

Results for assessment 53 53 53 DQA1*02:01,*04:01; DQB1*02:02,*04:02 44 4

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La 1	b Results for assessment 1 HLA DQ2 positive	Alleles of interest DQB1*02, DQB1*03:02	Interpretative comments The presence of HLA-DQ2 is associated with, but not diagnostic for, coeliac disease. HLA-DQ2 is present in about 21% of caucasians in the normal population	Comments	Received 09/04/2019	Tested 16/04/2019	Assessment Acceptable
1	2 DQ2: Positive, DQ8: Negative, DQA1*05: Negative	DQ2, DQ8, DQA1*05	This patient is negative for both the DQA1*05/DB1*02 heterodimer and DOR(DGB*103.02) which are present in over 95% patients with cooliac disease. HOR(DGB*103.02) which are present in over 95% patients with cooliac disease. HOR(DGB*103.02) which cooperates one half of the DGA1*05/DGB*102 horizon patients are patients. The DGA1*05/DGB*103.02 horizon patients are patients and patients are patients and patients are patients and patients are patients. This genotype has been associated with genetic susceptibility for coeliac diseases.		10/04/2019	12/04/2019	Acceptable
1	5 Not Tested				0000-00-00	0000-00-00	Not assessed
1	7 DOA1*0501 DOB1*02:01 (os) - DO2 Negative DOA1*05:05 02:01 DoB1*03:01 02:02 (time) - DO2 Negative DOA1*03:05 02:01 DOB1*03:02 - DO8 Negative	DOA119561 DOB119201 (de) DOA119568 0201 DB0193301 02:02 (trans) DOA119369 0201 DB0193302	The major association for Coeliac disease involves the haplotype: DOA1*05.01 #6 DOB1*02:01 (DO2) and a minority of cases with the haplotype: DOA1*03.01 #6* DOB1*03.02 (DO8), (Mature Reviews Immunology 2002.2:647) This patient is NEGATIVE for the DOA1*05.01-DOB1*02.01 (DO2) haplotype and DOA1*03-DB21*03.02 (DO8) haplotype. This patient has no genetic risk of having or developing coeliac disease.		11/04/2019	12/04/2019	Unacceptable
2	4 DQB1*02:02/06/10/11/12, *04:02/20/27/31; DQA1*02:01, *04:01/03N/04	DQB1*02, DQA1*05, DQB1*03:02	The patient possesses only one allele of the Coeliac Disease associated HLA-DQ2 molecule: DQB1*02 (DQA1*05 is not present). Patients with this genotype have a low risk of predisposition to Coeliac Disease. Other clinical indications are required for		10/04/2019	15/04/2019	Acceptable
2	5 DQB1*02:02; *04:02; DQA1*02:01; *04:01	DQ2	diagnosis. This patient is DQ2.2 positive, heterozygous. This patient is DQ2 positive which is		10/04/2019	18/04/2019	Acceptable
3	8 DQB1*02:02,*04:02 DQA1*02:01,*04:01	DQ8 DQB1*02 and DQB1*03:02	associated with Coeliacs Disease. This individual carries the DQB1*02:02 (DQ2) variant that has a weak association with		10/04/2019	18/04/2019	Acceptable
4	2 DQA1*02:01 DQA1*04:01/04:03N/04:04 DQB1*02:02/08/10/11/12/26/50/62/64/65/80/84/89/95/97/103/104	HLA-DQ	coeliac disease (Low risk). This patient is Heterozygous POSITIVE for HLA-DQ2 (but is DQA1*05 NEGATIVE) and NEGATIVE for HLA-DQ8 (DQB1*03:02). Patients with this genotype have a LOW RISK	1	11/04/2019	12/04/2019	Acceptable
	DQB1*04:2020-03404;104:3304;3505-0340;39043999997/103104 DQB1*04:02040-03404;104:3304:1804:3904:2404:25004:2604:29104:320- 3304:34/04:35/04:36N/04:37/04:39/04:40/04:41N/04:43/04:44/04:45/04:46N/04:47/04	4: :4	of predisposition to Coeliac disease				
7	8 Not tested				10/04/2019	0000-00-00	Not assessed
8	5 DQA1*05 negative DQB1*02 positive	DQA1*05 DQB1*02	This individual does not have the HLA-DQ variants associated with coeliac disease. This assay		10/04/2019	23/04/2019	Acceptable
	DQB1*0302 negative	DQB1*0302	tests for the presence of HLA-DQ2 (DQA1*05/DQB1*02) and HLA-DQ8 (DQB1*0302) which				
			are found in more than 97% of patients with coeliac disease. However, 2-3% patients with coeliac disease have a rare genotype that is not detected by this assay.				
8	6 Coeliac disease-associated HLA alleles present: DQB1*02:02 DQA1*02:01 HLADQ2 PRESENT - HLA-DQ2 2 HLADQ8 ABSENT	DQ2 and DQ8 associated DQ81* and DQA1*	HLA-DQ.2, which is associated with low genefic susceptibility for coeliac disease (CD), has been detected in heteroogous form in this patient. As 25-30% of the general population has one of the CD-associated HLA alleles encoding DQ2 and/or DQ8 and only 3% of these individuals develop coeliac disease, identification of a CD-associated HLA allele is not diagnostic of CD. The presence of DQ2 and/or DQ8 increases the listendor dark the patient has CD but a diagnosis must be based on clinical findings, serum antibody detection tests and/or intestinal biopsy.		10/04/2019	23/04/2019	Acceptable
8	7 positive hetrozygous for HLA-DQ2.2 and I² subunit HLA-DQ2.2/DQ2.5, rest negative	HLA-DQ2.2, HLA-DQ 2.5, HLA-DQ8, Beta subunit HLA-DQ2.2/DQ2.5	90-95% of Coeliac patients are HLA DQ2 or DQ8 positive (Husby S, et al. European Society for Pediatric Gastroenterology, Hepatology, and Nutrition guidelines for the diagnosis of coeliac disease. J Pediatr Gastroenterol Nutr 2012; 54:13660).		10/04/2019	24/04/2019	Acceptable
10	9 DQA1*05: negative DQB1*02:01 / DQB1*02:02: positive DQB1*03:02 (DQ8): negative	DQA1*05 DQB1*02:01 / DQB1*02:02 DQB1*03:02 (DQ8)	There is no associated risk for coeliac condition		10/04/2019	16/04/2019	Acceptable
11	3 Haemolysis of the sample. No result available	DQ2 and DQ8		Haemolysis of the sample. DNA concentration too low and bad quality. No result	09/04/2019	0000-00-00	Not assessed
12	3 Not Tested			available	0000-00-00	0000-00-00	Not assessed
12	4 Not Tested				0000-00-00	0000-00-00	Not assessed
12	6 DQA*05=NEG, DQB*02=POS, DQA*02=POS, DQA*03=NEG, DQB*0302=NEG	DQA*05, DQB*02, DQA*02, DQA*03, DQB*0302			10/04/2019	15/04/2019	Acceptable
12	7 HLA-DQ2* positive, HLA-DQ8* negative	HLA-DQA1* HIA-DQB1*			11/04/2019	16/04/2019	Acceptable
14	9 DQB1*02 positive; DQB1*03:02 negative; DQA1*05 negative 2 HLA-DQA1*05 absent HLA-DQB1*02 present HLA-DQB1*03:02 (DQB) absent 0 DQB1*02:02:04.02 DQA*02:01:04:01	DQB192: 03:02; DQA1*05 HLA-DQA1*05, HLA-DQB1*02 and HLA-DQB1*03:02 (DQ8)	Slightly increased risk for coellac disease PrÄdsence de Iåt "malfa le HLA-DQB1*02 (DQ2) mais absence des allÄ-les HLA- DQA1*05 et DQB1*03:02 (DQB.) Risque faible de prÄddisposition Ä la maladie ch*liaque. Absence of allelles DQB1*02:01-DQA*05:01 and DQB1*03:02-DQA*03:01		11/04/2019	29/04/2019 22/04/2019 18/04/2019	Acceptable
	4 HLA-DQA1*02:01, *04:01; HLA-DQB1*02:02,*04:02	DQ8 : DQB1*03:02-DQA*03:01 HLA-DQA1*, and HLA-DQB1* are typed to the 4-digit level to determine whether	haplotype DQA1*05:01. DQB1*02:01 : absence		10/04/2019		
	11275 G. 10, 10, 10, 10, 10, 10, 10, 10, 10, 10,	HLA-DQ2 is coded by DQA1*05:01, DQB1*02:01; HLA-DQ2 is coded by DQA1*05:05, DQB1*03:01 and DQA1*02:01, DQB1*02:02; HLA-DQ8 is coded by DQA1*03:01, DQB1*03:02	haplotypes DQA1*03:05, DQB1*03:01 and DQA1*02:01, DQB1*02:02 : absence haplotype DQA1*03:01, DQB1*03:02 : absence		1010412010	1110412010	/ toopiable
			The patient has HLA-D02 encoded by an haplotype not listed to be most at risk of ceilar disease. >95% of ceilar disease patients express HA-D02 encoded by D0A1*05.01, D0B1*02.01 and D0A1*02.01, D0B1*02.02. 5% of ceilar disease patients express HLA-D08 encoded by D0A1*03.01, D0B1*03.01, D0B1*03.02. HA-D02 or D08 are expressed in 39-40% of the Caucasian population. HA-D02 or D08 are expressed in 39-40% of the diagnosis of ceilar disease.				



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159 DQA1*03 negative, DQA1*05 negative, DQB1*02 positive and DQB1*03:02 negative		Absence of susceptibility phenotype for coeliac disease	11/04/2019		
173 DOA1*05NEGATIVE	HLA-DQA1*05, HLA-DQA1*03, HLA-DQB1*02, HLA-DQB1*03:02, HLA-DQB1*03:03	Low risk of cellac disease	17/04/2019	23/04/2019	Acceptable
176 DQA1*05-Pos DQB1*02-Pos DQB1*03:02-Neg	DQA1*05 DQB1*02 DQB1*03:02	Celiac tissue type examination: Positive for HLA-DQB1 * 02 (DQ2), HLA-DQA1 * 05 and negative for HLA-DQB1 * 03: 02 (DQ8). The genetic risk of Celiac disease is cresent	15/04/2019	16/04/2019	Unacceptable
201 DQA1*02:01 DQA1*04:01 DQB1*02:02 DQB1*02:02	DQB1 03.02	ревент.	10/04/2019	17/04/2019	Acceptable
219 DQB1*03:02: negative DQA1*05: negative DQB1*02: positive DQA1*02: positive	HLA-DQB1*03:02, HLA-DQA1*05, HLA-DQB1*02, HLA-DQA1*02	English translation: "HLA-DO2 is lesteded in the form of HLA-DO81*02 and HLA-DOA1*02. A small minority of coeliac patients has these alleles. The alleles are common in the general population. Coeliac disease is unlikely. However, the test dance can not exclude coeliac	24/04/2019	30/04/2019	Acceptable
Reported serotype: DQ2.2 223 DQA1*02 positive, DQA1*03 negative, DQA1*05 negative, DQB1*02 positive, DQB1*03:02 negative	DQA1*02, DQA1*03, DQA1*05, DQB1*02, DQB1*03:02	disease."	10/04/2019	16/04/2019	Acceptable
224 DQA1*05=NEG, DQB1*02=POS; DQB1*03:02 group (DQ8)= NEG 225 DQ2-positive, DQ8-negative	DQA1*02, DQA1*03, DQA1*05 DQB1*02, DQB1*03:02	The patient is HLA-DQ2.2 positive. Coeliac disease is associated with this HLA-type in 5%.	10/04/2019 15/04/2019		Acceptable Acceptable
245 DQB1*02,*04; DQA1*02,*04	DQB1*02+DQA1*05 DQB1*02 DQB1*03:02	Presence of DQ2 antigen not associated with an alpha chain at risk: low to very low risk to develop coeliac disease. This result alone does not confirm the diagnosis.	10/04/2019	17/04/2019	Acceptable
255 DQB1*02 pos	DQA1*05, DQB1*02, DQB1*0302	DQ2.5 neg DQ8 neg	10/04/2019	23/04/2019	Acceptable
263 DQA1*0201 present, DQA1*03 absent, DQA1*05 absent DQB1*02 present (heterozigous), DQB1*0302 absent	HLA-DQA1*0201, DQA1*03, DQA1*05 HLA-DQB1*02, DQB1*0302 Homozygous or heterozygous status for DQB1*02 only	The HLA-OQB1 *02 allele is present in heterozygosity. This condition, while compatible with the presence of cellac disease, has not been shown to significantly increase the risk of disease compared to the general population.	18/04/2019	23/04/2019	Acceptable
269 positive	HLA DQ2.2 HLA DQ2.5 HLA DQ8	HLA DQ2 2 positive, heterozygot HLA DQ2 5 negative HLA DQ8 negative	07/04/2019	17/04/2019	Acceptable
274 HLADQ2trans, haplotype2 carrier (detected:DQA1*02, DQB1*02, DRB1*07)	HLADQ2cis, HLADQ2trans, HLA DQ2trans hp1, HLADQ2trans hp2, HLA DQ8		12/04/2019	15/04/2019	Acceptable
276 DQA1*05 negative, DQB1*02 positive, DQB1*03:02 negative	DQA1*05 DQB1*02		10/04/2019	23/04/2019	Acceptable
278 Positive for genotype HLA-DQ2.2	DQB1*03:02 DQA1*02, DQA1*02/*0301, DQA1*03, DQA1*0302/03, DQA1*05, DQB1*02, DQB1*0	2 The genotype indicates a risk of developing coeliac disease	15/04/2019	25/04/2019	Acceptable
279 Not Tested			0000-00-00	0000-00-00	Not assessed
281 Positive association with coeliac disease.	DQA1*		09/04/2019	16/04/2019	Acceptable
DQA1*02:01-DQB1*02:02 type. 307 DQA1*02:01,*04:01 DQB1*02,*04	DGB1* DGA1*02 DGA1*05 DGA1*05 DGA1*03 DGB1*02	Presence of just the beta chain of the DQ2 dimer (DQB1*02 positive - DQA1*05 negative).	10/04/2019	12/04/2019	Acceptable
315 POSITIVE (DQB1*02, DQB1*04) 317 Positive for allel: DQA1x02, DQA1x02/x0301, DQB1x02, DQB1x02/x0302	DQB1*03:02 DQB1*02, DQB1*03:02 HLA DQA1 and HLA DQB1	Positive for HLA DQ 2.2			Acceptable Unacceptable
319 DQ2 Neg DQ8 Neg	DQA1*05 Neg DQB1*02 Pos DQB1*0302 Neg		10/04/2019	12/04/2019	Unacceptable
331 DQB1*02:02, DQB1*04:02 333 DQB1*02, DQA1*02, BRB1*07	DQB1*02:01, DQB1*03:02 DQA1*05, DQA1*02, DGA1*03, DQB1*02, DQB1*0301, DQB1*0302, DRB1*03,	Absence of DQB1*02:01; absence of DQB1*03:02	12/04/2019 04/04/2019	24/04/2019 18/04/2019	Acceptable Acceptable
338 HLA-DQB1*02:02;DQA1*02:01;DRB1*07	DRB1*11, DRB1*12, DRB1*07, DRB1*04 HLA-DQB1; DQA1 ;DRB1*03,*04,*07,*11	Presence of DQ2.2 (DR7&CDQ2) haplotype in heterozygous state. This condition has been shown to confer a certain, but lower risk to Celiac Disease.	11/04/2019	28/05/2019	Acceptable
339 Found DQB102 positive, but a genetic predisposition for Celiac Diasease is unlikely	DQA105, DQB102 and DQB10302		10/04/2019	16/04/2019	Acceptable
346 HLA-DQA1*05-NEG, HLA-DQB1*02-POS, HLA-DQB1*03:02P (DQ8)-NEG 347 HLA-DQ2.5-negative, HLA-DQ2.2-positive, HLA-DQ8-negative	DQA1*05, DQB1*02, DQB1*03:02P (DQ8) HLA-DQA1 / HLA-DQB1	Increased risk for the development of Coeliac Disease; determination of serological parameters or biopsy from the small intestine recommended.			Acceptable Acceptable
355 HLA-DQ2.2 Positive		parameters or pulpsy from the small intestine recommended. The patient has a genetic disposition to develop celiac disease. Analyzing for celiac antibodies in plasma is recommended.	10/04/2019	30/04/2019	Acceptable
359 Alleles positive: DQA1*02, DQA1*01/*04/*06, DQA1*02/*03:01, DQB1*02, DQB1*04/*05, alfa-subunit HLA-DQ2.2, beta-subunit HLA-DQ2.2/DQ2.5	DQA1*02, DQA1*03, DQA1*05, DQA1*01/*04/*06, DQA1*02/*03:01, DQA1*03:02/03, DQB1*02, DQB1*03:02, DQB1*03/*06, DQB1*04/*05, alfa-subunitHLA-DQ2.2, alfa-subunitHLA-DQ2.5, alfa-subunitHLA-DQ2.2/DQ2.5, beta-subunitHLA-DQ2.2/DQ2.5, beta-subunitHLA-DQ2.5, beta-sub		11/04/2019	17/04/2019	Acceptable
363 HLA DQ2.2 = Present ; HLA DQ2.5 = Absent ; HLA DQ8 = Absent	subunit HLA-DQ8 DQA1*02, DQA1*0201, DQA1*03, DQA1*0302/03, DQA1*05, DQB1*02,		09/04/2019	16/04/2019	Acceptable
413 Presence of DQ2 beta chain (DQA1*02:01; DQB1*02:02; DRB1*07)	DQB1*02/*0302 DQ2, DQ8 based on the results of DQA1*05:01; DQA1*05:05; DQA1*02:01; DQA1*05	0;	10/04/2019	15/04/2019	Acceptable
1350 DQ2.2 (DQ2 trans haplotype (Hp2) carrier)	DQB1*02:02; DQB1*02:01; DQB1*03:02; DRB1*. detected alleles (allelic groups):	detected HLA genotype is associated with the rare risk of coellac disease	10/04/2019	16/04/2019	Acceptable
	HLA-DQ2 cis (D\Q2.5); DQA1*05-DQB1*02-DRB1*03 HLA-DQ2 trans (DQ2.5); DQA1*05-DQB1*03:01-DRB1*11/DRB1*12 DQA1*02-DQB1*02-DRB1*07 HLA-DQ2 trans haplotype (Hp1) carrier: DQA1*05-DQB1*03:01-DRB1*11/DRB1*12				
	HLA-DQ2 trans haplotype (Hp2) carrier (DQ2.2): DQA1*02-DQB1*02-DRB1*07 DQ8:DQA1*03-DQB1*03:02-DRB1*04				



NARCOLEPS otal distributed							
Total submitte	d 21						
Reference nber acceptabl							
er unacceptabl	e 1						
Assessed 5	<u>2</u>						
1	b Results for assessment 1 DQB1*06:02 Negative	Alleles of interest DQB1*06:02	Interpretative comments HLA-DQB1 allele known to be associated with Narcolepsy is not present	Comments	Received 09/04/2019	16/04/2019	
	2 5 Not Tested						Not assessed Not assessed
1	7 DQB1*06:02- Negative 4 DQB1*02:02/06/10/11/12, *04:02/20/27/31; DQA1*02:01, *04:01/03N/04	DQB1*06:02 allele DQB1*06:02	This patient is NEGATIVE for the narcolepsy associated allele DQB1*06:02. HLA-DQB1*06:02 is associated with narcolepsy-cataplexy. This patient is NEGATIVE for HLA-DQB1*06:02.		11/04/2019 10/04/2019	12/04/2019	Acceptable
	5 DQB*06:02 NEGATIVE 8 DQB1*02:02, *04:02	DQB*06:02 DQB1*06:02	The patient does not carry the associated HLA alleles which confer susceptibility to Narcolepsy.		10/04/2019 10/04/2019	18/04/2019 18/04/2019	
4	2 DQB1*02:02/06/10/11/12/26/50/62/64/65/80/84/89/95/97/103/104 DQB1*04:02/04:04/04:11/04:13/04:18/04:19/04:23/04:24/04:25N/04:26/04:29/04:32/04: 33/04:34/04:35/04:36N/04:37/04:39/04:40/04:41N/04:43/04:44/04:45/04:46N/04:47/04:4	HLA-DQB1*06:02	DQB1*06:02 NEGATIVE. Narcolepsy is associated with the expression of the human leukocyte antigen (HLA) class II molecule DQB1*06:02.		11/04/2019	12/04/2019	Acceptable
7 8 8 8	6				10/04/2019 10/04/2019	23/04/2019 23/04/2019	Not assessed Not assessed Not assessed Not assessed
10	9 DQA1*01:02: negative	DQA1*01:02 DQB1*06:02	There is no associated risk for narcolepsy condition		10/04/2019	16/04/2019	Acceptable
11	DQB1*06:02: negative 3 Haemolysis of the sample. No result available	DQB1*06:02		Haemolysis of the sample. DNA concentration too low and bad quality. No result available	09/04/2019	0000-00-00	Not assessed
12	3				0000-00-00	0000-00-00	Not assessed
12	4				0000-00-00	0000-00-00	Not assessed
12 12	7 HLA-DQB1*06:02 negative	HLA-DQB1*06:02			10/04/2019 11/04/2019		
12	9 DQB1*06:02 negative	DQB1*06:02	Risk for narcolepsy not increased		22/04/2019		
	2 HLA-DQB1*06:02 absent 0 DQB1*02;04.	HLA-DQB1*06:02 DQB1*06:02.	Absence of allele DQB1*06:02.		11/04/2019 17/04/2019		
15	4 HLA-DQB1*02:02,*04:02	HLA-DQB1*06:02	Allele DQB1*06:02 : absence		10/04/2019	17/04/2019	Acceptable
			The HLA-DQB1*06:02 is found in 15-25% of the overall population and in 90-100% of narcolepsy patients.				
15 17 17 20 21	6 1	DQ81*06.02, DQA1*01.02	Absence of susceptibility phenotype for narcolepsy		17/04/2019 15/04/2019 10/04/2019	23/04/2019 16/04/2019 17/04/2019	Not assessed Not assessed Not assessed Not assessed Not assessed
	3 DQA1*01:02 negative, DQB1*06:02 negative	DQA1*01:02, DQB1*06:02			10/04/2019	16/04/2019	Acceptable
	4 DQA1*01:02=NEG, DQB1*06:02=NEG 5 DQB1*06:02-negative	DQB1*06:02	The patient donÂ't have the HLA-type thatÂ's associated with narcolepsy.		10/04/2019 15/04/2019		
24	5 DQB1*02,*04	DQB1*08:02	Absence of the susceptibility allele for narcolepsy-cataplexy DQB1'06:02. This allele is present in 12 to 38' of the general population, in 40 to 60% of patients with narcolepsy without cataplexy and in 18% of patients with diopathic hypersonnia. This result makes the diagnosis of narcolepsy-cataplexy unlikely but does not exclude the diagnosis.		10/04/2019	17/04/2019	Acceptable
25					10/04/2019		
26 26							Not assessed Not assessed
27	4				12/04/2019	15/04/2019	Not assessed
27 27	6 DQB1*06:02 negative 8	DQB1*06:02			10/04/2019 15/04/2019		Acceptable Not assessed
27		DQB1*			0000-00-00		
30	1 No known association with narcolepsia. 7	DQB1*			09/04/2019 10/04/2019		Not assessed
31 31	5 NEGATIVE (DQB1*02, DQB1*04)	DQB1*06:02			10/04/2019 15/04/2019		
31	9				10/04/2019	12/04/2019	Not assessed
33 33	1 DQB1*02:01, DQB1*03:02	DQB*06:02	Absence of DQB1*06:02				Unacceptable Not assessed
33	8				11/04/2019	28/05/2019	Not assessed
33 34							Not assessed Not assessed
34	7				16/04/2019	17/04/2019	Not assessed
35 35							Not assessed Not assessed
36	3				09/04/2019	16/04/2019	Not assessed
41 135	3				10/04/2019	15/04/2019	Not assessed Not assessed
135	U				10/04/2019	10/04/2019	NUL assessed



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801/2019							
ACTINIC PRURIGO Total distributed		Results for assessment					
Total submitted		3					
Reference		DRB1*08:01, DRB1*07:01					
Number acceptable		3					
Number unacceptable Assessed		- <u>V</u>					
ASSESSEU B	1	<u> </u>					
Lat	Results for assessment		Alleles of interest	Interpretative comments	Comments	Received	Tested Assessment
11 12	1						16/04/2019 Not assessed 12/04/2019 Not assessed
15							0000-00-00 Not assessed
17	7					11/04/2019	12/04/2019 Not assessed
24			DDD4404.07	The state of the population and the state of		10/04/2019	15/04/2019 Not assessed
25	DRB1*04:07 NEGATIVE		DRB1*04:07	This patient is negative for HLA DRB1*04 alleles which are associated with Actinic Prurigo		10/04/2019	18/04/2019 Acceptable
38	B DRB1*08:01, *07:01		DRB1*04:07	This patient does not carry DRB1*04:07, which confers susceptibility to Actinic Prurigo		10/04/2019	18/04/2019 Acceptable
42	2 DRB1*07:01/07:05/07:07/07:	:10N/07:11/07:13/07:15/07:16/07:19/07:21/07:24/07:25/	07:HLA-DRB1*04:07	This patient is HLA-DRB1*04:07 NEGATIVE. Actinic Prurigo is associated with the		11/04/2019	12/04/2019 Acceptable
	27/07:28/07:29/07:30/07:31/0	07:32/07:33/07:34/07:35/07:37/07:38/07:40/07:41/07:43	/0	expression of the human leukocyte antigen (HLA) class II molecule DRB1*04:07			
		9/07:51/07:52/07:53/07:55/07:56/07:58N/07:59/07:60/07					
	84/07:85/07:86/07:87N/07:88	7:72/07:73/07:74/07:75/07:77/07:78/07:79/07:81/07:82/0	Jr:				
		:50/08:55/08:64/08:77/08:78N/08:79/08:86					
78	В						0000-00-00 Not assessed
85 86							23/04/2019 Not assessed 23/04/2019 Not assessed
87							24/04/2019 Not assessed 24/04/2019 Not assessed
109							16/04/2019 Not assessed
113	3		Not assessed		Haemolysis of the sample. DNA concentration too low and bad quality. No result	09/04/2019	0000-00-00 Not assessed
123	,				available	0000 00 00	0000-00-00 Not assessed
124	4						0000-00-00 Not assessed
126							15/04/2019 Not assessed
127 129							16/04/2019 Not assessed 29/04/2019 Not assessed
142							22/04/2019 Not assessed
150	D					17/04/2019	18/04/2019 Not assessed
154 159						10/04/2019	17/04/2019 Not assessed 12/04/2019 Not assessed
173							23/04/2019 Not assessed 23/04/2019 Not assessed
176	6					15/04/2019	16/04/2019 Not assessed
201							17/04/2019 Not assessed
219 223							30/04/2019 Not assessed 16/04/2019 Not assessed
224							17/04/2019 Not assessed
225	5					15/04/2019	17/04/2019 Not assessed
245 255	Not Tested		NT	NT		10/04/2019	17/04/2019 Not assessed
263	3					18/04/2019	23/04/2019 Not assessed 23/04/2019 Not assessed
269	9					07/04/2019	17/04/2019 Not assessed
274							15/04/2019 Not assessed
276 278							23/04/2019 Not assessed 25/04/2019 Not assessed
279	9						0000-00-00 Not assessed
281	1					09/04/2019	16/04/2019 Not assessed
307 315	7					10/04/2019	12/04/2019 Not assessed 16/04/2019 Not assessed
317						15/04/2019	17/04/2019 Not assessed
319	9					10/04/2019	12/04/2019 Not assessed
331							24/04/2019 Not assessed
333 338							18/04/2019 Not assessed 28/05/2019 Not assessed
339	9						16/04/2019 Not assessed
346	6						23/04/2019 Not assessed
347 355	5						17/04/2019 Not assessed 30/04/2019 Not assessed
359	9						17/04/2019 Not assessed
363	3						16/04/2019 Not assessed
413	3					10/04/2019	15/04/2019 Not assessed
1350	J					10/04/2019	16/04/2019 Not assessed



d Diseases

		UK NEQAS for H&I \$	Scheme 8 - HLA Genotypi	ng for Coeliac and Other I	HLA	Asso	ociated D
B01/2019 BIRDSHOT RETINDEATHY Total distributed Total submitted Reference Number unacceptable Number unacceptable Assessed II	Results for assessment 7 7 7 A*02, A*29 7 ED						
Lab Results for assessment 11 HLA-A29 Positive	<u></u>	Alleles of interest A*29	Interpretative comments HLA-A affele known to be associated with but not diagnostic for birdshot choicretinopathy is present	Comments	Received 09/04/2019	Tested A 16/04/2019 A	
12 15 Not Tested 17			сиотменторацу із резеня		10/04/2019 0000-00-00 11/04/2019	12/04/2019 N 0000-00-00 N 12/04/2019 N	lot assessed lot assessed
24 25 A*29		A*29	The HLA A29 antigen associated with Birdshot Chorioretinopathy is present. The presence of a particular HLA antigen does not establish the diagnosis of any particular disease, but provides a probability statement for the possible existence of the disease in the patient.		10/04/2019 10/04/2019	15/04/2019 N 18/04/2019 A	cceptable
38 A*02, *29		A*29	This patient carries HLA-A*29 which confers susceptibility to Birdshot Retinopathy			18/04/2019 A	
302-44902-59012-5902-5902-5902-5902-5902-5902-5902-590	5700 (1980 - 1980 - 1980 - 2080 - 2550 - 258		This patient is HLAA'29 POSITIVE. Birdinfor retinochronidopathy is associated with the expression of the human leukocyte artigen (HLA) class I molecule A'29.		11/04/2019	12/04/2019 A	coeplable
78 85 86 87 109 113 123 124 126		Not assessed		Neemolysis of the sample. DNA concentration too low and bad quality. No result available	10/04/2019 10/04/2019 10/04/2019 10/04/2019 09/04/2019	0000-00-00 N 23/04/2019 N 23/04/2019 N 16/04/2019 N 0000-00-00 N 0000-00-00 N 0000-00-00 N	lot assessed lot assessed lot assessed lot assessed lot assessed lot assessed lot assessed
127 129					11/04/2019 22/04/2019	16/04/2019 N 29/04/2019 N	lot assessed
142 150 A*02;29		A*29.	Presence of allele A*29.		11/04/2019 17/04/2019	22/04/2019 N 18/04/2019 A	cceptable
154 159 173 176 201					10/04/2019 11/04/2019 17/04/2019 15/04/2019 10/04/2019	17/04/2019 N 12/04/2019 N 23/04/2019 N 16/04/2019 N 17/04/2019 N	lot assessed lot assessed lot assessed
219					24/04/2019	30/04/2019 N	lot assessed
223						16/04/2019 N	
224						17/04/2019 N	
225						17/04/2019 N	
245 A*02,*29		A*29	Presence of the A*29 susceptibility allele. The presence of the A*29 allele, associated with the clinical signs of the disease, is strongly in favor of the diagnosis of birdshot disease. The prevalence of A*29 in patients with birdshot is 90 to 100% according to published studies.			17/04/2019 A	
255						23/04/2019 N	
263						23/04/2019 N	
269						17/04/2019 N	
274 276						15/04/2019 N 23/04/2019 N	
278						25/04/2019 N	
279						0000-00-00 N	
281					09/04/2019	16/04/2019 N	
307		400			10/04/2019	12/04/2019 N	
315 POSITIVE (A*02, A*29) 317		A*29			10/04/2019 15/04/2019	16/04/2019 A 17/04/2019 N	
319					10/04/2019	12/04/2019 N	lot assessed
331					12/04/2019	24/04/2019 N	lot assessed
333					04/04/2019	18/04/2019 N	lot assessed
338					11/04/2019	28/05/2019 N	lot assessed
339					10/04/2019	16/04/2019 N	lot assessed
346					16/04/2019	23/04/2019 N	lot assessed

10/04/2019 15/04/2019 Not assessed



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801/201 DISEAS						
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Assessed 5	Z Z					
	b Results for assessment	Alleles of interest	Interpretative comments	Comments	Received	Tested Assessment
1	1 HLA-B51(5) Positive 2 HLA-B*51: Positive	B*51 HLA-B*51	HLA-B51(5) is associated with but is not diagnostic for Behcet's disease This patient is positive for HLA-B*51 (the HLA specificity associated with Behcet's		09/04/2019	16/04/2019 Acceptable 12/04/2019 Acceptable
	2 PLANE ST. POSITIVE	HDA'D ST	disease).		10/04/2015	12/04/2019 Acceptable
			HLA-B*51 is present in approximately 9% of the normal Caucasian population.			
			There or is present in approximately 5% or the normal deduction population.			
	5 Not Tested 7				11/04/2019	0000-00-00 Not assessed 12/04/2019 Not assessed
2	4				10/04/2019	15/04/2019 Not assessed
2	5 B*51	B*51	The HLA B51 antigen associated with Bechets Disease is present. The presence of particular HLA antigen does not establish the diagnosis of any particular disease, but	a .	10/04/2019	18/04/2019 Acceptable
			provides a probability statement for the possible existence of the disease in the			
3	8 HLA-B*44.*51	B*51	patient. This patient carries HLA-B*51 which confers susceptibility to Behcet's disease.		10/04/2019	18/04/2019 Acceptable
4	2 B*44:03/44:26/44:35/44:36/44:38/44:39/44:47/44:79/44:85/44:89/44:94/44:98/44:103	/ 48 *51	This patient is HLA-B*51 POSITIVE. Behcet's disease is associated with the		11/04/2019	12/04/2019 Acceptable
	:108N/44:111/44:115/44:122/44:125/44:128/44:141/44:147/44:154/44:155/44:157/44 9/44:161/44:164/44:165/44:167/44:175/44:178/44:180/44:182/44:183/44:186/44:188	:15 M4:	expression of the human leukocyte antigen (HLA) class I molecule B51.			
	189/44:192/44:198N/44:202/44:205/44:207/44:222/44:228/44:231/44:233/44:237N/4	4.2				
	39/44:250/44:252/44:258/44:278/44:280/44:281/44:284/44:286 B*51:01/51:03/51:04/51:09/51:11N/51:24/51:27N/51:28/51:30/51:32/51:33/51:35/51:	38/				
	51:39/51:41N/51:48/51:49/51:51/51:52/51:55/51:60/51:65/51:67/51:69/51:71/51:73/5	1:7				
	4/51:75/51:76/51:77/51:79/51:80/51:83/51:84/51:86/51:95/51:96/51:98N/51:99/51:10 1:102/51:105/51:107/51:109/51:110N/51:111/51:117/51:121/51:122/51:123/51:125/5	0/5				
	26/51:127/51:129/51:130/51:134/51:136/51:137/51:138/51:142/51:145/51:149N/51:1	51/				
	51:154/51:155/51:156/51:158/51:159/51:161/51:162/51:163/51:164/51:165/51:166/5 67/51:168/51:169/51:171/51:173Q/51:174/51:177/51:178N/51:183/51:184N/51:186/5					
	187/51:188/51:191/51:193/51:196/51:198/51:201/51:203/51:204/51:205/51:207/51:21	 09/				
	51:210/51:212/51:215/51:219/51:224/51:229/51:230/51:232/51:233/51:234/51:235N					
7	8 Not tested				10/04/2019	0000-00-00 Not assessed 23/04/2019 Not assessed
8	6				10/04/2019	23/04/2019 Not assessed
8 10						24/04/2019 Not assessed 16/04/2019 Not assessed
	3 Haemolysis of the sample. No result available	B51		Haemolysis of the sample. DNA concentration too low and bad quality. No result		0000-00-00 Not assessed
12	2			available	0000-00-00	0000-00-00 Not assessed
12	4				0000-00-00	0000-00-00 Not assessed
12	6 7				10/04/2019	15/04/2019 Not assessed 16/04/2019 Not assessed
12 12	9				22/04/2019	29/04/2019 Not assessed
14	2 HLA-B*51 present 0 B*44:51	HLA-B*51 B*51	Presence of allele B*51.		11/04/2019	22/04/2019 Acceptable 18/04/2019 Acceptable
15	4		Treatment of annie 5 of .		10/04/2019	17/04/2019 Not assessed
15	9 3 HLA-B*51 POSITIVE	HLA-B*51	High risk of Behħet Syndrome			12/04/2019 Not assessed 23/04/2019 Acceptable
17 20	6	TIDYO UT	riigi riak or beinger of raione		15/04/2019	16/04/2019 Not assessed
20	11				10/04/2019 24/04/2019	17/04/2019 Not assessed 30/04/2019 Not assessed
21 22	3				10/04/2019	16/04/2019 Not assessed
22	4				10/04/2019	17/04/2019 Not assessed 17/04/2019 Not assessed
	5 B*44,*51	B*51	Presence of the susceptibility allele HLA-B*51. The presence of this allele confers on			17/04/2019 Acceptable
			the wearer a risk multiplied by 6 to develop a BehÄŞet disease compared to the general population, 30-60% of affected patients have this allele. This result alone doe	6		
			not support the diagnosis of Behħet's disease. It must be associated with other	•		
			clinico-biological signs corresponding to the diagnostic criteria.			
25	5				10/04/2019	23/04/2019 Not assessed
26	3				18/04/2019	23/04/2019 Not assessed
26	0				07/04/2010	17/04/2019 Not assessed
27	4				12/04/2019	15/04/2019 Not assessed
27	6				10/04/2019	23/04/2019 Not assessed
27	8					25/04/2019 Not assessed
27	9					0000-00-00 Not assessed
28	1				09/04/2019	16/04/2019 Not assessed
30	7				10/04/2019	12/04/2019 Not assessed
		Dica				
31 31	5 POSITIVE (B*44, B*51) 7	B*51			10/04/2019	16/04/2019 Acceptable 17/04/2019 Not assessed
-						
31						12/04/2019 Not assessed
33	1				12/04/2019	24/04/2019 Not assessed
33	3				04/04/2019	18/04/2019 Not assessed
33	8				11/04/2010	28/05/2019 Not assessed
33	9				10/04/2019	16/04/2019 Not assessed
34	6				16/04/2019	23/04/2019 Not assessed
34	7					17/04/2019 Not assessed
35	5				10/04/2019	30/04/2019 Not assessed
35	9				11/04/2019	17/04/2019 Not assessed
36	3				09/04/2019	16/04/2019 Not assessed
41					10/04/2019	15/04/2019 Not assessed
135	<u> </u>				10/04/2019	16/04/2019 Not assessed



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801/ RHEUMATOID ARTHI Total distrib Total subm Refer Number accep Number unaccep Assesse	RITIS Results for assessment utted 2 titted 2 ence DRB1*08:01, DRB1*07:01 table 1 table -				
	Lab Results for assessment 11 12 15 17 24 25 38 42 78 85 86 87	Alleles of interest	Interpretative comments	Comments	Roceived Testad Assessment 0904/2019 10/2019 10/2019 Not assessed 1004/2019 12/24/2019 Not assessed 0000-00-00 0000-00-00 Not assessed 11/04/2019 12/04/2019 Not assessed 10/04/2019 18/04/2019 Not assessed 10/04/2019 18/04/2019 Not assessed 10/04/2019 12/04/2019 Not assessed 10/04/2019 23/04/2019 Not assessed 10/04/2019 23/04/2019 Not assessed 10/04/2019 23/04/2019 Not assessed 10/04/2019 24/04/2019 Not assessed 10/04/2019 16/04/2019 Not assessed 10/04/2019 16/04/2019 Not assessed 10/04/2019 Not assessed
	113 Haemolysis of the sample. No result available	DRB1*01:01, DRB1*01:02, DRB1*04:01, DRB1*04:04, DRB1*04:05, DRB1*04:08, DRB1*04:09, DRB1*04:10, DRB1*10, DRB1*14:02,		Haemolysis of the sample. DNA concentration too low and bad quality. No result available	09/04/2019 10/04/2019 Not assessed 09/04/2019 0000-00-00 Not assessed
	123 124 126 127 129 142 150 154 159 173 176 201 219 223 224	DRB1*14.06			0000-00-00 0000-00-00 Not assessed 0000-00-00 0000-00-00 Not assessed 1004/2019 I5/04/2019 Not assessed 11/04/2019 I5/04/2019 Not assessed 22/04/2019 Sy01-2019 Not assessed 22/04/2019 29/04/2019 Not assessed 11/04/2019 29/04/2019 Not assessed 11/04/2019 18/04/2019 Not assessed 11/04/2019 18/04/2019 Not assessed 10/04/2019 18/04/2019 Not assessed 15/04/2019 18/04/2019 Not assessed 15/04/2019 16/04/2019 Not assessed 15/04/2019 16/04/2019 Not assessed 10/04/2019 1/04/2019 Not assessed 10/04/2019 1/04/2019 Not assessed 10/04/2019 1/04/2019 Not assessed 15/04/2019 Not assessed 15/04/2019 Not assessed 10/04/2019 1/04/2019 Not assessed
	245 DRB1*07,*08	DRB1*04:01,*04:04.*04:05,*04:08 DRB1*10 DRB1*01:01,*01:02,*01:04 DRB1*14:06	No single or double dose shared epitope. The patient has no shared epitopes associa with a risk of developing rheumatoid arthritis. 20% of patients with rheumatoid arthritis do not have a shared epitope, so this result does not eliminate the diagnosis.	at S	10/04/2019 17/04/2019 Acceptable
	255 263 269 274 276 278 279 281 307 315 317 319 323 331 DRB1*07, DRB1*08 333 338 346 347 355 399 363 4413	DRB1*04	Absence of DRB1*04		10/04/2019 23/04/2019 Not assessed 18/04/2019 23/04/2019 Not assessed 07/04/2019 17/04/2019 Not assessed 07/04/2019 Not assessed 12/04/2019 Not assessed 12/04/2019 Not assessed 12/04/2019 Not assessed 15/04/2019 Not assessed 15/04/2019 Not assessed 09/04/2019 18/04/2019 Not assessed 10/04/2019 12/04/2019 Not assessed 10/04/2019 18/04/2019 Not assessed 10/04/2019 28/05/2019 Not assessed 10/04/2019 30/04/2019 Not assessed 10/04/2019 30/04/2019 Not assessed 10/04/2019 17/04/2019 Not assessed 10/04/2019 15/04/2019 Not assessed 10/04/2019 Not asses
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801/2019 DIABETES istributed submitted	Results for assessment 5					
Reference cceptable	DRB1*08:01, *07:01; DQA1*02:01,*04:01; DQB1*02:02,*04:02 4					
cceptable ssessed 🗹	<u>.</u>					
Lab 11 12 15 17 24 25 38 42 78 85 86 87 109		Alleles of interest	Interpretative comments	Comments	0000-00-00 11/04/2019 10/04/2019 10/04/2019 10/04/2019 11/04/2019 10/04/2019 10/04/2019 10/04/2019 10/04/2019	Tested Assessment 1604/2019 Not assessed 1204/2019 Not assessed 1804/2019 Not assessed 1804/2019 Not assessed 1204/2019 Not assessed
113	Haemolysis of the sample. No result available	DRB1*04/DQA1*03:01/DQB1*03:02, DRB1*03:01/DQA1*05:01/DQB1*02:01		Haemolysis of the sample. DNA concentration too low and bad quality. No result available	09/04/2019	0000-00-00 Not assessed
123 124 126 127	DQB1*02. *04: DQA1*02:01	DQB1'02 '03.01, '03.02 '03.03, '03.04, '04, '05.01, '05.03, '06.01, '06.02, '06.03,	Notificially for ink or protection accordant banks are:		0000-00-00 10/04/2019 11/04/2019	0000-00-00 Not assessed 0000-00-00 Not assessed 15/04/2019 Not assessed 16/04/2019 Not assessed 29/04/2019 Acceptable
		DQA1*02:01, *03.05 DRB1*04:01, *04:02, *04:03/6, *04:04, *04:05, *04:07	пециа нак (но нак от ргоесион аввосиате нарисурев)			
142 150 154	DRB1107.08 HLA-DRB1107.708, HLA-DQA1102.01, 104.01; HLA-DQB1102.02,104.02	DR3 : DRB1*03.01 - DR4 : DRB1*04.05. H.LA-DR3*1* is typed to the 2-dgall level and HLA-DQA1* and HLA-DQB1* are typed to the 4-dgall real to detect the following hapidopses : H.LA-DRB1*04, DQA1*03.01, DQB1*03.01 H.LA-DRB1*04, DQA1*03.01, DQB1*03.02	Absence of allA ¹ les DR3 OR DR4. halptoype DR3 D02 (DR811V3.0A)1V3.01, D081V0.201): absence halptoype DR4.0D2 (DR811V4.DA41V3.01, D081V0.302): absence The patient has no HLA halptoype associated with type 1 diabeties. The DR3, D02 and DR4.D08 halptoypes are found in 95% of type 1 diabetes patients. The HLA-DR3 and DR4 antigens are found 40% of the Caucasian population.		17/04/2019	22/04/2019 Not assessed 18/04/2019 Acceptable 17/04/2019 Acceptable
159 173 176 201 219 223 224 225		Susceptible:	Absence of alleles that may constitute a susceptibility or protective HLA haplotype to		17/04/2019 15/04/2019 10/04/2019 24/04/2019 10/04/2019 10/04/2019 15/04/2019	12/04/2019 Not assessed 23/04/2019 Not assessed 16/04/2019 Not assessed 7/04/2019 Not assessed 30/04/2019 Not assessed 17/04/2019 Not assessed 17/04/2019 Not assessed 17/04/2019 Not assessed 17/04/2019 Not assessed
240	3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0 3.0	DBB119301:DQA119301:DQB119301 DBB119401:DQA119301:DQB11930204 DBB119401:DQA119301:DQB11930204 DBB119401:DQA119301:DQB11930204 DBB119401:DQA119301:DQB11930204 DBB119401:DQA119301:DQB11930204 DBB119401:DQA119101:DQB119302	type I diabetes. This result does not constitute a criterion for excluding the disease.		13042515	770-1200 Peccepation
255 253 269 274 276 278 279 281 307 315 317 319 333 338 339 346 347 355 359 363 31350	DRB1'07, DRB1'08	DR81*03, DR81*04	Absence of DRB1*03 : Absence of DRB1*04		1804/2019 0704/2019 12/04/2019 12/04/2019 10/04/2019 10/04/2019 10/04/2019 10/04/2019 10/04/2019 10/04/2019 11/04/2019 11/04/2019 16/04/2019 16/04/2019 11/04/2019 11/04/2019 11/04/2019 11/04/2019 11/04/2019 10/04/2019 10/04/2019	2304/2019 Not assessed 17,04/2019 Not assessed 2504/2019 Not assessed 2504/2019 Not assessed 2504/2019 Not assessed 1604/2019 Not assessed 1604/2019 Not assessed 1604/2019 Not assessed 17,04/2019 Not assessed 12,04/2019 Not assessed 17,04/2019 Not assessed 15,04/2019 No



	UI	K NEQAS for H&I Scheme 8	- HLA Genotyping for Coellac and Ot	ner HLA Associated Diseases	
801/2019 OTHER Total distributed Total submitted Reference Number acceptable	Results for assessment 0 0				
Number unacceptable	:				
Assessed ☑	<u> </u>				
Assessed		Alleles of interest	Interpretative comments	Haemolysis of the sample. DNA concentration too low and bad quality. No result available	Received
355 359 363 413 1350					10/04/2019 30/04/2019 Not assessed 11/04/2019 17/04/2019 Not assessed 09/04/2019 16/04/2019 Not assessed 10/04/2019 15/04/2019 Not assessed 10/04/2019 16/04/2019 Not assessed