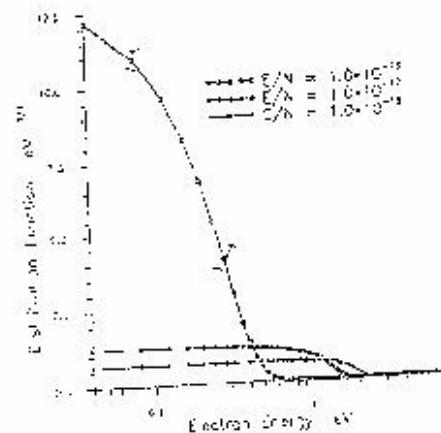
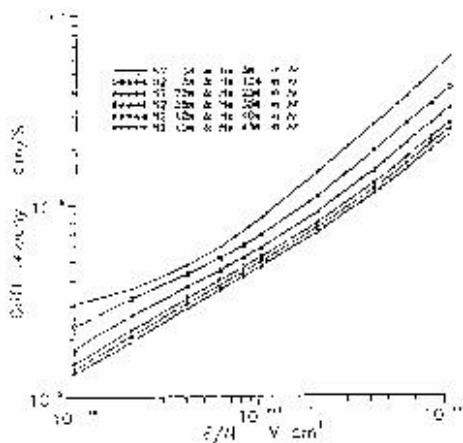


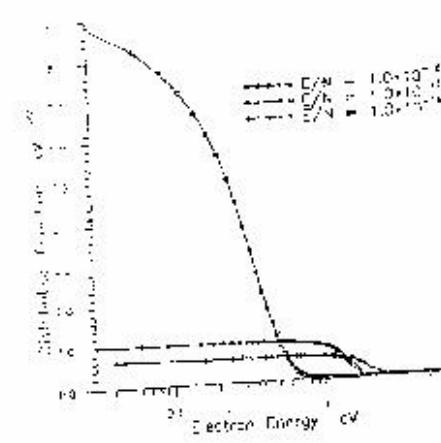
شكل (9) خصائص طاقة الالكترونات كـ $E/N$  لمزج من غاز (أرجون-هيليوم-نيتروجين).



شكل (6) التوزيع الالكتروني كـ $f(E)$  لطاقة الالكترون في مزج غاز (70% أرجون 15% هيليوم 15% نيتروجين).



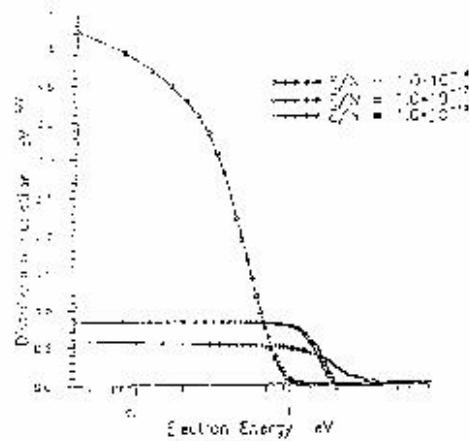
شكل (10) اتجاه سرعة الالكترونات كـ $E/N$  لمزج من غاز (أرجون-هيليوم-نيتروجين).



شكل (7) التوزيع الالكتروني كـ $f(E)$  لطاقة الالكترون في مزج غاز (80% أرجون 10% هيليوم 10% نيتروجين).

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شكل (8) التوزيع الالكتروني كـ $f(E)$  لطاقة الالكترون في مزج غاز (90% أرجون 5% هيليوم 5% نيتروجين).

## Abstract

Alkaline protease was purified from *B. stearothermophilus* ATCC25725 by several steps included heat treatment, DEAE-Cellulose ion exchange chromatography and gel filtration on Sepharose-6B column. Gel filtration resulted in separation of the enzyme preparation into protease I and protease II, the obtained purification folds were 5.9 and 9.8 respectively and recovery was 13 and 37 respectively.

The molecular weight of the purified protease I and Protease II were 13182 and 20892 daltons as determined by gel filtration and 56234 and 69502 daltons respectively as determined by SDS-PAGE.

The optimal pH for activity protease I and II on casein were 7 and 10 respectively while the protease I and protease II were most stable in pH range (6.5-7.5) and (8-10) respectively using casein as a substrate.

The maximum purified proteases activity was observed at 65 and 85 °C respectively. The protease I and II retained 100% activity at 60 and 65 °C for 30 min.