Table 1: Immune Cells and Cytokines in Synovium and Lymph Nodes

Table 1: Immune Cells and Cytokines in Synovium and Lymph Nodes Immune Cells Cytokines Released Cytokines Acting On				
	- Cytominos Holoacca	Them and Their		
		Functions		
MLS	TNF-α, IL-1β, IL-6,	TNF-α: Activates MLS		
WILO	TGF-β, IL-8, CCL-2	to adapt M1		
	β, 12 3, 332 2	phenotype and		
		promotes release of		
		TNF-α, IL-1β, IL-6. IL-		
		6: Sustains survival		
		and enhances		
		cytokine release. IL-		
		1β: Activates MLS to		
		release TNF-α and IL-		
		6. IL-17: Enhances		
		pro-inflammatory		
		cytokine production		
		$(TNF-\alpha, IL-1\beta, IL-6).$		
		IFN-y: Activates MLS		
		to M1 phenotype and		
		enhances MHC II		
		expression. GM-CSF:		
		Promotes survival and		
		expansion. TGF-β:		
		Shifts MLS to M2		
		phenotype. IL-10:		
		Suppresses M1		
		phenotype and		
		promotes tissue		
		repair.		
Recruited	TNF-α, IL-1β, IL-6, IL-	TNF-α: Activates M1		
Macrophages	12, IL-23, IL-8, CCL2,	phenotype and		
Madrophagoo	RANKL	induces pro-		
		inflammatory		
		cytokines (TNF-a, IL-		
		1β, IL-6). IL-6:		
		Sustains survival and		
		cytokine release. IL-		
		1β: Promotes		
		differentiation of		
		osteoclast precursors		
		and release of TNF-α,		
		IL-6. IL-17: Synergizes		

		to enhance TNF-α, IL-1β, IL-6 production. IFN-γ: Activates M1 phenotype and enhances MHC II expression. GM-CSF: Promotes survival and expansion. IL-10: Suppresses M1 phenotype and promotes tissue repair. TGF-β: Shifts macrophages to M2 phenotype. RANKL: Induces osteoclast differentiation with M-CSF. M-CSF: Essential for macrophage differentiation and osteoclast precursor response to RANKL.
Treg Cells	TGF-β	TGF-β: Promotes Treg cell differentiation and proliferation in low IL-6, IL-1β conditions. IL-6, TNF-α: Decrease Treg differentiation.
Osteoclast Cells		TNF-α: Amplifies osteoclast differentiation and prolongs survival. IL-1β: Potentiates RANKL/RANK signaling for differentiation. RANKL: Initiates differentiation from precursors. M-CSF: Prepares precursors for RANKL signaling and promotes survival and proliferation.

FLS (Fibroblast-like Synoviocytes)	IL-6, IL-8, IL-1β, CXCL12, CXCL9, CXCL10, CXCL11, VEGF, GM-CSF, M- CSF, RANKL	TNF-α: Promotes proliferation. IL-1β: Triggers MMP production, proliferation, and migration. IL-6: Enhances survival and resistance to apoptosis. IL-17:
		Synergizes with TNF-α for proliferation. TGF-β: Promotes differentiation from progenitor cells.
Th17 Cells	IL-17, IL-22, GM-CSF, TNF-α, IFN-γ	TGF-β: Critical for differentiation in presence of IL-6 or IL-1β. IL-6, TGF-β: Required for production. IL-23: Promotes survival and proliferation. IL-1β: Enhances pathogenicity and survival. IL-12: Converts to Th1-like phenotype. IFN-γ: Promotes plasticity and transition to Th1-like phenotype.

Table 2: Cytokines and Cells That Release Them

Cytokine/Chemokine	Cells That Release It
TNF-α	MLS, Recruited Macrophages,
	Th17 Cells
IL-1β	MLS, Recruited Macrophages, FLS
IL-6	MLS, Recruited Macrophages, FLS
TGF-β	MLS, Treg Cells
IL-8	MLS, Recruited Macrophages, FLS
CCL-2	MLS, Recruited Macrophages
IL-12	Recruited Macrophages
IL-23	Recruited Macrophages
RANKL	Recruited Macrophages, FLS,
	Osteoblasts, Synovial Fibroblasts,
	T Cells, B Cells

GM-CSF	MLS, FLS, Th17 Cells
M-CSF	FLS
IL-17	Th17 Cells
IL-22	Th17 Cells
IFN-γ	Th17 Cells
CXCL12	FLS
CXCL9	FLS, Th17 Cells
CXCL10	FLS, Th17 Cells
CXCL11	FLS, Th17 Cells
VEGF	FLS
CCL20	Th17 Cells

This tabular arrangement consolidates your collected data, categorizing cytokines/chemokines with the immune cells that release them and interact with them. Functions for each cytokine have been specified to aid in differential equation modeling.