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Topographic analysis and translation of human autologous monocytes: the imprinting of a distinct autologous monocyte subpopulation in the immune system. To analyze the topography, origin and fine localization of human autologous monocytes, we separated synovial fluid monocytes from eight rheumatoid arthritis (RA) patients by means of density sedimentation, and analyzed them by means of fluorescent microscopy and a new computerized image analysis method. A distinct subpopulation of cells, termed autologous monocytes, could be identified on the basis of their distinctive cell morphology. Positive staining of the nucleus was restricted to the marginal position of the cell, whereas the cytoplasm was bound by a perinuclear halo. These cells were visualized by fluorescent microscopy in a characteristic pattern in the synovial lining. The cell density within the joint revealed a peak in the RA synovium, which was significantly lower in the cellular fraction (p less than 0.001). No autologous monocyte could be detected in synovial fluid supernatant. Topographic analysis of monocytes in the synovial lining revealed considerable differences between individual patients. When the expression of various antig

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