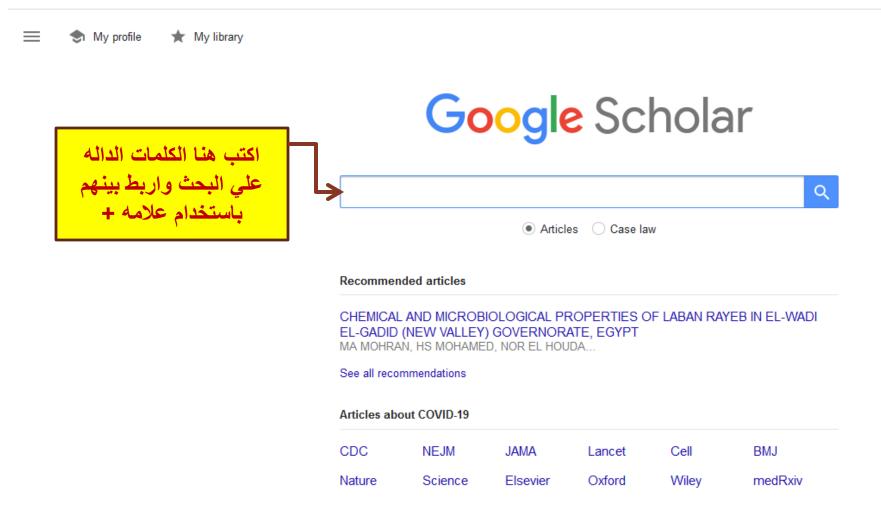


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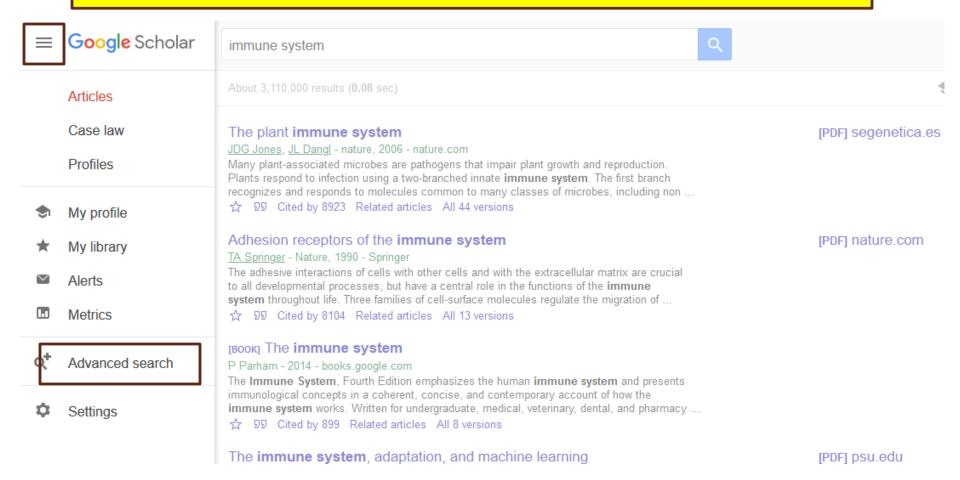
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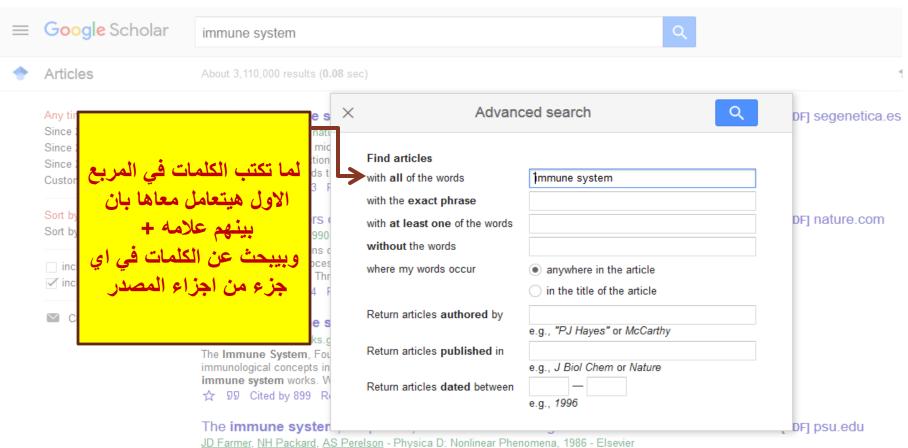




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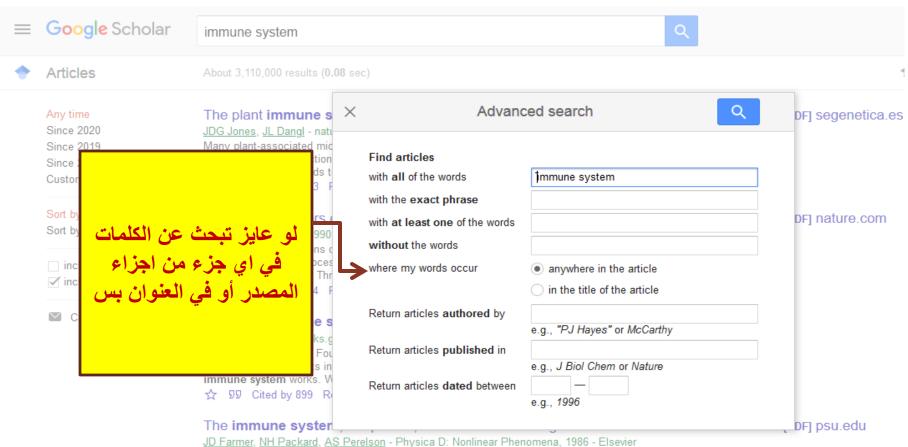
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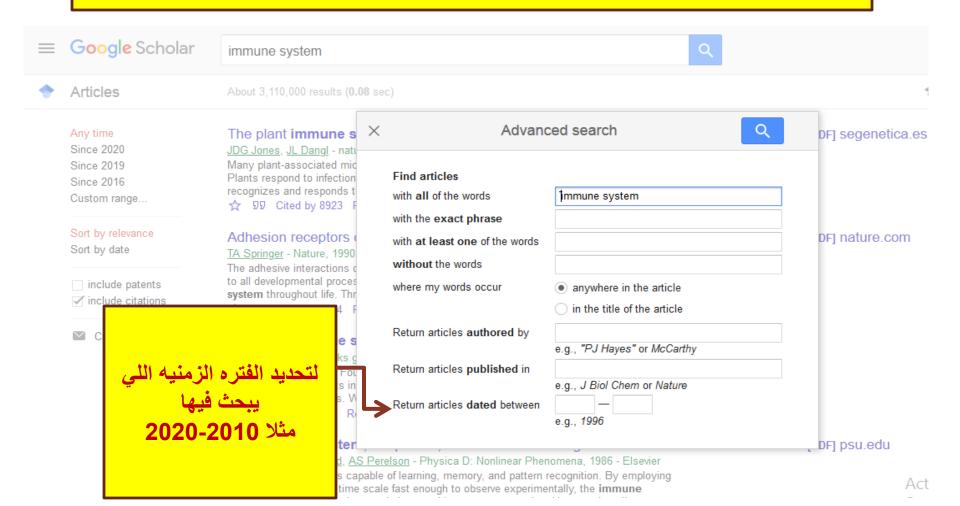
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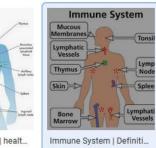


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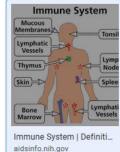
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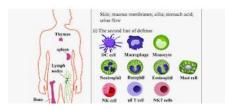
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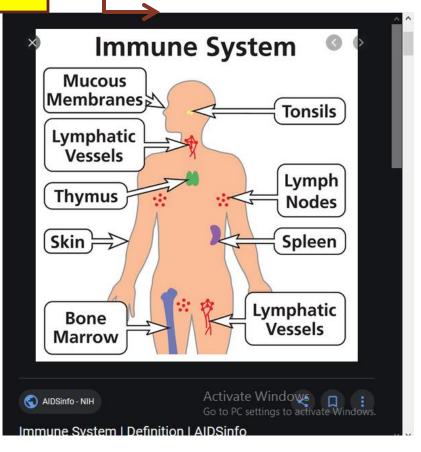


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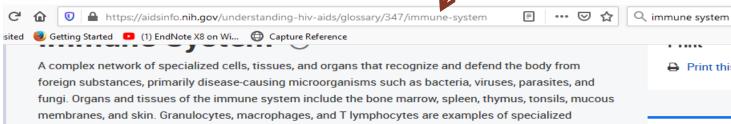
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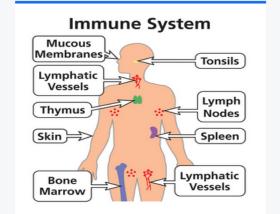
cells. The lymphatic vessels of the immune system carry immune cells, which converge in lymph nodes

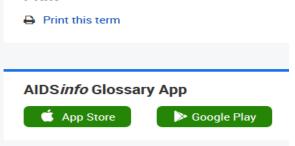
found throughout the body. A swollen lymph node often indicates an active immune response to a foreign

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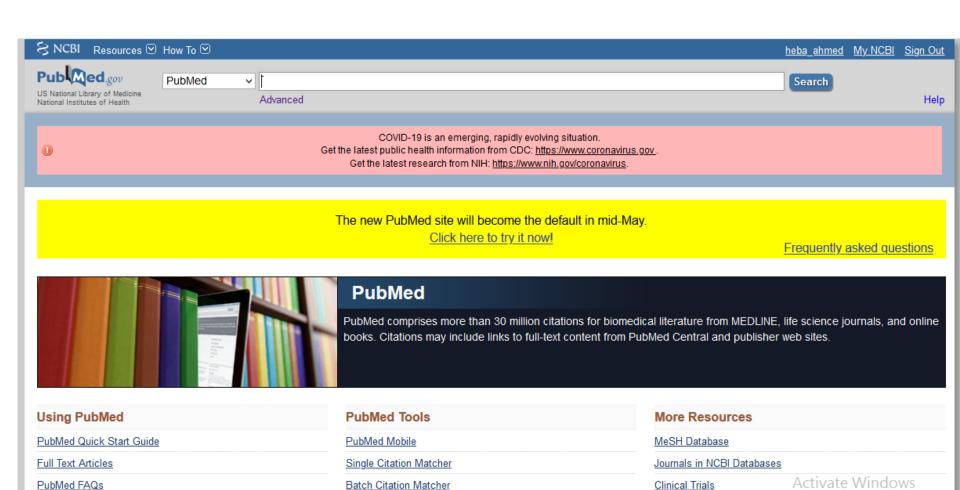


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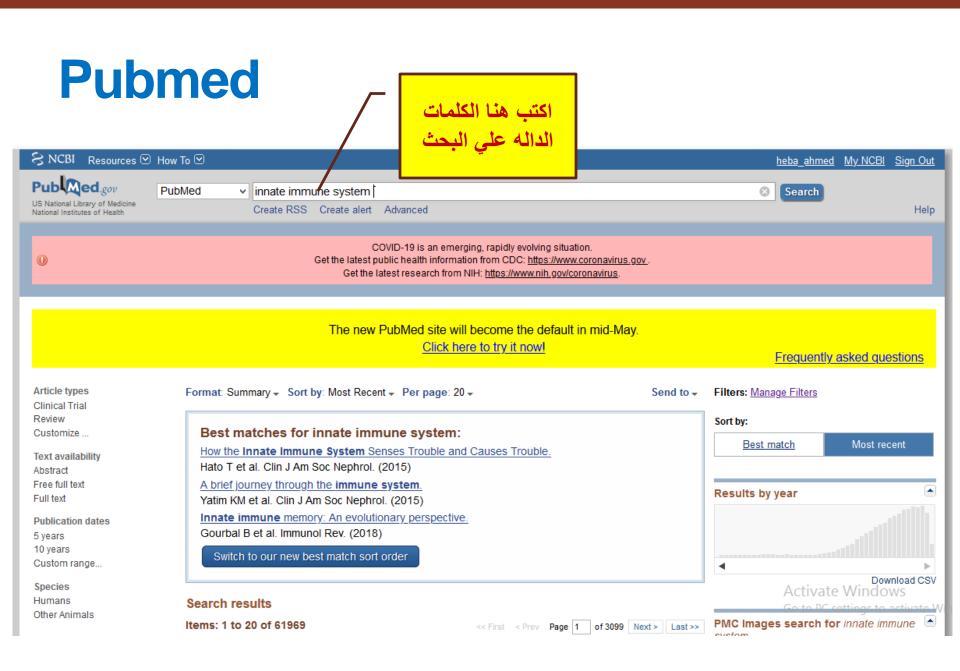


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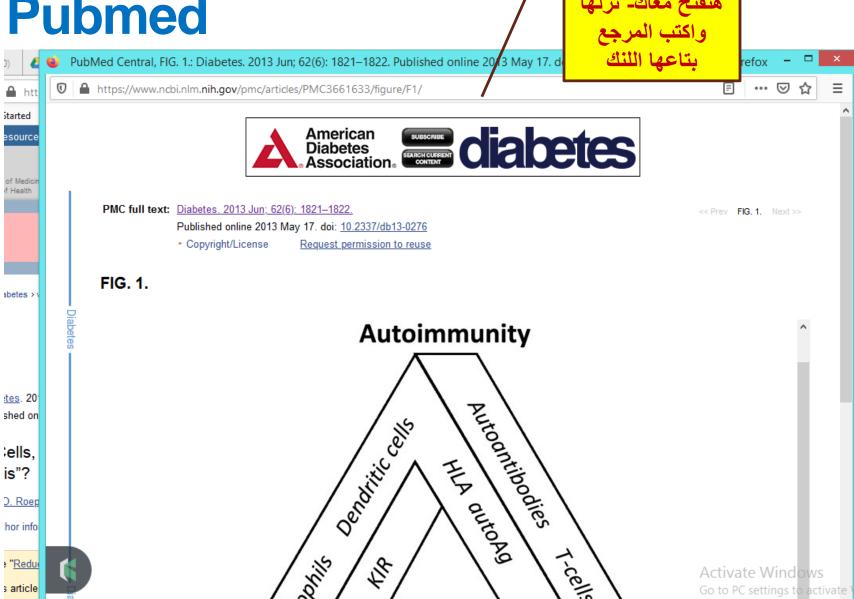
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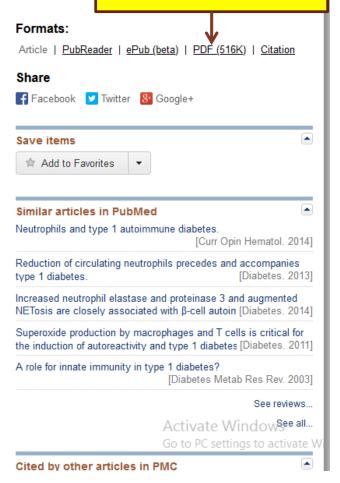
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In this issue of *Diabetes*, Valle et al. (2) present their perhaps surprising observation that slightly reduced circulating neutrophil counts associate with type 1 diabetes. The authors demonstrate that this phenomenon



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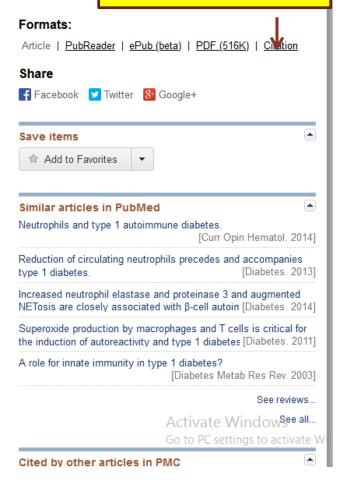
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Diet reversal and immune modulation show key role for liver and adipose tissue T cells in murine non-alcoholic steatohepatitis.

Van Herck MA¹, Vonghia L², Kwanten WJ³, Julé Y⁴, Vanwolleghem T³, Ebo D⁵, Michielsen PP³, De Man J⁶, Gama L⁷, De Winter BY⁶, Francque SM³.

Author information

Abstract

BACKGROUND & AIMS: Non-alcoholic steatohepatitis (NASH) is a multisystem condition, implicating liver and adipose tissue. Whereas the general involvement of the innate and adaptive immune system has been established, we aimed to define the exact role of the functionally diverse T-cell subsets in NASH pathogenesis through diet reversal and immunological modulation.

METHODS: Multiple experimental setups were used in 8-week-old C57BL/6J mice, including prolonged high-fat high-fructose diet (HFHFD) feeding, diet reversal (DR) from HFHFD to control diet, and administration of anti-CD8a and anti-IL-17A antibodies. Plasma ALT, glucose, and lipid levels were determined. Liver and adipose tissue was assessed histologically. Cytotoxic T (Tc), regulatory T (Treg), T helper (Th) 1, and Th17 cells were characterized in liver and visceral adipose tissue (VAT) via flow cytometry and RNA analysis.

RESULTS: HFHFD feeding induced the metabolic syndrome and NASH, which coincided with an increase in hepatic Th17, VAT Tc, and VAT Th17 cells and a decrease in VAT Treg cells. Although DR induced a phenotypical metabolic and hepatic normalization, the observed T-cell disruptions persisted. Treatment with anti-CD8a antibodies lowered Tc cell numbers in all investigated tissues and induced a biochemical and histological attenuation of the HFHFD-induced NASH. Conversely, anti-IL-17A antibodies decreased hepatic inflammation without affecting other features of NASH or the metabolic syndrome.

CONCLUSIONS: HFHFD feeding induces important immune disruptions in multiple hepatic and VAT T-cell subsets, refractory to diet reversal. In particular, VAT Tc cells are critically involved in NASH pathogenesis, linking adipose tissue inflammation to liver disease.



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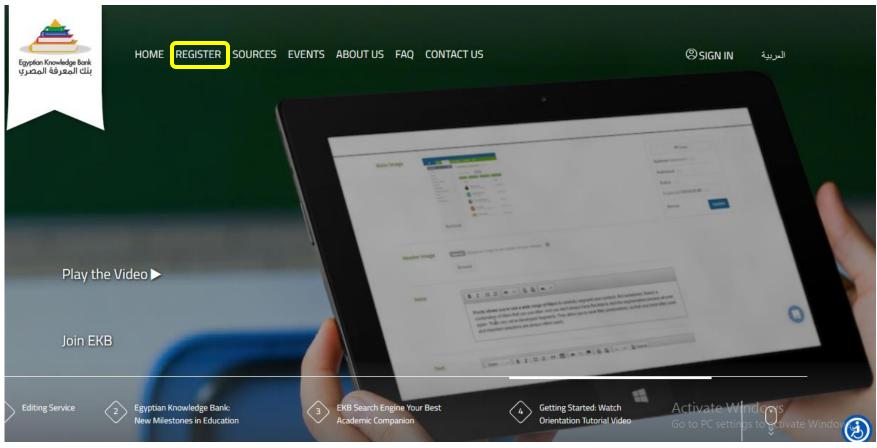
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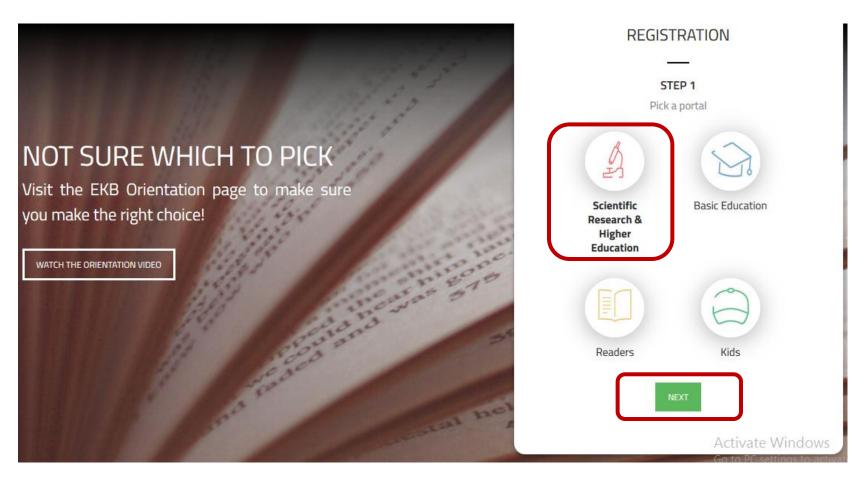
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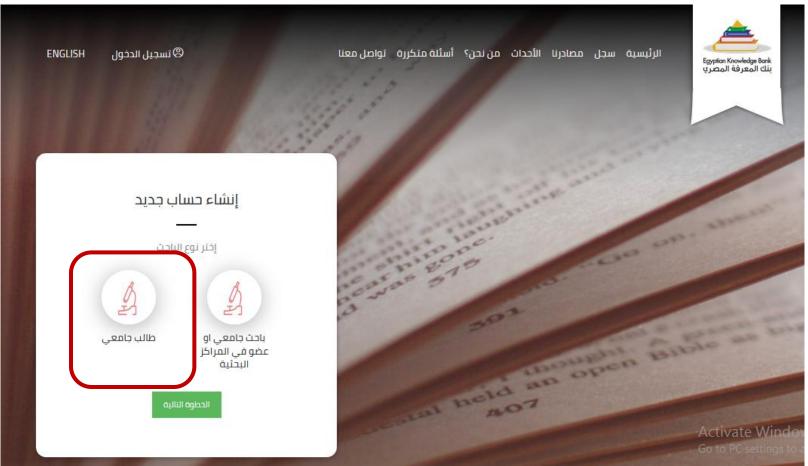
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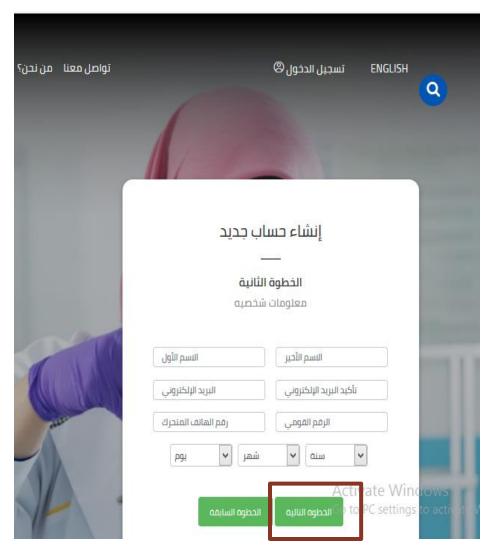
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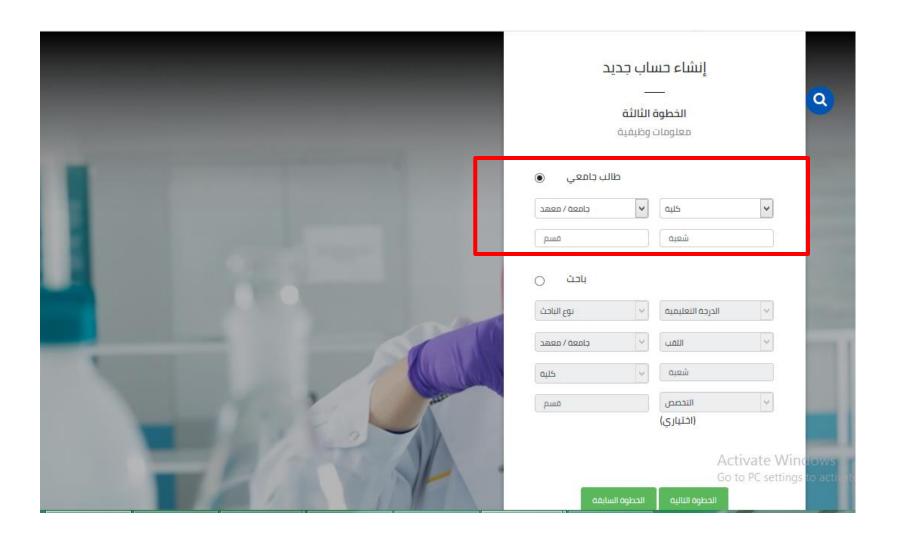


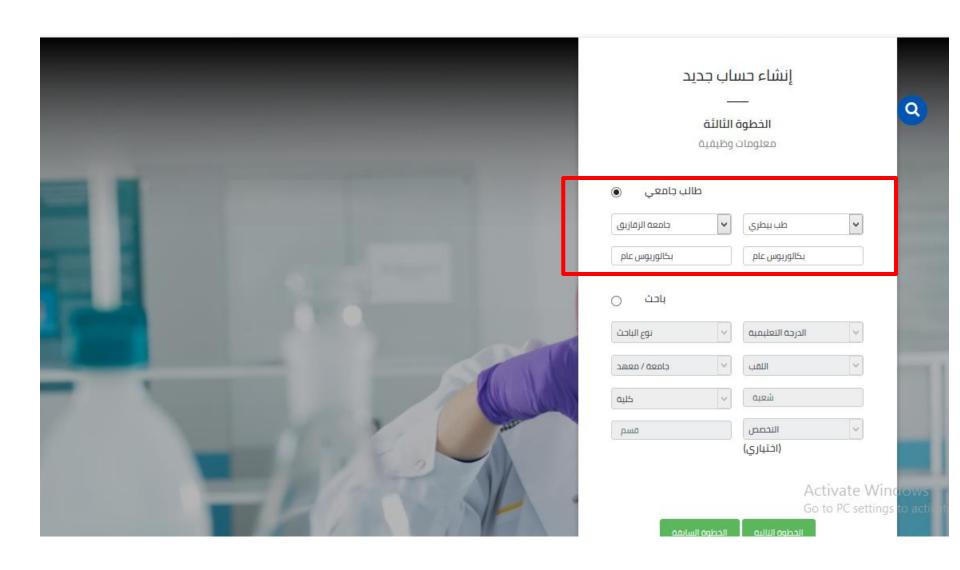




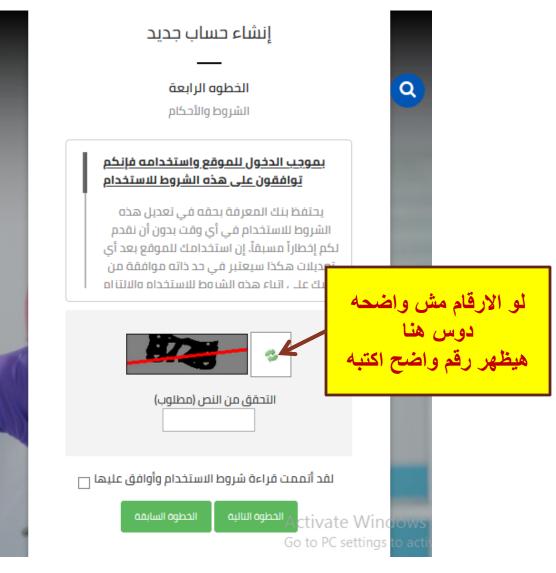


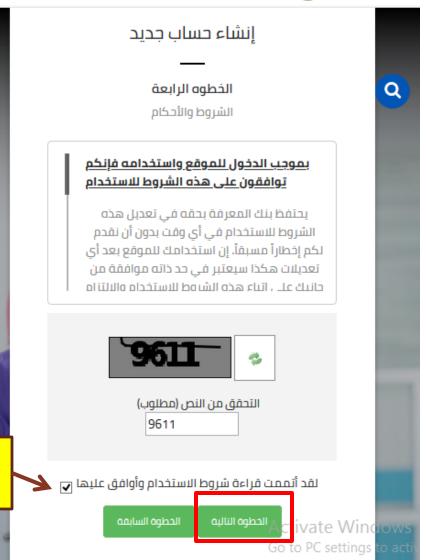
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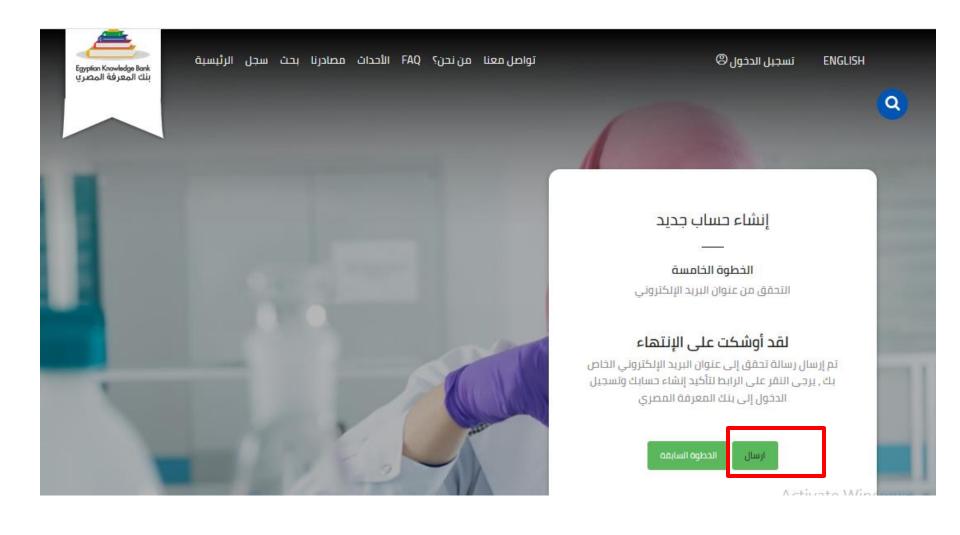








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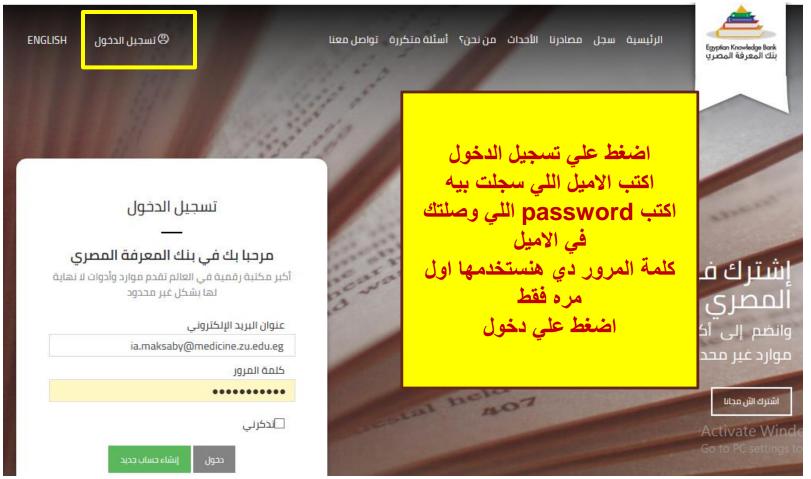
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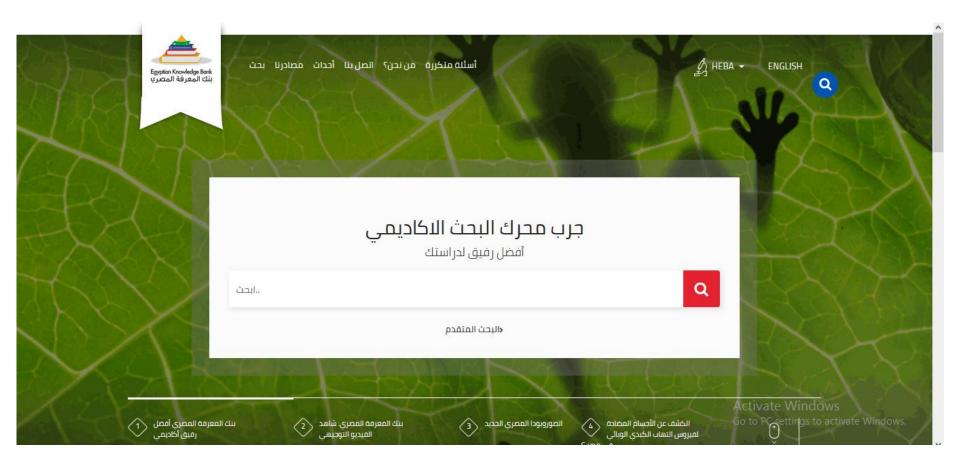
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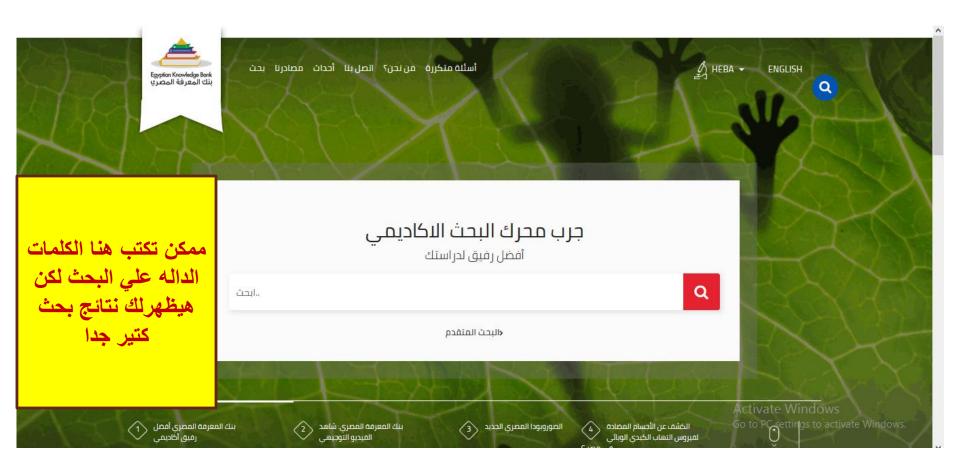
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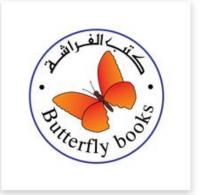
















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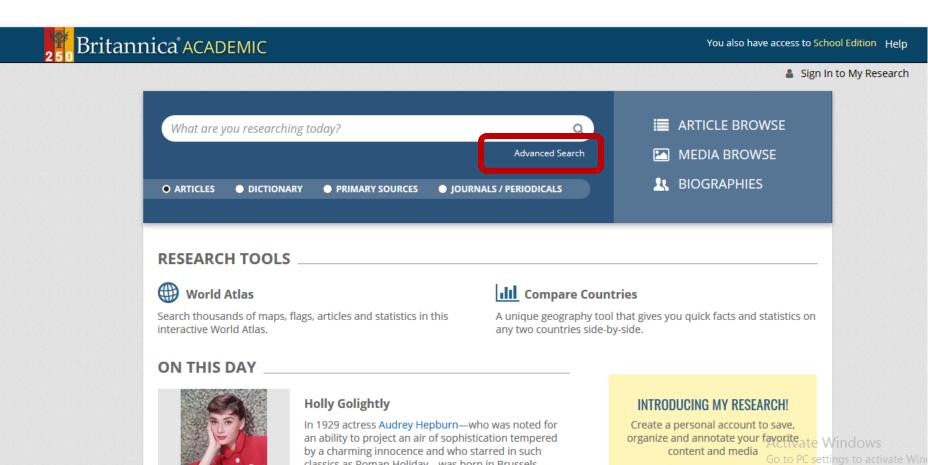
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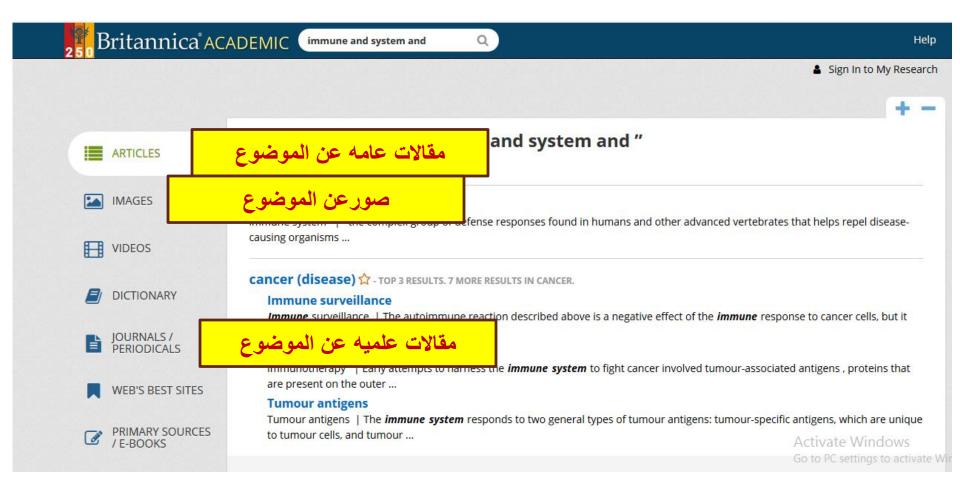
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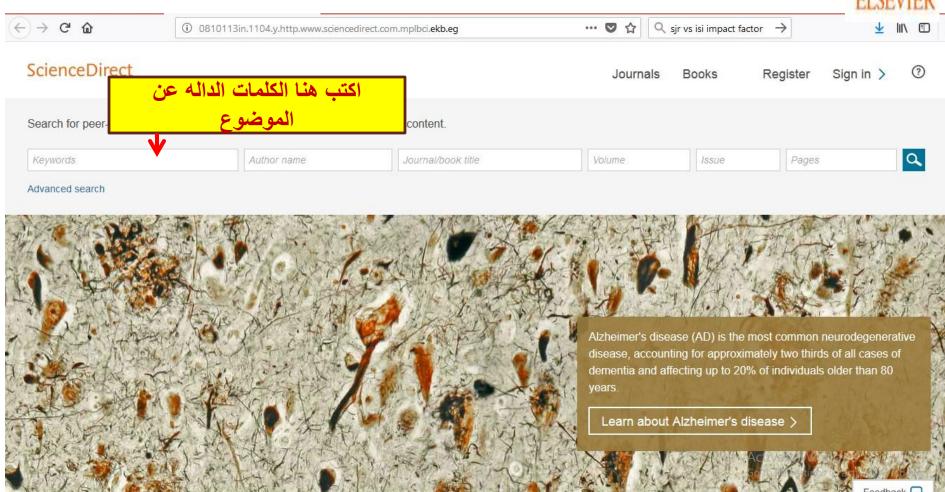
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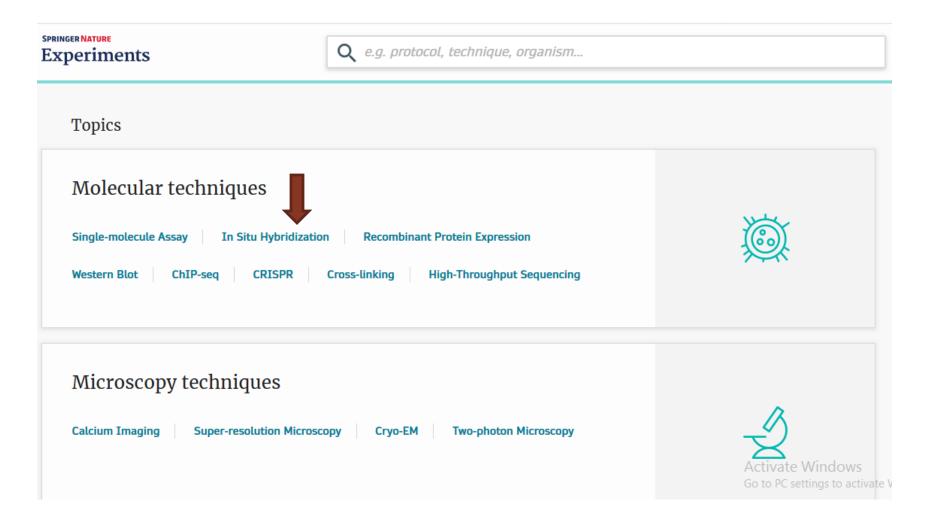
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- ح عدم الاعتماد علي مذكرة او كتاب القسم في اعداد البحث.
- جزء conclusion فقط هو الذي يتم فيه تلخيص ما فهمته من المحاضرات او الكتاب الخاص بالقسم وتكتب باسلوبك
 - ح الالتزام بعدد الصفحات والتنسيق

