Overexpression of PaNAC03, a stress induced NAC gene family transcription factor in Norway spruce leads to reduced flavonol biosynthesis and aberrant embryo development

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Background

In plants, the NAC [for NAM (no apical meristem), ATAF (Arabidopsis transcription activation factor), CUC (cup-shaped cotyledon)] family of transcription factors (TFs) is one of the largest plant TF gene families. The gene family is estimated to comprise 117 members in Arabidopsis thaliana and 144 and 161 respectively in rice and poplar [1, 2]. The NAC gene family in conifers appears to be at least as large as in Arabidopsis and might possibly even be expanded [3]. The boreal forest in the Northern hemisphere is dominated by conifers, many of which are economically and ecologically important. Still, relatively little is known about how conifers, and other gymnosperms, sense and respond to abiotic and biotic stress. General knowledge about inducible defence responses and their regulatory pathways are primarily derived from studies in angiosperm model plants, which in some cases can be extrapolated to gymnosperm systems [4–9], despite their evolutionary divergence [10]. A recent study showed that the accumulation of flavonoids and the gene induction pattern in the flavonoid pathway correlated to the level of resistance in Norway spruce to the root rot fungus Heterobasidion annosum (Fr.) Bref. sensu lato (hereafter referred to as H. annosum s.l.) [9]. H. annosum s.l. is a complex of five closely related species [11, 12] that have partly overlapping host ranges. These results indicated a differential control of defence responses between resistant and susceptible genotypes.

NAC TFs were first identified in forward genetic screens as key regulators of developmental processes [13–16]. NAC proteins have been shown to regulate central developmental processes such as embryo patterning and vascular patterning in both angiosperms and gymnosperms [15–18]. However, NAC proteins are also one of the most important groups of differentially regulated TFs in plant defence [19–21]. NAC TFs commonly possess a conserved DNA-binding NAC domain at the N-terminus, which includes nearly 160 amino acids that are divided into five subdomains (A–E) [22]. The C-terminal regions of NAC proteins are highly divergent [13, 22] and confer the regulatory specificity of transcriptional activation [1]. Based on the phylogeny of and domain structure in the NAC proteins it is possible to structure and organize the functional specificities of the conserved NAC domains and the divergent C-termini [1, 17, 22]. The NAC subgroups, e.g. subgroup III-3 in Arabidopsis, which contains the stress-related NAC proteins, ANAC019, ANAC055, ANAC072, ATAF1 and ATAF2, have common unique C-terminal motifs dominated by a negatively charged matrix with a few conserved bulky and hydrophobic amino acid residues that form the transactivation domains [1]. This group of paralogous Arabidopsis NAC genes show co-expression in response to stress hormones [20, 21, 23] and several members are known to act as regulators of plant responses to abiotic [19, 20, 23] and biotic [20, 24, 25] stressors. Transgenic plants overexpressing members of this subgroup (ATAF1, ATAF2, ANAC019 or ANAC055) show increased susceptibility to necrotrophic pathogens such as Botrytis cinerea or Fusarium oxysporum [20, 21, 24, 25] while an anac019 anac055 double mutation [21] or expression of an ATAF1 repressor construct [24] lead to enhanced resistance against B. cinerea. Taken together, this suggests that subgroup III-3 NAC transcription factors may be important transcriptional integrators between biotic and abiotic stress. A number of NAC TFs with similarity to Arabidopsis subgroup III-3 NACs among the differentially regulated TFs in recent transcriptome studies of spruce responses to biotic stress [9, 26] indicate that spruce orthologs of well-characterized Arabidopsis NACs control similar programmes in spruce and Arabidopsis not only in plant development [17, 18] but also in plant responses to stress.

The aims of this study were to: I) analyse the classification and stress-induced expression pattern of H. annosum s.l.-induced Norway spruce NAC TFs; II) investigate the downstream target genes of PaNAC03 in Norway spruce; III) investigate if PaNAC03 had the capacity to regulate the promoter PaLAR3, a gene in the downstream regulation module. To address the first aim we queried sequence databases to identify homologous sequences, identified the modular structure and phylogenetic placement of H. annosum s.l.-induced Norway spruce NACs. We also determined the expression patterns of the H. annosum s.l.-induced NAC TFs in response to different stressors. To investigate downstream target genes of PaNAC03 Norway spruce cell lines overexpressing PaNAC03 were constructed and their transcriptome was compared with the wild-type Norway spruce cell line to identify misregulated genes. To address our last aim we isolated the promoter of PaLAR3 and fused it to the GUS reporter gene and performed transactivation studies of PaNAC03 and PaLAR3 in Nicotiana benthamiana.

Methods

Sequence search and phylogeny

Six putatively unique transcripts (PUT) with similarity to angiosperm NAC transcription factors (Table 1) identified in previous RNAseq experiments [9, 26] were used to query the Norway spruce genome portal (http://congenie.org/) using Blastn [27] and TAIR (https://www.arabidopsis.org/) and Genbank using Blastx. The significant hits were downloaded and nucleotide and amino acid sequence alignments were made with Picea sequences from Genbank and P. abies 1.0 [3]. For phylogenetic analysis of the identified Norway spruce NAC genes additional Norway spruce gene models were downloaded
from the Norway spruce genome portal and subgroup III-1, III-2 and III-3 Arabidopsis NAC amino acid sequences were downloaded from TAIR. The sequences were trimmed to the conserved N-terminal region and aligned with the Clustal W algorithm in MEGA 5.0 [28]. Phylogenetic trees were created using the Neighbor-joining algorithm in the same program with 1000 bootstrap values, p-distance estimations as a statistical model, uniform substitution rates and an estimation based on partial sequences with a cutoff value of 95%.

Predicted subgroup III-3 Norway spruce NAC protein sequences were inspected for presence of a conserved N-terminal [22] and C-terminal domains [1]. The charge and hydrophobicity of the predicted proteins were estimated with EMBOSS Pepinfo software [29], the hydrophobicity of the predicted amino acid sequences was plotted using Kyte & Doolittles hydrophobicity index with a window of 11 amino acids. Sequence identity and similarity analysis of the full length and C-terminal regions of the identified Norway spruce NAC proteins was performed with the ident and sim functions of the Sequence manipulation suite [30].

### Determination of gene expression patterns

#### Biotic and abiotic stress

Thirty-year-old trees of eight independent Norway spruce genotypes which are part of a Swedish clonal forestry program and grow in a stand situated at Årdala, Sweden, (59°01’ N, 16°49’ E) [31] were inoculated with *H. annosum* s.l. The inoculation and sampling procedures are described in detail in Danielsson et al. [9]: Briefly, three ramets per genotype and two roots per ramet were used in the experiment. On one root, a wooden plug colonized by *H. annosum* s.s. (Sä 16–4) [32] was attached to an artificial wound on the root surface with Parafilm; the other root was wounded only and sealed with Parafilm. Phloem samples (ca 90 mm² pieces) for RNA extraction were harvested at the start of the experiment (0 days post inoculation) and at 5 and 15 days post inoculation (dpi) and preserved in RNA later (Ambion) for subsequent RNA extraction.

Total RNA was isolated according to Chang et al. [33]. Poly (A) + RNA was purified and amplified using MessageAmpIII (Ambion). Purified amplified RNA (aRNA, 1 μg) from each genotype were reverse transcribed with the iScript cDNA synthesis kit (Bio-Rad). The cDNA synthesis was diluted 1:1 in deionized water. Each genotype was used as an independent biological replicate.

#### Plant stress hormone treatments

To analyse the response of candidate genes to stress hormones and compare it to the response to *H. annosum* s.l., two-week-old Norway spruce seedlings (Rörby FP-65, 09 L022–1001) were transferred under axenic conditions to Petri plates with filter paper (five seedlings/plate), moistened with fertilized liquid media [34] and treated homogenized *Heterobasidion parviporum* (Rb175). For treatments with methyl jasmonate (MeJA) or methyl salicylate (MeSA) as previously described by Arnerup et al. [7]. Every treatment was performed in triplicate. After 72 h, seedlings were immediately frozen in liquid nitrogen and stored at −80 °C until further use. Total RNA was isolated according to Chang et al. [33] after DNAse I treatment one μg of total RNA was reverse transcribed with the iScript cDNA synthesis kit (Bio-Rad).

#### Somatic embryo maturation treatment

Samples for analysis of *PaNAC03* expression levels during embryo development, was a generous gift from Drs. Irena Molina and Malin Abrahamsson. Briefly, samples were collected from five sequential developmental stages (classification based on Zhu et al. [35]): +PGR (Proliferating cultures + Plant growth regulators (PGR) five days after subculture), —PGR (Proliferating cultures — PGR five days after subculture), EE (Early embryos differentiated after

### Table 1 Norway spruce subgroup III-3 NAC genes and their closest homolog in Arabidopsis thaliana

| Isogroup      | Gene       | Congenie (BlastN) | E-value | Best hit in NCBI | E-value | Locus     | Annotation   |
|---------------|------------|-------------------|---------|------------------|---------|-----------|--------------|
| isogroup00240a | PaNAC03    | MA_8980g0010      | 0       | ABK26029         | 0       | AT1G01720.1 | ATAF1        |
| isogroup00812b | PaNAC04    | MA_264971g0010    | 0       | AAC32123         | 0       | AT1G77450.1 | ANAC032      |
| isogroup02038a | PaNAC05    | MA_5115g0010      | 0       | ABK26029         | 1.00E-99| AT1G77450.1 | ANAC032      |
| isogroup05528a | MA_86256g0010 | 2.32E-144       | ABK26029 | 2.00E-145         | AT1G01720.1 | ATAF1        |
|               | MA_64687g0010 | MA_75192g0010    | ABK26029 | 2.00E-127         | AT1G01720.1 | ATAF1        |
|               | MA_103386g0010 | MA_103386g0010   | ABK26029 | 9.00E-145         | AT1G01720.1 | ATAF1        |
| isogroup02925b | MA_8533126g0010 | 2.19E-111       | ABB16510 | 5.00E-82          | AT1G25580.1 | SOG1         |
| isogroup05889b | MA_23113g0010 | 1.83E-18        | no hit   | No hit            | No hit   | No hit    |             |

*Induced in both wounding and inoculation treatments

*Induced only in response to inoculation treatment

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one week on maturation medium); LE1 and LE2 (late early embryos developed after two and three weeks on maturation medium, respectively). Three independent samples were collected for every stage and frozen in liquid nitrogen and stored at −80 °C until extraction. Total RNA were extracted with the Spectrum Plant Total RNA kit (Sigma Aldrich) after DNase I treatment one μg of total RNA was reverse transcribed with the Quanta cDNA synthesis kit (Quanta Biosciences).

Quantitative reverse-transcribed PCR (qPCR)
For analyses of gene expression levels an aliquot of cDNA equivalent to 25 ng of RNA was used per 20 μL of PCR reaction using SsoFast EVAGreen Supermix (Bio-Rad) and a final concentration of 0.5 μM of each primer. Primers were designed using Primer3 software (http://primer3.wi.mit.edu/) with a melting temperature (Tm) between 58 °C and 60 °C, and amplicon length between 95 and 183 bp (Additional file 1). The thermal-cycling condition parameters, run on an iQ™ Multicolor Real-Time PCR Detection System (Bio-Rad), were as follows: 95 °C for 30 s; 40 cycles of 95 °C for 5 s, 58 or 60 °C for 20 s. Each run was followed by a melt curve analysis to validate the specificity of the reaction. A linear plasmid standard curve was used to measure the PCR efficiency in each of the experiments, and primer pairs with efficiency lower than 95% were discarded. Two technical replicates were prepared for each sample.

The relative expression was calculated using the 2ΔΔCT-method [36, 37], transcript abundance was normalized to the reference genes phosphoglucomutase [38], eukaryotic translation initiation factor 4A (elF4A) [39] and elongation factor 1-α (ELF1α) [5]. The stability of reference gene expression was assessed with the Bestkeeper tool separately for every experiment [40]. Differential expression between treatments were tested with Kruskal-Wallis- and Mann-Whitney U-tests using the GraphPad Prism5 software (GraphPad Inc.).

Transformation of Norway spruce
Full-length cDNA sequences of PaNAC03 were obtained by amplification with the specific primers PaNAC03FL (Additional file 1), designed based on comparison of full-length or partial sequences of P. abies, P. glauca and P. sitchensis homologues, from a pool of cDNA from Norway spruce bark inoculated with H. annosum s.l. For the PCR reaction we used Dream-Taq Polymerase (Fermentas). AttB1 and attB2 adapters were added to the 1148 bp product by PCR using Dream-Taq Polymerase. The resulting PCR product was recombined into the pDONR/Zeo (ThermoFisher) vector followed by LR recombination into pMDC32 vector [41]. The resulting vector was verified by test-digestion and sequencing.

Cell lines constitutively expressing PaNAC03 were established by Agrobacterium-mediated transformation of Norway spruce somatic embryogenic cell line 95:61:21, as described by Minina et al. [42]. In brief, pMDC32:: PaNAC03 and pMDC32:: GUS [42] was transformed into the Agrobacterium tumefaciens C58C1 strain with the additional virulence plasmid pTOK47. Transformed bacteria were then grown overnight with the appropriate selection and collected by centrifugation and resuspended in infiltration buffer (10 mM MgCl2, 10 mM MES, pH 5.5, and 150 μM acetosyringone) to an OD600 of 10. Seven days old Norway spruce suspension cultures and Agrobacterium was mixed in a 5:1 ratio and acetosyringone was added to a final concentration of 150 μM. The co-cultivation was allowed to proceed for 4 h. Thereafter the cells were plated on a filter paper placed on the top of solidified proliferation medium with PGR [43] and incubated at room temperature in the darkness for 48 h. Then, filters were transferred on solidified proliferation medium with PGR containing 400 μg ml⁻¹ timentin and 250 μg ml⁻¹ cefotaxime and incubated under the same conditions for 5 days. Subsequently, filter papers were transferred onto fresh solidified proliferation medium with PGR containing 20 μg ml⁻¹ hygromycin, 400 μg ml⁻¹ timentin, and 250 μg ml⁻¹ cefotaxime and subcultured onto fresh medium every week. The transgenic calli were picked from the plates after a month and transferred to solidified proliferation medium with PGR containing 20 μg ml⁻¹ hygromycin, 400 μg ml⁻¹ timentin, and 250 μg ml⁻¹ cefotaxime. Transgenic lines were maintained on proliferation medium with PGR and 20 μg ml⁻¹ hygromycin.

Nine transgenic lines were selected for DNA and RNA extraction for verification of the insert and expression levels respectively. To verify the transformation, DNA was extracted by homogenizing and boiling a 3–5 mm diameter callus in an Eppendorf tube in 20 μl 0.5 M sodium hydroxide at 95 °C, quickly centrifuging and diluting 5 μl of the supernatant in 495 μl 10 mM Tris–HCl pH 8. Five μl of the dilution was used in a 25 μl PCR reaction using DreamTaq (Thermo Scientific) and Hyg primers (Additional file 1).

Total RNA was extracted by using a modified CTAB extraction protocol [33]. After DNase I treatment (Sigma-Aldrich) cDNA was synthesized from 1 μg of total RNA using the iScript cDNA synthesis kit (BioRad). Expression levels of PaNAC03 was tested by qRT-PCR by using an iQ5 Multicolor Real-Time PCR Detection System (BioRad) and SsoFast EvaGreen Supermix (BioRad) as stated previously and two independent lines (4.1 and 4.2) with expression levels 1.7 times higher than the WT cell line were selected for maturation initiation, RNA sequencing and chemical analysis. The initiation of somatic embryo maturation in the overexpression lines and the control line was done
according to the protocol described by Filonova et al. [44], briefly for each line pre-weighed pieces of callus was placed on half strength LP medium for a week before the explants were transferred onto the maturation medium, the maturation response was scored after four and six weeks on maturation medium, embryos resembling the LE2, ME1 and ME2 stages [35] were noted.

Transcriptome profiling of PaNAC03 overexpression lines

RNA extraction and Illumina sequencing The two selected overexpression (OE) lines, 4.1 and 4.2, along with the WT line (95:61:21) were incubated on solidified proliferation medium with PGR at room temperature in the darkness for six days and approx. 7 mm diameter large calli were picked from the lines and frozen in liquid nitrogen. The samples were ground in a mortar in liquid nitrogen and extracted by using the RNasy Plant Mini Kit (Qiagen) using the RLT buffer and following the manufacturer’s instructions, thereafter the samples were treated with DNase I (Sigma-Aldrich). Three biological replicates per line were used for Illumina sequencing. The RNA integrity was analysed by using the Agilent RNA 6000 Nano kit (Agilent Technologies Inc.). Sequencing libraries were prepared at the SNP&SEQ Technology Platform (SciLifeLab, Uppsala) using the TruSeq stranded mRNA sample preparation kit according to the manual TruSeq stranded mRNA sample preparation guide. Sequencing was done using HiSeq 2500, paired-end 125 bp read length, v4 sequencing chemistry.

Filtering, mapping and differential expression The raw sequences were filtered by a nesoni clip for the read pairs using Nesoni 0.128 (http://www.vicbioinformatics.com/nesoni-cookbook/index.html#) (See Additional file 2 for scripts used). To enable alignments to a reference database we constructed a Bowtie reference from the ‘Trinity contaminant free’ dataset downloaded from the Norway spruce genome portal (http://congenie.org/) using Bowtie2 version 2.2.4 (http://bowtie-bio.sourceforge.net/bowtie2/index.shtml). The clipped read pairs were aligned to Trinity using Tophat version 2.0.13 [45]. The resulting alignment files from Tophat were provided to cufflinks version 2.2.1 to produce an assembly for each sample. The assemblies were then merged using cuffmerge (included in the cufflinks package). We then applied the newer workflow by running cuffquant (http://cole-trapnell-lab.github.io/cufflinks/manual/) that calculates transcript abundances from the single assembly file and the aligned read files produced by the Tophat run which was run separately for each sample. Differential expression analysis was performed with cuffdiff [45, 46].

Chemical analysis of Norway spruce overexpression lines

Norway spruce OE lines (4.1 and 4.2) overexpressing the PaNAC03 gene and the wild-type cell line 95:61:21 were grown in liquid proliferation medium without PGR for two weeks. Thereafter, the cells were collected and flash frozen in liquid nitrogen after which the samples were freeze-dried. The freeze-dried samples were ground using a ball mill. Once pulverized, the sample-weight was noted. Specialised metabolite content was assessed with the method described by Hammerbacher et al. [47].

Transactivation of pPaLAR3 by PaNAC03

PaLAR3 transactivation by PaNAC03 The PaLAR3 promoter has two allelic forms, PaLAR3A and PaLAR3B. Both were amplified from genomic DNA using pPaLAR3A and pPaLAR3B primer sets (Additional files 1 and Additional file 3). After amplification, they were cloned into pJET1.2 plasmids using the CloneJET PCR cloning kit (Thermo scientific). From this plasmid, PCR products were amplified with the pPaLAR3A_2 and pPaLAR3B_2 primer sets (Additional file 1). These two PCR products were subsequently cloned into the destination plasmid pCF201 which was adapted from the pGA580 vector used for Agrobacterium transformation [48] by overlap extension PCR. To be able to do so, the destination plasmid was amplified into two separate PCR products. For the first PCR fragment the primers TetA2 forward and PU5 reverse were used and for the second PCR fragment GUS forward and TetA2 reverse were used (Additional file 1). All the PCR product fragments were purified with the GeneJet PCR purification kit (Thermo Scientific) as instructed by the manufacturer’s protocol. The promoter fragments were separately combined with these destination fragments and amplified in a three fragment overlap extension PCR using the method from (Bryksin and Matsumura 2010) with the adaptation PCR protocol: Initial denaturation at 98 °C for 2 min, followed by three cycles of denaturation at 98 °C for 15 s, annealing at 60 °C for 2 min and elongation at 72 °C for 5 min, then 14 cycles of denaturation at 98 °C for 15 s, annealing at 60 °C for 30 s, elongation at 72 °C for 5 min, then the final elongation at 72 °C for 10 min. Single mutations (Additional file 3) in the PaLAR3A promoter were created by two fragment overlap extension PCR. Mut_XbaI_F or Mut_KpnI_F were combined with the TETA2_reverse primer to make the first fragment and Mut_XbaI_R or Mut_KpnI_R were combined with TetA2 forward for the second fragment. The two corresponding fragments were combined in an OEPCR with the same PCR conditions as described above. A double mutation was created by using Mut_XbaI_KpnI primers with the corresponding TETA2 primers and the same method was repeated.
The newly formed plasmids were isolated with DpnI restriction endonuclease [49]. The restriction mix was incubated at 37 °C for 15 min and deactivated at 80 °C for 5 min. 1 μl of DpnI treated OE-PCR product was transformed into chemically competent E. coli cells (One Shot™ TOP10 Competent Cells, Invitrogen) and shake incubated for a minimum of 3 h at 37 °C. Colony PCR screen was performed with screening primers (Additional file 1). Positive clones were selected on agar plates with tetracycline (5 μg ml⁻¹), and plasmids were isolated with the GeneJet Plasmid Miniprep Kit (Thermo Scientific). Transformation of Agrobacterium tumefaciens (strain C58C1-RS with the helper plasmid pCH32) was done with the heat-thaw method as described [50]. Cells were plated on agar plates with tetracycline (5 μg ml⁻¹), kanamycin (5 μg ml⁻¹) and rifampicin (50 μl ml⁻¹) and transformants were selected with colony PCR using the same primers as for E.coli.

The transactivation experiment is an adapted version of the one described in (Leborgne-Castel et al. 1999). Four to six weeks old Nicotiana bethaminiana plants were grown under a 16-h photoperiod at 23 °C. Infiltration occurred as described in (Voinnet et al. 2003). The following 1:1 mixes of A. tumefaciens harboring the different effector and reporter constructs were prepared. After 72 h, leaf disks were taken and GUS expression and total protein were measured. The GUS colorimetric assay was described in a protocol in Wilson et. al. [51] where 20 μl of cleared extract were added to 250 μl GUS assay buffer as well as to GUS assay buffer with 6 mM 4-Nitrophenyl β-D-glucuronide (PNPG). The reaction was incubated overnight covered in aluminum foil. OD₄₀₅nm was measured in a microplate reader of the type Fluostar Optima. The GUS activity was determined in mol PNP per minute and gram protein. The protein concentration was determined by the Bio-Rad protein assay [52]. Student t-tests were performed to calculate significant changes based on 6–12 biological replicates per measurement.

Results
Norway spruce contain multiple clade III-3NAC transcription factor gene family members
The RNAseq dataset from the time course study of H. annosum s.s. inoculated Norway spruce [9, 26] contained six putatively unique transcripts (PUTs) with similarity to NAC TFs, all PUTs had at least one blastn hit in the P. abies genome v1.0 gene catalogue and a phylogenetic analysis of Norway spruce, rice, poplar and Arabidopsis protein sequences show that the Norway spruce genome has at least seven NAC gene models (Fig. 1) which fall within subgroup III-3 described by Jensen et al. [1]. We essentially see four clades within subgroup III-3, the predicted amino acid sequence of six of these genes, including PaNAC03- PaNAC05, form a sister group to a clade with members from all angiosperm species including ANAC032, ATAF1, ATAF2, ANAC102. The Norway spruce clade and two other clades, one of them specific to rice, are distinctly separated from the ANAC019, ANAC055, ANAC072, PNAC118 and PNAC120 protein sequences (Fig. 1). The six sequences in the Norway spruce clade share a higher amino acid similarity with each other than with MA_75192p0010, which clusters closer to the ANAC019, ANAC055, ANAC072, PNAC118 and PNAC120 branch (Additional file 4 and Additional file 5).

PaNAC03 (MA_8980g0010), PaNAC04 (MA_264971g0010), and PaNAC05 (MA_5115g0010) correspond to isogroup00240, isogroup00812 and isogroup02038 respectively (Table 1) identified in the time course study of the Norway spruce’s transcriptional responses to H. annosum s.s. [9, 26]. The predicted proteins from PaNAC03 and PaNAC04 share a maximum of 81% identity and 90% similarity in the conserved N-terminal domains and 59% similarity over the complete predicted protein sequence (Additional file 5). The two sequences cluster closely in the phylogeny together with three other potential NAC genes, all highly similar (Additional file 5). The third expressed Norway spruce clade III-3 like NAC, PaNAC05, clusters outside this group of highly similar NAC sequences (Fig. 1) and the protein share approximately 40% identity on amino acid level with the PaNAC03 and PaNAC04 proteins.

The conserved N-terminal A-E motifs [22] were present in all the identified Norway spruce NACs (Additional file 4). The C-terminal region is highly conserved between PaNAC04, MA_103386p0010 and MA_86256p0010 and is dominated by polar and charged amino acids (Additional file 4). PaNAC03 share a common C-terminal motif (SEKEE (V/I) QSSFRLE, Additional file 4) with all Norway spruce clade III-3 NACs except PaNAC05. The C-terminal motifs in Norway spruce subgroup III-3 NACs are different from the negatively charged matrix with a few conserved bulky and hydrophobic amino acid residues in Arabidopsis subgroup III-3 NACs [1].

Pathogen-induced expression of clade III-3-like Norway spruce NACs
We selected PaNAC03 and PaNAC04 for expression analysis as representatives of NACs responding to both wounding and inoculation (PaNAC03) and of NACs primarily responding to inoculation (PaNAC04) in the time
Fig. 1 (See legend on next page.)
course study of Norway spruce transcriptional responses to *H. annosum* s.s. [9, 26], as these PUTs were the most highly expressed in either category. The qRT-PCR analysis showed that *PaNAC03* is significantly induced in response to both inoculation and wounding treatments (*P* < 0.05 for both treatments) compared to the control although the induction level was significantly higher after inoculation compared to wounding at 5 dpi (*P* = 0.01) (Fig. 2a). *PaNAC04* was significantly induced at 5 dpi both after wounding and inoculation with *H. annosum* s.s. (*P* = 0.008 and *P* = 0.004 respectively) compared to the control. The qRT-PCR data also showed that *PaNAC04* transcript levels were significantly higher after inoculation compared to the wounding treatment at 15 dpi (*P* = 0.02) (Fig. 2b). The responsiveness of *PaNAC03* and *PaNAC04* to *H. parviporum* inoculation or to plant defence hormones (MeJA and MeSA) was tested in young seedlings. Both genes were significantly induced in response to MeJA and MeSA treatments (Figs. 3a and b) but only *PaNAC03* was significantly induced in response to fungal inoculation (Fig. 3a).

**PaNAC03 overexpression in Norway spruce leads to altered developmental and metabolite profiles**

Eight selected hygromycin-resistant lines were verified to be transformed with pMDC32:: *PaNAC03*. Five of these lines were shown to moderately overexpress *PaNAC03*, 1.2-2.2 times the WT line (Additional file 6). In the WT line, *PaNAC03* expression is at, or below, the detection limit during early embryo development and no truly quantifiable expression was detected until LE2 (3 weeks after ABA treatment) (Additional file 7). Two OE lines, 4.1 and 4.2, expressing *PaNAC03* at equal levels (1.7 times compared to WT) were tested for maturation capacity with a standard maturation protocol [44] (Filonova et al. 2000). Both lines formed distinct embryonal masses in response to ABA treatment albeit at a lower frequency than the WT line (t-test, *P* = 0.095 and *P* = 0.048 for OE line 4.1 and 4.2 respectively, Fig. 4a). However, the embryonal masses appeared to lack a normal protoderm and rarely developed into...
normal mature embryos (Fig. 4b). Thus, the proliferating OE lines 4.1 and 4.2 and the WT, were selected for transcriptome and metabolite profiling (Additional file 6). A small number of mature embryos with a reduced number of/or fused cotyledons, were obtained from the OE lines (Fig. 4c). The embryos from the OE lines showed a normal germination response after a standard desiccation treatment [44], but a significantly smaller fraction of the germinated embryos showed epicotyl formation and growth (Fig. 4c).

**A limited number of consistently misregulated genes are found in the overexpression lines**

The transcriptomes of the PaNAC03 OE lines and the WT line were sequenced with Illumina HiSeq sequencing generating 15.6-17.9 M reads per sample that passed Illumina’s chastity filter and between 15.4 and 17.8 M read pairs were kept after Nesoni filtering (Additional file 8). The overall read mapping rate from Tophat was 28–63% where most samples had around 60% mapping (Additional file 9).

The analysis of the RNA-seq data-set showed that compared to the WT line 4.1 and 4.2 had 1683 and 740 differentially regulated genes respectively, and 482 genes were consistently misregulated in both OE-lines (Fig. 5). Of these, 153 were consistently up-regulated in both 4.1 and 4.2 and 329 were consistently down-regulated in
strongly regulated gene in the consistently down regulated domain was a gene model, MA_10251997g0010, with similarity to the Arabidopsis transcription factor KANADI (AT5G16560.1) (Additional file 12). Four peroxidases associated with the GO term GO:0042542 were down regulated in the OE lines, three of these were class III peroxidases MA_195910g0010 (PabPrx132), MA_195775g0010 (PabPrx131) and MA_185755g0010 (PabPrx01) (Additional file 11).

**PanNAC03 overexpression lines show reduced levels of flavonoids**

Interestingly, three key genes in the flavonoid biosynthesis pathway were concomitantly down-regulated in the PanNAC03 OE-lines: a chalcone synthase, MA_1035 9605g0010, homologous to the Arabidopsis gene transparent testa 4 (TT4, AT5G13930), a flavonoid 3′-hydroxylase (F3′H, MA_10434709g0010) a possible homologue to the Arabidopsis gene transparent testa 7 (TT7, AT5G07990) and the previously described PaLAR3 gene (MA_10001337g0010) [47, 56] (Additional file 12). We only detected one consistently induced gene associated within the phenylpropanoid pathway, MA_10429470g0020, which encodes an isoflavone reductase with similarity to AT4G39230 which might be involved in lignin biosynthesis.

Given the concomitant down-regulation of Norway spruce homologs to key genes in the flavonoid pathway, we analysed the levels of specific specialized metabolites in the PanNAC03 OE-lines namely of the major stilbenes, the immediate catalytic products of PaLAR3, catechin and gallatechecin, and finally a number of flavonoids. The major stilbene in Norway spruce, astringin, showed no significant differences between the WT and the PanNAC03 OE-lines, neither did the flavonoids kaempferol or isorhamnetin (Fig. 6). However the levels of naringenin, apigenin, eriodictyol and catechin, gallicatechin and their dimers were all lower in OE-line 4.2 ($P<0.05$, One way-ANOVA) and line 4.1 ($0.1>P>0.05$, One-way ANOVA) (Fig. 6).

**PanNAC03 does not suppress the activity of the PaLAR3 promoter**

One of the consistently down regulated genes (PaLAR3, MA_10001337g0010) has been thoroughly studied before [47, 56] and the promoters from two different alleles, PaLAR3A and PaLAR3B, have been isolated, the promoters show a high overall similarity and they differ primarily by two indel-regions present in the PaLAR3A promoter only, containing two putative NAC binding sites (Nemesio-Gorriz 2016). The WT line, 95:61:21, exhibits almost no significant differences between the WT and the PanNAC03 OE-lines, neither did the flavonoids kaempferol or isorhamnetin (Fig. 6). However the levels of naringenin, apigenin, eriodictyol and catechin, gallicatechin and their dimers were all lower in OE-line 4.2 ($P<0.05$, One way-ANOVA) and line 4.1 ($0.1>P>0.05$, One-way ANOVA) (Fig. 6).
Fig. 6 Down-regulation of flavan-3-ol in PaNAC03 OE-lines. Quantification of flavonoids in the WT line 95:61:21 and the PaNAC03 OE-lines 4.1 and 4.2 after two weeks culture. The flavonoids were quantified according to Hammerbacher et al. (2014) \((N = 3)\) superscript letters indicate significant differences (Kruskal-Wallis test).
09g0010, by direct interaction with the promoter of these genes. To test this hypothesis, the PaLAR3A and PaLAR3B promoters were cloned into a GUS reporter vector and were used in a transactivation experiment in N. benthamiana leaves with and without PaNAC03, the basal activity of the promoters and effect of PaNAC03 on these promoters were quantified. The basal expression of the PaLAR3A and PaLAR3B promoters was similar (Figure 7a). However, co-expression of PaNAC03 strongly activated the PaLAR3A promoter (P < 0.05) but did not affect the activity of the PaLAR3B promoter (Fig. 7a), showing a different interaction of PaNAC03 with the two promoters.

To investigate if the two putative NAC binding sites in the indel-region unique to the PaLAR3A promoter (Additional file 3) were the targets for PaNAC03 causing the specific activation of the PaLAR3A promoter we mutated these sites constructing promoter pPaLAR3A_mut. The mutated promoter was cloned into the GUS reporter vector. Thereafter PaNAC03 was co-expressed with either pPaLAR3A_mut or the native pPaLAR3A in N. benthamiana leaves. Interestingly the basal activity of pPaLAR3A_mut was higher than that of the native PaLAR3A promoter (Fig. 7b) showing that these putative NAC binding sites can affect PaLAR3A expression, the deletion of the NAC binding sites did however not affect the transactivation of pPaLAR3A by PaNAC03, there was no significant difference in relative activity of pPaLAR3A_mut (1.5 +/- 0.8 times the promoter alone) or the native pPaLAR3A (2.6 +/- 1.8 times the promoter alone).

Discussion

In this study we identified seven gene models in the Norway spruce genome assembly v 1.0 that show homology to the stress-induced subgroup III-3 NACs in Arabidopsis [1]. Generally, the Norway spruce subgroup III-3 NAC gene family members are highly similar displaying the conserved N-terminal A-E motifs characterizing NAC domain proteins [22] and also a relatively conserved C-terminal region including a conserved C-terminal motif SEKEE (V/I) QSSFRL, a motif present also in most sequences amplified with the marker Sb29 [57]. The conserved C-terminal region in Norway spruce subgroup III-3 members is different from the C-terminal motifs Arabidopsis subgroup III-3 members and likely the Norway spruce promoters do not have a transactivation domain similar to the Arabidopsis members [1], and it is unclear if the function of Norway spruce subgroup III-3 NAC is similar to that of Arabidopsis subgroup III-3 members. The Norway spruce subgroup III-3 NACs share, at least partly, an element of stress inducibility with the Arabidopsis members based on the expression data available in the Norway spruce genome portal. Three of the Norway spruce subgroup III-3 gene models were identical to the PUTs PaNAC03, PaNAC04 and PaNAC05 from a de novo transcriptome assembly of the interaction between Norway spruce and H. annosum s.s. [9, 26]. A fourth gene

![Fig. 7 Transactivation of PaLAR3 promoters by PaNAC03. Figure a, transactivation of native PaLAR3A (open and light grey bars) and PaLAR3B promoters (open and light grey hashed bars) by PaNAC03 in N. benthamiana, the figure shows the results from one of three representative experiments (N = 12). Figure b shows one representative transactivation experiment with native PaLAR3A (open and light grey bar) and PaLAR3A_mut promoters (open and light striped bars) by PaNAC03 in N. benthamiana (N = 9). The mean +/- SE is indicated for each measurement. Asterisks indicate significant differences * = P < 0.05; ** = P < 0.01; *** = P < 0.001 (Mann-Whitney U-test)
model, MA_86256p0010, showed similarity to another PUT. Taken together, it shows that certain Norway spruce subgroup III-3 NACs, like their Arabidopsis orthologs, respond to biotic stress. Interestingly, the biotic stress responsive gene models to (PaNAC03, PaNAC04, PaNAC05 and MA_86256p0010) cluster together in the phylogeny with MA_103386p0010 and MA_64687p0010, which do not respond to H. annosum s.i. inoculation, forming a sister group to the Arabidopsis ANAC032, ATAF1, ATAF2 and ANAC102 proteins and to a group of rice NAC genes known to respond to abiotic stress [58]. This differentiation in terms of expression pattern between Norway spruce paralogs is consistent with the concept of subfunctionalization [59].

To confirm the responsiveness to biotic and abiotic stress, the expression of PaNAC03 and PaNAC04 was analysed in response to H. annosum s.s. infection or wounding in phloem of mature Norway spruce trees by qRT-PCR. Both genes showed significant induction in response to either treatment, but as previously reported for other Norway spruce genes, [5, 7, 9, 60], the induction was higher after inoculation than after wounding. However only PaNAC03 transcription was significantly induced in response to H. parviporum treatment. This discrepancy seen in the induction patterns of PaNAC04 between experiments could be an effect of several different factors; obviously different organs of conifers show different transcriptional responses to pathogens in seedlings, the responses appears to be more organ-specific than pathogen-specific [4] suggesting that the organ analysed, phloem versus seedling roots, could explain the differential induction between PaNAC03 and PaNAC04. Furthermore, the age of the host is another factor which may affect the manifested defence responses in conifers. The two fungal species, H. annosum s.s. and H. parviporum, are known to have different host preferences (as reviewed in [11]) and different capacity to produce phytotoxins [61], suggesting that their interaction with the host may differ and could lead to a differential regulation of PaNAC04. However, in our previous studies confronting Norway spruce plants of different ages with H. parviporum and H. annosum s.s. [5, 7, 9, 60] we have seen consistent induction of defence related genes with the two different fungal species over different age classes and tissues tested. Thus, the gene induction patterns could indicate that PaNAC03 is likely to have a closer association with the transcriptional responses to biotic stress in Norway spruce than PaNAC04. Therefore we selected PaNAC03 for functional analysis by overexpression in somatic embryogenic cultures.

The overall plan of embryo development is similar in angiosperms and gymnosperms despite their separation approximately 300 million years ago. There are, however, several distinct differences in the embryo development programme between the two plant lineages. In angiosperms the first tissue to differentiate during embryogenesis is the protoderm which is formed by periclinal divisions of cells of the early globular embryo [62]. The formation of the protoderm, which restricts cell expansion, is essential for the remaining developmental process [63]. In contrast, in gymnosperms the surface layer of the embryonal mass divides both periclinal and anticlinally. Nevertheless, the outer cell layer in the embryonal mass in Norway spruce embryos defines a functional protoderm [63, 64]. The developing embryonal masses in the PaNAC03 OE-lines masses appeared to lack the normal conifer “protoderm” i.e. a smooth outer surface, the ruggedness of the embryo surface were reminiscent of the phenotype of other transgenic Norway spruce lines with a disturbed protoderm formation [35, 63, 64]. The embryonal masses did generally not develop into mature cotyledonary embryos. A small number of mature, but aberrant looking, embryos were recovered from the OE-lines. These embryos showed a normal germination response but a significantly smaller fraction of the germinated embryos showed epicotyl formation and growth. Based on these observations we concluded that the embryo development programme is disturbed at a very early stage in the PaNAC03 OE-lines.

Among the 482 consistently misregulated gene models identified by transcriptome sequencing of the two PaNAC03 OE-lines we found a number of genes known to control various aspects of patterning and embryo development in Norway spruce. The most strongly induced gene model is MA_122121g0010 which encodes a HD-ZIP IV protein highly similar to PaHB2 [53] and the Arabidopsis genes GLABRA 2 and ANTHCYANINLESS2 [54, 55, 65] associated with patterning in Norway spruce and Arabidopsis while PaHB1, controlling protoderm formation [64], was slightly down regulated in the OE lines. PaHB2 is not expressed during early embryo development in WT-lines [53] and neither is MA_122121g0010. Thus, the misregulation of MA_122121g0010 may be the cause of the aberrant embryo morphology in the OE-lines. Other strongly upregulated gene models in the OE-lines with potential to cause of the aberrant embryo morphology encode HBK4 (MA_114226g0010), PaPIN1 (MA_100472g0010) and PaACT4 (MA_135063g0010), which all have been shown to control Norway spruce somatic embryo development [18, 66, 68], and specifically the differentiation of the shoot apical meristem and cotyledons [18, 69] processes which appears to be affected in the PaNAC03 OE lines. A gene model (MA_10251997g0010) with similarity to the Arabidopsis gene KANADI1 (AT5G16560.1) shows a three-fold lower expression in the PaNAC03 OE-lines. In Arabidopsis embryos KANADI1 is initially expressed in the central domain protoderm at the late globular embryo stage and appears to have a role in specifying the peripheral identity in
the developing Arabidopsis embryo in interplay with HD-ZIP III proteins [70–72]. It may be noteworthy that, as mentioned before, PaPIN1 and also a Norway spruce HD-Zip III gene model (MA_10427484g0010) with similarity to ATHB15 (AT1G52150) and appears to show contrasting induction levels compared to the KANADI-like gene in OE-lines; these expression patterns are reminiscent of the interaction between the KANADI1, HD-ZIP III and PIN1 in Arabidopsis [71, 73]. Taken together the maturation and RNA-seq data indicates that ectopic expression of PaNAC03 interferes with the protoderm formation and early embryo patterning through misregulation of transcriptional modules controlling these processes. This pleiotropic effect must be taken into consideration when further examining the consistently misregulated genes in PaNAC03 OE-lines as any gene misregulation may be an effect of the aberrant early embryo development.

About two thirds of the consistently misregulated genes were consistently repressed in the overexpression lines. The consistently repressed genes more commonly associated GO terms related to response to abiotic stimuli, stress responses and responses to hydrogen peroxide. There were three consistently down regulated class III peroxidases, including PaPrx01, among the consistently misregulated genes. Previously, PaPrx01 has been shown to respond to H. parviform treatments in Norway spruce cultures [74] and it is suggested to contribute to H2O2 production in suspension cultures of Norway spruce, indicating a potential role of PaNAC03 in redox homeostasis under stress in Norway spruce as H. parviform treatments in Norway spruce cultures appears to repress several peroxidases [74].

We observed consistent misregulation of three key genes in the flavonoid biosynthesis pathway in the overexpression lines, a CHS homologous to the Arabidopsis thaliana gene transparent testa 4 (TT4, AT5G13930), one F3’H homologous to the Arabidopsis gene transparent testa 7 (TT7, AT5G07990) and the previously described PaLAR3 gene [47]. Variation in the PaLAR3 locus associated with enhanced resistance to H. parviform and with increased accumulation of the catalytic product of the enzyme (+) catechin [56].

The concomitant misregulation of key genes in the flavan-3-ol pathway is associated with reduced levels of flavan-3-ols in the OE-lines; both naringenin and apigenin, which are products formed downstream of CHS but before steps catalysed by either F3’H or PaLAR3 were down regulated in the PaNAC03 overexpression lines (as indicated in Fig. 6). Eriodictyol, a catalytic product of F3’H was also reduced. The catalytic product of PaLAR3, (+)-catechin, was also significantly reduced in the OE lines. While other metabolites not directly associated with flavan-3-ol production accumulated to the same levels as in the WT line, showing that the down regulation of key members in the flavan-3-ol pathway lead to a specific reduction in these compounds. Although regulation of anthocyanin or proanthocyanin pathways by NAC TFs is not commonly reported in literature. The NAC TFs BL, controlling the blood red flesh phenotype in peach, and ANAC078 in Arabidopsis appear to control certain members of the anthocyanin or proanthocyanin pathways [75, 76]. The metabolite and transcriptome profiling of the OE lines appeared to indicate that PaNAC03 could act as a negative regulator of 3-flavanol production in Norway spruce, possibly by acting directly on the misregulated flavonoid biosynthesis genes. To test this possibility we co-expressed PaNAC03 with the promoter of either of the two alleles at the PaLAR3 locus [56] in N. bethamiana leaves; hypothesising that PaNAC03 would reduce PaLAR3 promoter activity if it acts as a repressor. However, in this system PaNAC03 strongly activated the promoter of the PaLAR3A allele suggesting that PaNAC03 does not act as a negative regulator of flavan-3-ol production by direct interaction with PaLAR3. However, the down-regulation of CHS transcription, encoding the rate-limiting step in flavonoid biosynthesis [77] might have had an effect on substrate availability for downstream metabolite biosynthesis, explaining the lower transcriptional and metabolite levels observed in our study. Transcript profiling also showed an up-regulation of an isoflavone reductase gene that could be involved in lignin biosynthesis [78]. Lignan and lignin biosynthesis directly compete for substrates used in the flavonoid pathway and might therefore also negatively regulate flavonoid biosynthesis, as has been observed in our PaNAC03 over-expressing lines. It is possible that the down-regulation of CHS, F3’H and PaLAR3 genes in PaNAC03 overexpressing lines could be mediated by another factor such as misregulation of an upstream regulatory gene or the interference of constitutive PaNAC03 expression with early embryo patterning. It should be noted that flavonols and the transparent testa mutants has been linked to auxin homeostasis and polar auxin transport [79, 80] in plants. Another possible explanation to the discrepancy between the overexpression and transactivation experiments is that PaNAC03 act in a heterodimer, as has been shown for other stress-responsive NACs [81, 82], with a currently unidentified TF to downregulate the CHS, F3’H and PaLAR3 genes in the OE-lines. This possibility could be tested by yeast two-hybrid screening of cDNA libraries from embryogenic cultures using PaNAC03 as a bait.

Conclusion
PaNAC03 and its orthologs form a sister group to well characterized stress-related angiosperm NAC genes and at least PaNAC03 is responsive to biotic stress and appear to act in the control of defence associated secondary metabolite production. However, the unexpected embryo
phenotype of the PaNAC03 OE lines emphasizes the still enigmatic connection between specialized metabolism and patterning in plants, raising questions on the role of subgroup III-3 NAC TFs in development and embryo patterning.

Additional files

- **Additional file 1:** Sequences of primers used in the study. (XLSX 11 kb)
- **Additional file 2:** Scripts used for Nesoni, tophat, cufflinks and cuffdiff. (XLSX 14 kb)
- **Additional file 3:** Figure S2. Schematic representation of the PaLAR3A promoter (Genbank accession no. KX674229.1) in black and the PaLAR3B promoter (KX674230.1) in red, the white regions in PaLAR3B promoter corresponds to deletions in the sequence compared to PaLAR3A promoter. The two NAC binding sites (TTTCGT) present in the region unique to the PaLAR3A promoter are indicated in yellow. In the PaLAR3A_mut promoter it is only these two sites which has been mutated in the remainder of the PaLAR3A promoter is intact. (XLSX 69 kb)
- **Additional file 4:** Clustal W Alignment of Norway spruce subgroup III-3 NAC proteins. The coloured boxes correspond to the conserved N-terminal motifs. A (light blue), B (pale green), C (pale red), D (black) and E (pale gold). The shaded residues indicate residues conserved in the C-terminal region. (XLSX 21 kb)
- **Additional file 5:** Amino acid identity and similarity in subgroup III-3 NAC proteins. Percent amino acid identity (above the diagonal) and similarity (below the diagonal) in the complete protein sequences (A) or the C-terminal part of the proteins (B). (DOCX 16 kb)
- **Additional file 6:** Relative expression of putative PaNAC3 overexpression lines. The relative expression was determined in relation to the untransformed wild type line 95:61:21. (DOCX 33.1 kb)
- **Additional file 7:** Transcriptional regulation of PaNAC3 in response to standard maturation treatment in the wild type line 95:61:21. (PDF 447 kb)
- **Additional file 8:** Table S7. RNAseq metrics after Nesoni filtering. (DOCX 19 kb)
- **Additional file 9:** Alignment summaries from tophat. (DOCX 11 kb)
- **Additional file 10:** Enriched GO terms among consistently down- or up-regulated genes in PaNAC3 overexpression lines. (TIF 45020 kb)
- **Additional file 11:** Consistently up-regulated genes in PaNAC3 overexpression lines. (DOCX 15 kb)
- **Additional file 12:** Consistently down-regulated genes in PaNAC3 overexpression lines. (DOCX 17 kb)

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Availability of data and materials

The short read data generated from the NAC OE lines is deposited in SRR5022423-SRR5022431 in BioProject PRJNA350779 at the NCBI. Further requests for materials should be addressed to ME (Malin.Elfstrand@slu.se).

Authors’ contributions

ME, IE and KL conceived the study. KD planned and executed the experimental work with some assistance by MNG. AH and MNG performed the chemical profiling of the NAC OE lines. JW planned and executed the transactivation study in N. benthamiana. KD, ME and MNG drafted the manuscript; all authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

Consent for publication

Not applicable.

Ethics approval and consent to participate

Not applicable.

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