

Supplemental Materials

Supplemental Table: Selected completed birth cohorts – relatives and general population now completed or in follow up*							
Program: Location	Population screened	Location	Number screened	Screening material	Screening assays	Rates of positive screens	Comments
BABYDIAB 1989-2000 (1)	Newborn children of those with T1D	Germany	2364	Cord blood then venous blood	ICA and RBA: IAA, GADA, IA-2A, ZnT8A and TTG AA	<ul style="list-style-type: none"> • AA+: 220(9%) • ≥2 AA+: 123(5%) 	<ul style="list-style-type: none"> • AA screening in cord blood, at 9 months and 2, 5, and 8 years • From 3 yrs, yearly oral glucose tolerance test monitoring if AA+
DAISY 1993-2004	Newborn general population (GP) and relatives <4 yrs	Colorado, US	Newborns : 32,114	Cord blood for HLA and serum for AA	<ul style="list-style-type: none"> • RBA and ECL: IAA, GADA, IA-2A, ZnT8A, tTGA 	<ul style="list-style-type: none"> • 1,424 GP newborns and 1,123 relatives identified and followed • AA+: 8% • ≥2 AA+: 5% 	<ul style="list-style-type: none"> • Genetically at-risk newborns based on HLA genotyping and relatives followed at 9, 15, 24 months and annually thereafter until age 20 y • AA+ followed q3-6 mo until 30 y
DEW-IT (2) 1995-2001 2010-2012	GP newborn blood spots	Washington, US	42000 blood spots tested (3)	Dried new born screening blood spots, then serum for AA	HLA genotyping; RBA: IAA, GADA, IA-2A, and later, ZnT8A	<ul style="list-style-type: none"> • 14.2% of children eligible for AA surveillance(3) • 3748 followed over time (3) • AA+: 173 (5%) • ≥2 AA+: 170 (5%) 	Cover letter and consent form mailed to Washington families. Consenting families received HLA genotyping of dried newborn blood spots followed by AA monitoring of at-risk individuals.
DiPiS (4) 2000-2004	GP newborns	Sweden	35688	Cord blood for HLA, blood spots for GADA and IA2A, serum for IAA and ZnT8	HLA genotyping; RBA: IAA, GADA, IA-2A, ZnT8A	<ul style="list-style-type: none"> • 7826 positive screens(3) • 4359 followed over time • AA+: 184 (4%) • ≥2 AA +: 100 (2%) 	<ul style="list-style-type: none"> • Children identified for surveillance based on risk score that include HLA genotype and environmental, demographic, and historical risk factors. • Positive screens with yearly follow up. Those with ≥2 AA+ followed every 3 months.
TEDDY (5) 2004-2010	Newborns in both relatives and GP	Clinical centers in US Finland, Germany, Sweden	424,788	Capillary blood spots	HLA genotyping; RBA: IAA, GADA, IA-2A, tTGA	21589 (0.05%) of screens with high-risk HLA; 8676 parents consented to follow up.	<ul style="list-style-type: none"> • High-risk newborns followed every 3-6 months for 15 yrs for AAs and T1D, with documentation of potential environmental contributors. • 90% without a known relative with T1D

Recent follow up data obtained from recent published references (3; 5) and personal communications (M Rewers)

GP=general population; AA- autoantibody; Study Acronyms: DAISY: Diabetes Autoimmunity Study in the Young; DEWIT: Diabetes Evaluation in Washington Study; DiPiS: Diabetes Prediction in Skane; TEDDY: The Environmental Determinants of Diabetes in Youth

References

1. Ziegler AG, Hummel M, Schenker M, Bonifacio E: Autoantibody appearance and risk for development of childhood diabetes in offspring of parents with type 1 diabetes: the 2-year analysis of the German BABYDIAB Study. *Diabetes* 1999;48:460-468
2. Wion E, Brantley M, Stevens J, Gallinger S, Peng H, Glass M, Hagopian W: Population-wide infant screening for HLA-based type 1 diabetes risk via dried blood spots from the public health infrastructure. *Ann N Y Acad Sci* 2003;1005:400-403
3. Anand V, Li Y, Liu B, Ghalwash M, Koski E, Ng K, Dunne JL, Jonsson J, Winkler C, Knip M, Toppari J, Ilonen J, Killian MB, Frohnert BI, Lundgren M, Ziegler AG, Hagopian W, Veijola R, Rewers M, Group TDS: Islet Autoimmunity and HLA Markers of Presymptomatic and Clinical Type 1 Diabetes: Joint Analyses of Prospective Cohort Studies in Finland, Germany, Sweden, and the U.S. *Diabetes Care* 2021;
4. Elding Larsson H: A Swedish approach to the prevention of type 1 diabetes. *Pediatric diabetes* 2016;17 Suppl 22:73-77
5. Bonifacio E, Weiss A, Winkler C, Hippich M, Rewers MJ, Toppari J, Lernmark A, She JX, Hagopian WA, Krischer JP, Vehik K, Schatz DA, Akolkar B, Ziegler AG, Group TS: An Age-Related Exponential Decline in the Risk of Multiple Islet Autoantibody Seroconversion During Childhood. *Diabetes Care* 2021;