

# Airbus A320-231, G-OOAC

## AAIB Bulletin No: 11/97 Ref: EW/C97/5/7 Category: 1.1

<b>Aircraft Type and Registration:</b>	Airbus A320-231, G-OOAC
<b>No &amp; Type of Engines:</b>	2 International Aero Engine V2500-A1 turbofan engines
<b>Year of Manufacture:</b>	1992
<b>Date &amp; Time (UTC):</b>	26 May 1997 at 0555 hrs
<b>Location:</b>	Bristol Airport
<b>Type of Flight:</b>	Public Transport
<b>Persons on Board:</b>	Crew - 7 - Passengers - 175
<b>Injuries:</b>	Crew - Nil - Passengers - Nil
<b>Nature of Damage:</b>	Damage to the No 3 brake pack
<b>Commander's Licence:</b>	Airline Transport Pilot's Licence
<b>Commander's Age:</b>	39 years
<b>Commander's Flying Experience:</b>	7,680 hours (of which 3,720 hours were on type) Last 90 days - 212 hours Last 28 days - 77 hours
<b>Information Source:</b>	AAIB Field Investigation

## History of the flight

After the aircraft arrived at Bristol, it was prepared for the next flight to Mahon. The oncoming commander confirmed from the Technical Log that there had been no unserviceabilities on the previous flight and that there were no 'Carried Forward Defects'; he also completed an external check which revealed no apparent defects.

The pre-start checks were normal as was the start and pushback from the stand. No abnormalities were noted and the aircraft was taxied to the active runway. The first officer had been designated as the handling pilot for the outbound sector and, after a short hold, he carried out a normal take-off. When safely airborne, the commander retracted the landing gear and the aircraft was established in the climb. The flight continued uneventfully until approximately 35 minutes later when, with the aircraft under Brest ATC control, the crew were advised by ATC that parts of a brake unit and some

hydraulic fluid had been found at Bristol. The commander then contacted Bristol ATC to clarify the message and it was confirmed that brake unit parts and hydraulic fluid had been found on the taxiway, at the runway hold area and on the runway; he was also advised that the debris was suspected to have come from his aircraft. The commander then made a comprehensive check of the aircraft displays and instrumentation, but there were no fault indications. Following radio contact with his company operations department, who confirmed that the debris had probably come from G-OOAC, the commander decided to divert to London Gatwick. He chose this destination as the most suitable considering his location, the company engineering support at Gatwick, the runway length and his experience of the airport.

The diversion was uneventful and the commander took over the handling duties for the approach to Runway 26 at Gatwick. He did not declare an emergency, but advised ATC of the situation and requested that the Airport Fire Service (AFS) meet the aircraft after landing. The cabin staff and passengers had been briefed before the approach. The commander made a gentle landing and then used full reverse thrust to retard the aircraft; autobrake had intentionally not been selected and gentle wheel braking was applied as the ground speed slowed to 60 kt. Once the aircraft had cleared the runway, the commander brought it to a halt and shut both engines down. The AFS were already positioned and confirmed that there was no excessive heat evident from the aircraft wheels; on the flight deck, the crew noted that the brake temperatures remained below 100°C. The aircraft was then towed to the parking stand where the crew and passengers disembarked normally. An initial examination revealed that the No 3 brake unit had failed.

### **Brake unit description and history**

The brake units fitted to this aircraft were of the 'carbon' type, with the rotors, stators and end discs (the heat pack) all made from a carbon/carbon matrix. Each disc is manufactured from stacked layers of Rayon fabric, and processed under a combination of heat and pressure such that a rigid, almost pure, carbon material is formed which has good structural strength resulting from the directionality of the original Rayon fibres. The type of brake unit fitted to this aircraft is shown in Figure 1 and comprised four rotor discs, three stator discs, and two pressure plates. All discs have wearing surfaces on both sides except the two pressure plates, which abut only against the first and last rotors. The rotors and stators are 'keyed', respectively, to the wheel hub and torque tube by integral metal capped lugs at their outer or inner edges. To generate braking torque, pressure is applied across the stack by a multiple piston hydraulic assembly, each piston acting through metal insulators and a thrust plate. Half of the pistons are associated with the parking brake system, the other half with the normal braking system. The aircraft hydraulic supplies to the brake units have hydraulic fuses installed to limit fluid loss in the event of a serious leak. The claimed advantages of 'carbon' disc brakes are that they are significantly lighter than steel based brakes and have a higher thermal capacity. They possess good friction characteristics and generally should exhibit long lives, particularly if used in a manner such that they become 'hot', as advocated by the manufacturer. At low temperatures they are prone to 'snatch', or operate roughly, and wear more quickly. In common with most brakes, high braking torque is often generated at low rotational speed. Other similar brake failures examined during this investigation had occurred at low speeds or on pushback from the stand. The failed unit from G-OOAC had been overhauled by the manufacturer and released to the operator on 19 December 1995. It entered service on 7 April 1996 at position No 3 where it remained until the failure. During that time it had accumulated 4,924 flight hours and 1,661 landings.

### **Initial examination**

Debris removed from the runway at Bristol and the damaged brake unit from the aircraft were transported to the AAIB where an initial examination was carried out in conjunction with a representative of the manufacturer. It was determined that the carbon pressure plate nearest to the pistons housing, the adjacent rotor and first stator disc were all missing from the heat pack, but it was apparent that the debris from the runway was composed of elements from these missing parts, with all but one of the stator clips having been recovered (Figure 2). The distribution of this debris on the runway indicated that disc failure had occurred at, or about, the point of brake release prior to the take-off roll, with the fragmentation and general disc damage having probably occurred during the take-off run. The severely damaged metal thrust plate and nine of the fourteen insulators, two of which had broken diametrically across the rivet hole, were also recovered. Distortion and damage also occurred to several of the hydraulic pistons as these had extended to full travel during the brake failure, releasing a quantity of hydraulic fluid. It was apparent that the failure had been initiated by detachment of one or more lugs, probably on the first stator, and that the resultant entrapped debris had caused major disruption of the affected discs as the wheel rotated during taxi and the take-off roll.

Initial visual inspection of the carbon discs in this brake unit showed all to exhibit evidence of oxidation, the severity of which increased towards the piston housing end of the brake. Generally, the inner regions of the discs had suffered to a greater extent than the outer. In these circumstances, oxidation is manifest by loss of the material (with the consequent reduction in load-carrying ability of the discs, particularly around the inner and outer lugs), with carbon combining with oxygen to form CO and CO<sub>2</sub> (carbon monoxide and carbon dioxide gases). This effect becomes evident above 426°C and is dependent both on time and temperature. The rate of material loss, however, increases rapidly with increasing temperature. Figure 3 shows a comparison between lightly and severely oxidised surfaces.

In order to counter oxidation, the exposed 'non-wearing' surfaces of each carbon disc are treated with an inhibitor or anti-oxidant (AO), in this case the manufacturer's proprietary material based on phosphoric acid (H<sub>3</sub>PO<sub>4</sub>). This is painted on by hand in the form of a transparent liquid and then baked at high temperature. Under magnification, it normally appears as a crystalline salt deposit which should completely cover the surface. Soon after the introduction of these brakes the AO compound used was modified, due to its poor performance, to a standard known as M-1 and this was the standard released for production in December 1992 and applied to the discs in the subject brake.

### **Detailed examination**

Discussions with the brake manufacturer and several UK based operators of the A320, who used the same brake units, revealed that there had been a series of similar failures affecting, particularly, the three main charter airlines operating A320 aircraft in the UK. One operator in particular had experienced 11 incidents of disc failure between August 1996 and May 1997 over a fleet of 7 aircraft, with landings achieved at failure varying between 857 and 1,318. In an attempt to further understand this problem, detailed examination was carried out on samples taken from discs in the failed brake and from two newly manufactured/unused discs (one treated and one untreated) which were supplied by the manufacturer. This examination was performed using a scanning electron microscope (SEM) and energy dispersive X-ray (EDX) analyses with the aim of obtaining data on the degree of protection afforded by the M-1 inhibitor on both used and new discs. This was to be assessed in terms of surface coverage, penetration into the matrix of the carbon material, the level of any contamination on the failed discs and the possible effects of elevated temperature. The manufacturer had indicated that certain contaminants such as sodium, potassium, calcium,

chromium, iron and seawater, can act in a catalytic manner to oxidise carbon and pointed out that potassium acetate is a major constituent of many commercially available runway de-icers. Phosphoric acid has an affinity with iron.

*In the figures referred to below, most SEM micrographs were taken using the Backscattered Electron Detector (BSD), which images regions of varying atomic weights as differences in grey level - 'heavier' elements appear brighter than 'lighter' ones. Such images have the notation 'Detector = QBSD' in the data zone. Other SEM micrographs were taken using the Secondary Electron (SE) detector, which gives a higher resolution image and shows good topography. These images have the notation 'Detector = SEI' in the data zone.*

Samples for examination were cut from two locations, i.e. the outermost and innermost surfaces from the following discs (see Figure 1), including at least one surface which had been protected by a statormetal clip:- i.e. disc 7, disc 5, the uninhibited unused stator (used as a reference) and the unused M-1 inhibited stator. All the samples, the external surfaces of which were to be examined, were mounted on SEM stubs such that the AO treated surfaces were uppermost. As they were electrically conductive, it was possible to examine them directly in the SEM. Samples to be sectioned were first 'potted' in a resin block, ground and polished, and then lightly 'sputter-coated' with platinum so that the entire sample became sufficiently electrically conductive to enable it to be imaged within the electron beam of the SEM.

Figure 4 and Figure 5 illustrate surface and sectional detail of the unused M-1 treated disc. It was apparent from these micrographs, which are representative of all the surfaces and sections examined, that the AO deposits (bright areas) did not completely cover the carbon material, indicated here as fibre bundles of different orientation, and that there was significantly more AO present on the inner surface than the outer. It was also evident from the sectional views that the carbon matrix was not homogeneous but contained voids, at least in the material close to the surface, and that there was significant, but not total, penetration of the AO into these voids with the better protection occurring at the inner surface.

Figure 6 shows similar SEM views from disc 7 (failed brake unit) and this was typical of the other discs examined from this brake. Incomplete coverage of the surfaces by the AO is apparent in addition to the effects of oxidation, i.e. rougher surfaces than seen on the unused disc. Figure 7 illustrates an area of AO that had become globular in form and this characteristic was seen in areas throughout the disc samples examined from the failed brake unit. An EDX scan of the elemental composition of these globules is shown in Figure 9 where, in addition to the main constituents of the AO, there are traces of iron, sodium, aluminium, silicon, and potassium. This disc did not appear to have been contaminated with hydraulic oil (Skydrol LD4), as had occurred to the broken discs close to the piston housing. Two spectrochemical analyses of the oil taken from the affected system (green) several months (21 February 1997) before and shortly after this incident (5 July 1997) are shown in Figure 10. Minor traces of the elements zinc, chromium, silicon and calcium were found to be present in the oil in the earlier sample and chromium and calcium in the later sample, but neither contained sodium or potassium as identified by the EDX scan of the globules. This also suggested that the sample shown in Figure 7 had not been contaminated with oil and that the elements detected represented a low level of 'in-service' contamination.

Four of the samples taken from the unused M-1 inhibited disc, after initial examination, were each then heated to a different temperature ranging from 300°C to 600°C in a pre-heated oven for 15 minutes. After cooling, each sample was re-examined and no significant change in the appearance of the surface or the character of the AO material was apparent. However, the sample heated to

300°C was then re-heated to 1000°C and re-examined, whereupon it was seen that in some areas the AO had taken up a globular form as shown in Figure 8. This sample also showed a different ratio of elements when compared with other samples. The levels of phosphorus (P) had decreased while those of aluminium (Al) and oxygen (O) had increased.

### **Additional information**

Due to concern arising from these brake failures, several measures have been introduced/proposed by the airlines concerned and the brake manufacturer, such as wheel removal on a regular basis for examination of the exposed areas of the outer surfaces of discs for units with in excess of 1,100 landings. As part of an on-going reliability assessment, one of the affected airlines had required crews to log post-landing peak brake temperatures as indicated on the Engine Indicating and Crew Alerting System (EICAS), some 5 minutes after the aircraft was parked, over a period of some 5 weeks. The results from this showed that, generally, indicated temperatures ranged between 200°C and 400°C but that, following approximately 5% of the documented landings, temperatures exceeded 400°C. The maximum temperature recorded during this period was 680°C following a landing with a tailwind where medium auto brake was used. These were sensed temperatures from a probe fitted to the pistons' housing; actual disc temperatures may have been higher.

In order to obtain a comprehensive assessment of brake operating conditions of that airline, the brake manufacturer has developed a 'data logger' known as the Brake Performance Monitoring System (BPMS), which is designed to fit into the standard electronic equipment rack without requiring modification. All information relevant to the brakes on the various data busses would be monitored and logged over a period of several weeks of revenue service. Approval to install this unit by the manufacturer and airline is currently being sought from the CAA.

### **Conclusions**

Whilst the examination of the samples from this failed brake unit was not exhaustive, and other failed brake unit discs were not examined in such detail, this investigation revealed apparent similarities between the new M-1 treated disc supplied by the manufacturer and disc samples from lightly oxidised areas of the failed brake unit. The distribution of the AO was extremely patchy, and the amounts on the inner and outer surfaces differed, leaving areas of 'non-wearing' carbon exposed to the atmosphere. Where the carbon had been heavily oxidised it was not possible to be certain of the extent of the original AO coverage but, if the new disc was typical of in-service discs, then oxidation would have occurred whenever the disc temperature rose above 426°C. The AO penetration into the material below the surface of this unused disc was 8.5mm. There were, however, many voids within the carbon of all disc sections examined, with only a proportion exhibiting evidence of AO penetration from the surface to provide any protection. It therefore seemed probable that when high brake temperatures were experienced oxidation could also have occurred below the protected surfaces if air were able to access these voids.

The heating tests on the samples showed that the general appearance and elemental content of the AO did not appear to change appreciably up to 600°C. The one sample initially heated to 300°C and subsequently to 1000°C, however, showed the AO to have partially adopted a globular form at the higher temperature, similar to that seen in areas on the surfaces of the discs from the failed brake. This suggested that the brake unit had been operated to temperatures significantly higher than for the onset of oxidation to occur, the extent of oxidation indicating that this had most likely been

occurring on a regular basis. This view was supported by the peak temperatures logged by flight crews over a period of normal revenue operation.

Some contamination of the AO was found on discs from the failed brake. A large amount of this contamination appeared to be iron, but sodium and potassium were also present in many areas, particularly on disc 7 (least damaged and apparently unaffected by hydraulic oil contamination), where there was more AO on the surfaces. Much of the globular formation of the AO was particularly rich in iron, one of the elements recognised to act in a catalytic manner in the oxidation of carbon. Thus the severe oxidation seen on these failed discs seems to have resulted from incomplete AO coverage on the exposed surfaces and penetration of the sub-surface voids, compounded by operation at disc temperatures in excess of 426°C. Contamination by 'catalytic' elements was found and these may have been a contributory factor in the failure mode of the discs. Since August 1996 the manufacturer has again modified the composition of the AO to further enhance its performance, this latest version being known as M-2, and data is currently being gathered from their various overhaul facilities with a view to identifying an expected improvement in disc condition and life. The operator which had experienced 10 brake failures advised that, of some 40 installed brake units on two aircraft fleets for which it had responsibility, there were currently only 3 units which were known to contain M-1 treated discs. Since the installation of M-2 treated heat packs, they reported a marked reduction in the presence of carbon oxidation and a gradual increase in the number of landings achieved to overhaul.