Amniotic fluid macrophages and the antenatal diagnosis of anencephaly and spina bifida

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Summary. The macrophage content of amniotic fluid has been measured and the upper limit of normal on an arbitrary scale is 41. Amongst 65 amniotic fluids collected for antenatal diagnostic studies before 22 weeks’ gestation there were eight which had macrophage counts ranging from 82 to 6226, three of these were shown to have anencephaly and two spina bifida. The reasons for three apparently false positives are as yet undetermined. Rhesus iso-immunized amniotic fluids were found to have macrophage counts of up to 276 and a possible explanation for this is considered. It is argued that an elevated amniotic fluid macrophage count may indicate a CNS defect or possibly other fetal abnormality.

The finding by Sutherland, Brock, and Scrimgeour (1973) of large numbers of macrophages in the amniotic fluid from two cases of anencephaly has been used by Nelson, Ruttiman, and Brock (1974) to make the antenatal diagnosis of a further case. The macrophage content of spina bifida and anencephalic amniotic fluids has been quantitated and compared with that from normal pregnancies.

Materials and methods

Amniotic fluids were obtained by amniocentesis for antenatal chromosome studies, α-fetoprotein estimation, during the management of Rhesus iso-immunized pregnancies and at induction of labour in cases where the fetus was known to have anencephaly.

Amniotic fluid cultures were established as previously described (Sutherland, Grace, and Bain, 1973). The culture vessel was a 50 mm Petri dish which contained five or six 6 x 22 mm glass coverslips. The culture medium was Ham’s F10 supplemented with 30% fetal calf serum. The day after cultures had been set up (usually about 20 hours later) one of the coverslips was aseptically removed from the Petri dish, washed well in phosphate buffered saline, fixed with methanol, stained with Giemsa, and mounted on a slide. The number of glass adherent cells in 10 low power fields (x 100) were counted and this number was adjusted to relate to 10 ml of amniotic fluid, ie, if the cells from 5 ml of amniotic fluid had been inoculated into the Petri dish then the number was multiplied by two. The resultant macrophage count (glass adherent cells/10 low power fields/10 ml of amniotic fluid) has been used to compare different samples of amniotic fluid. Amniotic fluid α-fetoprotein was measured according to Brock and Sutcliffe (1972).

Results

The macrophage counts for the 94 fluids studied are shown in Fig. 1. All the fluids classed as ‘normal’ had low macrophage counts (range 0–41), and there was no reason to suspect fetal CNS abnormality either because the infant had been delivered or because the amniotic fluid α-fetoprotein levels were not elevated. Four amniotic fluids from cases of hydramnios of unknown aetiology had macrophage counts within the normal range. A number of samples of amniotic fluid showing different degrees of blood staining had normal macrophage counts. Rhesus iso-immunized amniotic fluids had highly variable macrophage counts ranging up to 276. Where serial samples were obtained from these pregnancies the macrophage counts tended to decrease as gestation progressed.

Details of the cases with anencephaly or spina bifida are shown in Table I. In all except one of these fluids the macrophage count was elevated,
levels. These pregnancies were allowed to continue and the one which had the highest macrophage count went into premature labour at 33 weeks' gestation and resulted in a normal male infant. The placenta showed multiple recurrent infarctions, there was a small succenturiate lobe and the cord had many false knots. The two other similar pregnancies are continuing.*

The possibility that interfering with the amniotic fluid cultures on the day after they have been set up may affect their success was tested by comparing the times required to achieve cytogenetic results from the first 25 normal cultures on which macrophage counts were performed (mean 13.0 days) with the 20 similar cultures studied before macrophage counts were being done routinely (14.5 days). In all cases where a cytogenetic result was required after macrophage counts had been performed this was achieved. It was not possible to differentiate between amniotic fluids with normal and raised macrophage counts (except in those cases where the macrophage count was grossly elevated) by direct inspection of the cultures with the inverted microscope. Quantitation of stained material was necessary for this purpose and its preparation did not prejudice the success of the cultures in any way.

**Discussion**

The antenatal diagnosis of anencephaly and spina bifida can be achieved with a high degree of success using amniotic fluid α-fetoprotein measurements (Lancet, 1974). Ideally all samples of amniotic fluid, collected for any reason before mid-pregnancy, should have α-fetoprotein estimations carried out. The two cases of spina bifida in this report were diagnosed from amniotic fluids collected primarily for antenatal chromosome studies.

The nature and origin of the glass-adherent cells in the amniotic fluid must be considered before their clinical relevance is discussed. All the glass-adherent cells in amniotic fluid are probably not macrophages but erythrophagocytic properties

* These subsequently led to the birth of full-term normal infants.
There have been claims that the macrophages in the amniotic fluid are a reliable indicator of neural tube defects. However, more recent studies have suggested that macrophages may be present in normal amniotic fluid and that their presence does not necessarily indicate the absence of neural tube defects. This has led to the suggestion that the presence of macrophages in amniotic fluid should be considered as a marker of potential problems, rather than a definitive diagnosis.

Further studies of the macrophages in amniotic fluid are required to assess fully the usefulness of this parameter in antenatal diagnosis. The false positives, which suggest a potential problem, are of particular concern, as the diagnosis of neural tube defects is often made on the basis of other markers, such as alpha-fetoprotein levels.

REFERENCES


