

The role of the innate immune cells in the development of type 1 diabetes

December 19 2012

Julien Diana and Yannick Simoni of the "Immune Mechanisms in Type 1 Diabetes," Inserm/Université Paris Descartes, directed by Agnès Lehuen, have just published the results of their work on type 1 diabetes in the *Nature Medicine* journal.

The researchers reveal the role of the innate immune cells, especially the <u>dendritic cells</u>, that cause the activation of the killer <u>T-lymphocytes</u> whose action is directed against the p pancreatic cells. The results obtained in mice make it possible to consider new ways of regulating the auto-immune reaction generated by the innate immune cells.

Type 1 diabetes, or insulin-dependent diabetes, is an auto-immune disease characterised by the destruction of insulin-producing pancreatic β cells that are present in the Islets of Langerhans which are themselves in the pancreas. The peculiarity of this type of diabetes lies in the fact that the cells are destroyed by T lymphocytes that kill the patient's immune system. This is an auto-immune reaction. Much of the research has highlighted the role of auto-reactive T lymphocytes in the pancreatic β cells. Yet the mechanisms involved in the initial activation of the immune system that triggers the sequence of events leading to the death of the cells are still ill-defined.

The work of the team working on "<u>Immune mechanisms</u> of Type 1 diabetes" (Inserm/Université de Paris Descartes) in NOD (Non-obese diabetic) mice, the model used for studying Type 1 diabetes, reveals the essential role played by the cells of the innate immune system that were



not hitherto considered to be involved in diabetes. These cells were known to play a role in other auto-immune diseases such as lupus and psoriasis. In this study, researchers managed to describe the mechanisms initiating the activation of T lymphocytes attaching themselves to the pancreatic β cells.

The innate immune system is normally activated when an infection occurs. This is when the mobilisation and activation of <u>neutrophils</u> and dendritic cells is observed, constituting the first stages of the immune response. Abnormally in NOD <u>diabetic mice</u>, the natural physiological death of pancreatic β cells, occurring as the process progresses, involves an innate auto-immune response in the pancreas. Researchers have provided details of the introduction of a sequence of activation events in these innate immune cells in the Islets of Langerhans.

"We have observed in mice that treatment prevents activation of the innate immune cells, neutrophils and dendritic cells, warning of the onset of diabetes by inhibiting the appearance of auto-immune T responses aimed at the pancreas" explains Agnès Lehuen, head of the Inserm team.

These results show, for the first time, the important role played by innate immune cells in the sequence of events leading to the onset of Type 1 diabetes. Researchers continue to strive to understand how to regulate the auto-<u>immune reaction</u> produced by dendritic cells without compromising the <u>innate immune system</u>, an essential one in cases of infection. Several routes are being taken to attempt to regulate the production of the INF α alarm molecule that precedes activation of the killer T-cells, for example, by specifically targeting certain activation routes for the pDC dendritic cells.

"These therapeutic approaches are currently being tested in other autoimmune diseases such as lupus and psoriasis. Such innovative treatments could be useful in the prevention of <u>Type 1 diabetes</u>. It will first be



necessary, however, to perform studies in diabetic and pre-diabetic patients to be able to better understand how the innate <u>immune cells</u> function, something that has not been studied until recently in autoimmune diabetes," concludes Agnès Lehuen.

More information: Crosstalk between neutrophils, B-1a cells and plasmacytoid dendritic cells initiates autoimmune diabetes. Julien Diana et al., *Nature Medicine*.

Provided by Institut National de la Sante et de la Recherche Medicale

Citation: The role of the innate immune cells in the development of type 1 diabetes (2012, December 19) retrieved 5 May 2024 from <u>https://medicalxpress.com/news/2012-12-role-innate-immune-cells-diabetes.html</u>

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