT/talc (70/30) blends showed complete separation: extracted pure PBAT and residual pure talc (FTIR peaks: 1009  $\text{cm}^{-1}$  Si-O-Si, 670  $\text{cm}^{-1}$  Si-O-Mg). In contrast, silane-modified blends (e.g., PBAT/talc/VTMOS/Luperox 70/30/1/0.1) yielded only ~55 wt% extractable PBAT. FTIR of the non-extractable residue showed characteristic peaks of both PBAT (1715  $\text{cm}^{-1}$  C=O, 729  $\text{cm}^{-1}$  benzene, ~2950  $\text{cm}^{-1}$  -CH<sub>2</sub>-) and talc (shifted Si-O-Si peak at ~968  $\text{cm}^{-1}$ , 667  $\text{cm}^{-1}$  Si-O-Mg), confirming PBAT-talc copolymer formation via Si-O-Si bonds from hydrolyzed silane.

## 2.2. Film Processing and Mechanical/Barrier Properties

Modified composites and unmodified control were blown into films (~10  $\mu\text{m}$ ) using a single-screw blown film line (Killion, screw  $\varnothing 25.4$  mm, L/D=25:1, die gap 1.5 mm, melt temp ~150°C, screw speed 15 rpm). Tensile properties (ASTM D882) were measured. Results are summarized in Table 2-1 and Table 2-2.

Table 2-1 Tensile Properties (Machine Direction) of blown films

Sample	% Free Radical Initiator	Type and (% Amount of Silane)	Young's modulus (MPa)	Yield Stress (MPa)	Tensile Stress (MPa)	Break Elongation (%)
1	0	0	117.21	12.69	14.20	600
2	0.1	VTMOS (0.5)	391.97	39.99	39.99	300
3	0.1	VTMOS (1.0)	378.18	31.72	32.41	230
4	0.1	VMDMOS (1.0)	330.95	29.30	29.65	470

Table 2-2 Tensile Properties (Transverse Direction) of blown films

Sample	% Free Radical Initiator	Type and (% Amount of Silane)	Young's modulus (MPa)	Yield Stress (MPa)	Tensile Stress (MPa)	Break Elongation (%)
1	0	0	118.59	9.65	10.34	300
2	0.1	VTMOS (0.5)	349.56	22.75	22.75	270
3	0.1	VTMOS (1.0)	395.07	25.51	26.20	200
4	0.1	VMDMOS (1.0)	355.77	24.13	24.13	430

Based on materials properties and cost considerations, we choose sample #2 as a modified formula at all the further tests. Water vapor transmission rate (WVTR) test was conducted at 10 micron film under the condition at 38°C, 90% RH for 24 hours. The test results demonstrate that water retention was improved significantly in sample #2. The WVTR was decreased from 1132 g/(m<sup>2</sup>·24hr) (sample #1) to 382.7 g/(m<sup>2</sup>·24hr) (sample #2), which meets the mulch film standard requirement <800 g/(m<sup>2</sup>·24hr).

## 3. Aging Test

An aging test of biodegradable film samples (with/out grafting) was evaluated in accordance with the national standards to assess the mulch film's durability and resilience over a longer period of time through shorter periods of extreme weathering. The test conditions included an irradiance of 173.4 W/m<sup>2</sup> at 340 nm, a black panel temperature of 65  $\pm$  3 °C, a relative humidity of 50  $\pm$  10% RH, and a cyclic light/spray exposure for 100 hours. After aging, tensile tests were performed on aged films, only elongation results were recorded in table 2-3 per standard requirements. The aging test results show significantly improved resistance to environmental aging after grafting reaction. Sample #2 comfortably meets and exceeds the national standards for post-aging mechanical performance.

Table 3-1: Results of aging test

Orientation	Unit	Requirement	Post Aging Elongation	
			Sample #1	Sample #2
Machine Direction	%	$\geq 80$	11	160
Transverse Direction	%	$\geq 100$	29	230

#### 4. Biodegradation Test

Biodegradability under composting conditions was assessed according to ISO 14855 or equivalent standards. Test material (Sample #2 film) and positive control (semi crystalline cellulose) were mixed with mature compost in vessels maintained at  $58 \pm 2^\circ\text{C}$  and above 50% moisture levels. The total organic carbon (TOC) was around 12%, the pH was about 7.8, and the volatile solids content (VS) above 55%.

Evolved  $\text{CO}_2$  was monitored continuously by infrared analysis over 180 days. The degree of biodegradation was calculated as  $(\text{CO}_2 \text{ from test material}) / (\text{Theoretical evolved carbon dioxide (ThCO}_2)) \times 100\%$ , and disintegration was assessed visually and gravimetrically (mass loss). Biodegradation reached a plateau of around 90% mineralization by day 120, achieving an average of 92.8% biodegradation by day 180. Plant growth (ryegrass, cress) and earthworm survival tests in compost-amended soil showed no adverse effects compared to blank compost controls, meeting ecotoxicity requirements. As shown in Table 4-1, gravimetric analysis showed a 98.4% average disintegration in sample 2. Visual observation confirmed the progressive fragmentation and disappearance of our test material. Furthermore, all of the results of sample #2 met the ASTM D6400 requirements.

Table 4-1: Degree of Disintegration

Apparatus Number	Total dry solid in test sample input (g)	Total dry solids in the residual test sample (g)	Degree of disintegration (%)
#1	1019.0	17.1	98.3
#2	1019.0	15.9	98.4
Average			98.4

## 5. Field study for Cotton Production

A field study was conducted in China to evaluate the effectiveness of using grafted biodegradable mulch film (sample #2) to aid cotton planting, including its degradation process in the field. Mulch films were placed over Tahe 2 cultivar cotton, an early-to-mid-maturing upland cotton type that requires 120 days to mature. The mulch films were placed on machines and installed with standard practices.

### 5.1. Methods and Measurements

Three treatment groups were tested: 10.2  $\mu\text{m}$  thick biodegradable film (sample #2), 12.1  $\mu\text{m}$  thick PE film, and bare soil (control). A thinner BDM was chosen due to stronger tensile properties of the biodegradable films vs. PE mulch films, but both types of films were laid down with a cross-sectional width of 205 cm. A randomized block design was conducted with three replicates per treatment group. Traditional bare soil served as control.

Throughout the planting time, ground temperature was measured by using automatic temperature recording probes (accuracy  $\pm 0.2^\circ\text{C}$ ) which were buried at a depth of 10 cm. Data was recorded every 60 minutes at each of the three different locations per treatment group, then daily average temperatures were calculated. The state of degradation of the films was observed every 10 days post-application and classified by stage. The stages were determined by the size of cracks on the film: induction ( $\leq 2$  cm cracks), cracking (2–20 cm), fragmentation ( $> 20$  cm), disintegration (residues  $\leq 16$  cm<sup>2</sup>), and disappearance. Plant height, boll count, average single boll weight, and yield were measured as crop metrics during the harvest.

### 5.2. Results

Although biodegradable mulch films were slightly less effective than traditional PE films at increasing plant height and cotton boll weight, the number of cotton bolls produced per plant and the overall yield per hectare were higher for biodegradable films. Overall levels of productivity for the two products were very similar.

Table 5-1: Effect of different mulch film covering on cotton yield

	Plant Height (cm)	Cotton bolls per plant	Avg boll weight (g)	Cotton yield (kg/hectare)
PE Film	80.3 $\pm$ 0.5	6.42 $\pm$ 0.20	4.84 $\pm$ 0.10	5494.5 $\pm$ 62.5
BDM Film	79.8 $\pm$ 0.6	6.48 $\pm$ 0.13	4.82 $\pm$ 0.11	5518.6 $\pm$ 53.2
No Mulch Film	76.4 $\pm$ 0.6	3.20 $\pm$ 0.10	4.20 $\pm$ 0.10	2379.2 $\pm$ 84.3

Table 5-1 displays the statistical results of the cotton yields, comparing plant heights, the number of cotton bolls per plant, and weights of each boll for cotton grown with different films. Whereas cotton plants covered in mulch films both exhibit increases in height vs. uncovered plants, there is not a statistically significant difference in plant height using PE vs. BDM film. It may be observed that the presence of mulch films significantly affects the yield of cotton. The yield of biodegradable mulch film is the highest, though there is no significant difference in the yields of different mulch films. The yields of cotton cultivated on land with different mulch films are 2.3 times that of cotton cultivated on bare land. This is mainly reflected in the number of cotton bolls per covered plant (3.22–3.28 more than plants cultivated on bare land) and the 0.42–0.44 g increase in individual boll weight.

The results of the study demonstrated that when it came to soil temperature measurements, the biodegradable mulch film outperformed PE film, increasing soil temperature of 10 cm by 0.35–0.89°C during early growth (0–60 days) (Figure 5-1). There was no significant difference in terms of moisture retention and weed control.

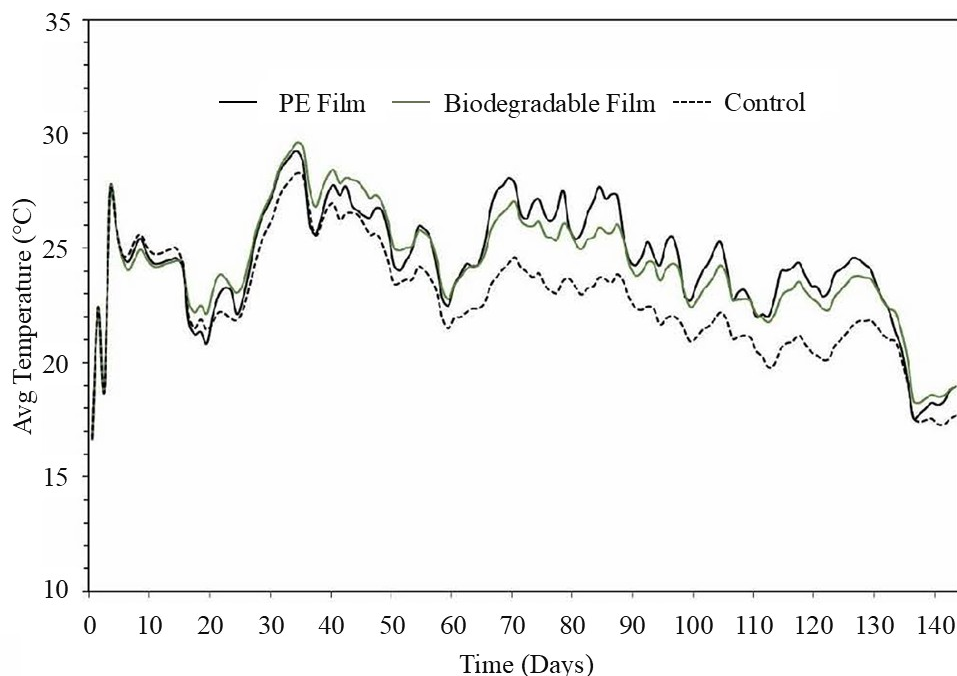


Figure 5-1: Changes in soil temperature in cotton fields with PE film, biodegradable film and with no film.

The disintegration speed of films were then matched with crop cycles (Figure 5-2). The degradation timelines were followed as planned for cotton growth needs. Cotton was sown on May 5. 120 days later, the mulch film surface had multiple (2-3 per square meter) natural cracks or holes with cross-sections below 2 cm, entering the induction period. 141 days after the cotton was sown, natural cracks with cross-sections between 2 cm and 20 cm began to appear, entering the cracking period. 152 days after the seeds were sown, it entered the fragmentation period, in which the film surface began to have natural cracks more than 20 cm long. 154 days after planting, it entered the disintegration period: the mulch film on the surface lost its flexibility and fragmented, with mulch film fragment averaging areas below 16 cm<sup>2</sup>. Finally, full degradation was achieved by harvest. It can be seen that after the biodegradable mulch film entered the induction period, the subsequent degradation time was relatively fast, and it only took one week to transition from the large cracking period to the fragmentation period. There was essentially no change in the visible structure of the PE film.





Figure 5-2: Disintegration of biodegradable mulch film for Tahe 2 cultivar cotton. Data was measured at three different sites (BDM01, BDM02, BDM03). Induction of degradation began by day 120 and full degradation occurred by harvest.

## 6. Conclusions

The study successfully developed and validated high-performance BDMs specifically engineered for long-growth-cycle crops like cotton. As demonstrated by field tests, reactive grafting of PBAT with talc filler using vinyl-silane coupling agents significantly enhanced key properties, such as mechanical strength, durability, moisture retention, and biodegradability. In particular, grafted PBAT achieved tensile properties comparable to PE film and demonstrated near-complete mineralization and disintegration under composting conditions within 180 days, with no ecotoxic effects. The optimized BDM provided equivalent agronomic support (soil warming, moisture retention, weed control) and crop yield to conventional PE film throughout the critical 120-150 day cotton growth period. Critically, it then degraded over a predictable timeline aligned with the crop cycle, entering disintegration shortly after boll opening and achieving full degradation post-harvest before the next planting season. This technology effectively resolves the major limitation of premature disintegration of existing BDMs for long-duration crops. Thus, farmers avoid yield losses associated with premature disintegration. By enabling customization of degradation kinetics through material formulation, it offers a viable, sustainable solution to replace persistent PE mulch, eliminating plastic pollution and retrieval costs without compromising agricultural productivity.

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