**AD NUMBER**

AD836618

**NEW LIMITATION CHANGE**

**TO**

Approved for public release, distribution unlimited

**FROM**

Distribution authorized to U.S. Gov’t. agencies and their contractors; Administrative/Operational Use; 08 JUN 1964. Other requests shall be referred to Department of The Army, Fort Detrick, Attn: Technical Release Branch/ TID, Frederick, MD 21701.

**AUTHORITY**

AMXFD ltr, 9 Feb 1972

THIS PAGE IS UNCLASSIFIED
DDC AVAILABILITY NOTICE

Reproduction of this publication in whole or in part is prohibited. However, DDC is authorized to reproduce the publication for United States Government purposes.

STATEMENT #2 UNCLASSIFIED

This document is subject to special export controls and each transmittal to foreign governments or foreign nationals may be made only with prior approval of Dept. of Army, Fort Detrick, ATTN: Technical Release Branch/TID, Frederick, Maryland 21701

DEPARTMENT OF THE ARMY
Fort Detrick
Frederick, Maryland
USE OF IMMUNO-FLUORESCENCE FOR THE DETECTION OF
THE SENSITIZATION OF MEGAKARYOCYTES

[Following is a translation of an article
by J. Duheille and H. Herbeuval in the
French-language journal Comptes Rendus
Soc. Biol., (Proceedings of the Society
of Biologists), Vol 157, Paris, 1963,
pages 150-154.]

The presence of anti-thrombocyte antibodies has been
demonstrated in the case of various diseases [Note: J. Dausset,
Clinical and Biological Immuno-hematology (Paris: Flammarion
1956); P. Miescher and K. O. Vorlander, Clinical and Experi-
mental Immuno-hematology (Paris: Flammarion, 1959).], and
it would be useful to find out whether this immunological
process affects only the platelets or whether its action al-
so extends to the megakaryocytes. Such a study would help
us to find out whether the thrombopenias observed in these
diseases are due only to the peripheral destruction of the
sensitized platelets or whether the production of these plate-
lets is upset from its very outset.

The use of immuno-fluorescent techniques has just
brought us the first inklings of a solution to this problem.
It is now possible, thanks to the work of Coons ([Note:] A.
Vol 102, page 49), to objectivize, with the help of a fluo-
rescent immune serum against human globulins, the fixing on
a determined cell of autoantibodies, or at least substances
having antigenic properties identical to those of human serum
globulins. This is—the method, following McKenna and Pisci-
otta ([Note:] Blood, 1962, Vol 19, page 654.), that we
have applied to the study of megakaryocyte platelet sensitiza-
tion.

-1-
Material and Methods

We examined four patients suffering from purpuras of different origins ([Note:] We thank Mr. Neimann, who very kindly allowed us to examine the patients at the Children's Medical Clinic in Nancy), which could be divided into two acute infantile purpuras, of which one showed serous anti-platelet antibodies, one purpura accompanying a panocytopenia, and one non-thrombopenic purpura.

Medullary fluid obtained by sternal puncture from each patient was spread in thin smears on microscopic slides, either immediately or else after collection on a non-formaldehyde preserving fluid and triple washing, as in our technique for the study of leucocyte sensitization ([Note:] J. Duheille, H. Herbeuval, F. Bellut and Y. Badonnel, C. R. Soc. Biol., 1962, Vol 154, page 2093).

The dried smears were fixed for one minute in a bath of ethyl alcohol at a temperature of 95 degrees, then rehydrated in three baths of packed physiological serum (ClNa 0.14 M., phosphates of Na 0.01 M, pH = 7.2), over a period of fifteen minutes.

They were then covered with fluorescent serum against human globulins (immune serum from a goat, joined to fluorescein iacthiocyanate, after a modified Riggs technique [see last note] and adsorbed on powdered organs immediately before use). The antigenic-antibody reaction is allowed to continue in a damp enclosure at laboratory temperature over a period of thirty minutes, then the preparations are cleansed of any excess of fluorescent antiserum by four baths of packed physiological solution, each lasting five minutes. This length of bath represents the minimum, and in one case lengthening it to twenty-four hours noticeably improved the contrast of the preparations by eliminating a non-specific fluorescence due to traces of unlinked fluorescein. The preparations were mounted in packed physiological serum under sealed glass covers, and examined under the microscope both in ultraviolet light and in phase contrast (Wild apparatus). They lasted hardly twenty-four hours and therefore it is valuable to record photographically the results obtained.

Results

1. Reading the preparations. On the smears of unwashed marrow the interstices separating the cells were filled with dried plasma, fixing the marked immune serum and brilliantly outlining the cell contours. The preliminary washing of the marrow suppressed this phenomenon and the background
of the preparation appeared uniformly dark. In both cases most of the elements that appeared did not fix the marked immune serum, and were distinguished only by a weak auto-fluorescence. Some cells stood out on this dark background, identifiable by phase contrast either as plasmocytes or as eosinophilous granulocytes since the fluorescent serum still contained, in spite of adsorption on powdered organs, traces of unlinked fluorescein.

When the megakaryocytes are sensitized the fluorescent immune serum attaches itself to the cytoplasm, to which it gives an intense luminosity, clearly differentiated from the scarcely observable luminosity of the other medullary elements. The nucleus, on the other hand, is only slightly fluorescent and its outline shows up in negative on the brilliant cytoplasmic background (figure 1). It is easy to identify these cells by passing from ultraviolet lighting to phase contrast: The important dimensions, the granular appearance of the cytoplasm, the multilobed contours of the nucleus, all signal the fact that they belong to the megakaryocyte series (figure 2). The youngest units in the megakaryocyte mass, corresponding to little promegakaryocyte or to megakaryoblasts, usually escape sensitization as far as one can determine from figures 1 and 2.

In negative preparations, as in the marrow of healthy subjects, the megakaryocytes show only a weak fluorescence, scarcely more distinct than that of the other cells and not to be compared with that of the sensitized megakaryocytes. It is possible that an improvement in the immunological specificity of the marked anti-serum would get rid of this slight residual fluorescence.

2. Clinical correlations. Out of the four medullary samples we examined, two (coming from two cases of acute infantile thrombopenic purpura) presented a very clear megakaryocyte sensitization. One of these patients, with a severe attack, had received many blood transfusions in the three months preceding the examination; the presence of immune anti-megakaryocytio-platelet antibodies cannot, therefore, be ruled out. But such antibodies, likely to sensitize heterologous megakaryocytes, would on the other hand (according to all probability) have no effect on those of the patient himself. We are therefore led to think that the observed sensitization is really the work of auto-antibodies. The second patient, whose serum contained anti-platelet antibodies discernable by ordinary techniques, had also developed antibodies directed against his own megakaryocyte line.

The two negative samples corresponded one to a purpura secondary to a pancytopenia and the other to a purpura where no examination could reveal the participation of the megakaryocyte-line.
Discussion and Conclusion

Our results differ on several points from those published by McKenna and Pisciotta: while these authors observed megakaryocyte sensitization only in cases of chronic idiopathic thrombocytic purpura, our two positive cases involved children suffering from acute thrombocytic purpura. It is possible that, in these two cases, it was a matter of an initial episode announcing the onset of a chronic purpura, although the isolated nature of this attack and its development to total recovery hardly plead in favor of such a hypothesis. It is also possible that the different nature of the serum against human globulins that was used was responsible, in part, for this disparity: Our anti-serum came from hyper-immunized goats while that of McKenna and Pisciotta was a horse immune serum. A rabbit immune serum gave these authors no results. It could be that the goat immune serum usually shows up a globulin other than the 2-globulin revealed by the horse immune serum and characteristic of chronic idiopathic thrombocytic purpura. The nature of the globulin shown up by our study therefore merits further study. We can assume, however, that fibrin is not concerned since the positiveness of the reaction persists even after collection of the marrow on anticoagulant and multiple washings.

All the same, it is not possible, given these results only, to attribute a pathogenic role to the anti-megakaryocyte auto-antibodies thus revealed: they could be only the secondary signs of underlying processes, for, in one of our cases, we have seen them persist in spite of the complete and perfect clinical recovery of the patient; they could in this case have the value only of an immunological scar.

Although the limited number of cases studied in this preliminary work does not yet allow us to appreciate fully the value of this new method of detecting megakaryocyte sensitizations, the first results obtained let us nevertheless see numerous possibilities offered by this technique in the field of immuno-hematology ([Note:] Work carried out with the help of the National Institute of Health and the National Social Security Fund).

(Laboratory of Medical Pathology [Director: M. R. Herbeuval], Faculty of Medicine, Nancy.)
Figure 1. Thrombopenic purpura. Sternal bone marrow, unwashed, colored by a fluorescent immune serum against human globulins. In the center, two sensitized megakaryocytes, strongly fluorescent, and a nonfluorescent megakaryoblast. Plasmatic fluorescence in the interstices separating the other cells.

Figure 2. The same field in phase contrast.

- END -

-5-