

**AKTIVITAS ANTIINFLAMASI DAN POTENSI ANTIKANKER
FORMULASI NANOKRISTAL EKSTRAK DAUN KELOR DAN
CIPLUKAN SECARA *IN VITRO***

SKRIPSI

Diajukan sebagai salah satu syarat untuk memperoleh gelar Sarjana Sains
Program Studi Biologi



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**PROGRAM STUDI BIOLOGI
FAKULTAS PENDIDIKAN MATEMATIKA DAN ILMU PENGETAHUAN ALAM
UNIVERSITAS PENDIDIKAN INDONESIA
2025**

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Sebuah Skripsi yang diajukan untuk memenuhi salah satu syarat memperoleh
gelar Sarjana Sains pada Fakultas Pendidikan Matematika dan Ilmu Pengetahuan
alam

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Agustus 2025

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Dengan ini saya menyatakan bahwa skripsi dengan judul "Aktivitas Antiinflamasi dan Potensi Antikanker Formulasi Nanokristal Ekstrak Daun Kelor dan Ciplukan secara *In Vitro*" ini beserta seluruh isinya adalah benar-benar karya saya sendiri. Saya tidak melakukan penjiplakan atau pengutipan dengan cara-cara yang tidak sesuai dengan etika ilmu yang berlaku dalam masyarakat keilmuan. Atas pernyataan ini, saya siap menanggung risiko/sanksi apabila dikemudian hari ditemukan adanya pelanggaran etika keilmuan atau ada klaim dari pihak lain terhadap keaslian karya saya ini.

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KATA PENGANTAR

Puji syukur penulis panjatkan ke hadirat Allah SWT karena berkat rahmat dan karunia-Nya, penulis dapat menyelesaikan penyusunan skripsi ini dengan judul "Aktivitas Antiinflamasi dan Potensi Antikanker Formulasi Nanokristal Ekstrak Daun Kelor dan Ciplukan secara *In Vitro*". Skripsi ini disusun sebagai salah satu syarat untuk memperoleh gelar Sarjana pada Program Studi Biologi, Fakultas Pendidikan Matematika dan Ilmu Pengetahuan Alam, Universitas Pendidikan Indonesia.

Penulis menyadari bahwa dalam proses penyusunan ini masih terdapat banyak kekurangan, baik dalam hal isi maupun penyajian. Oleh karena itu, penulis membuka diri terhadap masukan, kritik, dan saran yang membangun demi kesempurnaan karya ini di kemudian hari. Akhir kata, semoga skripsi ini dapat memberikan manfaat khususnya bagi penuliss dan umumnya bagi pembaca serta menjadi sumbangsih kecil dalam pengembangan ilmu pengetahuan di bidang biomedik.

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ABSTRAK

Stres oksidatif memicu inflamasi kronis yang dapat menyebabkan kerusakan DNA dan perkembangan kanker. Pengobatan konvensional sering menimbulkan efek samping seperti gangguan pencernaan, resistensi obat, dan toksisitas sistemik. Penelitian ini bertujuan mengevaluasi potensi kombinasi nanokristal ekstrak daun kelor dan ciplukan (NKEKC) sebagai agen antioksidan, antiinflamasi, dan antikanker secara *in vitro*. Nanokristal dibuat dari ekstrak serbuk kelor dan ciplukan, kemudian dikarakterisasi PSA dan ZPA. Hasil menunjukkan ukuran partikel NKEK dan NKEC masing-masing 323 nm dan 671 nm, dengan zeta potensial -40.5 mV dan -52.3 mV. Uji DPPH menunjukkan NKEKC mampu menghambat radikal bebas hingga 55,06% pada konsentrasi 4 mg/mL. Pada uji denaturasi protein, NKEKC menghambat denaturasi hingga 98,39% pada konsentrasi 1 mg/mL. Uji sitotoksitas pada sel MCF-7 menunjukkan viabilitas sel 87,48% pada konsentrasi 1 mg/mL, namun belum mencapai IC50. Selain itu, NKEKC mampu menginduksi penuaan sel (*senescence*) pada sel MCF-7 sebesar 53,47% pada konsentrasi 0,25 mg/mL dan meningkat hingga 87,78% pada 4 mg/mL. Analisis siklus sel NKEKC terhadap sel MCF-7 menunjukkan bahwa NKEKC dapat menginduksi penghentian siklus sel pada fase G0 dan S. Berdasarkan hasil tersebut, dapat disimpulkan bahwa ekstrak kelor dan ekstrak ciplukan tergolong partikel nano yang memiliki aktivitas antioksidan dan antiinflamasi yang baik. NKEKC berpotensi sebagai agen antikanker melalui mekanisme induksi penuaan sel.

Kata kunci: stres oksidatif, inflamasi, kanker, kelor, ciplukan, nanokristal

ABSTRACT

Oxidative stress triggers chronic inflammation, which can lead to DNA damage and cancer development. Conventional treatments often cause side effects such as digestive disorders, drug resistance, and systemic toxicity. This study aims to evaluate the potential of a combination of nanocrystals from Moringa oleifera and Physalis angulata leaf extracts (NKEKC) as antioxidant, anti-inflammatory, and anticancer agents in vitro. Nanocrystals were prepared from powdered extracts of Moringa and Physalis and characterized using particle size analysis and zeta potential analysis. The results showed that the particle sizes of NKEK and NKEC were 323 nm and 671 nm, respectively, with zeta potentials of -40.5 mV and -52.3 mV. NKEKC showed a particle size of 1115 nm and a zeta potential of -22.2 mV. The DPPH assay indicated that NKEKC could inhibit free radicals up to 55.06% at a concentration of 4 mg/mL. In the protein denaturation assay, NKEKC inhibited protein denaturation by 98.39% at 1 mg/mL. Cytotoxicity testing on MCF-7 cells showed a cell viability of 87.48% at 1 mg/mL, indicating that NKEKC has not yet reached the IC50 value. Moreover, NKEKC induced cellular senescence in MCF-7 cells by 53.47% at 0.25 mg/mL, increasing to 87.78% at 4 mg/mL. NKEKC cell cycle analysis of MCF-7 cells showed that NKEKC can induce cell cycle arrest in the G0 and S phases. Based on these results, it can be concluded that Moringa and Physalis extract are classified as nanocrystals with good antioxidant and anti-inflammatory activity. NKEKC also has potential as an anticancer agent through the induction of cellular senescence.

Keywords: oxidative stress, inflammation, cancer, kelor, ciplukan, nanocrystal

DAFTAR ISI

LEMBAR PENGESAHAN	i
LEMBAR PERNYATAAN	ii
KATA PENGANTAR.....	iii
UCAPAN TERIMA KASIH	iv
ABSTRAK	vi
<i>ABSTRACT</i>	vii
DAFTAR ISI	viii
DAFTAR TABEL.....	x
DAFTAR GAMBAR	xi
DAFTAR LAMPIRAN	xii
BAB I PENDAHULUAN	1
1.1 Latar Belakang	1
1.2 Rumusan Masalah	5
1.3 Pertanyaan Penelitian.....	5
1.4 Tujuan Penelitian	6
1.5 Batasan Penelitian	6
1.6 Manfaat Penelitian	6
1.7 Struktur Penulisan Skripsi.....	7
BAB II AKTIVITAS TERAPEUTIK DAUN KELOR DAN CIPLUKAN SEBAGAI ANTIOKSIDAN, ANTIINFLAMASI DAN ANTIKANKER SECARA <i>IN VITRO</i>	9
2.1 Inflamasi dan Mekanismenya	9
2.2 Induksi Karsinogenesis akibat Inflamasi Kronis.....	10
2.3 Antioksidan dan Peranannya	11
2.4 Kandungan dan Potensi Medis Kelor (<i>Moringa oleifera</i>).....	13
2.5 Kandungan dan Potensi Medis Ciplukan (<i>Physalis angulata</i>).....	14
2.6 Kanker Payudara dan Pengendalian Kanker	16
2.7 Potensi Kelor dan Ciplukan sebagai Antiinflamasi dan Antikanker	19
2.8 Nanokristal dalam Dunia Kesehatan	21
BAB III METODOLOGI PENELITIAN.....	24
3.1 Jenis Penelitian.....	24
3.2 Desain Penelitian.....	24

3.3 Waktu dan Lokasi Penelitian.....	24
3.4 Populasi dan Sampel	25
3.5 Alur Penelitian.....	25
3.6 Alat dan Bahan	26
3.7 Prosedur Penelitian.....	26
3.7.1 Ekstraksi Daun Kelor dan Daun Ciplukan	26
3.7.2 Pembuatan Nanokristal dan Karakterisasi	26
3.7.3 Pembuatan Formulasi Nanokristal Ekstrak Kelor dan Nanokristal Ekstrak Ciplukan	26
3.7.4 Uji Aktivitas Antioksidan.....	26
3.7.5 Uji Aktivitas Antiinflamasi	27
3.7.6 Preparasi Sel dan Larutan Stok NKEKC	28
3.7.7 Uji Sitotoksitas sel MCF-7 dengan perlakuan Nanokristal	29
3.7.8 Uji <i>Senescence</i> Sel MCF-7	30
3.7.9 Uji Aktivitas Siklus Sel	31
3.7.10 Analisis statistik	32
BAB IV TEMUAN DAN PEMBAHASAN	33
4.1 Karakterisasi Nanokristal.....	33
4.2 Analisis Aktivitas Antioksidan.....	35
4.3 Aktivitas Antiinflamasi	38
4.4 Analisis Aktivitas Antikanker	41
4.4.1 Analisis Aktivitas Sitotoksik	41
4.4.2 Aktivitas <i>Senescence</i> Sel MCF-7	43
4.4.3 Analisis Siklus Sel	46
4.5 Hubungan Aktivitas Antioksidan dan Antiinflamasi Formulasi Nanokristal Ekstrak Daun Kelor dan Ciplukan	49
4.6 Potensi Formulasi Nanokristal Ekstrak Daun Kelor dan Ciplukan sebagai Agen Antioksidan, Antiinflamasi, dan Antikanker	50
BAB V SIMPULAN, IMPLIKASI, DAN REKOMENDASI	52
5.1 Simpulan	52
5.2 Implikasi.....	52
5.3 Rekomendasi	52
DAFTAR PUSTAKA	54
LAMPIRAN.....	66

DAFTAR TABEL

Tabel 4.1. Hasil Karakterisasi PSA dan ZPA.....	33
Tabel 4.2. Aktivitas Antioksidan NKEKC.....	35
Tabel 4.3. Hasil Aktivitas Antiinflamasi oleh NKEKC	38
Tabel 4.4. Aktivitas Antikanker NKEKC dengan Uji Sitotoksik	42
Tabel 4.5. Aktivitas <i>Senescence</i> Sel MCF-7 oleh NKEKC	43
Tabel 4.6 Aktivitas Penghentian Siklus Sel MCF-7 oleh NKEKC	46

DAFTAR GAMBAR

Gambar 2.1 Siklus Sel	18
Gambar 3.1. Alur Penelitian	25
Gambar 3.2 Skema plat 96 sumur (<i>96 well plate</i> untuk uji antikosidan DPPH ..	27
Gambar 3.3 Skema plat 96 sumur (<i>96 well plate</i>) untuk uji sitotoksik	30
Gambar 3.4 Skema plat 12 sumur (<i>12-Well plate</i>) untuk uji <i>senescence</i> sel	31
Gambar 4.1. Aktivitas inhibisi denaturasi protein oleh NKEKC	39
Gambar 4.2 Sel yang tidak terwarnai tidak mengalami <i>senescence</i> (a), sel yang terwarnai mengalami <i>senescence</i> (b)	44
Gambar 4.3 Kontrol Negatif (perbesaran 100x)	45
Gambar 4.4 Konsentrasi NKEKC 4.0 mg/mL (perbesaran 100x)	45
Gambar 4.5 Konsentrasi NKEKC 2.0 mg/mL (perbesaran 100x)	45
Gambar 4.6 Konsentrasi NKEKC 1.0 mg/mL (perbesaran 100x)	45
Gambar 4.7 Konsentrasi NKEKC 0.5 mg/mL (perbesaran 100x)	45
Gambar 4.8 Konsentrasi NKEKC 0.25 mg/mL (perbesaran 100x)	45
Gambar 4.9 Hasil Analisis Flowsitometri Siklus Sel MCF-7	48
Gambar 4.10 Peran NKEKC sebagai Antioksidan, Antiinflamasi, dan Antikanker	51

DAFTAR LAMPIRAN

Lampiran 1 Alat dan Bahan.....	66
Lampiran 2 Hasil Karakterisasi Nanokristal Ekstrak Kelor.....	68
Lampiran 3 Hasil Karakterisasi Nanokristal Ekstrak Ciplukan	69
Lampiran 4 Hasil Uji NKEKC	70
Lampiran 5 Hasil Analisis Statistik Uji Antioksidan	72
Lampiran 6 Hasil Analisis Statistik Uji Antiinflamasi	75
Lampiran 7 Hasil Analisis Statistik Uji <i>Senescence</i>	81
Lampiran 8 Hasil Analisis Statistik Uji Siklus Sel.....	82
Lampiran 9 Hasil Analisis Statistik Korelasi Spearman's.....	84
Lampiran 10 Dokumentasi Penelitian.....	85

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