



	<b>Experiment title:</b> Uppsala (II) BAG, LS-1665 (T. Alwyn Jones BAG) Cellulases; EG3	<b>Experiment number:</b> LS 1665
<b>Beamline:</b> ID 14:EH4	<b>Date of experiment:</b> from: 28 Apr 2000 to: 30 Apr 2000	<b>Date of report:</b>  <i>Received at ESRF:</i>
<b>Shifts:</b> 1.5 (EG3: 1.5)	<b>Local contact(s):</b> Julien Lescar	

**Names and affiliations of applicants** (\* indicates experimentalists):

T. Alwyn Jones, Uppsala University, alwyn@xray.bmc.uu.se  
Sherry L. Mowbray, Swedish Univ. Agric. Sciences, mowbray@alpha2.bmc.uu.se  
Jerry Ståhlberg, Swedish Univ. Agric. Sciences, Jerry.Stahlberg@molbio.slu.se  
\* Mats Sandgren, Uppsala University, mats@alpha2.bmc.uu.se  
\* Inés Muñoz, Swedish Univ. Agric. Sciences, ines@alpha2.bmc.uu.se  
\* Torsten Unge, Uppsala University, torsten@alpha2.bmc.uu.se  
\* Seved Löwgren, Uppsala University, seved@alpha2.bmc.uu.se

**Report:**

**Protein Name EG3: Endoglucanase 3 from *T. reesei***

**Brief background and outline of project**

(Cel12A, GH family 12). EG3 is a minor component in the cellulase system, but may nevertheless play an important role. It differs from the other cellulases in that it does not contain the additional cellulose binding module and linker typical for many cellulases. It is also smaller than other fungal celolytic enzymes and subsequently appears to have a larger ability to penetrate the cellulose substrate. We have recently solved the structure and now aim at obtaining ligand complexes with wild-type and mutated enzymes for understanding catalytic mechanism, substrate binding and specificity. Very large crystals have been obtained that diffract to very high resolution (0.9-1.0 Å) and which could hopefully yield structures at atomic resolution.

**Datasets collected on ID14:EH4**

Two datasets were collected at ID14:EH4 on protein crystals from two new *T. reesei* EG3 apo catalytic inactive mutants. Both datasets were of high quality and fairly high completeness. The space-group of these datasets were the same as previously datasets P21, but the cell parameters were new and the datasets had to be solved by molecular replacement methods. Both these catalytic inactive mutants structures have now been solved and refined. Neither of these structures did contain a ligand in the active site but these structures will be used in a

manuscript, in progress right now, where we try to explain the catalytic mechanism of the enzyme.

### **Data statistics for the datasets:**

#### Dataset1:

Ligand: No

Mutation: A

Resolution: 40-1.7A

Space group: P21

Cell: a:62.49, b:77.55, c:83.41,  $\beta$ : 90,  $\alpha$ :98.46,  $\gamma$ :90

Completeness: 98%

#### Dataset 2:

Ligand: No

Mutation: B

Resolution: 30-1.2A

Space group: P21

Cell: a:43.1, b:103.8, c:45.33,  $\beta$ : 90,  $\alpha$ :118.4,  $\gamma$ :90

Completeness: 90%

