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Genomic and Transcriptomic Characterization of Atypical Recurrent Flank Alopecia in the Cesky Fousek

Silvie Neradilová 1, Alexandria M. Schauer 2,3, Jessica J. Hayward 4, Magdalena A. T. Brunner 2,3, Magdalena Bohutínská 5, Vidhya Jagannathan 6, Laurie B. Connell 7, Adam R. Boyko 4,†, Monika M. Welle 2,3,† and Barbora Černá Bolfíková 1,*,†

1 Department of Animal Science and Food Processing, Faculty of Tropical AgriSciences, Czech University of Life Sciences Prague, 16500 Prague, Czech Republic; neradilova@ftz.czu.cz
2 Institute of Animal Pathology, Vetsuisse Faculty, University of Bern, 3012 Bern, Switzerland; alexandria.schauer@vetsuisse.unibe.ch (A.M.S.); magdalena.holcova@natur.cuni.cz (M.M.W.)
3 DermFocus, University of Bern, 3012 Bern, Switzerland
4 Department of Biomedical Sciences, College of Veterinary Medicine, Cornell University, Ithaca, NY 14853, USA; jjh276@cornell.edu (J.J.H.); boyko@cornell.edu (A.R.B.)
5 Department of Botany, Faculty of Science, Charles University, 12800 Prague, Czech Republic; monika.welle@vetsuisse.unibe.ch (M.M.W.)
6 Institute of Genetics, Vetsuisse Faculty, University of Bern, 3012 Bern, Switzerland; vidhya.jagannathan@vetsuisse.unibe.ch
7 Molecular & Biomedical Sciences and School of Marine Sciences, University of Maine, Orono, ME 04469, USA; laurie.b.connell@maine.edu
* Correspondence: bolfikova@ftz.czu.cz
† These authors contributed equally to this work.

Abstract: Non-inflammatory alopecia is a frequent skin problem in dogs, causing damaged coat integrity and compromised appearance of affected individuals. In this study, we examined the Cesky Fousek breed, which displays atypical recurrent flank alopecia (aRFA) at a high frequency. This type of alopecia can be quite severe and is characterized by seasonal episodes of well demarcated alopecic areas without hyperpigmentation. The genetic component responsible for aRFA remains unknown. Thus, here we aimed to identify variants involved in aRFA using a combination of histological, genomic, and transcriptomic data. We showed that aRFA is histologically similar to recurrent flank alopecia, characterized by a lack of anagen hair follicles and the presence of severely shortened telogen or kenogen hair follicles. We performed a genome-wide association study (GWAS) using 216 dogs phenotyped for aRFA and identified associations on chromosomes 19, 8, 30, 36, and 21, highlighting 144 candidate genes, which suggests a polygenic basis for aRFA. By comparing the skin cell transcription pattern of six aRFA and five control dogs, we identified 236 strongly differentially expressed genes (DEGs). We showed that the GWAS genes associated with aRFA are often predicted to interact with DEGs, suggesting their joint contribution to the development of the disease. Together, these genes affect four major metabolic pathways connected to aRFA: collagen formation, muscle structure/contraction, lipid metabolism, and the immune system.

Keywords: dog; GWAS; canine alopecia; atypical recurrent flank alopecia; Cesky Fousek; RNA-seq; differential gene expression; skin biopsies

1. Introduction

The domestic dog (Canis lupus familiaris) has recently become a frequently used model organism for genomic medicine research. Over the last two decades, approximately 350 diseases have been identified that have the same or very similar underlying pathways in both dogs and humans [1–3]. The intensive selection during breed development, combined with a limited number of founding individuals, has resulted in long genomic regions of
linkage disequilibrium within breeds. This makes association studies in dogs simpler and more straightforward than in humans [1,2]. In order to fix desirable traits, closely related founding individuals were usually bred together, resulting in high levels of inbreeding. Thus, newly established breeds often go through population bottlenecks followed by strong genetic drift that, together with inbreeding, decreases genetic diversity and increases homozygosity in populations. This leads to the fixation of undesirable traits and keeps harmful recessive alleles or dominant alleles with incomplete penetrance, resulting in late-onset and complex diseases within populations [4,5]. Genome-Wide association studies (GWAS) have proven to be a very effective tool for discovering the single nucleotide polymorphisms (SNPs) associated with a disease, even with a very low number of sampled individuals [6].

One undesirable trait observed frequently in dogs and less often in other mammalian species is alopecia or hypotrichosis, which results in the absence of the hair coat or a severely reduced number of hairs on some body regions or the entire body. Successful hair growth is dependent on the finely tuned interactions of signaling molecules and transcription factors, as well as a fully functional stem cell compartment within the hair follicle (HF). This interplay orchestrates the HF morphogenesis during the embryonic stage and the postnatal hair cycle (HC), and its coordination is dependent on complex interactions between signals of the follicular and dermal microenvironment, systemic factors, and environmental factors [7–11]. The HC is a lifelong process during which the HF undergoes periodic stages of growth (anagen), regression (catagen), and quiescence (telogen) [12,13]. This process relies on follicular stem cells (SCs) located in their niche [14,15]. In early anagen, cell proliferation leads to the expansion of the HF downward before the cells engulf the dermal papilla to form the hair bulb. The hair bulb is composed of matrix cells, which are highly lineage-restricted proliferative precursors. These matrix progenitors subsequently become differentiated post-mitotic cells that constitute the layers of the companion layer, the inner root sheath and the hair shaft (reviewed in [14,15]). During catagen, the next phase of the HC, the lower portion of the follicle regresses. Lastly, a state of proliferative quiescence ensues while the HF is in telogen. At any point during the next HC, but mostly during anagen, the old club hair, which is firmly anchored to the outer root sheath by trichilemmal keratin during telogen, is shed in a process called exogen [16]. Disruption of the HC results in alopecia due to either HC arrest or impaired hair quality, leading to the postnatal onset of noninflammatory alopecia.

Noninflammatory alopecia is a relatively common problem in dogs. Some forms of alopecia have a clear hereditary cause [17–21], and affected dogs are born without hair. In other forms, a hereditary background is suspected because of a clear breed predisposition, and these cases typically exhibit later onset of the clinical phenotype. Examples of such disorders are alopecia X [22–26] and recurrent flank alopecia (RFA) [27,28]. The cause of these disorders remains unknown.

Typical RFA is characterized by recurrent episodes of well demarcated, hyperpigmented areas of alopecia that affect several canine breeds, including Boxers, Rhodesian Ridgebacks, and Airedale Terriers [27–31]. The hair on the affected parts will usually regrow within a few weeks with recurrent loss the following year [31–33]. RFA is not correlated with the sex of the individual [34] and always affects adults [27]. Previous studies have shown that the onset of RFA is influenced by the photoperiod [31,34]. Histologically, RFA is associated with severely dilated infundibula filled with abundant keratin and shortened, narrowed, and often distorted proximal HF parts, resulting in the typical “witch’s feet” appearance [27].

An atypical form of RFA (aRFA) is anecdotaly described in Vizslas, German Wire-haired Pointers, and the Cesky Fousek (CF). In this atypical form, the alopecia may still wax and wane but is more severe and affects the flanks, the sacral area, the thighs, the base of the tail, and sometimes the ears and nose (Figure 1).
2. Materials and Methods

2.1. Blood Sample Collection

Altogether, 216 samples (non-affected n = 116, affected n = 100) were collected (189 from the Czech Republic and 27 from Cesky Fousek North America (CFNA)); 72 males and 144 females (Table S1). Relatives were not excluded from the dataset. The blood draws were done in cooperation with the Czech Cesky Fousek Breeding Club (KCHCF) and CFNA.
during 2016–2019. Blood samples were shipped to the Cornell Veterinary Biobank, where DNA extraction was performed by standard salt precipitation and the DNA was then stored at −20 °C. The level of severity was determined by a responsible member of the breeding club (KCH CF). In the past, the club has developed a protocol for the identification of all aRFA levels and this protocol was also followed in our study. The Affliction of individuals was marked during sample collection, and their affected status and severity was updated throughout the study’s duration.

2.2. Biopsy Sample Collection

Six-millimeter punch biopsies were taken under local anesthesia for histological evaluation and RNA extraction (seven control dogs and seven affected dogs). From the seven control dogs, two 6 mm punch biopsies were taken from sites close to each other. From the seven dogs affected with aRFA, two neighboring biopsies were taken from a completely alopecic site and two from a distant, fully haired area (shoulder). One biopsy from each site was fixed and stored in buffered Formafix 10%® (Formafix AG, Hittnau, Switzerland). The second biopsy from each site was collected in RNA later (Invitrogen, CA, USA) and stored at −20 °C until RNA extraction was performed. Based on the histological evaluation of the samples and the diagnosis of hypothyroidism in one of the dogs, we had to exclude some samples from further analysis. A comprehensive list of animals used for this study can be found in Table S2.

2.3. Genotyping

Genotyping was performed on a semi-custom 220k CanineHD array (Illumina, CA, USA), currently available as the Embark genetic test [www.embarkvet.com, accessed on 19 February 2022]. In total, 216 samples were genotyped: 47 samples in 2016 with 214,582 markers and 169 samples in 2018–2019 with 239,490 markers. The number of markers differs due to the upgrading of the genotyping array. The positions of the markers are listed in CanFam3.1.

PLINK [36,37] datafiles were generated and the data were checked for errors in sex and genotype missingness. All samples had a genotyping rate higher than 95% so no samples were excluded at this point. Only markers with minor allele frequency (MAF) higher than 0.05 were included in the analyses. Data from the two different arrays were merged and discordant SNPs between duplicate samples, in accordance with a previous study [38], were removed from the datasets. Moreover, in order to avoid the resulting bias caused by the non-balanced sex of the individuals entering the analysis (Table S1), we also filtered out the chromosome Y retrocopies [39] and sex-associated markers (a total of 96 SNPs). After filtering, the number of SNPs used for GWAS was 140,024.

Principal component analysis (PCA) was performed prior to the GWAS analyses in order to (i) check for population structure between the Czech Republic and USA samples, (ii) look for any batch effects due to the two genotyping arrays used, and (iii) identify any individual outliers. PCA was run on unlinked SNPs only, using PLINK command–indep 50 5 2, in the program EIGENSTRAT in the EIGENSOFT v5.0.1. package [40,41]. PCA plots were visualized in R i386 3.6.1 [42].

The population stratification of the data was corrected in GEMMA by including a relatedness matrix, calculated from genotypes, as a random effect. We calculated the genomic inflation factor (λ), based on p-values, in the R package snpStats [43]. Lambda inflation factor compares the median test statistic and expected null distribution and it detects the normality of the data distribution with a value of 1.0 representing no inflation. Manhattan and quantile–quantile plots of p-values were constructed in R. The significance thresholds for the GWAS analyses were set on Bonferroni correction on unlinked SNPs (using –indep 100 10 10 in PLINK). LD plots were created from LD analyses run in PLINK and using Matplotlib library in the Jupyter notebook [44,45].
2.4. Case/Control GWAS

A Genome-wide association study (GWAS) was conducted using a linear-mixed model in GEMMA v0.98.1 [46]. In total, 213 individuals out of 216 were used for the case/control GWAS study—three affected individuals were excluded from this analysis due to an unusual manifestation of alopecia (alopecia on the head), resulting in a dataset of 96 affected and 117 control individuals (Table S1).

2.5. Quantitative GWAS (QGWAS) and Additional GWAS Analyses

In order to discover variants with a direct association with aRFA level, we performed a QGWAS analysis with 216 individuals. The three animals with the unusual manifestation of aRFA on their heads were included in this analysis. All individuals were divided into one of six phenotypic categories. The number of individuals in each category and the code of each category are: healthy (n = 111; code “0”), head affection (n = 3; code “0.1”), level 1 aRFA (n = 6; code “0.25”), level 2 aRFA (n = 28, code “0.5”), level 3 aRFA (n = 49; code “0.75”), level 4 aRFA (n = 19; code “1”). Each sample was assigned to the corresponding category according to the aRFA level: Head affection—the individual loses hair on the top of the head, ears, and sometimes the top of the nose. Level 1—the individual loses hair on the ears only (can enter the breeding program); Level 2—the individual loses hair on the body sides up to the size of approximately $10 \times 10$ cm; Level 3—hair loss on the body sides up to approximately $10 \times 25$ cm; Level 4—hair loss on the body sides up to approximately $10 \times 40$ cm; Level 5—hair loss on the body sides larger than $10 \times 40$ cm (this level was not represented in our dataset).

Moreover, we conducted four additional GWAS analyses to identify specific variants that were associated with level 2 aRFA (28 individuals), level 4 aRFA (19 individuals), age of onset before 2 years of age (26 individuals), and age of onset at 6–8 years of age (20 individuals). The last two groups were also affected by level 2 aRFA or worse. We did not consider level 1 aRFA as “affected”. The control group for all the above-mentioned groups was composed of 35 individuals aged 10+ years in which the chances of developing aRFA were very low. The settings of the allelic and genotyping frequencies were the same as in the main case/control GWAS analysis.

2.6. Haplotype Identification

Prior to haplotype identification, we divided the genotyping data by chromosomes using PLINK v.1.9 [47,48] and subsequently phased each chromosome of interest, based on the case/control GWAS results, in SHAPEIT.v2.r837 [49]. The settings were left at their default levels with 7 MCMC burn-in iterations, 8 pruning iterations, and 20 main iterations. The number of conditioning states (K) was left at 100, the --window size setting was 2 Mb, and the genetic map was not provided, leaving the --rho value at its default (0.0004). Each phased chromosome was then transferred to PED/MAP format and run in PLINK v1.07 [37] in order to estimate the haplotypes in both the case and control groups of individuals. The setting was set to --hap-window from 1 to 10 SNPs to obtain all one-, two-, and up to ten-SNP windows across the dataset (respecting the chromosome boundaries).

2.7. Histopathological Analysis

Formalin-fixed biopsies were processed for routine histological analysis by embedding in paraffin, microtoming (3 µm), and staining with hematoxylin and eosin (H&E). The samples were blinded and a histopathological analysis was conducted to characterize specific histological features and patterns associated with this alopecic disease. Histological evaluation was also utilized to include or exclude samples not suitable for RNA extraction based on histological findings that might influence gene expression (e.g., secondary lesions such as inflammation). Based on this, we excluded the biopsies from one control dog, two lesional sites of alopecic dogs, and three normal skin sites from affected dogs from the analysis, resulting in a total of six samples from control animals (B2, B5, B6, B12, B13 and B14), five samples of alopecic skin (B3, B7, B9, B10, B11), and four samples of normal
skin from affected dogs (B3, B9, B10, B11). Factors for exclusion included an endocrine imbalance in one of the alopecic dogs and pronounced inflammation in the biopsy of one control dog, three biopsies of normal skin of alopecic dogs, and one biopsy of alopecic skin of an affected dog.

2.8. RNA Extraction and RNA-Seq Experiments

RNA extraction and cDNA sequencing experiments were conducted according to the protocol outlined previously in [25,50]. All 11 samples were of high quality with a RIN > 9. After sequencing, the Illumina BCL output files with base calls and qualities were converted to FASTQ file formats and demultiplexed.

All reads that passed quality control were mapped to the canine reference genome (CanFam3.1) by STAR aligner version 2.5.3.a, as described in [25,50]. The alignment of RNA-seq reads from each sample was summarized by the number of splice arrangements per sample. The read abundance was calculated using the count software HTseq [51] and an NCBI annotated GTF (release 103) file.

2.9. Differential Expression Analysis

The R DESeq2 package [52] was used for differential expression analysis as described in [25]. For each gene, normalized read counts were fit to a generalized linear model (GLM) with the design formula where the condition was the factor of interest in two states: control and affected. Transcripts were considered to be differentially expressed with a Benjamini and Hochberg false discovery rate (FDR) of <0.01. The differentially expressed genes (DEGs) were mapped to biological networks using an open-source, open access, and manually curated pathway database called Reactome [https://reactome.org/ version 71; accessed on 12 January 2021]. Separate lists of upregulated and downregulated genes were uploaded separately into the database and were analyzed and matched with known biological processes and pathways.

2.10. Protein–Protein Interaction Analysis

We searched for potential functional associations among our GWAS and differentially expressed candidate genes using the STRING database [53], following the approach described in the study of Bohutinská et al. [54]. We were able to retrieve predicted protein–protein interactions for 132 out of 144 GWAS candidates and for 144 out of 236 strongly DEGs (exceeding the Log2FC value of +/-2). We used the ‘multiple proteins’ search in Canis lupus, with text mining, experiments, databases, co-expression, neighborhood, gene fusion, and co-occurrence as information sources. We used a minimum confidence of 0.4 and retained only 1st shell associations (proteins that are directly associated with the candidate protein: i.e., immediately neighboring network circles).

3. Results

3.1. Population Genetic Structure of aRFA Affected and Control Individuals

We first inquired whether aRFA individuals appear evenly distributed among populations. Even though a certain level of genetic differentiation between the Czech and the North American populations exists [55], the case and control samples were evenly distributed across the whole dataset (Figure S1a); thus, we decided to use all samples for the GWAS analyses. We also checked for a possible batch effect since the samples were genotyped in different years and the genotyping array had been updated. No batch effect was identified, as shown in Figure S1b.

3.2. Case/Control GWAS

To identify genetic variants associated with aRFA, we performed GWAS, using the presence/absence of aRFA as a predictor of phenotype. Our main within-breed case/control GWAS analysis revealed a significant association with aRFA on chromosome 19 ($p = 1.08 \times 10^{-6}$) (Table 1; Figure 2). Of the top ten SNPs, six are located on chromosome 8 and five of these
are within the region 43,341,000–43,490,000 bp. Genotypes and their frequencies for the top SNPs on chromosomes 19 and 8 are shown in Table 2. The lambda value ($\lambda = 1.01$) shows that the stratification correction worked well. The significance threshold was based on the Bonferroni correction (alpha = 0.1; cut-off = $1.16 \times 10^{-6}$). Only the chr19 association can be considered significant, while the other identified variants are considered suggestive.

The distribution of genotypes for the chromosome 19 association shows that 59% of controls are of genotype AA while only 27% of cases are of the same genotype, and nearly 19% of cases are GG compared to only 6% of controls. For the chromosome 8 association, the highest proportion of cases (70%) have the genotype GG compared to only 44% of controls.

**Table 1.** Case/control GWAS results showing chromosome (Chr), SNP name, position (bp), allele frequency, and raw $p$-value for the top twenty SNPs. One significant SNP was identified on chromosome 19. The interrupted line represents the Bonferroni cut-off.

|Chr| SNP Name                  | Position (bp) | Allele Freq | $p$-Value       |
|---|----------------------------|---------------|-------------|-----------------|
|19 | BICF2G630255452            | 47,856,573    | 0.333       | $1.08 \times 10^{-6}$ |
|8  | BICF2P465820               | 43,487,284    | 0.262       | $3.10 \times 10^{-5}$ |
|8  | TIGRP2P114211_rs8542415    | 434,942,31    | 0.262       | $3.10 \times 10^{-5}$ |
|8  | BICF2S23110497             | 25,810,719    | 0.205       | $3.30 \times 10^{-5}$ |
|36 | BICF2P1194573              | 28,584,717    | 0.271       | $6.72 \times 10^{-5}$ |
|30 | BICF2G630401492            | 26,273,661    | 0.326       | $8.07 \times 10^{-5}$ |
|8  | BICF2P361090               | 43,341,287    | 0.233       | $8.90 \times 10^{-5}$ |
|8  | BICF2P543725               | 43,371,261    | 0.233       | $8.90 \times 10^{-5}$ |
|8  | BICF2S23137831             | 43,418,611    | 0.233       | $8.89 \times 10^{-5}$ |
|13 | BICF2P281837               | 63,012,417    | 0.057       | $9.61 \times 10^{-5}$ |
|6  | BICF2P742566               | 35,078,147    | 0.309       | $9.86 \times 10^{-5}$ |
|8  | BICF2P177234               | 43,520,222    | 0.235       | $1.10 \times 10^{-4}$ |
|41 | BICF2S23546044             | 18,45,101     | 0.310       | $1.11 \times 10^{-4}$ |
|8  | TIGRP2P114933_rs9187625    | 46,799,348    | 0.493       | $1.24 \times 10^{-4}$ |
|31 | BICF2P1368177              | 7,605,782     | 0.104       | $1.27 \times 10^{-4}$ |
|31 | BICF2S2443709              | 7,615,165     | 0.104       | $1.27 \times 10^{-4}$ |
|13 | BICF2G630745860            | 61,855,230    | 0.149       | $1.28 \times 10^{-4}$ |
|14 | BICF2G630521203            | 10,825,554    | 0.061       | $1.36 \times 10^{-4}$ |
|30 | TIGRP2P370921_rs8763952    | 26,977,673    | 0.233       | $1.36 \times 10^{-4}$ |
|8  | BICF2P1102123              | 43,411,814    | 0.255       | $1.54 \times 10^{-4}$ |

**Table 2.** Genotypes of the two top SNPs from the case/control GWAS for chromosomes 19 and 8. The highest proportions in each phenotype group are underlined.

|Chr (SNP) | Genotype | No. Controls | % Controls | No. Cases | % Cases |
|----------|----------|--------------|------------|-----------|---------|
|chr19 (BICF2G630255452) | AA       | 69           | 59.0       | 26        | 26.8    |
|          | GA       | 41           | 35.0       | 53        | 54.6    |
|          | GG       | 7            | 6.0        | 17        | 18.6    |
|chr8 (BICF2P465820) | AA       | 16           | 13.7       | 1         | 1.0     |
|          | AG       | 50           | 42.7       | 27        | 27.8    |
|          | GG       | 51           | 43.6       | 68        | 70.1    |
Although the significance level of the case/control GWAS was not met, we considered these suggestive associations relevant as well. The significance threshold was based on the Bonferroni correction (cut-off = 1.16 × 10^−6). The lambda value is shown in the QQ plot.

3.3. Quantitative GWAS and Additional GWAS Analyses

Quantitative GWAS (QGWAS) and additional GWAS analyses were performed in order to find possible variants associated with the aRFA level of severity represented in our dataset, as well as the age of aRFA onset. A QGWAS analysis of six phenotypic categories showed that seventeen of the twenty top SNPs were on chromosome 8 (Table 3; Figure S2). Seven of the top ten SNPs overlapped with those identified in the case/control GWAS. Although the significance level of the case/control GWAS was not met, we considered these suggestive associations relevant as well. The significance threshold was based on the Bonferroni cut-off (alpha = 0.05) for all analyses mentioned in this section.

Table 3. Quantitative GWAS results showing associations with six phenotypic categories. Chromosome (Chr), SNP name, position (bp), allele frequency, and raw p-value for the top twenty SNPs. No variant reached the significance cut-off, and thus these variants are considered suggestive only.

| Chr | SNP Name          | Position (bp) | Allele Freq | p-Value     |
|-----|-------------------|---------------|-------------|-------------|
| 8   | BICF2P361090      | 43,341,287    | 0.235       | 4.81 × 10^−6 |
| 8   | BICF2P543725      | 43,371,261    | 0.235       | 4.81 × 10^−6 |
| 8   | BICF2S23137831    | 43,418,611    | 0.235       | 4.81 × 10^−6 |
| 8   | BICF2P465920      | 43,487,284    | 0.263       | 5.56 × 10^−6 |
| 8   | TIGRP2P114211_sn8542415 | 43,494,231 | 0.263       | 5.56 × 10^−6 |
| 8   | BICF2S23110497    | 25,810,719    | 0.207       | 5.81 × 10^−6 |
| 8   | BICF2P177234      | 43,520,222    | 0.236       | 7.36 × 10^−6 |
| 19  | BICF2G630255452   | 47,856,573    | 0.335       | 9.05 × 10^−6 |
| 8   | TIGRP2P114933_rs9187825 | 46,799,348 | 0.405       | 1.36 × 10^−5 |
| 8   | BICF2S23235533    | 15,314,523    | 0.251       | 1.53 × 10^−5 |
| 8   | BICF2S2921051     | 15,005,970    | 0.260       | 1.56 × 10^−5 |
| 8   | BICF2P1102123     | 43,411,814    | 0.256       | 1.85 × 10^−5 |
| 8   | BICF2P1109401     | 43,462,069    | 0.256       | 1.85 × 10^−5 |
| 8   | BICF2P146090      | 43,425,535    | 0.256       | 1.85 × 10^−5 |
| 8   | BICF2P396875      | 43,463,543    | 0.256       | 1.85 × 10^−5 |
| 8   | BICF2P755461      | 43,441,286    | 0.256       | 1.85 × 10^−5 |
| 8   | BICF2P762487      | 43,454,904    | 0.256       | 1.85 × 10^−5 |
| 8   | BICF2S2932019     | 46,809,268    | 0.493       | 2.18 × 10^−5 |
| 31  | BICF2P1366177     | 7,605,782     | 0.102       | 2.58 × 10^−3 |
| 31  | BICF2S2443709     | 7,615,165     | 0.102       | 2.58 × 10^−3 |
The genotypes of each phenotypic group for the top SNP (chr8, BICF2P361090) are presented in Table 4. Some groups consist of low sample numbers, and thus we cannot draw any definite conclusions (“head” and “L1”). Groups “healthy” and “L2” show the ratio of individuals with AA and CA genotypes close to 50%, while groups “L3” and “L4” show that most individuals carry the genotype AA. The CC genotype exhibits a comparatively lower frequency in all groups (Table 4).

Table 4. Genotypes for the top SNP (chr8, BICF2P361090) from the QGWAS for each of the six phenotypic categories, with percentages shown in parentheses. The highest proportions in each phenotype group are underlined.

| Chr (SNP) | Genotype | Healthy (%) | Head (%) | L1 (%) | L2 (%) | L3 (%) | L4 (%) |
|-----------|----------|-------------|----------|--------|--------|--------|--------|
| chr8 (BICF2P361090) | AA | 52 (47) | 1 (33) | 4 (67) | 14 (50) | 43 (88) | 15 (79) |
|           | CA | 46 (41) | 2 (67) | 1 (17) | 13 (46) | 5 (10) | 4 (21) |
|           | CC | 13 (12) | 0 | 1 (17) | 0 | 1 (2) | 0 |
|           | missing | 0 | 0 | 0 | 1 (4) | 0 | 0 |
| total     | 111 | 3 | 6 | 28 | 49 | 19 |

For the additional GWAS analyses, we found associations with aRFA onset before 2 years of age (Table 5, Figure S3) and level 4 aRFA (Table S3, Figure S4). The top SNP in both analyses was on chromosome 21 (BICF2G630640798) with raw $p = 5.01 \times 10^{-7}$ and $p = 1.28 \times 10^{-6}$, respectively (Table 5, Figures S3, S4 and Table S3). Moreover, in the analysis of the early onset before 2 years of age (Table 5), the result shows a stronger association ($p = 5.01 \times 10^{-7}$; Bonferroni cut-off = $5.8 \times 10^{-7}$; Table 1). The results of the GWAS analyses of individuals older than 6 years and level 2 aRFA showed no significant associations. The average genomic inflation factor for all four additional GWAS analyses was 1.02 (range 1.00–1.05).

Table 5. Additional analysis (age of onset before 2 years of age) GWAS results showing chromosome (Chr), SNP name, position (bp), allele frequency, and raw $p$-value for the top twenty SNPs. One significant SNP was identified on chromosome 21. The interrupted line divides significant association from the rest. The significance threshold based on the Bonferroni correction was set to $5.8 \times 10^{-7}$.

| Chr | SNP Name | Position (bp) | Allele Freq | $p$-Value |
|-----|----------|---------------|-------------|-----------|
| 21  | BICF2G630640798 | 47,085,771 | 0.221 | $5.01 \times 10^{-7}$ |
| 23  | BICF2S23432401 | 11,113,618 | 0.377 | $5.75 \times 10^{-6}$ |
| 37  | TIGRP2P420015_rs8709645 | 20,114,103 | 0.262 | $6.48 \times 10^{-6}$ |
| 8   | chr8_59707832 | 59,707,832 | 0.221 | $8.66 \times 10^{-6}$ |
| 37  | BICF2G630131116 | 25,669,986 | 0.148 | $1.07 \times 10^{-5}$ |
| 15  | BICF2G630419811 | 59,659,531 | 0.434 | $1.22 \times 10^{-5}$ |
| 23  | BICF2P438054 | 11,110,146 | 0.352 | $1.30 \times 10^{-5}$ |
| 21  | BICF2G630641744 | 46,513,869 | 0.254 | $1.60 \times 10^{-5}$ |
| 27  | BICF2G630139626 | 42,94,734 | 0.205 | $1.92 \times 10^{-5}$ |
| 17  | chr17_40427743 | 40,427,743 | 0.426 | $2.87 \times 10^{-5}$ |
| 21  | BICF2S23427379 | 46,584,445 | 0.270 | $3.07 \times 10^{-5}$ |
| 20  | BICF2P1328442 | 55,962,058 | 0.320 | $3.49 \times 10^{-5}$ |
| 27  | BICF2P675588 | 34,378,821 | 0.295 | $3.70 \times 10^{-5}$ |
| 23  | BICF2G630386401 | 13,368,971 | 0.459 | $4.20 \times 10^{-5}$ |
| 18  | BICF2G630693935 | 34,387,737 | 0.484 | $4.30 \times 10^{-5}$ |
| 27  | BICF2G630139599 | 4,253,386 | 0.180 | $4.40 \times 10^{-5}$ |
| 27  | BICF2G630139609 | 4,266,185 | 0.180 | $4.40 \times 10^{-5}$ |
| 27  | BICF2G630139630 | 4,299,688 | 0.180 | $4.40 \times 10^{-5}$ |
| 27  | BICF2G630139642 | 4,318,805 | 0.180 | $4.40 \times 10^{-5}$ |
| 27  | BICF2S23028384 | 4,247,215 | 0.180 | $4.40 \times 10^{-5}$ |
3.4. Haplotype Identification

In order to adequately extend the area on chromosomes where the candidate genes could be located, we conducted a haplotype analysis. Based on the results of the case/control GWAS we looked closely at the haplotype distribution on chromosomes 19, 8, 30, and 36. Table 6 shows the results for the most significant haplotypes and the most significant haplotypes containing the most significant SNPs (from the case/control GWAS). On chr19, 4443 SNPs passed filtering and the most significant haplotype, consisting of the motif ATGGTCA (\( p = 2.09 \times 10^{-11} \)), was found in 84% of cases and 54% of controls. A single-base (A or G; \( p = 2.03 \times 10^{-6} \)) haplotype containing the top chr19 SNP from the case/control GWAS study was found, with the A-haplotype in 54% of cases and 77% of controls, and the G-haplotype in 45% of cases and 24% of controls. On chr8, 6105 SNPs passed filtering and the most significant haplotype (AAG; \( p = 7.31 \times 10^{-8} \)) was found in 75% of cases and only 49% of controls. A haplotype containing the suggestive chr8 SNP from the case/control GWAS was found (GGG; \( p = 2.25 \times 10^{-7} \)) in 85% of cases and 62% of controls. On chr30, 3922 SNPs passed filtering and the most significant haplotype (GCGA; \( p = 5.04 \times 10^{-6} \)) was found in 16% of cases and 36% of controls. A haplotype containing the suggestive chr30 SNP from the case/control GWAS has the motif ATACAGGA (\( p = 1.45 \times 10^{-5} \)) and was found in 22% of cases and 41% of controls. On chr36, 2746 SNPs passed filtering and the most significant haplotype (CC; \( p = 3.5 \times 10^{-5} \)) was found in 37% of cases and 19% of controls. This haplotype also contains the suggestive chr36 SNP from the case/control GWAS.

Table 6. Haplotypes for chromosomes revealed by the case/control and quantitative GWAS and subsequent haplotype analysis. For each chromosome we show the most significant haplotype and a haplotype containing the most significant or suggestive SNP for each chromosome (marked by *).

| Chr | bp            | Haplotype  | % Cases | % Controls | p-Value     |
|-----|---------------|------------|---------|------------|-------------|
| 8   | 43,341,287–43,356,221 | AAG        | 75.0    | 49.4       | 7.31 \times 10^{-8} |
| 8 * | 43,463,820–43,494,231 | GGG        | 85.0    | 62.4       | 2.25 \times 10^{-7} |
| 19  | 19,807,697–20,172,164 | ATGGTCA    | 84.4    | 53.9       | 2.09 \times 10^{-11} |
| 19 *| 47,856,573   | A          | 54.7    | 76.5       | 2.03 \times 10^{-6} |
| 19 *| 47,856,573   | G          | 45.3    | 23.5       | 2.03 \times 10^{-6} |
| 30  | 26,126,946–26,143,675 | GCGA       | 15.8    | 35.5       | 5.04 \times 10^{-6} |
| 30 *| 26,245,545–26,328,881 | ATACAGGA  | 21.5    | 41.3       | 1.45 \times 10^{-5} |
| 36 *| 28,573,704–28,584,717 | CC         | 36.7    | 18.8       | 3.53 \times 10^{-5} |

3.5. Candidate Genes Identified by GWAS

Using the results of the different GWAS analyses with subsequently constructed LD plots (Figure S5) and the abovementioned haplotype analyses, we identified 144 potential candidate genes within a 2–4 MB window using the most significant and several suggestive SNPs on each chromosome (11 genes on chr19, 61 genes on chr8, 60 genes on chr30, and 11 genes on chr36) (Figure S5 and Table S4). Given the nature of aRFA and the available scientific information regarding RFA and other non-inflammatory alopecic disorders in dogs, we focused mainly on genes associated with circadian rhythm and keratin metabolism. We identified eight genes that met these criteria (CSNK2A1, PIF1, RORA, TCF12, FUT8, ZFP36L1, RNF111, SNX22) [www.genecards.org; www.pathcards.genecards.org, accessed on 19 February 2022]. The mRNA expression of four out of the 144 GWAS candidate genes have been previously associated with different HC stages (telogen—GULP1, anagen—PCLAF, PIF1, TLN2) [56]. A spreadsheet summarizing all GWAS candidate genes is shown in Table S4.

3.6. Histopathological Phenotyping and Sample Selection

To identify a precise histological phenotype of aRFA, we examined skin biopsies of aRFA affected and control dogs. The histological phenotype in all biopsies from the control...
dogs as well as samples of unaffected haired skin from alopecic dogs was histopathologically unremarkable (Figure 3A). HFs were predominantly in anagen, and the inferior portion of the HFs extended deep into the panniculus. Few follicles were in telogen or kenogen. Sebaceous glands appeared normal, and the epidermis was unremarkable.

All biopsies from the affected skin of alopecic animals displayed similar features previously described in typical RFA cases (Figure 3B–F). Anagen follicles were absent. Infundibuli were moderately to severely dilated, sometimes appeared long, and were filled with abundant orthokeratotic keratin, which was laminar to compact and extended into the openings of the secondary follicles, resulting in a “witch’s feet”-like appearance (Figure 3E). The follicular parts proximal to the infundibula were shortened and limited to the dermis (Figure 3B–E). In some sections, rare telogen (Figure 3C) or kenogen (Figure 3F) follicles could be identified but often only the outer root sheath was visible and a definitive follicular stage could not be assigned (Figure 3B–D). Atrophic follicles were present in some biopsies (Figure 3F). A mild distortion of the HFs could be observed (Figure 3B–F). The sebaceous glands appeared multifocally prominent. The epidermis was mildly hyperplastic and covered by mild to moderate basket-weave, orthokeratotic keratin. Excessive pigmentation was not seen.

![Figure 3. Histological representation of biopsy samples from control skin from unaffected dogs (A) and affected dogs (B–F). Note numerous anagen hair follicles in A identified by the presence](image-url)
of numerous hair bulbs (black arrows). In aRFA, infundibuli (gray arrows) are moderately to severely dilated (B, C, E) and are filled with abundant keratin, which extends into the openings of the secondary follicles, resulting in a “witch’s feet”-like appearance (E). The follicular parts proximal to the infundibuli are shortened and limited to the dermis (B–E). A few telogen follicles (C, white arrow) or kenogen follicles (F, white arrow) can be identified. Follicular atrophy may be seen (F, black cross). A mild distortion of the HF is observed (B–F). All samples are stained with hematoxylin and eosin (H&E) and the scale bars represent 200 microns.

3.7. RNA Sequencing Analysis

Single-end sequencing of the fifteen RNA libraries produced a mean number of 37,377,469 million (M) reads per sample on average (range: 31,686,907–43,209,034 M). The mean percentage of reads uniquely mapped to the genome was 90.19%, ranging from 87.50–91.59%. Among those, 80.84% on average mapped to the annotated canine transcriptome (range: 75.65–86.25%), resulting in 26 M counts per sample on average.

To identify genes that were differentially expressed in alopecic and healthy skin, we conducted a transcriptome analysis. A PCA plot was constructed based on gene expression profiles (controls n = 6; normal skin of affected dogs n = 4, alopecic skin of affected dogs n = 5) and demonstrates distinct clustering of samples from control dogs and biopsies of unaffected skin from affected dogs compared to alopecic skin of dogs with aRFA (Figure 4). Based on these clear clustering results, we combined samples from control dogs and healthy skin samples from dogs affected by aRFA (n = 10) and compared those with samples of affected skin from dogs with aRFA (n = 5) for further analysis.

Figure 4. Principal component analysis (PCA) of samples demonstrating clustering based on expression profiles plotted against the two most variable components (PC1 and PC2). Samples from control animals (red) and normal skin from affected animals (green) cluster together, whereas samples from alopecic skin from affected animals (blue) are clearly separated from the clusters representing normal skin but show a higher inter-group variability.

We identified a total of 1435 DEGs with an adjusted p-value of <0.01. Of these, 669 genes (46.6%) were downregulated, whereas 766 (53.4%) genes were upregulated in alopecic skin samples from affected dogs (Table S5). Of all deregulated genes, 135 were strongly upregulated with a log2fold change of at least 2 and 101 genes were strongly downregulated with a log2fold change of at least −2 (Table S5). Twenty-five of the DEGs have previously been associated with HF morphogenesis or HC in the literature (Table 7). With the exception of DLX5, LGR6, and NFATC2IP, all of the HC-associated genes were downregulated and most of them were associated with the WNT or SHH (Sonic Hedgehog)
pathway. Only four genes could be considered strongly downregulated, however, on the lower end of the scale (Log2FC ranging between $-2.00$ and $-2.89$; Table 7).

Table 7. Differentially expressed genes associated with a role in hair follicle (HF) morphogenesis and the hair cycle (HC) identified by transcriptome analysis comparing unaffected skin of control dogs and dogs affected with aRFA (n = 11) with alopecic skin of dogs with aRFA (n = 5).

| Gene Symbol | Full Gene Name | Described Function | Signaling Pathway | BaseMean | Log2FC |
|-------------|----------------|--------------------|-------------------|----------|--------|
| CTNNB1      | catenin beta   | promotes HF growth | WNT               | 20,928.210 | $-0.498$ |
| CUX1        | Ctx1, cut like homeobox 1 | inhibitor of HF differentiation | NOTCH | 1600.827 | $-0.895$ |
| DLX1        | distal-less homeobox 1 | HF cycling and differentiation | WNT | 48.951 | $-2.120$ |
| DLX2        | distal-less homeobox 2 | HF cycling and differentiation | TGF-b | 32.891 | $-1.825$ |
| DLX3        | distal-less homeobox 3 | HF cycling and differentiation | WNT | 1315.187 | $-1.154$ |
| DLX5        | distal-less homeobox 5 | HF cycling and differentiation | BMP | 85.608 | 1.026 |
| FGF5        | fibroblast growth factor 5 | catagen induction | FGF | 124.674 | $-2.896$ |
| FOXE1       | forkhead box E1 | governs HF stem cell (SC) niche | SHH | 475.559 | $-1.575$ |
| FOXC1       | forkhead box N1 | HF development, HS differentiation | WNT, BMP, SHH | 1199.807 | $-1.476$ |
| FZD2        | frizzled class receptor 2 | receptor WNT pathway | WNT | 155.689 | $-0.948$ |
| FZD3        | frizzled class receptor 3 | receptor WNT pathway | WNT | 412.223 | $-0.978$ |
| GLI1        | GLI family zinc finger 2 | HF SC related transcription factor | SHH | 541.121 | $-0.927$ |
| HHIP        | hedgehog interacting protein | HF organogenesis | SHH | 185.270 | $-2.185$ |
| HOXC13      | homeobox C13 | HS differentiation | WNT | 1064.553 | $-1.734$ |
| JAG1        | Jagged 1 | HF maintenance | Notch | 5310.336 | $-0.668$ |
| LEP1        | lymphoid enhancer binding factor 1 | HS differentiation | WNT | 957.142 | $-1.636$ |
| LGR4        | leucine rich repeat containing G protein-coupled receptor 4 | delays HC; inhibits activation of follicular SCs | WNT | 4786.889 | $-0.464$ |
| LGR5        | leucine rich repeat containing G protein-coupled receptor 5 | follicular SC marker; anagen initiation | WNT | 1028.425 | $-1.430$ |
| LGR6        | leucine rich repeat containing G protein-coupled receptor 6 | SC associated marker | WNT | 336.586 | 1.145 |
| LHX2        | LIM homeobox 2 | HF differentiation, SC associated marker | WNT | 645.982 | $-1.377$ |
| MSX2        | Msh homeobox 2 | HS differentiation | BMP | 295.450 | $-1.454$ |
| NCAM1       | neural cell adhesion molecule 1 | expressed in dermal papilla | FGF | 267.400 | $-1.642$ |
| NEAT2C1P    | nuclear factor of activated T cells 2 interacting protein | aging of HF stem cells | SHH | 1056.715 | 0.332 |
| SHH         | Sonic hedgehog | HF development and cycling | SHH | 51.377 | $-2.002$ |
| SMO         | Smoothened | HF development and cycling | SHH | 781.216 | $-0.858$ |

An analysis of the deregulated genes showed that some (n = 12) were associated with vitamin D and steroid hormone metabolism; however, of these only HSD3B2 could be considered strongly upregulated (Table 8).
Table 8. Differentially expressed genes associated with either vitamin D or steroid hormone metabolism comparing unaffected skin of control dogs and dogs affected with aRFA (n = 11) and alopecic skin of dogs with aRFA (n = 5).

| Gene Symbol | Full Gene Name | Function | BaseMean | Log2FC |
|-------------|----------------|----------|----------|--------|
| CYP27B1     | cytochrome P450 family 27 subfamily B member 1 | activates vitamin D3 | 209.288 | −1.650 |
| CYP2R1      | cytochrome P450 family 2 subfamily R1 | major vitamin D25-hydroxylase | 228.196 | 0.980 |
| CYP39A1     | Cytochrome P450 Family 39 Subfamily A Member 1 | 7-alpha hydroxylation of 24-hydroxycholesterol | 753.872 | −0.818 |
| CYP51A1     | cytochrome P450 family 51 subfamily A member 1 | cholesterol biosynthesis | 1933.523 | 0.712 |
| DHCR7       | 7-Dehydrocholesterol reductase | converts 7-dehydrocholesterol (substrate for vitamin D formation cholesterol) | 2100.759 | 0.719 |
| ESR2        | estrogen receptor 2 | nuclear receptor, expressed in the HF in outer root sheath, dermal papilla, matrix cells and in the bulge | 180.354 | −1.212 |
| HSD17B2     | 17β-Hydroxysteroid dehydrogenase 2 | inactivation of estrogens and androgens: converts estradiol to estrone, testosterone to androstenedione, and androstenediol to DHEA; activates the weak progestogen 20α-hydroxyprogesterone into the potent progestogen progesterone | 2720.505 | 1.061 |
| HSD17B6     | 17β-Hydroxysteroid dehydrogenase 6 | androgen catabolism: convert 3 alpha-adiol to dihydrotestosterone and androsterone to epi-androsterone. | 310.914 | 0.762 |
| HSD17B7     | 17β-Hydroxysteroid dehydrogenase 7 | biosynthesis of estrogen and cholesterol | 1341.347 | 0.687 |
| HSD3B2      | hydroxy-delta-5-steroid dehydrogenase, 3 beta- and steroid delta-isomerase 2 | biosynthesis of all classes of hormonal steroids | 154.717 | 2.406 |
| RXRG        | retinoid X receptor gamma | increases transcriptional function of VDR | 147.075 | −1.132 |
| VDR         | vitamin D receptor | nuclear transcription factor, absence leads to defects in HF regeneration and alopecia | 3347.023 | −1.211 |

3.8. Functional Classification of DEGs

The online Reactome pathway analysis revealed that amongst the genes downregulated in the skin biopsies, there is an overrepresentation of pathways involved in the organization and assembly of the extracellular compartment and signal transduction. For
the latter specifically, 17 downregulated genes identified were involved in the SHH and WNT signaling pathways (Table 7). Pathway analysis of the upregulated genes identified that 251 genes (33%) were involved in metabolism generally, whereas 135 (18%) of the upregulated genes were specifically related to the metabolism of lipids. Among the most relevant pathways identified for up and downregulated genes, no common pathways were found (Tables S6 and S7, respectively).

3.9. Protein–protein Interaction Analysis

To discover functional connections between GWAS candidate genes and RNA-seq deregulated genes, we conducted a protein–protein interaction analysis in STRING. The GWAS candidate genes showed enriched protein–protein interactions. Specifically, they were predicted to be connected in six clusters of more than three proteins each (Figure S6), altogether comprising 96 interaction edges (significant enrichment of interactions; observed N edges = 96, expected N edges = 20, \( p\)-value < 1.0 \( \times \) 10\(^{-16}\)). We identified very limited overlap between the 144 GWAS candidate genes and the 1435 DEGs in the skin biopsies. Only 11 identical genes (SLC25A3, GTDC1, ARG2, PAPLN, RAD51B, RDH11, HACD3, LACTB, SNX1, TPM1, TRIP4) were identified in the GWAS study and RNA-seq experiment and their overlap was not significant (\( p\)-value = 0.61, Fisher’s exact test; Table S8). Moreover, no functional connections between the 11 identical genes were found. Due to the lack of overlap between GWAS candidates and DEGs, we speculated that they may be linked at the level of functional processes. This would mean that genetic changes identified in the GWAS analysis resulted in gene expression shifts of the interacting protein partners. Thus, we sought evidence that GWAS candidate genes d with the DEGs, which would identify the molecular processes related to aRFA. We took advantage of protein interaction information from the STRING database, which provides an estimate of proteins’ joint contributions to a shared function [53]. For each GWAS candidate, we searched for the presence of STRING interactors with proteins coded by DEGs. To focus only on genes with a likely stronger impact on aRFA, we limited our list of DEGs to those strongly deregulated in the skin biopsies, exceeding the Log2FC value of +/− 2. Following this approach, we found that out of the 144 GWAS candidates, 40 were predicted to interact with at least one of the 236 strongly DEGs. In fact, thirteen interacted with more than one strongly DEG (Figure 5).

Several large STRING clusters were identified in this analysis (Figures 5 and S7). The largest of these clusters were centered on collagen formation, muscle structure/contraction, the immune system, and lipid metabolism. Some of these cohorts were already seen among the most significant functional pathways of deregulated genes (Tables 7 and 8). Taken together, STRING analysis supports our hypothesis of distinct molecular functions being involved in the pathogenesis of aRFA.
Figure 5. Interactions of GWAS candidate genes (green) and STRING-associated strongly differentially expressed genes colored by their level of expression. We used only medium confidence associations and higher (increasing thickness of lines connecting genes indicates greater confidence). Colorful bubbles represent the metabolic pathways common for each cluster of genes.

4. Discussion

Atypical recurrent flank alopecia in the Cesky Fousek is a disease that is influenced by genetic factors. Pedigree analysis shows that aRFA is more prevalent in some families than in others. Dostál et al. [57] suggest that aRFA in the Cesky Fousek is a recessive disease with incomplete penetrance. They conducted a simple statistical analysis of the offspring of parents that were affected, healthy, or a combination of both [57]. It is believed that the incomplete penetrance is dependent on environmental factors, such as housing and nutrition.

In this study, we identified two significant associations with aRFA in the Cesky Fousek (chr19 and chr21) and other suggestive associations on chromosomes 8, 36, and 30. The suggestive associations were used for the candidate genes’ identification because these
variants may sometimes help us to get more complete information about the connection of the genes to the phenotype [58]. The significance of the GWAS-identified variants in our study is comparable to other GWAS studies of complex genetic diseases in dogs (e.g., lymphoma, elbow dysplasia, mast cell tumor) [2], suggesting that aRFA likely has a polygenic inheritance. Even though we used a within-breed design, our dataset was rather small for a complex disease GWAS. Follow-up analysis using a larger sample size is needed to confirm these findings. Interestingly, the region identified in our study on chr19 maps to chr2 (144,837,140–147,020,527bp) in the human genome (hg38), and this region has been associated with male pattern baldness [59]. However, the significance of this association was relatively low ($p = 5.65 \times 10^{-10}$; 181 out of 287 associated regions), and thus there is a possibility that the overlap might have happened by chance.

In total, we identified 144 GWAS candidate genes based on significant and suggestive associations on chromosomes 19, 21, 8, 30, and 36. Genotype analysis for the chr19 variant (BICF2G630255452) did not reveal a clear pattern between cases and controls; nevertheless, most of the control individuals were of genotype AA while most of the cases were of genotype GG and GA, suggesting the G allele is associated with higher risk for aRFA. The frequency of genotypes on chr8 (BICF2P361090) identified by QGWAS suggest that the CC genotype is associated with a lower risk of aRFA occurrence. Proportionally, more individuals severely affected by disease (level 3 aRFA and level 4 aRFA) were of genotype AA, while mildly affected (level 2 aRFA) and healthy individuals were of genotype CA. The genotype could be a contributing factor and, along with environmental factors, may influence the severity of the disease.

There are several limitations to the current GWAS study. One of them is the uncertainty of the development of aRFA in individuals from the control group. aRFA may manifest later in life; therefore, some individuals may later be reclassified. Ideally, we would use only animals aged 10+ years; however, due to the small population size of the breed this was not possible. We believe that the lack of an age threshold does not strongly affect the results of our study, because the average age of aRFA onset for our affected group was 3.9 years and out of 117 controls only five were slightly under this average age of manifestation. Another possible limitation is that the causative variants might be fixed in the population or at a high frequency; thus, the GWAS method would not detect them in a similar manner to the detection of obsessive compulsive disorder in Doberman Pinchers [60]. In this case, the identified significant and suggestive associations may only be modifiers of the causative variants. We also need to consider that the Bonferroni cut-off for the case/control GWAS is rather high. There is a discussion about multiple testing corrections and which cut-off threshold to use in order to find truly significant results [61]. Although there are several concerns about GWAS in general, it is still considered the best method for detecting associations between SNPs and hereditary diseases, which can lead to the identification of possible causative genes and variants [61]. In order to overcome the limitations of the current study, it is necessary to conduct future research with more individuals, ideally from multiple dog breeds.

Transcriptome analysis revealed 1435 deregulated genes and the vast majority of these genes are also present within microdissected anagen and/or telogen HFs [56]. Only 43 (3%) of the deregulated genes in the alopecic skin biopsies of dogs with aRFA have not been identified in microdissected HF, suggesting that these genes are derived from the HF macroenvironment. The HF macroenvironment is gaining more and more attention and it is well known that the cyclical regeneration of the HF is not only controlled by factors derived from the follicular microenvironment but also from the dermal macroenvironment [7,62–64].

Among the deregulated genes in our study, 25 (1.7%) genes can be clearly identified as being involved in HF development, specific HC stages, follicular SCs, or the HC. Most of them are related to the WNT and SHH pathways, which are known to be important for anagen induction, promotion, and differentiation [7,14,65–68]. Recently, Alopecia X, another noninflammatory alopecic disorder with a presumed hereditary background,
was also connected to altered WNT and SHH pathways [25]. Specifically, in our study, thirteen genes (CTNNB1, CUX1, DLX2, DLX3, CTNNB1, HOXC13, FOXN1, FZD2, FZD3, LEF1, LHX2, LGR4, and LGR5) encoding transcription factors or signaling molecules of the WNT signaling pathway were all downregulated in affected skin samples. Equally, genes associated with the SHH pathway, including FOXE1, GLI1, HHIP, SHH, and SMO, were also downregulated. Among the genes involved in inhibiting anagen induction and in the BMP signaling pathway [14], only DLX5 was upregulated. Furthermore, MMP7, an antagonist of the WNT pathway, was also upregulated, indicating that there is likely an active inhibitory component of the WNT pathway involved in the HC arrest observed in aRFA [69]. Interestingly, genes associated with the HF stem cell niche, namely GLI2, LGR5, LHX2, and FOXE1 are all downregulated suggesting that impaired SC function is associated with the development of aRFA [14,70,71]. It has been shown that HC is dependent on a fully functional SC compartment [25,72] and it is possible that an altered SC compartment might be responsible for the extreme short follicles seen in histology. In conclusion, the findings of the RNA-seq experiments are compatible with the results of the histopathological examination showing a lack of anagen HFs and shortened follicles. The exact HC stage reflected by these shortened follicles cannot be assigned morphologically. Eventually, the downregulated genes associated with follicular SCs result in a completely dysfunctional HC that does not allow clear HC stages to be defined. This would also be supported by the histological observation of numerous HFs that have a dystrophic appearance. It is, however, still unclear whether the deregulated genes are the cause or the consequence of the HC arrest.

Eleven identical genes were identified in both the GWAS and the transcriptome analysis. While these concordant findings might imply that they represent the core genes involved in aRFA, we see two reasons that this is not necessarily the case. First, these genes were not predicted to interact, suggesting that they have roles in very different molecular processes. Second, although the 11 genes were classified as differentially expressed based on the expression significance, their actual expression shift was negligible (Log2FC ranging between 0.31 and 1.04), suggesting only a subtle impact of their differential expression on the organism.

In addition to the abovementioned DEGs associated with the WNT, SHH, and BMP signaling pathways 236 DEGs identified in the skin biopsies were predicted to interact with 40 genes of the GWAS study using the STRING database (Figure 5). They were associated with collagen formation, muscle structure/contraction, Lipid metabolism, and immune metabolic pathways (Figure 5). We identified significantly more interactions than expected in the network (enrichment p-value < 1.0 × 10^{-16}). In Figure 5, several regulators of different molecular processes are present, such as DNA repair (PCLAF aka KIAA0101), cell cycle (CCNB2), cellular and intracellular trafficking (SLC10A1, HERC1, RAB11A), cell signaling (ITGAV, RGS6, PSEN1), etc. Some of these regulators might be responsible for the altered function of the interacting DEGs.

Lipids have been implicated in three possible mechanisms for disrupting hair growth [73]. Two of the mechanisms are unlikely in aRFA but one mechanism suggested in this review, namely that an inherently altered lipid metabolism state may be linked to the HC by affecting signaling proteins involved in the SHH or WNT pathways, is a possibility [74]. This assumption is supported by the fact that several genes encoding for molecules involved in these pathways are deregulated in the skin biopsies of dogs with aRFA. If these signaling pathways are impaired, due to, for example, a sterol precursor accumulation, the induction and promotion of the anagen HC phase is impossible, resulting in alopecia [75]. It was found that obesity had a negative impact on HF SCs and can cause hair thinning [76,77]. In our study, 30 out of 100 affected individuals (30%) were identified as overweight, while only 10 out of 116 controls were overweight (11.6%; Table S1). Obesity could be a contributing factor to aRFA in some individuals. Amongst other lipids, cholesterol is of particular importance for the skin. It is crucial for keratinocyte differentiation, has an important barrier function, and is a precursor for steroid hormone synthesis in the skin [78].
Interestingly, in the skin biopsies of dogs with aRFA seven deregulated genes are encoding proteins, mainly enzymes, involved in sex hormone or cholesterol biosynthesis.

Cholesterol (7-Dehydrocholesterol) is also a precursor of vitamin D3 (cholecalciferol) under UV radiation [79]. Vitamin D3 is important for the skin. A mutation of Vitamin D receptors has been previously connected to Alopecia Totalis and a knockout of vitamin D receptors in mice stopped the initiation of the new HC [63,78,80–82]. In our study, we identified several downregulated genes associated with the vitamin D metabolism in affected skin samples (Table 8). Vitamin D has been shown to play an essential role in the biosynthesis of estradiol in mice and pigs [83]. It is well known that keratinocytes are the primary source of vitamin D and its active metabolite is processed in the skin, supporting a local deregulation of the estrogen metabolism partially mediated by vitamin D [81]. Thus, it might be interesting to explore the role of cholesterol on the HC further to identify new drugs targeting the control of cholesterol in the skin. Interestingly, genes associated with sex hormone metabolism were also downregulated in the skin biopsies of dogs affected by aRFA. The degree to which sex hormone biosynthesis and metabolism, which involves the hormones and enzymes of the complex hypothalamic–pituitary–gonadal axis, is associated with vitamin D metabolism and is involved in the pathogenesis of aRFA remains to be further explored. A disrupted sex hormone metabolic pathway and deregulated vitamin D metabolism have also been identified in another alopecic disorder with a most likely hereditary cause [25]. Future studies evaluating the nutritional and hormonal status of affected vs. control dogs would be helpful to gain more insights into the role played by these pathways.

The muscle structure/contraction metabolic pathway that was identified as another relevant pathway affecting aRFA might be associated with the arrector pili muscles (APM). The APM has recently gained attention since it inserts close to the SC region of the HF, has been associated with impaired HF cycling [84] in humans, and is associated with impaired SCs function with age [85]. Conversely, the SCs of the HFs express genes facilitating the formation of tendons and ligaments and establish a niche for smooth muscle myoblasts that create the APM [86,87]. The overexpression of these genes result in a poor vascular and nerve supply of the SC niche and contributes to SCs quiescence [87]. Our results show mostly a strong upregulation of genes in the muscle structure/contraction pathway. Moreover, genes encoding follicular SCs in the skin biopsies were downregulated further, indicating that in dogs with aRFA this pathogenetic mechanism may also be involved.

Disrupted immune system metabolism might be a contributing factor to aRFA occurrence. We identified genes involved in the immune system cluster (Figure 5) that are involved in glucocorticoid regulation, enzyme and ion cellular transport, and inflammatory response [www.genecards.org, accessed on 19 February 2022]. While the inflammation seen in the skin biopsies of two individuals is most likely caused by an impaired epidermal barrier and does not have a primary genetic cause, an altered glucocorticoid regulation, enzyme function, or ion cellular transport might be associated with the HC.

When we identified genes (out of the 144 GWAS candidates) connected to the preliminarily chosen metabolic pathways that could have a connection with aRFA, we discovered four genes controlling the circadian rhythm (RORA, PIF1, TCF12, CSNK2A1) and four keratin-associated genes (FUT8, ZFP36L1, RNF111, SNX22) (Table S4). The genes controlling the circadian rhythm might be associated with the seasonality of the disease. Besides circadian rhythm metabolism, melatonin metabolism has also been discussed as one of the causative factors of RFA [27]. In the protein–protein analysis of interacting GWAS genes and DEGs shown in Figure 5, Kyneurine 3-Monoxygenase (KMO) and its paralog Coenzyme Q6 (COQ6) were identified. Interestingly, KMO is part of the nicotinamide adenine dinucleotide (NAD) biosynthesis II (from tryptophan) pathway and the tryptophan utilization superpathway, and is thus directly connected to the melatonin degradation pathway [www.pathcards.genecards.com, accessed on 19 February 2022]. Although it has not been shown for dogs with aRFA, more than 50% of dogs with RFA respond well to melatonin treatment [27].
5. Conclusions

In this comprehensive study, we investigated the genetics and the differential gene expressions associated with aRFA in the Cesky Fousek using a unique combination of techniques. We performed a genome-wide association study on 216 individuals, RNA-seq experiments from skin biopsies of 11 dogs, and examined the histopathological phenotype of dogs with aRFA. This is the first complex genomic study of canine alopecia in dogs using such an extensive sample size. Histologically, we found that aRFA is similar to RFA and compatible with an impaired HC. The mRNA of genes associated with the initiation and promotion of the HC, as well as of genes encoding for follicular stem cell markers, were mostly downregulated. These findings explain the lack of anagen follicles in the skin of affected individuals. In total, we identified 144 candidate genes from the GWAS analysis (including both the significant and suggestive associations) and 236 strongly deregulated genes from the RNA-seq analysis. Using only the significant GWAS candidate genes (n = 12), we did not discover a direct functional connection to the strongly deregulated genes. However, using the suggestive GWAS candidate genes we discovered four major metabolic pathways associated with aRFA—collagen formation, muscle structure/contraction, lipid metabolism, and the immune system. The findings from our study suggest that aRFA has a complex genetic inheritance that warrants further study.

Given the limitations of the GWAS analyses, further genetic studies involving independent and larger cohorts, including multiple breeds, are needed in order to validate our findings and determine the specific variants that contribute to aRFA risk.

Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/genes13040650/s1,
Figure S1: Two PCA plots of all individuals collected for the study (n = 216). The PCA plot shows genetic distances between individuals. Figure (a) shows relationships between affected (red) and healthy (blue) individuals and their geographic origin (circular points—individuals from the Czech Republic; triangle points—individuals from the US club CFNA). Note that individuals kept in the US but born in the Czech Republic were placed in the “CZ” group. Figure (b) shows relationships between three plates of samples genotyped in 2016 (plate 1), 2018 (plate 2), and 2019 (plate 3). Figure S2: Manhattan and quantile–quantile plot of quantitative GWAS. The top SNP $p$-value does not reach the significance threshold (Bonferroni correction = $5.8 	imes 10^{-7}$; shown as a purple line); however, many SNPs were identified on chr8 (colored in green) in close proximity to each other. We consider these associations suggestive. The lambda value shows that stratification correction worked well. Figure S3: Manhattan and quantile–quantile plot of GWAS analysis of individuals affected up to two years of age. Identified variants could have an effect on the time and/or severity of the affect. One variant can be considered significant (chr21, BICF2G630640798, raw p-value $= 5.01 	imes 10^{-7}$) as it reaches the significance threshold (Bonferroni correction $= 5.8 	imes 10^{-7}$; shown as a purple line), which is located on the gene ANO3. The lambda value shows that stratification correction worked well. Figure S4: Manhattan and quantile–quantile plot of GWAS analysis of individuals who were severely affected (level 4 aRFA). The chromosomes carrying the top three SNPs are shown in green. The significance threshold was based on the Bonferroni correction ($5.8 	imes 10^{-7}$; shown in a purple line). The top SNP does not reach the significance level; however, it is the same SNP as in the early onset GWAS. The lambda value shows that stratification correction worked well. Figure S5: Linkage disequilibrium (r2) plots for top SNP on chromosomes 19 and 8 from the case/control GWAS. The plots show LD among the top SNP on the chromosome and other SNPs within 2Mb or 4Mb (black—no LD; blue—weak LD, green—intermediate LD, yellow—moderate LD; red—strong LD). Figure S6: Predicted protein–protein interactions among the 144 GWAS candidate genes’ products. The GWAS gene candidates showed enriched protein–protein interactions. Specifically, they were predicted to be connected in six clusters of more than three proteins. Each cluster is connected by lines of different thickness that represent the strength of evidence of functional connection between genes. The color of each gene has not been changed from the STRING database output. Figure S7: Interactions of GWAS candidate genes (green) and STRING-associated strongly differentially expressed genes colored by their level of expression. Only medium- and high-confidence associations were used (increasing thickness of lines connecting genes indicates greater confidence). Colorful bubbles represent the metabolic pathways common for each cluster of genes. Couples and smaller clusters of interacting
genes are also shown. Table S1: List of sampled animals for GWAS study. The table contains number of samples; date of birth (DOB); information about aRFA status (severity L2 and more; Y/N); severity of the affection (‘0’—healthy individual; ‘L1’—ears affected; ‘L2’—body sides up to $10 \times 10$ cm affected; ‘L3’—body sides up to $10 \times 25$ cm affected; ‘L4’—body sides up to $10 \times 40$ cm affected; ‘head’—affected only on the head; age of aRFA onset (in years). Information about body condition was gathered using surveys ‘O’—overweight, ‘JR’—just right, ‘NA’—information not available. Table S2: Summary of patient data from animals included in the study for transcriptome analysis. Animals were included or excluded from RNA-sequencing analysis based on presence of consistent histological features and clinical picture. Table S3: Results of one of the additional GWAS analyses for two groups of individuals—healthy individuals of age 10+ and severely affected alopecic individuals (level 4 aRFA). The top SNP on chromosome 21 is the same SNP as in the analysis of early affect (Table 5); however, the significance level is not met. Table S4: List of all genes identified by GWAS analysis and haplotype analysis. The known function for each candidate gene (from www.genecards.org accessed on 19 February 2022) and a metabolic super pathway the gene is a part of (from pathcards.genecards.org) are stated. Yellow represents genes controlling keratin metabolism. Orange represents genes associated with the circadian rhythm. Table S5: Results of differential expression analysis investigating biopsies of alopecic skin of dogs with aRFA and control biopsies. Significantly differentially expressed genes sorted by Log2 fold change. BaseMean = mean of normalized read counts across all samples; LfcSE = standard error of the log2FoldChange; Stat = the log2FoldChange divided by lfcSE. Strongly downregulated genes are red ($n = 101$) while strongly upregulated genes are green ($n = 135$). Table S6: Top ten most relevant pathways identified using Reactome pathway analysis on downregulated genes. Table S7: Top thirteen most relevant pathways identified using Reactome pathway analysis on upregulated genes. Table S8: Fisher’s exact test of eleven overlapping genes from the GWAS study and the RNA-seq study. The overlap was found to be non-significant.

Author Contributions: Conceptualization, S.N., L.B.C., A.R.B., M.M.W. and B. Č.B.; data curation, S.N., A.M.S., J.J.H., M.A.T.B., V.J. and A.R.B.; formal analysis, S.N., A.M.S., J.J.H., M.A.T.B., M.B., V.J. and A.R.B.; funding acquisition, S.N., A.M.S., L.B.C. and B.C.B.; investigation, S.N., A.M.S., M.A.T.B. and A.R.B.; resources, S.N., L.B.C., M.M.W., and B.C.B.; supervision, A.R.B., M.M.W. and B.C.B.; writing—original draft, S.N., A.M.S. and J.J.H.; writing—review and editing, A.M.S., J.J.H., M.A.T.B., M.B., V.J., L.B.C., A.R.B., M.M.W. and B.Č.B. All authors have read and agreed to the published version of the manuscript.

Funding: This research was funded by The Czech University of Life Sciences, grant numbers CIGA 20185006 and IGA 20213106; the Czech Cesky Fousek Breeding Club (KCHČF); Cesky Fousek North America; the Ministry of Agriculture of the Czech Republic, grant no. 84 (24.4.2017); Nadace “Nadání Josefa, Marie a Ždenky Hlávkových”; Nadace Český literární fond; and the SPEZKO grant 2018 of the VETSUISSE faculty for Alexandria Marie Schauer.

Institutional Review Board Statement: Blood samples and biopsies were taken in the Czech Republic under the certificate of competency according to § 17 of the Act No. 246/1992 coll. on Protection of Animals No. CZ02355 and each owner signed an informed consent form. Blood samples in the USA were collected in accordance with the Cornell University Institutional Animal Care and Use Committee (IACUC) protocol #2005-0151, with owner informed consent.

Data Availability Statement: The data will be available on the Dryad platform (https://datadryad.org/stash/ accessed on 19 February 2022) if this paper is accepted.

Acknowledgments: The authors are especially grateful to the dog owners for contributing samples for use in this project and all sponsors for contributing to the funding of this research. We would also like to thank Vandana Raghavan for assisting with the genotyping and data curation, as well as the technicians at the histology lab of the Institute of Pathology of the Vetsuisse Faculty in Bern for providing high quality slides. Thanks also to Tosso Leeb from the Institute of Genetics from the Vetsuisse Faculty in Bern for offering his lab space and for fruitful discussions.

Conflicts of Interest: The authors declare no conflict of interest. The funders had no role in the design of the study; in the analyses or interpretation of data; in the writing of the manuscript; or in the decision to publish the results.
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